No. GN 44, Wednesday, 23 November 1988

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GOVERNMENT NOTICES

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Special Gazettes Nos S 351, S 352, and S 353 are herewith

The date of publication of this Gazette is 23 November 1988.

ADVERTISING RATES FOR **GOVERNMENT NOTICES**

These rates will be:

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Notices submitted for publication will be charged at the rate of \$260.00 per typeset page plus \$200.00 per issue. Material supplied as camera-ready copy and magnetic tape (altered and unaltered) will be charged at the respective Government Notices rates.

Variation of closing times

CHRISTMAS/NEW YEAR PERIOD

Commonwealth of Australia Gazette

The last regular Gazette for 1988 will be the Public Service issue to be published on 22 December 1988 with normal closing times. There will be no regular issues of the Gazette on 27 December 1988, 28 December 1988, 29 December 1988, 3 January 1989, 4 January 1989 and 5 January 1989.

The Government Notices Issue of 11 January 1989 will have normal closing times.

Friday, 6 January 1989 at 10.00 a.m.

Departments are requested to note the dates upon which regular issues will not appear and to make every effort to avoid the need for Special Issues during the holiday period by making arrangements for all necessary administrative and executive material to be gazetted by Wednesday 21 December 1988. Where possible all other material should be held over until the first regular issue of 1989.

N.N.-8878067

GENERAL INFORMATION

IMPORTANT COPYRIGHT NOTICE

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Government Notices issues, published each Wednesday, containing all legislation, proclamations, special information and government departments notices and are sold at \$5.95 each or on subscription of \$290.00 (50 issues), \$150.00 (25 issues) or \$75.00 (12 issues).

NOTICES FOR PUBLICATION and related correspondence should be addressed to:

Gazette Officer, Australian Government Publishing Service, GPO Box 4007, Canberra ACT 2601. Telephone (062) 95 4656

or lodged at AGPS, Government Printing Office Building, Wentworth Avenue, Kingston. Notices are accepted for publication in the next available issue, unless otherwise specified.

Except where a standard form is used, all notices for publication must have a covering instruction setting out requirements. A typewritten original or good copies are to be provided, wherever possible double-spaced, with a margin surrounding the typewritten matter. Copy is to be confined to one side of the paper, sheets are to be of uniform size (preferably A4), numbered consecutively and fastened securely together. Dates, proper names and signatures particularly are to be shown clearly.

Copy will be returned unpublished if not submitted in accordance with these requirements.

CLOSING TIMES. Notices for publication should be lodged at AGPS, Government Printing Office Building, unless otherwise specified, by the following times (except at holiday periods for which special advice of earlier closing times will be given).

Government Notices Gazette all copy: Friday at 10.00 a.m. in the week before publication.

PRIVATE NOTICES

The rates of charge and conditions applying to acceptance of copy for private notices are as follows:

- (a) minimum charge up to 125 words \$58.00;
- (b) each 25 words (or part thereof) thereafter \$12.00.

Remittances must be forwarded with a copy of the notice for publication unless prior credit approval has been granted for account customers. Account customers are reminded that payment is due immediately on presentation of invoice. Should payment not be received within twenty-eight days of the invoice date, credit privileges will be withdrawn.

Notices received without payment or from account customers whose credit privileges have been withdrawn will be returned unpublished.

SUBSCRIPTIONS are payable in advance and are accepted for a maximum period of one year. All subscriptions are on a firm basis and refunds for cancellations will not be given. Rates include surface postage in Australia and overseas. Other carriage rates are available on application.

AVAILABILITY. The Gazette may be purchased by mail from:

Mail Order Sales, Australian Government Publishing Service, GPO Box 84, Canberra ACT 2601

or over the counter from Commonwealth Government Bookshops at:

 Adelaide:
 55 Currie St, tel. (08) 237 6955

 Brisbane:
 294 Adelaide St, tel. (07) 229 6822

 Canberra:
 70 Alinga St, tel. (062) 47 7211

 Hobart:
 162 Macquarie St, tel. (002) 23 7151

 Melbourne:
 347 Swanston St, tel. (03) 663 3010

Perth: 200 St George's Tce, tel. (09) 322 4737 Sydney: 120 Clarence St, tel. (02) 29 1940

Commonwealth Acts and Statutory Rules, Australian Capital Territory Ordinances and Regulations, and other Commonwealth Government publications may also be purchased at these addresses.

ALL REMITTANCES should be made payable to: Collector of Public Moneys, Australia Government Publishing Service.

OTHER ISSUES OF THE GAZETTE

Public Service issues contain notices concerning administrative matters, including examinations, vacancies, transfers and promotions within the Australian Public Service and the Services of the Australian Postal Commission, Australian Telecommunications Commission, Commonwealth Teaching Service and Defence Force appointments etc. These issues are published weekly at 10.30 a.m. on Thursday, and sold at \$8.95 each or on subscription of \$395.00 (50 issues), \$206.00 (25 issues) or \$103.00 (12 issues).

Business issues, published each Tuesday, containing Notices under the Co-operative Companies and Securities Scheme, Bankruptcy Act and Private Notices and sold at \$3.95 each or on subscription of \$220.00 (50 issues), \$116.00 (25 issues) or \$58.00 (12 issues).

Special issues include notices which require urgent publication. All costs associated with producing Specials will be borne by the responsible department or authority. A limited number of Special Gazettes will be made available for sale from the Commonwealth Government Bookshop, Canberra, on the day of publication. General distribution of these notices will be by their inclusion in the next published issue of the Government Notices Gazette or Business Gazette as well as in the next published issue of the series of the Gazette in which the notice would normally have been published.

Tariff concessions issues contain notices of tariff concessions proposed, granted or revoked in accordance with the provisions of Part XVA of the *Customs Act 1901*. These issues are published each Wednesday and are sold at \$1.95 or on subscription only at \$115.00 for 50 issues including surface postage.

Periodic issues contain lengthy notices of a non-urgent nature, including the following: certificates of Australian citizenship; registered tax agents; authorised celebrants; unclaimed deposits and moneys; Australian Public Service conditions of entry and advancement; appointments to the Australian Public Service; holders of import licences and tariff quotas. Issues are made at irregular intervals as required, at individual prices according to size. Advice of availability is given in the Government Notices, Business and Public Service issues immediately following the day of publication. Periodic issues are not available on subscription, but standing orders are accepted for all selected issues. Purchasing and Disposals issues of the Gazette provide information on Commonwealth purchases and disposals and

other matters of general interest to persons buying from or

selling to the Commonwealth. These issues are published each Wednesday and sold at \$3.95 or on subscription of \$200.00 including postage for 50 issues.

Index issues contain references to entries in the Government Notices issues and entries in the Orders in Council, Notices under the Superannuation Act, Notices under the Public Service Act, and Determinations under the Public Service Act sections of the Public Service issues. Index issues are published quarterly, are available over the counter from Commonwealth Government Bookshops and are supplied without charge to annual subscribers to the Government Notices issues.

N.N.-8878068

ISSUE OF PERIODIC GAZETTES

The following Periodic issues of the Gazette have been published.

Copies may be purchased from Commonwealth government bookshops or by mail from the relevant address given on the front page of this Gazette.

Gazette number	Date of publication	Subject	
PI	18.1.88	Tariff Quotas—Transfer of Quota Allocations—1 January 1987 to 30 November 1987	
P2	5.2.88	Australian Customs Service—Import Licences	
P3	19.2.88	Tariff Quotas—Motor vehicles multiple period tender quota allocations— 1 December 1987 to 31 January 1989	
P4	22.2.88	Tariff Quotas—Motor vehicles multiple period tender quota allocations— 1 October 1987 to 31 March 1989	
P5	23.2.88	Tariff Quotas—Motor vehicles tender quota allocations—I October 1987 to 31 March 1989	
P6	26.2.88	Tariff Quotas—Textile, clothing, footwear quota allocations—1 January 1988 to 28 February	
P7	16.3.88	Wildlife Protection (Regulation of Exports and Imports) Act 1982	
P8	14.3.88	Tariff Quotas—1988 Base Quota Allocations—Listing of tariff quota holders	
P9	24.3.88	Customs (Import Licensing) Regulations Exception Notice No. M68	
P10	6.4.88	Tariff Quotas—Base Motor Vehicles Quota Allocations—Listing of tariff quota holders	
P11	27.4.88	Tariff Quotas—Quota transactions for the period 871201 to 880331	
P12	13.5.88	Tariff Quotas—Textiles, clothing and footwear allocations of 1988 residua tender by ballot	
P13	27.5.88	Great Barrier Reef Marine Park Regulations (1987)	
P14	24.6.88	Australian Capital Territory, Unclaimed Moneys Ordinance 1950, Petroz NL (formerly Offshore Oil NL)	
P15	24.6.88	Tariff Quotas—Cheese Quota Allocations—1 July 1988 to 30 June 1989— Listing of Tariff Quota Holders	
P16	24.6.88	Tariff Quotas—Developing Country Quota Allocations—1 July 1988 t 28 February 1989—List of Quota Holders	
P17	1.7.88	Wildlife Protection (Regulation of Exports and Imports) Act 1982	
P18	30.6.88	Life Insurance Act 1945—Return of Unclaimed Moneys as at 31 December 1987	
P19	15.7.88	Amendment No. 1 to the National Health and Medical Research Counci Food Standards Code	
P20	30.6.88	Statement of Unclaimed Money Deposits and Money for Year Ended 31.12.87	
P21	22.7.88	Import Licences—Issued under the Customs (Import Licensing) Regulations for used, second hand or disposals earthmoving, construction and materials handling machinery and equipment—January to June 1988	
P22	8.8.88	Great Barrier Reef Marine Park Act 1975—Permits granted, refused, suspended or revoked	
P23	18.8.88	Tariff Quotas—Quota Transactions for the period 1 April 1988 to 30 June 1988	
P24	12.9.88	Wildlife Protection (Regulation of Exports and Imports) Act 1982	
P25	31.8.88	National Memorials Ordinance 1928—Determination of Nomenclature	
P26	26.10.88	Tariff Quotas—Textiles, Clothing and Footwear Base quota allocations for the Period 1 December 1988 to 28 February 1989—List of quota Holders	
P27	27.10.88	Tariff Quotas—Quota Transactions for the Period 1 July 1988 to 30 September 1988	
P28	11.11.88	Great Barrier Reef Marine Park Act 1975 Particulars of Permits Granted Refused Suspended or Revoked for the Period 1 July 1988 to 30 September 1988	

Special Information

PROCLAMATION

Commonwealth of Australia N. M. STEPHEN Governor-General

By His Excellency the Governor-General of the Commonwealth of Australia

I, SIR NINIAN MARTIN STEPHEN, Governor-General of the Commonwealth of Australia, acting with the advice of the Federal Executive Council and under subsection 358A (3) of the Crimes Act 1900 of the State of New South Wales in the application of that Act to the Australian Capital Territory by virtue of subsection 6 (1) of the Seat of Government Acceptance Act 1909, hereby declare each of the following enactments to be a corresponding law:

Search Warrants Act 1985, of the State of New South Wales:

- (b) Part 11A of the Crimes Act 1958, of the State of Victoria.
- (c) Criminal Investigation (Extra-territorial Offences) Act 1987, of the State of Western Australia;
- (d) Criminal Investigation (Extraterritorial Offences) Act 1984, of the State of South Australia;
- (e) Criminal Investigation (Extra-Territorial Offences) Act 1987, of the State of Tasmania; and
- (f) Criminal Investigation (Extra-territorial Offences) Act 1985, of the Northern Territory.
- (L.S.) GIVEN under my hand and the Great Seal of Australia on 14 November 1988.

By His Excellency's Command,

LIONEL BOWEN

Attorney-General

GOD SAVE THE QUEEN!

N.N.-8878070

NOTICES UNDER THE INDEPENDENT AIR FARES COMMITTEE ACT 1981

COST ALLOCATION REVIEW

EAST-WEST AIRLINES (OPERATIONS) LIMITED

In accordance with the provisions of section 13 of the Independent Air Fares Committee Act 1981, the Minister for Transport and Communications has requested the Independent Air Fares Committee to conduct a Cost allocation review in relation to air services provided by East-West Airlines (Operations) Limited over all its routes. A previous such review for the airline was completed in July 1985, and copies of the Committee's report are available from the IAFC office.

The members of the Committee are Mr K. F. Wraith (Chairman), Dr D. W. Zink, A.M. and Mr A. Oliver, C.M.G.

For this cost allocation review, the Committee plans to hold public hearings during February and March 1989 in Brisbane, Cairns, Hobart and Sydney. Notices will be placed in the Gazette and relevant daily newspapers in December 1988 giving information about the dates and locations of the hearings.

Purpose of review

When conducting a cost allocation review, the Committee is required by the Act to determine the manner in which all costs incurred by the operator in providing regular public transport air services are to be recorded from the 'flag-fall' and 'distance' components of passenger fares. As part of a review, the Committee also takes public evidence concerning guidelines and principles to be applied in approving domestic air fares.

In the circumstances of this Cost allocation review, the Committee does not propose to completely re-examine all matters, but rather to concentrate on establishing what changes are appropriate to the previous cost allocations and fare setting guidelines so far as East-West is concerned.

Public Submissions

Interested persons, organisations and associations are invited to submit in writing any views they consider should be taken into account in the Cost allocation review for East-West Airlines. They should also indicate whether they wish to give oral evidence at the hearings.

An information paper, which includes details of matters the Committee wishes to consider during the review, can be obtained by telephoning the office of the IAFC (03) 629 7174 or making a request in writing to the Secretary at the address set out below.

The closing date for written submissions is Friday, 20 January 1989. Two copies of all submissions would be appreciated, and should be addressed to the Secretary, Independent Air Fares Committee, GPO Box 94A, Melbourne Vic. 3001. Facsimile (03) 629 1853. Telex 38145.

Operator and reference

Section of Act

Date notified

Determination

AIRLINES OF TASMANIA (C4/88)

25

16.11.88

In response to an application by the operator under section 25 of the IAFC Act, determination of the following one-way economy air fares effective from 16 November 1988:

Launceston-King Is

\$93

being the same as those approved by Transport Tasmania.

Decision

EAST-WEST AIRLINES (D199/88)

17 (4)

10.11.88

Approval to offer, during the period 29 April 1989 to 28 April 1990, fixed price holiday discount air fares at discounts ranging from nil to 32.5 per cent off the approved economy air fares at 1.11.88 depending on route and season, and available under specified conditions as proposed by the operator including that fares must be used in conjunction with authorised package holidays.

Approved on basis of estimates provided by the operator that the discount fares will generate additional traffic and improve profitability.

ANSETT AIRLINES OF AUSTRALIA (D200/88)

17 (4)

10.11.88

Approval to offer a 30 per cent discount return fare during periods of low activity from 19 November 1988 to 18 November 1989 inclusive, under specified conditions as proposed by the operator including seven days advance purchase and three nights minimum stay away requirements, and limited seat availability on specific flights over selected routes.

Approved on basis of estimates provided by the operator that the discount fare will generate additional traffic and improve profitability. The Committee intends to closely monitor operating results of this fare.

AUSTRALIAN AIRLINES (D202/88)

17(4)

10 11 88

Approval to offer an Australian Originating Add-on fare at 25 per cent discount (was 30 per cent) off the normal economy fare from 1 January 1989 to 31 October 1990 under specified conditions as proposed by the operator available to passengers ticketed on international promotional air fares for travel commencing and terminating in Australia.

Approved on basis of estimates provided by the operator that the discount fare will generate additional traffic and improve profitability.

AIR NSW (D201/88)

17 (4)

16.11.88

Approval to offer, during the period 29 April 1989 to 27 April 1990, fixed price holiday discount air fares at 21.0 per cent off the approved economy air fares as at 1 September 1988, under specified conditions as proposed by the operator. A surcharge level of 6.0 per cent on the economy fare as at 1.9.88 is applicable at specified peak periods, but the air fare component cannot be higher than the relevant economy fare in force at the time of ticketing. The fares must be used in conjunction with an authorised package holiday.

Approved on basis of estimates provided by the operator that the discount fares will generate additional traffic and improve profitability.

N.N.-8878071

NOTICE OF CESSATION OF A STATUTORY LIEN IN RESPECT OF CERTAIN AIRCRAFT

NOTICE is hereby given that pursuant to section 75 (1) of the Civil Aviation Act 1988, a statutory lien vested in the Authority ceased to have effect in respect of each of the aircraft described hereunder.

Lien No.	Aircraft description and registration mark	Date on which the Lien ceased to have effect
00474	Piper PA-30, VH-KNT	26 October 1988
00698	Beech 58, VH-PRH	3 November 1988

Dated this 17th day of November 1988.

K. L. CLAYTON Registrar of Statutory Liens

N.N.-8878072

Legislation

Acts of Parliament assented to

IT IS HEREBY NOTIFIED, for general information, that His Excellency the Governor-General, in the name of Her Majesty, assented, on 8 November 1988 to the undermentioned Acts passed by the Senate and the House of Representatives in Parliament assembled, viz.:

No. 86 of 1988—An Act relating to the prevention and settlement of certain industrial disputes, and for other purposes (Industrial Relations Act 1988).

No. 87 of 1988-An Act to enact certain transitional provisions, and to repeal certain Acts and amend certain Acts, in consequence of the enactment of the Industrial Relations Act 1988, and for other purposes (Industrial Relations (Consequential Provisions) Act 1988).

A. R. BROWNING

Clerk of the House of Representatives

N.N.-8878073

Regulations

AUSTRALIAN CAPITAL TERRITORY

NOTIFICATION OF THE MAKING OF A REGULATION

NOTICE is hereby given that the undermentioned Regulation of the Australian Capital Territory has been made. Copies of the Regulation may be purchased at the Commonwealth government bookshop, 70 Alinga St, Canberra City Australian Capital Territory.

Ordinance under which the Regulation was made	Description of Regulation	Number and year of Regulation
Gaming Machine Ordinance 1987	Gaming Machine Regulations (Amendment)	1988 No. 20

Orders

NOTIFICATION OF THE MAKING OF ORDERS UNDER THE EXPORT CONTROL (ORDERS) REGULATIONS

NOTICE is hereby given that the undermentioned orders under the Export Control (Orders) Regulations have been made. Copies of the orders can be purchased over the counter from the Commonwealth government bookshop at 70 Alinga St, Canberra City ACT 2600, or by mail from Mail Order Sales, GPO Box 84, Canberra 2601.

Number of Orders	Description of Orders		
No. 17 of 1988	Export Control (Fresh Fruits and Vegetables) Orders		

NOTIFICATION OF THE MAKING OF ORDERS UNDER THE CIVIL AVIATION REGULATIONS

NOTICE is hereby given that an amendment to Civil Aviation Orders Part 105, was made on 14 November 1988 and is effective from 18 November 1988.

AD/Cessna 303/7: NLG Actuator attach fitting.

Copies of the Order are available for inspection and may be purchased over the counter from the Civil Aviation Authority, 607 Swanston St, Carlton South Vic. 3053 or by mail from the Civil Aviation Authority, Publications Centre, GPO Box 1986, Carlton South Vic. 3053.

N.N.-8878076

By-laws

AUSTRALIAN TELECOMMUNICATIONS COMMISSION

NOTIFICATION OF THE MAKING OF BY-LAWS

NOTICE is hereby given that the Australian Telecommunications Commission has made the undermentioned By-laws on 9 November 1988.

N.N.-8878075

Copies can be obtained from the Corporate Secretary, Telecom Australia, 17th Floor, 199 William St, Melbourne Vic. 3000.

Act under which By-laws were made	Description of By-laws
Telecommunications Act 1975	Telecommunications (General) By-laws—Amendment No. 52

CA03T040 CR\$

N.N.--8878077

FEDERAL AIRPORTS CORPORATION

NOTIFICATION OF THE MAKING OF BY-LAWS

NOTICE is hereby given that the undermentioned By-laws have been made. Copies of the By-laws can be purchased from the Commonwealth Government Bookshop, 70 Alinga St, Canberra City ACT.

Act under which the By-laws were made	No. of By-laws (or form of individual for By-laws)	Description of By-laws
Federal Airports Corporation Act 1986	2.1	Part I—Preliminary Amended
1980	15D.1—15E.2	Part IVC Regulation of building and engineering works on federal airports
	15F.1—15F.5	Part IVD Regulations for the security of federal airports

CA03F082 CR\$

N.N.--8878078

CORRIGENDUM

GAZETTE NO. GN 43, 16 NOVEMBER 1988

FEDERAL AIRPORTS CORPORATION

Insert '1986' in column 1 after the word 'Act'.

Delete from column 2 '3.1' and delete the wording in column 3 opposite that entry.

Insert after the entry in column 2 '3.2-3.13' and the wording in column 3 opposite that entry, words and figures as follows:

in column 2 '8.3A

in column 3

48.4

Removal of abandoned vehicles

Additional reference to 8.3A

9.1

Additional reference to 8.3A'

Delete from column 3 the word and symbol 'Part IVA' (second occurring) and substitute the word and symbol 'Part IVR'.

Determinations

Commonwealth Teaching Service Act 1972

NOTICE OF THE MAKING OF DETERMINATION UNDER SECTIONS 20 AND 23

NOTICE is hereby given that the following determination has been made under sections 20 and 23 of the Commonwealth Teaching Service Act. Copies can be obtained from the Department of Industrial Relations, 1 Farrell Pl, Canberra City ACT (062) 43 7154.

Number and year of Determination	Description of Determination	Date made
No. 4 of 1988	Amendment to Determination 44—Canberra Institute of the Arts Second tier salary increases	19.10.88

N.N.-8878079

Government Departments

Administrative Services

AUSTRALIAN ELECTORAL COMMISSION

I HAVE ascertained and set out in the schedule for each State and Territory the number of electors enrolled in each Division as at the date indicated and for each State and the Australian Capital Territory have determined the average divisional enrolment and the extent to which the number of electors enrolled in each Division differs from the average divisional enrolment.

THE SCHEDULE

New South Wales as at 23 September 1988:

Average Divisional Enrolment 70 033

Division	Enrolment	average divisional enrolment	Wentworth Werriwa	66 71
————————	Enroiment	enroimeni		
Banks	67 953	-2.97	State Total	3 57
Barton	71 346	1.87		
Bennelong	70 312	0.40	Victoria as at 23 Se	ptember 1988:
Berowra	70 622	0.84	Average Divisional	-
Blaxland	68 141	-2.70	Average Divisional	Linoinicit 70
Bradfield	68 468	-2.23		
Calare	71 307	1.82		
Charlton	69 665	-0.53		
Chifley	70 007	-0.04	Division	Enrol
Cook	69 679	-0.51		
Cowper	75 116	7.26	Aston	72
Cunningham	66 496	-5.05	Ballarat	73
Dobell	74 071	5.77	Batman	64
Dundas	68 293	-2.48	Bendigo	73
Eden-Monaro	70 647	0.88	Bruce	65
Farrer	71 759	2.46	Burke	77
Fowler	69 569	-0.66	Calwell	72
Gilmore	68 524	-2.15	Casey	72
Grayndler	69 508	-0.75	Chisolm	69
Greenway	66 444	-5.12	Corangamite	74
Gwydir	69 195	-1.20	Corio	69
Hughes	70 835	1.15	Deakin	68
Hume	69 429	-0.86	Dunkley	69
Hunter	70 787	1.08	Flinders	73
Kingsford-Smith	70 825	1.13	Gellibrand	68
Lindsay	70 927	1.28	Gippsland	73
Lowe	69 519	-0.73	Goldstein	69
Lyne	75 141	7.29	Henty	67
Macarthur	67 979	-2.93	Higgins	66
Mackellar	70 259	0.32	Holt	6
Macquarie	67 758	-3.25	Hotham	70
Mitchell	71 096	1.52	Indi	74

% Deviation from

Division	Enrolment	% Deviation from average divisional enrolment
Newcastle	67 147	-4.12
New England	71 748	2.45
North Sydney	67 677	-3.36
Page	70 636	0.86
Parkes	67 722	-3.30
Parramatta	69 000	-1.48
Phillip	70 675	0.92
Prospect	75 492	7.79
Reid	70 799	1.09
Richmond	68 297	-2.48
Riverina-Darling	65 385	-6.64
Robertson	73 915	5.54
St George	70 348	0.45
Shortland	71 877	2.63
Sydney	69 545	-0.70
Throsby	66 596	-4.91
Warringah	69 298	-1.05
Wentworth	66 501	-5.04
Werriwa	77 358	10.46
State Total	3 571 693	

Divisio n	Enrolment	% Deviation from average divisional enrolment
Aston	72 592	3.31
Ballarat	73 148	4.10
Batman	64 862	-7.69
Bendigo	73 303	4.32
Bruce	65 937	-6.16
Burke	77 691	10.57
Calwell	72 149	2.68
Casey	72 919	3.78
Chisolm	69 460	-1.15
Corangamite	74 940	6.65
Corio	69 076	1.69
Deakin	68 473	-2.55
Dunkley	69 119	1.63
Flinders	77 314	10.03
Gellibrand	68 923	1.91
Gippsland	73 535	4.65
Goldstein	69 511	-1.07
Henty	67 161	-4.42
Higgins	66 844	-4.87
Holt	67 778	-3.54
Hotham	70 657	0.56
ndi	74 904	6.60

		~ D : .:			Of Davistian from
Division	Enrolment	% Deviation from average divisional enrolment	Division	Enrolment	% Deviation fron average divisiona enrolmen
Isaacs	68 354	-2.72	Perth	71 148	-1.54
Jagajaga	64 346	-8.43	Stirling	68 831	-4.7
			Swan	69 091	-4.39
Kooyong	68 193	-2.95		73 902	
Lalor	78 330	11.48	Tangney	13 902	2.2
La Trobe	71 607	1.91		222.424	
McEwen	73 433	4.51	State Total	939 431	
McMillan	69 052	-1.73			
Mallee	71 597	1.89	South Australia as	at 29 September 198	8:
Maribyrnong	66 793	-4.94	Average Divisional	Enrolment 72 857	
Melbourne	69 110	-1.65	Average Divisional	Emonitent 72 037	
Melbourne Ports	67 103	-4.50			01 D
Menzies	69 424	-1.20			% Deviation fron
Murray	71 138	1.24	_		average divisiona
Scullin	63 381	-9.80	Division	Enrolment	enrolmen
Streeton	69 970	-0.42			
Wannon	70 401		Adelaide	74 245	1.92
		0.19	Barker	72 101	-1.04
Wills	67 829	-3.47	Bonython	72 893	0.0
			Boothby	73 795	1.29
State Total	2 740 357			70 062	
			Grey		-3.84
Queensland as at 26 Se	entember 1988		Hawker	71 694	-1.60
	-		Hindmarsh	72 895	0.03
Average Divisional En	rolment /1 122		Kingston	71 877	-1.3
			Makin	71 872	-1.35
		% Deviation from	Mayo	75 455	3.57
		average divisional	Port Adelaide	73 232	0.51
Division	Enrolment	enrolment	Sturt	72 163	0.95
			Wakefield	74 860	2.75
Bowman Brisbane	73 707 68 472	3.63 -3.73	State Total	947 144	
Capricornia	68 950	-3.05	State of Tasmania	as at 23 September I	088.
Dawson	71 275	0.22		•	700.
Fadden	66 564	-6.41	Average Divisional	Enrolment 60 8/5	
Fairfax	74.015	4.07			
Fisher	77 765	9.34			% Deviation from
Forde	66 403	-6.64			average divisiona
Griffith	68 490	-3.70	Division	Enrolment	enrolmen
Groom	72 408	1.81	Division	2 nr onne n	
Herbert	73 468	3.30	D.	/A ##A	
Hinkler	69 721	-1.97	Bass	60 778	-0.16
Kennedy	70 127	-1.40	Braddon	61 371	0.8
Leichhardt	70 800	-0.45	Denison	60 349	-0.80
			Franklin	60 689	-0.3
Lilley	68 836	-3.21	Lyons	61 189	0.53
McPherson	82 301	15.72			
Maranoa	69 498	-2.28	State Total	304 376	
Moncrieff	76 054	6.93	State I otal	304 370	
Moreton	67 113	-5.64	Australian Camital	Territory as at 23 Se	ntember 1088
Oxley	67 056	-5.72	•	•	premoci 1300.
Petrie	72 522	1.97	Average Divisional	Enrolment 83 481	
Rankin	69 052	-2.91	-		
Ryan	74 130	4.23			% Deviation from
Wide Bay	68 206	-4.10			average divisiona
			Division	Enrolment	enrolmen
State Total	1 706 933		Conhess	81 748	2.01
State of Western Austr	ralia as at 27 Sen	tember 1988	Canberra Fraser	81 /48 85 214	-2.00 2.00
	ao at a' oop	1700.	FIASET	a3 214	/ ()

State of Western Australia as at 27 September 1988: Average Divisional Enrolment 72 264

Division Enro		Deviation from verage divisional enrolment
Brand	74 733	3.42
Canning	59 760	-3.47
Cowan	76 042	5.23
Curtin	71 339	-1.28
Forrest	72 538	0.38
Fremantle	73 368	1.53
Kalgoorlie	72 311	0.07
Moore	75 343	4.26
O'Connor	71 025	-1.71

ivisional irolment -2.0885 214 2.08 Fraser Territory Total 166 962 Northern Territory as at 27 September 1988: Average Divisional Enrolment 75 231 Total for Australia 10 452 127

> COLIN A. HUGHES Electoral Commissioner

The Arts, Sport, the Environment, **Tourism and Territories**

AUSTRALIAN CAPITAL TERRITORY

Trading Hours Ordinance 1962

APPLICATION OF ORDINANCE

Under section 6 (1) of the Trading Hours Ordinance 1962 'the Ordinance', I, hereby declare that the Ordinance shall not apply in the part of the Australian Capital Territory being the area occupied by the store known as Grace Bros on levels 1, 2 and 3 of the Westfield Shopping Centre located on Block 1, Section 52, Division of Belconnen for the period of time starting at 6.00 p.m. and finishing at 8.00 p.m. on Wednesday, 30 November 1988.

Dated 16 November 1988.

WILLIAM JOHN HARRIS

ACT Administration Delegate of the Minister of State for the Arts and Territories

N.N.-8878080

AUSTRALIAN CAPITAL TERRITORY

Canberra Theatre Trust Ordinance 1965

APPOINTMENT

Under section 6 (1) of the Canberra Theatre Trust Ordinance 1965 I appoint:

Anthony Lewis Hayward Joan Margaret Taggart

to be Trustees of the Canberra Theatre Trust for the period of three years commencing on the date of this instrument. Dated 10 November 1988.

WILLIAM JOHN HARRIS

ACT Administration Delegate of the Minister of State for the Arts and Territories

N.N.--8878081

AUSTRALIAN CAPITAL TERRITORY

City Area Leases Ordinance 1936

INSTRUMENT OF APPROVAL UNDER SECTION 10

In pursuance of section 10 of the City Area Leases Ordinance 1936, I, PETER ROBERT GRIFFITHS being the officer for the time being occupying an office to the occupant of which the Minister has by instrument in writing under section 12C of the Seat of Government (Administration) Act 1910 delegated his powers under section 10 of the said Ordinance hereby approve of Peter Neil Gowland and Valerie Mary Gowland ('the Applicant') carrying on the profession trade occupation or calling of venison and game distributors and ecological consultant ('the business') on Block 27, Section 25, Division of Hawker known as 9 Mataranka St, Hawker ('the land') subject to the following conditions relating to the use of the land being observed by the Applicant in carrying on the business:

- (1) that this approval will remain valid only while the Applicant continues to be a bona fide resident of the land;
- (2) that the Applicant will ensure that the conduct of the business does not cause an annoyance, a nuisance or danger and is not offensive to any tenants or occupiers of adjoining lands;
- (3) that the Applicant will not erect or permit or suffer to be displayed or erected upon the land or any building thereon any advertising sign or hoarding whatever without the consent in writing of the

- National Capital Development Commission and the Building Controller;
- (4) that the Applicant will conduct the business strictly in accordance with the application made by the Applicant under section 10 of the City Area Leases Ordinance unless otherwise stipulated in this Instrument;
- (5) that no person other than the Applicant will conduct or in any way carry on the business on the land without the prior approval of the Minister;
- (6) that the Applicant will ensure that all residence and business related vehicles are parked within the confines of the land;
- (7) that the Applicant will not employ any assistants for the purpose of conducting or carrying on the business on the land without the prior approval of the Minister;
- (8) that deliveries to the land by commercial vehicles be restricted to two per week;
- (9) that this approval relates to the provision of office accommodation and garaging of a refrigerated vehicle;
- (10) that this approval will terminate on the thirty-first day of October 1989 or on such earlier date as the Minister determines in accordance with condition 11;
- (11) upon any failure to comply with any or all of the foregoing conditions the Minister or his Delegate under the said Ordinance may give written notice requiring the Applicant to show cause within a period of fourteen days why this approval should not be revoked. At the expiration of this period the Minister or his Delegate may revoke the approval.

Dated this 9th day of November 1988.

P. R. GRIFFITHS

Delegate of the Minister of State for the Arts and Territories

N.N.-8878082

AUSTRALIAN CAPITAL TERRITORY

City Area Leases Ordinance 1936

INSTRUMENT OF APPROVAL UNDER SECTION 10

In pursuance of section 10 of the City Area Leases Ordinance 1936, I, PETER ROBERT GRIFFITHS being the officer for the time being occupying an office to the occupant of which the Minister has by instrument in writing under section 12C of the Seat of Government (Administration) Act 1910 delegated his powers under section 10 of the said Ordinance hereby approve of Ronald Edward Cochrane ('the Applicant') carrying on the profession trade occupation or calling of medical practitioner ('the business') on Block 8, Section 1, Division of Melba known as 103 Kingsford Smith Drv, Melba ('the land') subject to the following conditions relating to the use of the land being observed by the Applicant in carrying on the business:

- (1) that this approval will remain valid only while the Applicant continues to be a bona fide resident of the land:
- (2) that the Applicant will ensure that the conduct of the business does not cause an annoyance, a nuisance or danger and is not offensive to any tenants or occupiers of adjoining lands;
- (3) that the Applicant will not erect or permit or suffer to be displayed or erected upon the land or any building thereon any advertising sign or hoarding whatever without the consent in writing of the National Capital Development Commission and the Building Controller;
- (4) that the Applicant will conduct the business strictly in accordance with the application made by the Applicant under section 10 of the City Area Leases Ordinance unless otherwise stipulated in this Instrument;

- (5) that no person other than the Applicant will conduct or in any way carry on the business on the land without the prior approval of the Minister;
- (6) that the Applicant will ensure that all residence and business related vehicles are parked within the confines of the land:
- (7) that the Applicant will not without the prior approval of the Minister employ any more than two assistants for the purpose of conducting or carrying on the business on the land and those assistants shall be employed only in the capacity of a nurse and receptionist;
- (8) that the business will only be conducted on the land between the hours of 9.00 a.m. and 6.00 p.m. Monday to Friday and 9.00 a.m. and 12.00 noon Saturdays except in the case of an emergency;
- (9) that this approval will terminate on the thirty-first day of October 1989 or on such earlier date as the Minister determines in accordance with condition 10;
- (10) upon any failure to comply with any or all of the foregoing conditions the Minister or his Delegate under the said Ordinance may give written notice requiring the Applicant to show cause within a period of fourteen days why this approval should not be revoked. At the expiration of this period the Minister or his Delegate may revoke the approval.

Dated this 9th day of November 1988.

P. R. GRIFFITHS

Delegate of the Minister of State
for the Arts and Territories

N.N.-8878083

AUSTRALIAN CAPITAL TERRITORY

City Area Leases Ordinance 1936

INSTRUMENT OF APPROVAL UNDER SECTION 10

In pursuance of section 10 of the City Area Leases Ordinance 1936, I, PETER ROBERT GRIFFITHS being the officer for the time being occupying an office to the occupant of which the Minister has by instrument in writing under section 12C of the Seat of Government (Administration) Act 1910 delegated his powers under section 10 of the said Ordinance hereby approve of David Lindsay Barnes ('the Applicant') carrying on the profession trade occupation or calling of plumber and drainer ('the business') on Block 2, Section 17, Division of Scullin known as 27 Levien St, Scullin ('the land') subject to the following conditions relating to the use of the land being observed by the Applicant in carrying on the business:

- that this approval will remain valid only while the Applicant continues to be a bona fide resident of the land;
- (2) that the Applicant will ensure that the conduct of the business does not cause an annoyance, a nuisance or danger and is not offensive to any tenants or occupiers of adioining lands:
- (3) that the Applicant will not erect or permit or suffer to be displayed or erected upon the land or any building thereon any advertising sign or hoarding whatever without the consent in writing of the National Capital Development Commission and the Building Controller;
- (4) that the Applicant will conduct the business strictly in accordance with the application made by the Applicant under section 10 of the City Area Leases Ordinance unless otherwise stipulated in this Instrument;
- (5) that no person other than the Applicant will conduct or in any way carry on the business on the land without the prior approval of the Minister;

- (6) that the Applicant will ensure that all residence and business related vehicles are parked within the confines of the land;
- (7) that the Applicant will not employ any assistants for the purpose of conducting or carrying on the business on the land without the prior approval of the Minister;
- (8) that only one room in the house be used for the conduct of the business;
- (9) that the business will only be conducted on the land seven days per week;
- (10) that all equipment and materials shall be stored in the carport only;
- (11) that this approval will terminate on the thirtieth day of September 1989 or on such earlier date as the Minister determines in accordance with condition 12;
- (12) upon any failure to comply with any or all of the foregoing conditions the Minister or his Delegate under the said Ordinance may give written notice requiring the Applicant to show cause within a period of fourteen days why this approval should not be revoked. At the expiration of this period the Minister or his Delegate may revoke the approval.

Dated this 9th day of November 1988.

P. R. GRIFFITHS

Delegate of the Minister of State for the Arts and Territories

N.N.-8878084

AUSTRALIAN CAPITAL TERRITORY

City Area Leases Ordinance 1936

INSTRUMENT OF APPROVAL UNDER SECTION 10

In pursuance of section 10 of the City Area Leases Ordinance 1936, I, PETER ROBERT GRIFFITHS being the officer for the time being occupying an office to the occupant of which the Minister has by instrument in writing under section 12C of the Seat of Government (Administration) Act 1910 delegated his powers under section 10 of the said Ordinance hereby approve of Ricky Allen Flavell and Laurelle June Flavell ('the Applicant') carrying on the profession trade occupation or calling of freelance graphic design and publishing ('the business') on Block 1, Section 282, Division of Kambah known as 12 Kathleens Park, 17 Pinkerton Cct, Kambah ('the land') subject to the following conditions relating to the use of the land being observed by the Applicant in carrying on the business:

- (1) that this approval will remain valid only while the Applicant continues to be a bona fide resident of the land:
- (2) that the Applicant will ensure that the conduct of the business does not cause an annoyance, a nuisance or danger and is not offensive to any tenants or occupiers of adjoining lands;
- (3) that the Applicant will not erect or permit or suffer to be displayed or erected upon the land or any building thereon any advertising sign or hoarding whatever without the consent in writing of the National Capital Development Commission and the Building Controller;
- (4) that the Applicant will conduct the business strictly in accordance with the application made by the Applicant under section 10 of the City Area Leases Ordinance unless otherwise stipulated in this Instrument:
- (5) that no person other than the Applicant will conduct or in in any way carry on the business on the land without the prior approval of the Minister;
- (6) that the Applicant will ensure that all residence and business related vehicles are parked within the confines of the land;

- (7) that the applicant may employ the equivalent of one full-time assistant in the conduct of the business on the land between 8.30 a.m. and 5.30 p.m., with no more than one person assisting in the business on the land at any one time:
- (8) that Clause 7 does not prevent the Applicant accepting up to two school students per year to be engaged in the business to gain work experience for a maximum of two weeks per year per student;
- (9) that the hours of operation for the business do not extend beyond 11.00 p.m.
- (10) that the Applicant will conduct the business strictly by appointment, and that such appointments will be organised to ensure that not more than two clients are in attendance at any one time;

- (11) that this approval will terminate on the thirty-first day of August 1989 or on such earlier date as the Minister determines in accordance with condition 12:
- (12) upon any failure to comply with any or all of the foregoing conditions the Minister or his Delegate under the said Ordinance may given written notice requiring the Applicant to show cause within a period of fourteen days why this approval should not be revoked. At the expiration of this period the Minister or his Delegate may revoke the approval.

Dated this 9th day of November 1988.

P. R. GRIFFITHS Delegate of the Minister of State for the Arts and Territories

N.N.--8878085

COMMONWEALTH OF AUSTRALIA

Wildlife Protection (Regulation of Exports and Imports) Act 1982

Section 11

DECLARATION OF APPROVED INSTITUTIONS

1, JOHN DERRICK OVINGTON, the Designated Authority under subsection 18 (1) of the Wildlife Protection (Regulation of Exports and Imports) Act 1982, in pursuance of subsection 11 (1) of the Act, hereby declare each of the organisations specified in Column 2 of the Schedule, in an item in the Schedule, to be an approved institution in relation to the class, or classes, of specimens specified in Column 3 of the Schedule in that item.

Dated this eighteenth day of November 1988.

J. D. OVINGTON Designated authority

SCHEDULE

Column I Item	Column 2 Name and country of approved institution	Column 3 Approved class, or classes, of specimens
1	Lionel E. Campion Little Moe River Road Yarragon Vic. 3823	Cervus elaphus
2	Zoo Berlin Hardenbergplatz 8 D-1000 Berlin 30 West Germany	Dacelo spp
3	Geoffrey A. Hemmings 44 Spurwood Rd Warrimoo NSW 2774	Agapornis spp
4	Keith Voss 'Olveston', 22 Hastings Rd Hawera, South Taranaki New Zealand	Agapornis spp
5	San Diego Zoo 2920 Zoo Drv San Diego, CA 92103, USA	Saguinus imperator

N.N.-8878086

COMMONWEALTH OF AUSTRALIA

Wildlife Protection (Regulation of Exports and Imports) Act 1982

Section 12

DECLARATION OF APPROVED ZOOLOGICAL ORGANISATIONS

I, JOHN DERRICK OVINGTON, the Designated Authority under subsection 18 (1) of the Wildlife Protection (Regulation of Exports and Imports) Act 1982, in pursuance of subsection 12 (1) of that Act, hereby declare each of the zoological organisations specified in column 2 of the Schedule, in an item in the Schedule, to be an approved zoological organisation in relation to the class, or classes, of specimens specified in Column 3 of the Schedule in that item.

Dated this eighteenth day of November 1988.

J. D. OVINGTON Designated Authority

SCHEDULE

Column 1	Column 2	Column 3
Item	Name and country of 200	Approved class, or classes, of specimens
1	Zoo Berlin Hardenbergplatz 8 D-1000 Berlin 30 West Germany	Dacelo spp
2	San Diego Zoo 2920 Zoo Drive San Diego, CA 92103, USA	Saguinus imperator

COMMONWEALTH OF AUSTRALIA

Wildlife Protection (Regulation of Exports and Imports)
Act 1982

Section 44

NOTICE

I, GRAHAM FREDERICK RICHARDSON, the Minister of State for the Arts, Sport, the Environment, Tourism and Territories in pursuance of subsection 44 (1) of the Wildlife Protection (Regulation of Exports and Imports) Act 1982, hereby notify that 1 am considering giving the following person authority under section 44 of the Act to export over a period of six months, the specimens specified below, on condition that prior to export of each consignment, the exporter obtains the permission of the Director of National Parks and Wildlife Service or his nominee.

The person and specimens are as follows:

Person

Specimens

H. N. Mighell PO Box 185 Caustis blakei—cut stems collected from 5ha of private property and 20ha of State Forest under licence.

Tin Can Bay Old 4580

In accordance with paragraph 44 (1) (f) of that Act, I invite interested persons to lodge with me comments in writing on the desirability of giving this authority. Such comments should be lodged at the following address not later than ten days after the date of publication of this Notice:

The Director Australian National Parks and Wildlife Service GPO Box 636

Canberra ACT 2601

Attention: Wildlife Protection Section Dated this 10th day of November 1988.

GRAHAM RICHARDSON

Minister of State for the Arts, Sport, the Environment, Tourism and Territories

N.N.-8878088

.COMMONWEALTH OF AUSTRALIA

Wildlife Protection (Regulation of Exports and Imports) Act 1982

Section 44

NOTICE

I, GRAHAM FREDERICK RICHARDSON, the Minister of State for the Arts, Sport, the Environment, Tourism and Territories in pursuance of subsection 44 (1) of the Wildlife Protection (Regulation of Exports and Imports) Act 1982, hereby notify that I am considering giving Andrew Edinger, 14 Lake St, Esperance WA 6450, authority under section 44 of the Act to export over a period of six months, consignments of shells of native Australian molluses on condition that prior to export of each consignment, the exporter obtains the permission of the Director of Australian National Parks and Wildlife Service or his nominee.

In accordance with paragraph 44 (1) (f) of that Act, I invite interested persons to lodge with me comments in writing on the desirability of giving this authority. Such comments should be lodged at the following address not later than ten days after the date of publication of this Notice:

The Director Australian National Parks and Wildlife Service GPO Box 636 Canberra ACT 2601

Attention: Wildlife Protection Section

Dated this 9th day of November 1988.

GRAHAM RICHARDSON Minister of State for the Arts, Sport, the Environment, Tourism and Territories

N.N.--8878089

COMMONWEALTH OF AUSTRALIA

Wildlife Protection (Regulation of Exports and Imports) Act 1982

Section 44

NOTICE

I, SENATOR GRAHAM RICHARDSON, the Minister of State for the Arts, Sport, the Environment, Tourism and Territories, in pursuance of subsection 44 (1) of the Wild-life Protection (Regulation of Exports and Imports) Act 1982, hereby notify that I am considering renewing the authority issued to B. Gunderson of Ravensbourne, Qld, under section 44 of the Act to export over a period of six months fronds of Macrozamia miquelii harvested in Queensland from areas of State Forests 451 and 915 on condition that prior to export of each consignment, the exporter obtains the permission of the Director of National Parks and Wildlife Service or his nominee.

In accordance with paragraph 44 (1) (f) of that Act I invite interested persons to lodge with me comments in writing on the desirability of giving this authority. Such comments should be lodged at the following address not later than ten days after the date of publication of this notice:

The Director Australian National Parks and Wildlife Service

GPO Box 636 Canberra ACT 2601

Attention: Wildlife Protection Section Dated this 9th day of November 1988.

GRAHAM RICHARDSON

Minister of State for the Arts, Sport, the Environment, Tourism and Territories

N.N.-8878090

AUSTRALIAN CAPITAL TERRITORY

Electricity and Water Ordinance 1988

DETERMINATION

The Australian Capital Territory Electricity and Water Authority (ACTEW) under section 48 of the Electricity and Water Ordinance 1988, makes the following determination fixing the charge that is to be payable to ACTEW for the bulk supply of water to Queanbeyan City Council for the financial year 1988-1989.

1. Charge

For the sale of water to Queanbeyan City Council—16.6 cents per kilolitre.

N.N.-8878091

PHYSIOTHERAPISTS BOARD OF THE AUSTRALIAN CAPITAL TERRITORY ELECTION OF DEPUTY CHAIRMAN OF THE

BOARD Pursuant to paragraph 6 (2) (b) of the Health Professions Roards (Procedures) Ordinance 1981, it is hereby notified

Pursuant to paragraph 6 (2) (b) of the Health Professions Boards (Procedures) Ordinance 1981, it is hereby notified that Patricia Levick was on 15 November 1988, unanimously elected by the Physiotherapists Board to be Deputy Chairman of the Board for twelve months from and including 21 November 1988.

Dated the sixteenth day of November 1988.

N. BAXTER Chairman of the Board

Attorney-General

PUBLICATIONS CLASSIFIED UNDER THE ACT CLASSIFICATION OF PUBLICATIONS ORDINANCE 1983

FOR WEEK ENDING 4.11.88

Publications classified under the A.C.T. Ordinance would attract the following classifications under the N.S.W. Indecent Articles and Classified Publications Act 1975 and the N.T. Classification of Publications Act 1979: U/R—Unrestricted; Category I - Restricted; Category 2-Direct Sale; Refused-Includes Child Pornography (CP) may not be sold.

Title	Edition	Author/Publisher	Decision flagging
Busen	No. 29	Pleasure Verlagsgesellschaft GmbH, Germany	Category 2
Busen	No. 29	Pleasure Verlags, W. Germany	Category 2
Club International	Vol. 17 No. 11 (C) 1988	Paul Raymond Publications, UK	Category 1
Club Pour Hommes	No. 20, May 1988	Paul Raymond Publications Ltd, UK	Refused
Easyriders (Australian)	No. 186, Vol. 18, December 1988	Paisano Publications, USA	Unrestricted
Escort	Vol. 8, No. 11, October 1988	Paul Raymond Publications, UK	Category 1
Fling	Vol. 30 No. 4, Issue 151, November 1988	Relim Publishing, USA	Category 2
Gallery Fantasies	Vol. 3, No. 4, Winter 1988	Montcalm Publishing Co., USA	Category 1
Gentlemans Companion	Vol. 8, No. 10, December 1988	Gentlemas Companion Magazine, USA	Category 2
Hot Dreams (The best of human digest)	Vol. 4, No. 6, December 1988	Thomaston Publications, USA	Category 2
Inside Foxy Lady	Vol. 7 No. 32	Verlag, Teresa Orlowski, W. Germany	Category 2
Knave	Vol. 20, No. 11 (C) 1988	Galaxy Publications, UK	Category 1
Live (Best of)	Vol. 8 No. 8 Winter 1988 collectors edition No. 5	Live Periodicals Inc., USA	Category 2
Lolitas Anal	No. 3	Silwa Film, W. Germany	Category 2
Mayfair	Vol. 23, No. 11	Fisk Publishing Co., UK	Unrestricted
Penthouse (French Edition)	No. 46, November 1988	Filipacchi Editee Par editions, France	Category 2
Penthouse (The Girls of)	November/December 1988	Penthouse International, USA	Category 1
Penthouse Forum	Vol. 18, No. 4, December 1988	Forum International, USA	Category 1
Penthouse Variations	Vol. 10, No. 12, November 1988	Viva International, USA	Category 1
Playboy (US)	Vol. 35, No. 12, December 1988	Playboy, USA	Unrestricted
Playboy's Book of Lingerie	November/December 1988	Playboy Press, USA	Unrestricted
Playgirl	Vol. XVI, No. 7 (Vol. 16, No. 7) December 1988	Playgirl Inc., USA	Unrestricted
Seventeen Special	No. 17	Coer'est Holland	Category 2
Seventeen Special	No. 17 (C) 1988	Coer'est Holland	Category 2
Sex Bizarre	No. 46, October 1988	Color-Climax Corp., Denmark	Category 2
Super Neger	No. 6, March 1988	COQ International, Denmark	Category 2
Topsy Special	No. 6, September 1988	Silva Film, West Germany	Category 2

- (1) Brought to the attention of the Victorian Government.
- (2) Brought to the attention of the SA Government.
- (3) Brought to the attention of the WA Government.
- (CP) Child Pornography.
- (*) Decision subject of an appeal.

Purpose

Justified

Gratuitous

FILM CENSORSHIP BOARD

OCTOBER 1988

Films examined in terms of the Customs (Cinematograph Films) Regulations and State's film censorship legislation are listed below.

Low

Frequent

Explicitness/Intensity

Medium

High

An explanatory key to reasons for classifying non-'G' films appears hereunder: Frequency

Infrequent

		Ingrequent		jueni	LU	v Median	i ilign	Justij	ieu Oraiuitous
S (Sex)		i i i	f f f		1 1 1	m m m	h h h	j j j	g g g
*** Films Board of Revie	ew Decis	ion							
Title	Produc	er		Country		Submitted length (mins)	Applicant		Reason for decision
For general exhibition—	'G' Suita	ble for all a	ges						
Blowpipes and Bulldozers		dell/P. Tait		Australia		60	Gaia Films		
Matter of Heart Story of Jiro, The	Kenl	iitney Gakushu kyusha Kiner yo Productioi	na j	USA Japan		102 106	Ronin Film Japan Infor and Cult		
Parental guidance-'PG'	' Parenta	l guidance re	ecomn	ended for	chi	ldren under 15	years		
Campus Man	P. Fow	rler/J. Landa	u 1	USA		93	Vestron Pic	tures	L (i-l-g) V (i-l-g) O (adult concepts)
Contradictions	D. Kna	aus	4	Australia		57	Film Austra	ılia	O (adult concepts)
Crossing Delancey	M. No	zik	Ţ	USA		96	Village Roa Corporat		O (sexual allusions adult concepts)
Deciduous Tree (main title not shown in English)		Kobun pany/Kindas Kyokai		Japan		102	Japan Infor	mation ure Centre	O (nudity adult concepts)
La Leggenda del Santo Bevitore	V. Ďe Cicu	Leo/R. tto/M. Ceccl /V. Cecchi		Italy		130	Ente Autor Gestione Cinema		O (adult concepts)
Mother vs Mother	Lau Ti		1	Hong Kong	3	96	Vacole		L (i-l-g) O (adult concepts)
Ngati	J. O'Sh	nea	1	New Zeala	nd	91	Australian Institute		L (f-l-j)
Qualcuno in Ascolto (High Frequency)	G. Pico Leop	cioli/G. pardi]	Italy		105	Ente Autor Gestione Cinema	omo di	V (i-l-j) O (adult concepts)
Things Change	M. Ha	usman	Ţ	USA		100	Fox Colum Distribute		L (i-l-j) O (adult concepts)
Without a Clue Young Einstein (edited version)		divant lous/W. Ross loach		UK Australia		106 91	Filmpac Ho Village Roa Corporat	dshow	V (i-m-j) O (adult concepts)
For mature audiences	M' Reco	mmended as	suital	ole for pers	ons	15 years and	over		
Consuming Passions	W. Cai			UK .		98	Vestron Pic	tures	O (sexual allusions adult concepts)
Dark Night	Lo Wa	i	1	Hong Kong	3	97	Golden Ree	el Films	S (f-m-j) O (adult theme)
Deceivers, The Distant Thunder	I. Merc R. Sch			UK/India Canada/US	SA	102 113	Vestron Pic United Inte		V (f-m-g) L (f-m-g)
Distant Voices, Still Lives	J. How	arth	Ī	U K		81	Pictures Dendy Cine	ema	V (i-m-g) O (adult concepts) L (i-m-g)

Title	Producer	Country	Submitted length (mins)	Applicant	Reason for decision
Evil Angels	V. Lambert	Australia	121	Village Roadshow Corporation	L (i-m-j) O (adult
Fatal Love	R. Wong	Hong Kong	82	Chinatown Cinema	concepts) V (i-m-g) S (i-m-g)
Film d'Amore e d'Anarchia	R. Cardarelli	Italy	123	Ente Autonomo di Gestione per il Cinema	O (nudity, sexual allusions)
Heart to Hearts	S. Shin	Hong Kong	94	Chinatown Cinema	O (adult concepts)
How to Pick Girls Up!?	Lau Tin Chi	Hong Kong	98	Vacole	V (i-m-g) O (adult concepts)
Imagine—John Lennon	D. Wolper/A. Solt	USA	105	Village Roadshow Corporation	L (i-m-j) O (adult concepts
lo e Mia Sorella (Me and My Sister)	M. Cecchi Gori/ V. Cecchi Gori	Italy	111	Ente Autonomo Di Gestione Per il Cinema	S (i-m-j) O (adult concepts
L'Ami De Mon Amie	M. Menegoz	France	100	Newvision Film Distributors	L (i-m-j)
Mignon E' Partita (Mignon has Left)	L. Pescarolo/G. De Laurentiis	Italy/France	99	Ente Autonomo Di Gestione Per Il Cinema	O (adult concepts)
Mimi Metallurgico	Euro International	Italy	115	Ente Autonomo Di Gestione Per Il Cinema	O (adult concepts V (i-m-j)
P.I. Private Investigations	S. Colin/ S. Sighvatsson	USA	91	Vestron Pictures	V (i-m-g) L (i-m-g)
Pasqualino Settebellezze (A.K.A. Seven Beauties)	Medusa Distribuzione	Italy	120	Ente Autonomo Di Gestione Per II Cinema	O (adult concepts V (i-m-j) S (i-m-j)
Prayer for the Dying,	P. Snell	UK/USA	107	Vestron Pictures	L (i-m-j) V (i-m-j)
Spellbinder	J. Wizan/B. Russell	USA	96	United International Pictures	L (i-m-g) O (occul-
Sposi (Brides and Bridegrooms)	A. Avanti/ C. Bonivento	Italy	97	Ente Autonomo Di Gestione Per II Cinema	L (i-m-g) O (adult
U2—Rattle and Hum	M. Hamlyn	USA	99	United International Pictures	L (i-m-g)
Un Complicato Intrigo Di Donne Vicoli e Delitti	F. Lucisano	Italy	107	Ente Autonomo Di Gestione Per II Cinema	V (i-m-j) L (f-m-j) O (adult concepts
Vroom	P. Lister	UK	88	Hoyts Distribution	L (i-m-j) O (adult concepts)
For Restricted Exhibition	on-'R' Persons 2 years	of age and over	, and under 18	years of age, are not ad	•
Bet on Fire (main title not shown in English)	Not shown	Hong Kong	95	Chinatown Cinema	V (i-m-g)
Blue Jean Cop (a)	J. Harman	USA	95	Hoyts Distribution	V (i-m-g) O (adult concepts anti-socia theme)
Blue Velvet	F. Caruso	USA	119	Hoyts Distribution	S (i-m-j) V (i-m-j L (f-m-g
Call Me	J. Quill/K. Martel	USA	98	Vestron Pictures	L (f-m-g) S (i-m-g)
Crazy Love	E. Provoost/ A. Keysman	Belgium	87	Dendy Cinema	S (i-m-j) O (adult concepts
Travolti Da un Insolito Destino	R. Cardarelli	Italy	114	Ente Autonomo Di Gestione Per Il Cinema	S (i-m-j) O (sexua concepts)

⁽a) Blue Jean Cop was inadvertently omitted from 'Films Registered Without Deletions' For Restricted Exhibition—'R' on September 1988 list.

Title	Producer	Country	Submitted length (mins)	Applicant	Reason for decision
Special Conditions:					
Fatto Di Sangue Tra Due Uomini	H. Colombo	UK	118	Ente Autonomo Di Gestione Per II Cinema	
La Fine Del Mondo Nel Nostro Solito Letto	G. Shiva	Italy	108	Ente Autonomo Di Gestione Per II Cinema	

N.N.-8876533

FILM CENSORSHIP BOARD

WEEK ENDING 28 OCTOBER 1988

Classifications assigned to films for sale/hire pursuant to the Australian Capital Territory Classification of Publications Ordinance 1983; the Northern Territory Classification of Publications Act 1985; the New South Wales Film and Video Tape Classification Act 1984; the Queensland Censorship of Films Act 1947-1984; the Tasmania Classification of Publications Act 1984; the Western Australia Video Tapes Classification and Control Act 1987.

An explanatory key to reasons for classifying non-'G' films appears hereunder:

	Frequency		Explicitness/Intensity			Purpose	
	Infrequent	Frequent	Low	Medium	High	Justified	Gratuitous
S (Sex)	i	f	1	m	h	i	g
V (Violence)	i	f	1	m	h	j	ğ
L (Language)	i	f	1	m	h	j	g

^{*} Films Board of Review

Decision

^{**} Code reasons unavailable for films originally classified before 1972

Title	Producer	Country	Submitted length (mins)	Applicant	Reason for decision
'G'-Suitable for general	l exhibition				
Australia—Take a Bow	B. Morris	Australian	101	Hoyts Polygram Video	
Monster Madness	Halcon Days Productions	USA	30	Majestic Products	
Never Give a Sucker An Even Break/ You're Telling Me	E. Cline/E. Kenton	USA	69	Marieth Investments	
Pro Football Funnies	M. Ehuoin	USA	30	Majestic Products	
Skydiving—The Ultimate Sport	Norman Kent Productions	USA	30	Majestic Products	
Truckin'—The 1987 European Truck Racing Organization Championship Season	Hay Fisher Productions	UK	55	Majestic Products	
Ultimate Home Fitness Club	Not shown	Australia	49	Rob Gild Productions	
'PG'—Parental guidance	required for those unde	er 15			
Bagdad Cafe	E. Adlon/P. Adlon	USA/West Germany	91	Communications and Entertainment	L (i-l-j) O (adult concepts)
Bingo Bongo	M. Cecchi Gori/V. Cecchi Gori	Italy	102	RCA/Columbia Pictures/Hoyts Video	O (adult concepts)
Boggy Creek II	C. Pierce	USA	94	Key United Home Video	L (i-l-j) V (i-l-g)
Bonanza—The Next Generation	D. Dortort	USA	93	Taft Hardie Group	V (i-l-j) O (adult concepts)
'M'-Mature (not recon	mended for viewing by	persons under 1	5)		
Against Her Will (a)	D. Karew	USA	90	CBS/Fox Video	V (i-m-g) O (adult concepts)
Australian Made	J. McLean	Australia	79	Hoyts Polygram Video	L (i-m-g)

Title	Producer	Country	Submitted length (mins)	Applicant	Reason for decision
Beach Balls	M. Leipzig	USA	75	RCA/Columbia Pictures/Hoyts Video	L (i-m-g) S (i-m-g)
Buy & Cell	F. Yablans	USA/Italy	91	Roadshow Home Video	L (i-m-g) V (i-m-g) O (nudity)
Cat Squad: Python Wolf	D. Salven	Canada	93	Filmpac Holdings	V (f-m-g)
Daddy	R. Florio/H. Frey	USA	93	Communications and Entertainment	O (adult concepts)
Distant Thunder	R. Schaffel	Canada/USA	113	United International Pictures	L (f-m-g) V (i-m-g)
Dracula A.D. 1972	J. Douglas	UK	92	Corporate Video	V (i-m-j)
Evil Angels	V. Lambert	Australia	121	Warner Home Video	L (i-m-j) O (adult concepts)
Horror House on Highway Five	Vistar International Productions	USA	89	Popular Home Products	L (i-m-g) V (f-m-g) O (drug use)
Imagine—John Lennon	D. Wolper/A. Solt	USA	105	Warner Home Video	L (i-m-j) O (adult concepts)
In Dangerous Company	R. Preuss/R. Newell	USA	91	Roadshow Home Video	V (i-m-g) O (i-m-g) S (i-m-g)
Jack's Back	T. Moore/C. Elwes	USA	97	CBS/Fox Video	L (i-m-g) O (adult concepts) V (i-m-j)
My Stepmother is an Alien	Not shown	Not shown	98	Hoyts Distribution	O (sexual allusions)
Philip Marlowe Private Eye—Blackmailers Don't Shoot/Spanish Blood	J. Slan	Canada	97	CBS/Fox Video	V (i-m-g) O (drug use)
Sonney and Jed	R. Loyola	Italy	91	Popular Home Products	L (f-m-g) V (i-m-g) O (adult concepts)
Trident Force	Anna Films International	Not shown	86	Roadshow Home Video	V (i-m-g) L (i-m-g) O (sexual allusions)

(a) Title changed from Hostage shown on 26 July 1988 gazette list.

'R'-Restricted (not to be sold or hire or delivered to minors or displayed in a public place unless container bears

Electric Blue	U.K.	57	Video Ray	O (exploitative nudity) S (i-m-g)
Not shown	U.S.A.	64	Mareith Investments	S (f-m-g)
K. Russell	U.K.	93	Outland Promotions	O (horror adult concepts)
M. Golan/Y. Globus	USA/ Yugoslavia	90	Hoyts Distribution	S (i-m-g) O (adult concepts)
D. Kellogg/G. Moskowitz	U.S.A.	41	Palace Entertainment Corporation	O (exploitative nudity)
M. Trikilis	U.S.A.	27	Palace Entertainment Corporation	O (exploitative nudity)
	Not shown K. Russell M. Golan/Y. Globus D. Kellogg/G. Moskowitz	Not shown U.S.A. K. Russell U.K. M. Golan/Y. Globus USA/ Yugoslavia D. Kellogg/G. Moskowitz U.S.A.	Not shown U.S.A. 64 K. Russell U.K. 93 M. Golan/Y. Globus USA/ Yugoslavia 90 D. Kellogg/G. Moskowitz 41	Not shown U.S.A. 64 Mareith Investments K. Russell U.K. 93 Outland Promotions M. Golan/Y. Globus USA/ Yugoslavia D. Kellogg/G. Moskowitz M. Trikilis U.S.A. 41 Palace Entertainment Corporation Your Corporation Palace Entertainment Corporation 7 Palace Entertainment

'X'-Extra Restricted (not to be sold or hired or delivered to minors or displayed except in a restricted publications area and bearing prescribed markings)

Behind Ginger	Vivid Video	U.S.A.	28	Capital Duplicators	S (f-h-g)
Blonde, The	Not shown	U.S.A.	68	Mareith Investments	S (f-h-g)
Brat on the Run	P. Thomas	U.S.A.	81	Leisuremail	S (f-h-g)
Crocodile Blondee Part	The Jellyroll	U.S.A.	83	Leisuremail	S (f-h-g)
II .	Company				

Title	Producer	Country	Submitted length (mins)	Applicant	Reason for decision
Debbie Does Dallas IV	M. Curtis	U.S.A.	73	Leisuremail	S (f-h-g)
Nightmare on Porn Street	Moonlight Entertainment	U.S.A.	74	Leisuremail	S (f-h-g)
Sex Asylum II—Sheer Bedlam	B. Seven	U.S.A.	87	Leisuremail	S (f-h-g) O (mild fetish)
Surf's Up	Ocean Productions	U.S.A.	85	Leisuremail	S (f-h-g)
Refused Classification					
Pop's Oasis	H. Hope/N. Hope	U.S.A.	94	Hoyts Distribution	O (gratuitous sexual violence)

N.N.--8876534

Community Services and Health

COMMONWEALTH OF AUSTRALIA NATIONAL HEALTH ACT 1953 PHARMACEUTICAL BENEFITS DETERMINATION UNDER SECTION 93

I, JOHN STEWART DEEBLE, Acting First Assistant Secretary, Health Benefits Division, Department of Community Services and Health and Delegate of the Minister of State for Community Services and Health, pursuant to section 93 of the National Health Act 1953, hereby make the following Determination:

- This Determination shall come into operation on the first day of December 1988.
- The Determination under section 93 of the National Health Act made on 18 July 1988 with effect from 1 August 1988 is hereby revoked.
- The pharmaceutical benefits referred to in this Determination shall be those contained in the Schedule to this Determination.
- A medical practitioner is authorised for the purpose of section 93 of the Act to supply any of the pharmaceutical benefits obtained by the medical practitioner, or by another medical practitioner for whom the first-named medical practitioner is temporarily acting as a locum, in pursuance of this Determination.
- Subject to this Determination, the maximum quantity or number of units of a pharmaceutical benefit that may be obtained by a medical practitioner in any one month for the purpose of section 93 of the Act is the number or quantity specified in the Schedule to this Determination in relation to the pharmaceutical benefit.
- Where a medical practitioner has obtained a pharmaceutical benefit for the purpose of section 93 of the Act, that medical practitioner is not entitled to obtain a further quantity or number of units of that pharmaceutical benefit for that purpose whilst in possession of a quantity or number of units of the pharmaceutical benefit obtained by the medical practitioner equal to or greater than the maximum quantity or number of units allowed for the pharmaceutical benefit by paragraph 5.
- A medical practitioner is not entitled to obtain a pharmaceutical benefit for the purpose of section 93 of the Act more often than twice in any two months.

THE SCHEDULE

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Maximum quantity
Adrenaline Injection B.P.	Adrenaline, 1 in 1,000, 1 mL	5
Aminophylline Injection B.P.	Ampoule, 250 mg in 10 mL	. 5
Atropine Sulphate Injection B.P.	Ampoule, 600 micrograms in 1 mL	5
Chlorpromazine Injection B.P.	Ampoule, 50 mg in 2 mL	10
OR		
Haloperidol B.P.	Injection, 5 mg in 1 mL amp.	10
Diazepam Injection B.P.	10 mg in 2 mL	5
Digoxin Injection B.P.	Ampoule, 500 micrograms in 2 mL	5
Diphtheria and Tetanus Vaccine, Adsorbed B.P.	Injection, 0.5 mL amp.	10
Diphtheria and Tetanus Vaccine, Adsorbed B.P., Diluted	Injection, 0.5 mL amp.	10
Ergometrine Injection B.P.	Ampoule, 250 micrograms in 1 mL	5
Erythromycin Ethyl Succinate B.P.	Injection, 100 mg (base) in 2 mL	5
Frusemide Injection B.P.	Ampoule, 20 mg in 2 mL	5
Glucose Intravenous Infusion B.P.	Ampoule, 5 g in 10 mL	5
Hydrocortisone Sodium Succinate B.P.	Injection set containing equivalent of 100 mg hydrocortisone and 2 mL solvent	2
OR		
Hydrocortisone Sodium Succinate B.P.	Injection set containing equivalent of 250 mg hydrocortisone and 2 mL solvent	1
OR		
Dexamethasone Sodium Phosphate B.P.	Injection, 1 mL amp. containing equivalent of 4 mg dexamethasone phosphate	5
Lignocaine Hydrochloride B.P.	Injection, 100 mg in 5 mL	4
OR		
Lignocaine Hydrochloride Injection B.P.	Syringe, disposable, 300 mg in 3 mL	1

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Metoclopramide Injection B.P.	Ampoule, 10 mg in 2 mL	10
OR		
Prochlorperazine Edisylate	Injection, 12.5 mg in 1 mL	10
OR		
Prochlorperazine Injection B.P.	Ampoule, 12.5 mg in 1 mL	10
Morphine Sulphate Injection B.P.	Ampoule, 15 mg in 1 mL	5
OR		
Morphine Sulphate Injection B.P.	Ampoule, 30 mg in 1 mL	5
Naloxone Hydrochloride	Injection, 2 mg in 5 mL disposable injection set	2
Pethidine Injection B.P.	Ampoule, 100 mg in 2 mL	5
Procaine Penicillin Injection B.P.	Syringe, disposable, 1.5 g	10
QR		
Benzylpenicillin Potassium B.P. with any determined brand of Water for Injections	<pre>Injection, 600 mg vial (with ampoule, 2 mL)</pre>	10
Promethazine Hydrochloride Injection B.P.	Ampoule, 50 mg in 2 mL	10
Terbutaline Sulphate B.P.	<pre>Injection, 100 micrograms in 1 mL amp.</pre>	5
OR		
Terbutaline Sulphate B.P.	<pre>Injection, 500 micrograms in 1 mL amp.</pre>	5
Verapamil Hydrochloride Injection B.P.	Ampoule, 5 mg in 2 mL	5
Dated this 16th	day of Nonember.	1988.
be 5		

JOHN DEEBLE

ACTING FIRST ASSISTANT SECRETARY

HEALTH BENEFITS DIVISION

DELEGATE OF THE MINISTER OF STATE FOR COMMUNITY SERVICES AND HEALTH

COMMONWEALTH OF AUSTRALIA NATIONAL HEALTH ACT 1953 PHARMACEUTICAL BENEFITS DECLARATION UNDER SUB-SECTION 85(2AA)

I, JOHN STEWART DEFENLE, Acting First Assistant Secretary, Health Benefits Division, Department of Community Services and Health and Delegate of the Minister of State for Community Services and Health, pursuant to sub-section 85(2AA) of the National Health Act 1953, hereby make the following Declaration:

- 1. This Declaration shall come into operation on 1 December 1988.
- 2. The drugs and medicinal preparations contained in the Schedule to this Declaration shall cease to be drugs and medicinal preparations to which Part VII of the National Health Act 1953 applies.

THE SCHEDULE

Atropine Methonitrate B.P.

Benzocaine B.P. with Adrenaline B.P.

Cholera Vaccine B.P. Cloxacillin Sodium B.P. with Purified Water B.P.

Dapsone Tablets B.P.

Epicillin

Ergotamine Tartrate B.P. with Caffeine B.P. and Cyclizine Hydrochloride B.P.

Ergotamine Tartrate B.P. with Caffeine Citrate B.P.C. 1959 and Diphenhydramine Hydrochloride B.P.

Flucloxacillin Sodium B.P. with Purified Water B.P.

Natamycin

Norethisterone Acetate B.P. with Ethinyloestradiol B.P. Ornidazole

Phenytoin Injection B.P.

Plague Vaccine B.P. 1973

Pyridostigmine Injection B.P.

Sodium Calciumedetate B.P.

Sodium Salicylate B.P.

Stilboestrol Tablets B.P.

Sulphamethizole B.P.

"Teric" N8 with Lauryldimethylbenzyl Ammonium Chloride

Trimetaphan Camsylate B.P. 1968

Typhoid Vaccine B.P.

Typhoid Vaccine, Oral

Typhus Vaccine B.P.

Anthrax Antiserum

Anthrax Vaccine

Antirabies Immunoglobulin Injection B.P.

Bacillus Calmette-Guerin Vaccine B.P.

Bacillus Calmette-Guérin Vaccine, Percutaneous B.P.

Black Snake Antivenom

Botulinum Antitoxin B.P.

Box Jellyfish Antivenom Brown Snake Antivenom

Death Adder Antivenom

Formaldehyde Solution B.P.

Funnel-Web Spider Antivenom

Haemodialysis Solutions B.P.

Haemodialysis Solutions B.P.C. 1973

Haemodialysis Solutions, Concentrated B.P.

Haemodialysis Solutions, Concentrated B.P.C. 1973

Intraperitoneal Dialysis Solutions, Concentrated

Pituitary Gonadotrophin, Human

Pituitary Growth Hormone, Human

Rabies Vaccine B.P.

Snake Bite Antivenom, Polyvalent

Sodium Hypochlorite Solution, Strong B.P.

Sodium Hypochlorite Solution 52.5 mg per mL

Stone Fish Antivenom

Taipan Antivenom

Tiger Snake Antivenom

Tuberculin Purified Protein Derivative B.P.

Tuberculin Purified Protein Derivatives, Atypical

Dated this

day of November 1988.

JOHN DEEBLE

ACTING FIRST ASSISTANT SECRETARY

HEALTH BENEFITS DIVISION

DELEGATE OF THE MINISTER OF STATE FOR COMMUNITY SERVICES AND HEALTH

N.N.-8878561

COMMONWEALTH OF AUSTRALIA NATIONAL HEALTH ACT 1953 PHARMACEUTICAL BENEFITS DETERMINATION UNDER SUB-SECTION 84C(7)

I, JOHN STEWART DEEBLE, Acting First Assistant Secretary, Health Benefits Division, Department of Community Services and Health and Delegate of the Minister of State for Community Services and Health, pursuant to sub-section 84C(7) of the National Health Act 1953, hereby make the following Determination:

- This Determination shall come into operation on the first day of December 1988.
- 2. The Determination under sub-section 84C(7) of the National Health Act made on 1 November 1986 with effect from 1 November 1986, as amended, is, in this Determination, referred to as the Principal Determination.
- Sub-paragraphs 10(a), 10(b) and 10(c) of the Principal Determination are amended by omitting "\$0.62" (wherever occurring) and substituting "\$0.63".
- 4. Sub-paragraphs 10(a), 10(c), 20(d), 38(c) and 38(d) of the Principal Determination are amended by omitting "\$0.90" (wherever occurring) and substituting "\$0.91".

Dated this

16th

day of Monember

1988.

JOHN DEEBLE

ACTING FIRST ASSISTANT SECRETARY

HEALTH BENEFITS DIVISION

DELEGATE OF THE MINISTER OF STATE FOR COMMUNITY SERVICES AND HEALTH

N.N.-8878562

COMMONWEALTH OF AUSTRALIA

National Health Act 1953

PHARMACEUTICAL BENEFITS

DECLARATION UNDER SUB-SECTION 85 (2)

No. PB 5 of 1988

I, JOHN STEWART DEEBLE, Acting First Assistant Secretary, Health Benefits Division, Department of Community Services and Health and Delegate of the Minister of State for Community Services and Health, pursuant to sub-section 85 (2) of the National Health Act 1953, hereby make the following Declaration:

- 1. This Declaration shall come into operation on 1 December 1988.
- 2. Declaration No. PB 3 under sub-section 85 (2) of the Act made with effect from 1 August 1988, as amended by Declaration No. PB 4 with effect from 1 August 1988, is hereby revoked.
- In this Declaration:

"the Act" means the National Health Act 1953;

"approved pharmacist" has the same meaning as in sub-section 84 (1) of the Act;

"ready-prepared pharmaceutical benefit" means a drug or medicinal preparation in respect of which there is in force a determination under sub-section 85 (6) of the Act;

"extemporaneously-prepared pharmaceutical benefit" means a pharmaceutical benefit other than a ready-prepared pharmaceutical benefit.

- 4. Part VII of the Act applies in relation to each of the drugs and medicinal preparations as are specified in Schedule 1.
- 5. A medicinal preparation composed of a compound that includes a pharmaceutical benefit specified in column 1 of Schedule 2, other than a compound specified in column 2 of that Schedule opposite to that pharmaceutical benefit, is not a medicinal preparation to which Part VII of the Act applies, unless that pharmaceutical benefit is also specified in Schedule 3, in which case the provisions of paragraphs 7 and 8 apply.
- 6. Part VII of the Act does not apply in relation to a medicinal preparation composed of a compound that includes a ready-prepared pharmaceutical benefit, other than Water for Injections or a pharmaceutical benefit specified in column 1 of Schedule 2.
- 7. Part VII of the Act applies in relation to medicinal preparations composed of one or more of the drugs or medicinal preparations specified in Schedule 3.
- 8. Part VII of the Act applies in relation to medicinal preparations composed of one or more of the drugs or medicinal preparations specified in Schedule 3 with the addition of one or more of the substances specified in Schedule 4.
- 9. The substances specified in Schedule 4 are additives for the purposes of paragraph 85 (2) (b) of the Act.
- 10. Part VII of the Act applies in relation to each of the drugs and medicinal preparations as are specified in Schedule 5.
- 11. The drugs and medicinal preparations specified in Schedule 5 are additional pharmaceutical benefits made available under arrangements provided for by section 100 of the Act.
- 12. Where circumstances are specified in column 2 of Schedule 1 or Schedule 3 opposite the name of a pharmaceutical benefit specified in column 1 of either of those Schedules, that pharmaceutical benefit is a relevant pharmaceutical benefit for the purposes of section 88A of the Act.
- 13. Where circumstances are specified in column 2 of Schedule 3 opposite the name of a pharmaceutical benefit specified in column 1 of that Schedule, those circumstances are also specified in relation to any medicinal preparation containing that pharmaceutical benefit.
- 14. The following circumstances are specified in relation to each relevant pharmaceutical benefit for the purposes of section 88A of the Act:
 - (a) Where a class of persons is specified in column 2 of Schedule 1 or Schedule 3—that the pharmaceutical benefit is to be supplied for the treatment of a person included in that class of persons;
 - (b) Where a disease or condition is specified in column 2 of Schedule 1 or Schedule 3—
 - (i) if sub-sub-paragraph (ii) does not apply—that the pharmaceutical benefit is to be supplied for the treatment of that disease or condition in relation to any person; or
 - (ii) if the disease or condition is specified in relation to a specified class of persons—that the pharmaceutical benefit is to be supplied for the treatment of that disease or condition in a person included in that class of
 - (c) Where a purpose is specified in column 2 of Schedule 1 or Schedule 3—that the pharmaceutical benefit is to be supplied for that purpose;
 - (d) Where it is specified in column 2 of Schedule 1 or Schedule 3 that compliance with authority procedures set out in sub-paragraph 14 (d) is required—
 - (i) that an application for the written authority of the Secretary in relation to the supply of the pharmaceutical benefit has been made by a medical practitioner, on a form approved by the Secretary, and has been forwarded to the Secretary by that medical practitioner or on behalf of that medical practitioner by a person other than an approved pharmacist, and the Secretary has approved the application; or
 - (ii) where the medical practitioner believes that the supply of the pharmaceutical benefit is required urgentlythat the medical practitioner has prepared an Authority Prescription on a form approved by the Secretary and has:
 - (A) issued the original and duplicate of the form to the person for whom the pharmaceutical benefit is to be supplied;
 - (B) on the departmental copy of the form declared that the circumstances specified in column 2 of Schedule 1 or Schedule 3 apply and signed that copy in the medical practitioner's handwriting; and

- (C) forwarded the departmental copy to reach the Secretary within 7 days from the date on which the original and duplicate were issued to the person for whom the pharmaceutical benefit is to be supplied.
- 15. Where a medical practitioner makes an application under sub-paragraph 14 (d) and the Secretary approves the application, the Secretary shall record the approval on a numbered authority and—
 - (a) where, in the approval, the Secretary requires the medical practitioner to vary the prescription for the supply of the pharmaceutical benefit—return the approval to the medical practitioner; or
 - (b) in any other case—return the approval to the medical practitioner or, if the medical practitioner so requests, forward the approval to the person to whom the pharmaceutical benefit is to be supplied.

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SCHEDULE 1

Ready-prepared pharmaceutical benefits

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Acetazolamide B.P.	_
Acetazolamide Sodium	_
Acetazolamide Tablets B.P.	_
Acetylcysteine	Bronchiectasis
,,	Cystic fibrosis (mucoviscidosis)
Acrylic Resin	Ileostomy or colostomy conditions
Acyclovir	In respect of the eye ointment:
	Eye infections caused by herpes simplex virus
	In respect of the tablet, 200 mg:
	In compliance with authority procedures set out in sub paragraph 14 (d):
	Moderate to severe initial genital herpes, complicated
	by severe pain, systemic symptoms or urinary
	retention
	In respect of the tablet, 400 mg: In compliance with authority procedures set out in sub
	paragraph 14 (d):
	Treatment of patients with herpes zoster with severe
	pain who are over 60 years of age and in whom the duration of rash is less than 72 hours
Adrenaline B.P.	
Adrenaline Hydrochloride	_
Adrenaline Injection B.P.	_
"Albumaid XP"	Phenylketonuria
"Albumaid XPXT"	Tyrosinaemia
"Alfaré"	In compliance with authority procedures set out in sub paragraph 14 (d): Biliary atresia
	Chyloascites
	Chylothorax
	Cystic fibrosis Enterokinase deficiency
	Intolerance to both milk protein and soya protein Severe diarrhoea of greater than two weeks duration in
	infants under the age of 4 months
Allopurinol B.P.	
Allopurinol Tablets B.P.	_
Alprenolol Tablets B.P.	_
Aluminium Hydroxide and Magnesium Carbonate Co-dried Gel	-
Aluminium Hydroxide, Dried B.P.	_
Aluminium Hydroxide, Dried B.P. with Magnesium Hydroxide B.P.	_
Aluminium Hydroxide, Dried B.P. with Magnesium Trisilicate B.P. and Magnesium Hydroxide B.P.	_
Aluminium Hydroxide Mixture B.P.	
Aluminium Hydroxide Mixture B.P. with Light Kaolin B.P. or Light Kaolin (Natural) B.P.	_
Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P.	_
Aluminium Hydroxide Mixture B.P. with Magnesium	
Hydroxide B.P. and Oxethazaine Amantadine Hydrochloride	The treatment of Parkinson's disease caused otherwise that by treatment with a drug
Ambenonium Chloride	_,
Amiloride Hydrochloride Tablets B.P.	_
Aminacrine Hydrochloride B.P. 1968	_
Aminoglutethimide	In compliance with authority procedures set out in sub-
, anniogration made	paragraph 14 (d): Post-menopausal metastatic breast cancer in patients wh have failed to respond adequately to endocrin manipulation

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
"Aminogran Food Supplement"	Phenylketonuria
"Aminogran Mineral Mixture"	Phenylketonuria
Aminophylline B.P.	
Aminophylline Injection B.P.	_
Aminophylline Tablets B.P.	_
Amiodarone Hydrochloride	In compliance with authority procedures set out in sub- paragraph 14 (d): For the continuing treatment of severe refractory cardiac arrhythmias where treatment with amiodarone hydro- chloride was initiated in a hospital (in-patient or out- patient)
Amitriptyline Tablets B.P.	_
Ammonium Chloride B.P.	-
Amoxycillin Capsules B.P.	_
Amoxycillin Sodium	
Amoxycillin Trihydrate B.P.	_
Amoxycillin Trihydrate B.P. with Potassium Clavulanate	_
Amoxycillin Trihydrate B.P. with Potassium Clavulanate and Purified Water B.P.	Ξ
Amoxycillin Trihydrate B.P. with Purified Water B.P.	
Amphotericin B.P.	_
Amphotericin Lozenges B.P.	_
Ampicillin Capsules B.P.	_
Ampicillin Sodium B.P.	_
Ampicillin Trihydrate B.P. with Purified Water B.P.	_
Amylobarbitone Sodium B.P.	Epilepsy
Antazoline Phosphate with Naphazoline Hydrochloride	- · · ·
Antazoline Sulphate with Naphazoline Nitrate B.P.	_
Aspirin B.P.	_
Aspirin Tablets B.P.	_
Aspirin Tablets, Dispersible B.P.	
Atenolol	_
Atropine Eye Ointment B.P.	_
Atropine Sulphate B.P.	_
Atropine Sulphate Injection B.P.	_
Atropine Sulphate Tablets B.P.	
Auranofin	_
Aurothioglucose	
Azathioprine Tablets B.P.	_
Baclofen Tablets B.P.	_
"Banish" Beclomethasone Dipropionate B.P.	Ileostomy or colostomy conditions In respect of the capsules and the sprays, metered aerosol, 50 micrograms per dose, 200 doses, 20.4 g and 100 micrograms per dose, 200 doses, 20.4 g: Asthma In respect of the spray, metered aerosol, 250 micrograms per dose, 200 doses, 20.4 g: In compliance with authority procedures set out in subparagraph 14 (d): Severe chronic asthma not responding to lower doses of beclomethasone dipropionate aerosol
Bendrofluazide Tablets B.P.	— — — — — — — — — — — — — — — — — — —
Benzathine Penicillin B.P.	
Benzathine Penicillin B.P. with Procaine Penicillin B.P.,	
Benzylpenicillin Potassium B.P. and Water for Injections	
Benzhexol Tablets B.P.	_
Benzoin Tincture, Compound B.P.	Ileostomy or colostomy conditions
Benztropine Injection B.P.	_
Benztropine Tablets B.P.	

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Benzyl Benzoate Application B.P.	
Benzyl Benzoate B.P.	_
Benzylpenicillin Potassium B.P.	_
Betamethasone Acetate with Betamethasone Sodium Phosphate B.P.	Alopecia areata For local intra-articular or peri-articular infiltration Granulomata, dermal Keloid Lichen planus hypertrophic Lichen simplex chronicus Lupus erythematosus, chronic discoid Necrobiosis lipoidica Uveitis
Betamethasone Dipropionate	_
Betamethasone Tablets B.P.	
Betamethasone Valerate B.P.	_
Betamethasone Valerate Cream B.P.	atom.
Betamethasone Valerate Ointment B.P.	***
Bethanechol Chloride	_
Biperiden Hydrochloride	_
Bisacodyl B.P.	Any disease or condition in a paraplegic or quadriplegic patient
Bisacodyl Suppositories B.P.	Any disease or condition in a paraplegic or quadriplegic patient For use by patients who are receiving long-term extensive nursing care in hospitals or nursing homes For use by patients for whose care a Commonwealth Domiciliary Nursing Care Benefit is approved
Bisacodyl Tablets B.P.	Any disease or condition in a paraplegic or quadriplegic patient For use by patients who are receiving long-term extensive nursing care in hospitals or nursing homes For use by patients for whose care a Commonwealth Domiciliary Nursing Care Benefit is approved
Bismuth Subcitrate	_
Bleomycin Sulphate	Germ cell neoplasms Lymphoma Squamous cell carcinoma
Bromocriptine Mesylate Capsules B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): Acromegaly, prior to surgery or radiotherapy or where surgery or radiotherapy is inappropriate Parkinson's disease Pathological hyperprolactinaemia where appropriate sur- gery or radiotherapy is not indicated or has already been used with incomplete resolution
Bromocriptine Mesylate Tablets B.P.	Urgent suppression of physiological lactation In compliance with authority procedures set out in sub- paragraph 14 (d): Acromegaly, prior to surgery or radiotherapy or where surgery or radiotherapy is inappropriate Parkinson's disease Pathological hyperprolactinaemia where appropriate sur- gery or radiotherapy is not indicated or has already been used with incomplete resolution
Bumetanide	_
Busulphan Tablets B.P.	_
Butyl Monoester Polymer with Ethanol B.P.	Ileostomy or colostomy conditions
Butyl Monoester Polymer with Isopropyl Alcohol B.P.	Ileostomy or colostomy conditions
Calciferol Tablets, High-Strength B.P.	Hypocalcaemia Hypoparathyroidism Osteomalacia following gastrectomy, severe steatorrhoea or renal failure Vitamin D-resistant rickets

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Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Calcitonin (Human)—Synthetic	In compliance with authority procedures set out in sub- paragraph 14 (d): Proven active Paget's disease of bone, or hypercalcaemia, in patients unable to tolerate both pork and salmon calcitonin or who are resistant to treatment with either pork or salmon calcitonin
Calcitonin (Pork) B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): Proven active Paget's disease of bone causing pain or disability Treatment initiated in a hospital (in-patient or out- patient) of hypercalcaemia For the continuation of treatment of patients already established on calcitonin under arrangements provided for by section 100 of the Act
Calcitriol	In compliance with authority procedures set out in sub- paragraph 14 (d): Hypocalcaemia due to renal disease Hypoparathyroidism Hypophosphataemic rickets Vitamin D-resistant rickets
Calcium Carbonate B.P.	Hypocalcaemia Osteoporosis Proven malabsorption
Calcium Carbonate B.P. with Calcium Lactate-Gluconate	Hypocalcaemia Osteoporosis Proven malabsorption
Calcium Folinate	Antidote to folic acid antagonists
Calcium Glubionate	_
Captopril	In compliance with authority procedures set out in sub- paragraph 14 (d): For the continuing treatment of severe refractory cardiac failure where treatment with captopril was initiated in a hospital (in-patient or out-patient) Hypertension where other drug therapy has failed or is inappropriate
Carbachol B.P. 1973	
Carbamazepine B.P.	_
Carbamazepine Tablets B.P. Carbimazole Tablets B.P.	
Carboplatin	In compliance with authority procedures set out in sub- paragraph 14 (d): Advanced stage ovarian carcinoma
Carmellose Sodium B.P. with Pectin and Gelatin B.P.	Ileostomy or colostomy conditions
Cefotaxime Sodium	In compliance with authority procedures set out in sub- paragraph 14 (d): Infections where positive bacteriological evidence con- firms that cefotaxime sodium is an appropriate thera- peutic agent Septicaemia, suspected or proven
Ceftriaxone Sodium	In compliance with authority procedures set out in sub- paragraph 14 (d): Infections where positive bacteriological evidence con- firms that ceftriaxone sodium is an appropriate thera- peutic agent Septicaemia, suspected or proven
Cephalexin B.P. with Purified Water B.P.	
Cephalexin Capsules B.P.	
Cephalothin Sodium B.P.	_
Cephazolin Sodium	In compliance with authority procedures set out in sub- paragraph 14 (d): Infections where positive bacteriological evidence con- firms that cephazolin sodium is an appropriate thera- peutic agent Septicaemia, suspected or proven
Charcoal, Activated B.P.	Ileostomy or colostomy conditions

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Chloral Hydrate B.P.	
Chlorambucil Tablets B.P.	
Chloramphenicol B.P.	
Chloramphenicol B.P. with Polymyxin B Sulphate B.P.	_
	Bacterial meningitis
Chloramphenicol Capsules B.P.	Intracranial bacterial infections Intraocular infections Rickettsioses Typhoid Other serious infections where positive bacteriological evidence confirms that chloramphenicol is the only appropriate antibiotic
Chloramphenicol Eye Drops B.P.	_
Chloramphenicol Eye Ointment B.P.	
Chloramphenicol Palmitate Mixture B.P.	Bacterial meningitis Intracranial bacterial infections Intraocular infections Rickettsioses Typhoid Other serious infections where positive bacteriological evidence confirms that chloramphenicol is the only appropriate antibiotic
Chloramphenicol Sodium Succinate B.P.	Bacterial meningitis Intracranial bacterial infections Intraocular infections Rickettsioses Typhoid Other serious infections where positive bacteriological evidence confirms that chloramphenicol is the only appropriate antibiotic
Chlorhexidine Gluconate	_
Chlormethiazole Capsules B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): Short-term alcohol or drug withdrawal therapy in a hos- pital or approved centre Status epilepticus
Chlormethiazole Edisylate B.P.	Short-term alcohol or drug withdrawal therapy in a hospital or approved centre Status epilepticus
Chloroquine Phosphate Tablets B.P.	_ · ·
Chloroquine Sulphate B.P.	
Chlorothiazide Tablets B.P.	_
Chlorpromazine Elixir B.P.	Chorea
•	Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness Severe conduct disorders in patients under the age of 16
	years
Chlorpromazine Injection B.P.	Chorea Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness Severe conduct disorders in patients under the age of 16 years

years

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Chlorpromazine Suppositories B.P.	Chorea
	Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia
	Senile dementia
	Malignant neoplasia (late stage) Radiation sickness
	Severe conduct disorders in patients under the age of 16 years
Chlorpromazine Tablets B.P.	Chorea Hyperactive states of organic or toxic delirium
	Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia
	Malignant neoplasia (late stage) Radiation sickness
	Severe conduct disorders in patients under the age of 16 years
Chlorpropamide Tablets B.P.	_
Chalestone mine	— In compliance with outbooks proceedures set out in out-
Cholestyramine	In compliance with authority procedures set out in sub- paragraph 14 (d): Bile salt malabsorption Hypercholesterolaemia Pruritus associated with partial biliary obstruction not responding to other therapy Severe diarrhoea associated with pelvic irradiation
Choline Theophyllinate B.P.	
Chorionic Gonadotrophin B.P.	In respect of the injection set containing 1 ampoule powder for injection 5,000 units and 1 ampoule solvent ImL: In compliance with authority procedures set out in sub-paragraph 14 (d): For use with bromocriptine mesylate in infertility associated with hyperprolactinaemia
	In respect of the other injection sets: In compliance with authority procedures set out in sub-
	paragraph 14 (d): For the treatment of anovulatory infertility in females under 41 years of age with no more than 2 live children by their present union For the treatment of infertility in males due to hypogonadotrophic hypogonadism For the treatment of infertility in males associated with
	isolated luteinizing hormone deficiency For the treatment of males who have combined deficiency of human growth hormone and gonadotrophins and in whom the absence of secondary sexual characteristics indicates a lag in maturation For the treatment of males over the age of 16 years who show clinical evidence of hypogonadism or delayed puberty
Cimetidine	In respect of the tablets, 200 mg and 400 mg: In compliance with authority procedures set out in sub- paragraph 14 (d): Duodenal ulcer (including pyloric and stomal ulcers), proven by current or prior x-ray, endoscopy or surgery Gastric ulcer, proven by x-ray, endoscopy or surgery within the previous two years Scleroderma oesophagus Severe ulcerating (erosive) oesophagitis, proven by en- doscopy and unresponsive to other measures Zollinger-Ellison syndrome In respect of the tablet, 800 mg:

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
	In compliance with authority procedures set out in sub- paragraph 14 (d): Duodenal ulcer (including pyloric and stomal ulcers) proven by current or prior x-ray, endoscopy or surgery Gastric ulcer, proven by x-ray, endoscopy or surgery within the previous two years Zollinger-Ellison syndrome
Ciprofloxacin Hydrochloride	In compliance with authority procedures set out in sub- paragraph 14 (d): Serious infections for which no other antimicrobial agent is appropriate
Clindamycin Capsules B.P.	Gram-positive coccal infections where the infection cannot be safely and effectively treated with penicillin or a derivative of penicillin
Clindamycin Palmitate Hydrochloride with Purified Water B.P.	Gram-positive coccal infections where the infection cannot be safely and effectively treated with penicillin or a derivative of penicillin
Clioquinol B.P.	
Clofibrate Capsules B.P.	Hypertriglyceridaemia Hypercholesterolaemia
Clomiphene Tablets B.P.	Anovulatory infertility Patients undergoing in-vitro fertilisation
Clomipramine Hydrochloride B.P.	Cataplexy associated with narcolepsy Obsessive compulsive disorders
Clonazepam	=
Clonidine Hydrochloride Tablets B.P.	
Clopamide	_ .
Clotrimazole B.P.	_
Cloxacillin Capsules B.P.	_
Cloxacillin Sodium B.P.	_
Codeine Phosphate B.P. with Aspirin B.P.	_
Codeine Phosphate B.P. with Paracetamol B.P.	žia.
Codeine Phosphate Tablets B.P. Colchicine Tablets B.P.	
Colestipol Hydrochloride	In compliance with authority procedures set out in sub- paragraph 14 (d): Hypercholesterolaemia
Colistin Sulphate B.P. with Neomycin Sulphate B.P.	
Colistin Sulphomethate Sodium B.P.	_
Copper Sulphate B.P.C. 1973	
Cortisone Tablets B.P.	_
Co-trimoxazole Mixture, Paediatric B.P.	_
Co-trimoxazole Tablets B.P.	
Cyclopenthiazide Tablets B.P.	_
Cyclophosphamide B.P.	_
Cyclophosphamide Tablets B.P.	_
Cyproheptadine Tablets B.P.	Prevention of migraine
Cytarabine B.P.	_
Danazol	In compliance with authority procedures set out in sub- paragraph 14 (d): Endometriosis, proven by visual means Hereditary angio-oedema Menorrhagia, intractable primary
Dantrolene Sodium	Treatment of chronic spasticity
Debrisoquine Tablets B.P.	— · · · · · · · · · · · · · · · · · · ·
"De-Lact Infant"	Acute gastro-enteritis complicated by lactose intolerance in
	infants under the age of 12 months Proven lactose intolerance in infants under the age of 12 months
Demeclocycline Capsules B.P.	Syndrome of inappropriate antidiuretic hormone secretion caused otherwise than by treatment with a drug
Desipramine Tablets B.P.	_

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Desmopressin	
Dexamethasone B.P.	
Dexamethasone B.P. with Framycetin Sulphate B.P. and Gramicidin	=
Dexamethasone Sodium Metasulphobenzoate with Framycetin Sulphate B.P. and Gramicidin	_
Dexamethasone Sodium Phosphate B.P.	In respect of the injection, 5 mL vial containing equivalent of 120 mg dexamethasone phosphate: For use in a hospital In respect of the other injections:
Dexamethasone Tablets B.P.	_
Dexamphetamine Tablets B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): For the treatment of a patient under the age of 21 years and who is hyperkinetic as a result of brain damage Narcolepsy
Dextran 40 Intravenous Infusion B.P. with Glucose B.P. Dextran 40 Intravenous Infusion B.P. with Sodium Chloride B.P.	
Dextran 70 Intravenous Infusion B.P. with Glucose B.P. Dextran 70 Intravenous Infusion B.P. with Sodium Chloride B.P.	
Diazepam B.P.	_
Diazepam Injection B.P.	_
Diazepam Tablets B.P.	-
Diazoxide Injection B.P.	_
Dichlorphenamide Tablets B.P.	
Diclofenac Sodium	_
Dienoestrol B.P.	-
Difenoxin Hydrochloride with Atropine Sulphate B.P.	_
Diflunisal Tablets B.P. "Digestelact"	In compliance with authority procedures set out in sub- paragraph 14 (d): Lactose intolerance
Digoxin Elixir, Paediatric B.P.	Lactose intolerance
Digoxin Injection B.P.	_
Digoxin Tablets B.P.	
Dihydroergotamine Mesylate B.P.	-
Dihydrotachysterol B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): Hypoparathyroidism Osteomalacia following gastrectomy, severe steatorrhoea or renal failure Vitamin D-resistant rickets
Diloxanide Furoate Tablets B.P.	
Diltiazem Hydrochloride	In compliance with authority procedures set out in sub- paragraph 14 (d): Angina where treatment with verapamil hydrochloride or nifedipine has failed or is inappropriate
Dimercaprol Injection B.P.	
Dimethicones B.P. Diphenoxylate Hydrochloride B.P. with Atropine Sulphate	lleostomy or colostomy conditions
B.P.	
Diphtheria and Tetanus Vaccine, Adsorbed B.P., Diluted	_
Diphtheria Antitoxin B.P.	
Diphtheria, Tetanus and Pertussis Vaccine, Adsorbed B.P.	_
Diphtheria Vaccine, Adsorbed B.P.	-
Diphtheria Vaccine, Adsorbed B.P., Diluted	-
Dipivefrine Hydrochloride	-

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Disodium Etidronate	In compliance with authority procedures set out in sub- paragraph 14 (d): Active Paget's disease of bone when calcitonin has been found to be unsatisfactory due to lack of efficacy of unacceptable side effects
Disopyramide Capsules B.P.	unacceptable side effects —
Disopyramide Phosphate Capsules B.P.	
Docusate Sodium B.P. with Bisacodyl B.P.	Any disease or condition in a paraplegic or quadriplegic
·	patient For use by patients who are receiving long-term extensive nursing care in hospitals or nursing homes For use by patients for whose care a Commonwealth Dom iciliary Nursing Care Benefit is approved
Domperidone	-
Dothiepin Capsules B.P.	-
Dothiepin Hydrochloride B.P.	
Doxepin Capsules B.P.	_
Doxepin Hydrochloride B.P.	aprile.
Doxorubicin Hydrochloride	autoria.
Doxycycline Capsules B.P.	-
Doxycycline Hydrochloride B.P.	In respect of the capsules, 50 mg (base) (containing enterior coated pellets) and the tablets, 50 mg (base): Bronchiectasis in patients over the age of 8 years Chronic bronchitis in patients over the age of 8 years Severe acne In respect of the other capsules and tablets:
Dydrogesterone Tablets B.P.	In compliance with authority procedures set out in sub paragraph 14 (d): Endometriosis, proven by visual means
Econazole Nitrate B.P.	
Econazole Nitrate Cream B.P.	_
Econazole Nitrate Pessaries B.P.	<u> </u>
Ecothiopate Iodide B.P.	_
Enalapril Maleate	In compliance with authority procedures set out in sub- paragraph 14 (d): For the continuing treatment of severe refractory cardiac failure where treatment with enalapril maleate was initiated in a hospital (in-patient or out-patient) Hypertension where other drug therapy has failed or is inappropriate
Ergometrine and Oxytocin Injection B.P.	_
Ergometrine Injection B.P.	
Ergometrine Tablets B.P.	
Ergotamine Tablets B.P.	_
Ergotamine Tartrate B.P. with Caffeine B.P.	
Erythromycin B.P.	wante
Erythromycin Estolate B.P.	_
Erythromycin Ethyl Succinate B.P.	_
Erythromycin Ethyl Succinate B.P. with Purified Water B.P.	_
Erythromycin Lactobionate	_
Erythromycin Stearate	<u></u>
Erythromycin Stearate Tablets B.P.	
Erythromycin Tablets B.P.	_
Ethacrynic Acid Tablets B.P.	Patients who are hypersensitive to other oral diuretics
Ethinyloestradiol Tablets B.P.	
Ethosuximide Capsules B.P.	_
Ethosuximide Elixir B.P.	_
Ethyloestrenol Tablets B.P.	. In compliance with authority procedures set out in sub paragraph 14 (d): Osteoporosis Patients on long-term treatment with corticosteroids

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Ethynodiol Diacetate B.P. with Ethinyloestradiol B.P.	
Etretinate	In compliance with authority procedures set out in sub- paragraph 14 (d): Darier's disease Erythrokeratoderma Pityriasis rubra pilaris Severe congenital ichthyosis (lamellar, bullous and sex linked) Severe intractable psoriasis Severe lichen planus Severe palmo-plantar keratoderma
Famotidine	In compliance with authority procedures set out in sub- paragraph 14 (d): Duodenal ulcer (including pyloric and stomal ulcers), proven by current or prior x-ray, endoscopy or surgery Gastric ulcer, proven by x-ray, endoscopy or surgery within the previous two years Zollinger-Ellison syndrome
Felodipine	In compliance with authority procedures set out in sub- paragraph 14 (d): Treatment where less costly alternative therapy is inappropriate
Fenoterol Hydrobromide B.P.	_
Ferrous Aminoacetosulphate	_
Ferrous Gluconate B.P.	_
Ferrous Gluconate Tablets B.P.	_
Ferrous Sulphate, Dried B.P.	_
Ferrous Sulphate, Dried B.P. with Folic Acid B.P.	_
Flecainide Acetate	In compliance with authority procedures set out in sub- paragraph 14 (d): For the continuing treatment of cardiac arrhythmias where treatment with flecainide acetate was initiated in a hospital (in-patient or out-patient)
Fluclorolone Acetonide B.P.	-
Flucloxacillin Capsules B.P.	_
Flucloxacillin Magnesium B.P. with Purified Water B.P.	_
Flucloxacillin Sodium B.P.	_
Flucytosine Tablets B.P.	Invasive or systemic fungal infections such as candidiasis, cryptococcosis and chromoblastomycosis
Fludrocortisone Tablets B.P.	_
Flumethasone Pivalate with Clioquinol B.P.	_
Fluocortolone Pivalate and Fluocortolone Hexanoate Cream B.P.	_
Fluocortolone Pivalate and Fluocortolone Hexanoate Ointment B.P.	_
Fluorometholone	-
Fluorouracil Injection B.P.	
Fluoxymesterone Tablets B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): Aplastic anaemias, proven Carcinoma of the breast Hypogonadism Osteoporosis
Fluphenazine Decanoate Injection B.P.	Chorea Hyperactive states of organic or toxic delirium Major pyschoses including: Major organic pyschoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness

Column 1	Column 2
	Circumstances (if any) specified for the purposes of section
Pharmaceutical benefit	88A of the Act
Fluphenazine Tablets B.P.	Chorea Hyperactive states of organic or toxic delirium Major psychoses including: Major organic pyschoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness
Folic Acid B.P.	Radiation sickness
Folic Acid Tablets B.P.	_
Fosfestrol Sodium B.P.	Carcinoma of the prostate
Framycetin Sulphate B.P.	
Frusemide B.P.	_
Frusemide Injection B.P.	_
Frusemide Tablets B.P.	In respect of the 500 mg strength: Renal failure In respect of the other strengths:
Fusidic Acid Mixture B.P.	In compliance with authority procedures set out in sub- paragraph 14 (d): For use in combination with another antibiotic in the treatment of serious staphylococcal infections confirmed by bacteriological evidence
Gas-gangrene Antitoxin, Mixed B.P.	_
Gentamicin Eye Drops B.P.	Invasive ocular infection Perioperative use Suspected pseudomonal eye infection
Gentamicin Injection B.P.	—
Glibenclamide Tablets B.P.	_
Gliclazide	_
Glipizide	_
Glucagon Hydrochloride	_
Glucose and Ketone Indicator—Urine	_
Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P.	_
Glucose Indicator—Blood	_
Glucose Indicator—Urine	_
Glucose Intravenous Infusion B.P.	_
Glyceryl Trinitrate Changel Trinitrate Tableta B.B.	
Glyceryl Trinitrate Tablets B.P. Griseofulvin Tablets B.P.	— Kerion
	Recalcitrant tinea infections
Haloperidol B.P. Haloperidol Solution B.P.	Chorea Hyperactive states of organic or toxic delirium Major psychoses including: Major organic pyschoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness Chorea
The second of th	Hyperactive states of organic or toxic delirium Major psychoses including: Major organic pyschoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness

Column I	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Haloperidol Tablets B.P.	Chorea Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness
Heparin Injection B.P. (Calcium Salt)	-
Heparin Injection B.P. (Sodium Salt)	_
Hexamine Hippurate Hexamine Mandelate	_
Homatropine Hydrobromide B.P.	_
Hydralazine Hydrochloride Tablets B.P.	_
Hydrochlorothiazide B.P. with Amiloride Hydrochloride B.P.	_
Hydrochlorothiazide B.P. with Triamterene B.P.	_
Hydrochlorothiazide Tablets B.P.	_
Hydrocortisone B.P.	_
Hydrocortisone Acetate B.P.	In respect of the rectal foam: Proctitis Ulcerative colitis In respect of the other forms:
Hudroparticone Acatota D.D. with Nacarrain Culphate D.D.	_
Hydrocortisone Acetate B.P. with Neomycin Sulphate B.P.	
Hydrocortisone Acetate Cream B.P. Hydrocortisone Acetate Injection B.P.	Alopecia areata
	For local intra-articular or peri-articular infiltration Granulomata, dermal Keloid Lichen planus hypertrophic Lichen simplex chronicus Lupus erythematosus, chronic discoid
Hydrocortisone Acetate Ointment B.P.	_
Hydrocortisone Sodium Succinate B.P.	In respect of the injection set containing equivalent of 500 mg hydrocortisone and 4 mL solvent: Any disease or condition in a patient receiving treatment in a hospital In respect of the other injection sets:
Hydroxocobalamin Injection B.P.	Pernicious anaemia and other proven vitamin B12 deficiencies Post-gastrectomy treatment Sub-acute combined degeneration of the cord
Hydroxychloroquine Tablets B.P.	_
Hydroxyurea Capsules B.P.	
Hyoscyamine Hydrobromide with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P.	_
Hyoscyamine Sulphate B.P. with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P.	
Hypromellose 4500 B.P.	_
Hypromellose 4500 B.P. with Dextran 70	_
Ibuprofen Tablets B.P.	
Idoxuridine B.P.	
Idoxuridine Eye Drops B.P.	-
Imipramine Hydrochloride B.P.	_
Imipramine Tablets B.P.	_
Indapamide	-
Indomethacin Capsules B.P.	- .
Indomethacin Suppositories B.P.	-
Influenza Vaccine (Split Virion), Inactivated B.P.	For prophylaxis of persons at special risk of adverse con- sequences from infections of the lower respiratory tract

Column I	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Insect Allergen Extract—Honey Bee Venom	_
Insect Allergen Extract—Paper Wasp Venom	_
Insulin Injection, Acid B.P.	_
Insulin Injection, Biphasic B.P.	_
Insulin Injection, Isophane B.P. Insulin Injection, Isophane B.P. and Insulin Injection, Neutral B.P.	_
Insulin Injection, Neutral B.P.	_
Insulin Injection, Protamine Zinc B.P.	_
Insulin Isophane, Human (Synthetic)	Diabetics exhibiting lipoatrophy, lipohypertrophy, local allergy or immunologic resistance
Insulin Isophane, Human (Synthetic) and Insulin Neutral, Human (Synthetic)	Newly diagnosed insulin dependent diabetics Diabetics exhibiting lipoatrophy, lipohypertrophy, local allergy or immunologic resistance
Insulin Neutral, Human (Synthetic)	Newly diagnosed insulin dependent diabetics Diabetics exhibiting lipoatrophy, lipohypertrophy, local allergy or immunologic resistance
Insulin Zinc Suspension B.P.	Newly diagnosed insulin dependent diabetics
•	-
Insulin Zinc Suspension (Amorphous) B.P.	_
Insulin Zinc Suspension (Crystalline) B.P. Insulin Zinc Suspension (Crystalline), Human (Synthetic)	Dishatian ambibiting lineatasanha lineabannastasanha Jasah
Insulin Zinc Suspension, Human (Synthetic)	Diabetics exhibiting lipoatrophy, lipohypertrophy, local allergy or immunologic resistance Newly diagnosed insulin dependent diabetics Diabetics exhibiting lipoatrophy, lipohypertrophy, local
	allergy or immunologic resistance Newly diagnosed insulin dependent diabetics
Intraperitoneal Dialysis Solutions B.P.C. 1973	_
Ipratropium Bromide	-
Iron Dextran Injection B.P.	-
Iron Polymaitose Complex	
Isoconazole Nitrate	_
Isoniazid Tablets B.P.	_
Isopropyl Monoester Polymer with Isopropyl Alcohol B.P. Isosorbide Dinitrate Tablets B.P.	Ileostomy or colostomy conditions
Isotretinoin	In compliance with authority procedures set out in sub- paragraph 14(d): Severe cystic acne not responsive to other therapy
Kaolin, Light B.P. or Light Kaolin (Natural) B.P. with Pectin	-
Ketoconazole	In compliance with authority procedures set out in sub- paragraph 14(d): Systemic or deep mycoses where other forms of therapy have failed
Ketoprofen B.P.	
Ketoprofen Capsules B.P.	_
Labetalol Hydrochloride Tablets B.P.	-
Lactulose Solution B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Hepatic coma or pre-coma (chronic portosystemic encephalopathy)
	Terminal malignant neoplasia
Lauramine Oxide with Octoxinol Leuprorelin Acetate	Ileostomy or colostomy conditions In compliance with authority procedures set out in sub- paragraph 14 (d): Advanced cancer of the prostate
Levodopa and Carbidopa Tablets B.P.	_
Levodopa B.P. with Benserazide	
Levodopa Tablets B.P.	_
Levonorgestrel B.P.	
Levonorgestrel B.P. with Ethinyloestradiol B.P.	_

Column I	Column 2 Circumstances (if any) specified for the purposes of section
Pharmaceutical benefit	88A of the Act
Lignocaine Hydrochloride B.P.	_
Lignocaine Hydrochloride Injection B.P.	-
Lincomycin Injection B.P.	_
Lindane B.P.	
Lindane Cream B.P.	
Liothyronine Tablets B.P.	_
Lithium Carbonate B.P. Lithium Carbonate Tablets B.P.	_
"Locasol New Formula"	In compliance with authority procedures set out in sub-
Locasor New Formula	paragraph 14(d): Hypercalcaemia in children under the age of 2 years Osteopetrosis
"Lofenalac"	Phenylketonuria
Loperamide Hydrochloride	-
Lypressin	In compliance with authority procedures set out in sub-
	paragraph 14(d): Treatment where less costly alternative therapy is inappropriate
"Maxamaid RVHB"	Pyridoxine non-responsive homocystinuria
"Maxamaid XP"	Phenylketonuria
"Maxamum XP"	Phenylketonuria
Medroxyprogesterone Acetate B.P. Medrysone Mefenamic Acid Capsules B.P. Mefruside Megestrol Acetate B.P. Melphalan Tablets B.P. Menopausal Gonadotrophin, Human	In respect of the injections, 50 mg in 1 mL vial and 500 mg in 2.5 mL vial, the oral suspension and the tablets 100 mg, 200 mg and 250 mg: Breast cancer Endometrial cancer Renal cell cancer In respect of the injection, 150 mg in 1 mL vial: Breast cancer Endometrial cancer Endometriosis Renal cell cancer In respect of the tablet, 10 mg: In respect of the tablet, 500 mg: Advanced breast cancer Dysmenorrhoea Menorrhagia Breast cancer In compliance with authority procedures set out in subparagraph 14(d): For the treatment of anovulatory infertility in females
	under 41 years of age with no more than 2 live childrer by their present union For the treatment of infertility in males due to hypogon- adotrophic hypogonadism
Mercaptopurine Tablets B.P.	_
Metformin Tablets B.P.	_
Methacycline Hydrochloride B.P.C. 1973	_
Methadone Injection B.P.	Severe disabling pain not responding to non-narcotic analgesics
Methadone Tablets B.P.	Severe disabling pain not responding to non-narcotic analgesics
Methdilazine Hydrochloride	Prevention of migraine
Methenolone Acetate	In compliance with authority procedures set out in sub- paragraph 14(d): Osteoporosis Patients on long-term treatment with corticosteroids
Methotrexate B.P.	•
Methotrexate Injection B.P.	

	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Methotrexate Tablets B.P.	_
Methsuximide	In compliance with authority procedures set out in sub- paragraph 14(d): Treatment where less costly alternative therapy is inappropriate
Methyclothiazide	_
Methyldopa Tablets B.P.	_
Methylphenobarbitone B.P.	Epilepsy
Methylprednisolone Acetate Injection B.P.	For local intra-articular or peri-articular infiltration
Methylprednisolone Sodium Succinate	
Methyl Salicylate Liniment A.P.F.	_
Methyltestosterone Tablets B.P.	In respect of the 5 mg strength: Hypogonadism in patients under the age of 16 years In respect of the other strengths: Carcinoma of the breast Hypogonadism Klinefelter's syndrome (seminiferous tubule dysgenesis)
Methysergide Tablets B.P.	_
Metoclopramide Hydrochloride B.P.	-
Metoclopramide Injection B.P.	
Metoclopramide Tablets B.P.	_
Metolazone	_
Metoprolol Tartrate	_
Metronidazole Benzoate	_
Metronidazole B.P.	Prophylaxis in large bowel surgery Treatment, in a hospital, of acute anaerobic sepsis
Metronidazole Suppositories B.P.	_
Metronidazole Tablets B.P.	-
Mexiletine Hydrochloride Capsules B.P.	_
Mianserin Hydrochloride Tablets B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Depressive illness in persons with cardiovascular disease bladder neck dysfunction or glaucoma
Miconazole	
Miconazole Nitrate B.P.	_
Miconazole Nitrate Cream B.P.	_
Minocycline Hydrochloride	_
Minoxidil	In compliance with authority procedures set out in sub- paragraph 14(d): For the continuing treatment of severe refractory hyper- tensive disease where treatment with minoxidil was initiated in a hospital (in-patient or out-patient) For the continuing treatment of a patient who has al- ready received, for more than 6 months, therapy with minoxidil for severe refractory hypertension
Misoprostol	In compliance with authority procedures set out in sub- paragraph 14(d): Duodenal ulcer (including pyloric and stomal ulcers) proven by x-ray, endoscopy or surgery within 3 months of application Gastric ulcer, proven by x-ray, endoscopy or surgery within 3 months of application
Mithramycin	For use in a hospital for: Inoperable testicular neoplasm Symptomatic hypercalcaemia and hypercalcinuria associated with advanced neoplasia
Mitozantrone Hydrochloride	Metastatic or locally advanced breast cancer
Morphine Sulphate B.P. with Tacrine Hydrochloride	Severe intractable pain
M. II. O.I. I. I. I. I. I. D.D.	*****
Morphine Sulphate Injection B.P.	
Morphine Sulphate Tablets B.P.	Severe disabling pain not responding to non-narcotic analgesics

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Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Mustine Hydrochloride B.P.	
Nalidixic Acid Tablets B.P.	For use as a urinary antiseptic in patients with neurogenic bladder
	Urinary tract infections where current clinical and bacteri- ological evidence confirm that nalidixic acid is an appro- priate therapeutic agent
Naloxone Hydrochloride	_
Nandrolone Decanoate Injection B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Carcinoma of the breast Osteoporosis Patients on long-term treatment with corticosteroids
Nandrolone Phenylpropionate Injection B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Carcinoma of the breast Osteoporosis Patients on long-term treatment with corticosteroids
Naphazoline Hydrochloride	
Naproxen B.P.	_
Naproxen Tablets B.P.	_
Neomycin Sulphate Eye Drops B.P.	_
Neomycin Tablets B.P.	Acute leukaemia during induction of remission with chemotherapy Bowel sterilisation preparatory to major surgery Encephalopathy, hepatic
Neomycin Undecenoate with Bacitracin Zinc B.P.	Encephatopathy, nepatic
Neostigmine Injection B.P.	Ξ
Neostigmine Tablets B.P.	-
Niclosamide Tablets B.P.	_
Nicotinic Acid Tablets B.P.	
Nifedipine	_
Nitrazepam Tablets B.P.	
Nitrofurantoin B.P.	_
Nitrofurantoin Mixture B.P.	-
Nitrofurantoin Tablets B.P.	_
Norethisterone Acetate B.P.	Carcinoma of the breast Carcinoma of the prostate
Norethisterone B.P. with Ethinyloestradiol B.P.	_
Norethisterone B.P. with Mestranol B.P.	-
Norethisterone Tablets B.P.	-
Norfloxacin	Acute bacterial enterocolitis Urinary tract infection
Nortriptyline Hydrochloride B.P.	_
Nortriptyline Tablets B.P. "Nutramigen"	In compliance with authority procedures set out in sub- paragraph 14(d):
	Allergy to both cows' milk and soy protein in children under the age of 2 years Cystic fibrosis (mucoviscidosis) Galactosaemia Glycogen storage disease due to glucose-6-phosphatase deficiency in children under the age of 2 years
	Homocystinuria Lactose intolerance
Nystatin B.P.	
Nystatin Mixture B.P.	_
Nystatin Ointment B.P.	_
Nystatin Pessaries B.P.	_
Nystatin Tablets B.P.	_
"Odorgon"	Ileostomy or colostomy conditions
•	Ileostomy or colostomy conditions —

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Oestrone B.P.C. 1954	_
Olsalazine Sodium	In compliance with authority procedures set out in sub-
	paragraph 14(d): Inflammatory bowel disease involving the colon in pa- tients with proven hypersensitivity to sulphonamides
Orphenadrine Hydrochloride Tablets B.P.	Parkinsonism
Oxazepam Tablets B.P.	_
Oxprenolol Hydrochloride B.P.	_
Oxprenolol Tablets B.P.	_
Oxycodone Hydrochloride	Severe disabling pain not responding to non-narcotic analgesics
Oxycodone Pectinate	Severe disabling pain not responding to non-narcotic analgesics
Oxymetholone Tablets B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Aplastic anaemias, proven Myelosclerosis
Oxytocin Injection B.P.	_ ·
Pancreatin B.P.	Cystic fibrosis (mucoviscidosis) Following pancreatico-duodenectomy Pancreatic steatorrhoea
Pancrelipase	Cystic fibrosis (mucoviscidosis) Following pancreatico-duodenectomy Pancreatic steatorrhoea
Papaveretum B.P.C. 1973	_
Papaveretum B.P.C. 1973 with Hyoscine Hydrobromide B.P. Paracetamol B.P.	_ _
Paracetamol Tablets B.P.	_
Paraffin, Soft White B.P.	Ileostomy or colostomy conditions
Paraffin, Soft White B.P. with Liquid Paraffin B.P.	_
Penicillamine Tablets B.P.	Acute heavy metal intoxication Cystinosis Cystinosis with calculus formation
	Cystinuria with calculus formation Haemoglobinuria, paroxysmal cold Severe rheumatoid arthritis Wilson's disease (hepatolenticular degeneration)
Perhexiline Maleate	In compliance with authority procedures set out in sub paragraph 14(d): Angina not responding to other therapy
Pericyazine	Chorea
	Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states
	Schizophrenia Senile dementia
	Malignant neoplasia (late stage) Radiation sickness Severe conduct disorders in patients under the age of 10
Dakiding Introduct D.D.	years
Pethidine Injection B.P. Pethidine Tablets B.P.	In compliance with authority procedures set out in sub paragraph 14(d):
Phenelzine Tablets B.P.	Disabling pain associated with proven malignant neoplasia Depressive illness resistant to treatment with either tricyclic antidepressants or electroconvulsive therapy Phobic states
Phenethicillin Capsules B.P.	_
Phenethicillin Potassium B.P. with Purified Water B.P. Phenethicillin Tablets B.P.	_
Phenindione Tablets B.P.	Patients stabilised on phenindione for more than 6 months
Phenobarbitone Injection B.P.	Epilepsy
-	

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Phenoxybenzamine Capsules B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): Neurogenic urinary retention Phaeochromocytoma
Phenoxymethylpenicillin (Benzathine Salt)	_
Phenoxymethylpenicillin (Hydrabamine Salt)	_
Phenoxymethylpenicillin Potassium Capsules B.P.	-
Phenoxymethylpenicillin Potassium Tablets B.P.	_
Phensuximide B.P.C. 1973	In compliance with authority procedures set out in sub- paragraph 14(d): Treatment where less costly alternative therapy is
Observation that a third with side D.D.	inappropriate
Phenylephrine Hydrochloride B.P.	-
Phenytoin B.P.	
Phenytoin Capsules B.P.	-
Phenytoin Mixture B.P.	_
Phytomenadione Injection B.P.	
Pilocarpine	In compliance with authority procedures set out in sub- paragraph 14(d): Chronic glaucoma when other miotics are not tolerated
Pilocarpine Hydrochloride B.P.	
Pindolol B.P.	_
Pindolol Tablets B.P.	_
Piperazine Oestrone Sulphate	_
Piroxicam	pane.
Pizotifen Malate	_
PK AID I"	Phenylketonuria
'PK AID II"	Phenylketonuria
Pneumococcal Vaccine, Polyvalent	Splenectomised persons over the age of 2 years Persons with Hodgkin's disease Persons at high risk of pneumococcal infections
Polygeline	
Polyisobutylene	Ileostomy or colostomy conditions
Polymyxin B Sulphate B.P. with Bacitracin B.P. 1968 and Neomycin Sulphate B.P.	
Polymyxin B Sulphate B.P. with Bacitracin Zinc B.P. and Neomycin Sulphate B.P.	_
Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Gramicidin	
Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Hydrocortisone B.P.	_
Polyvinyl Alcohol	_
Polyvinyl Alcohol with Povidone B.P.	The same of the same of the same
Polyvinylpyrrolidone and Vinyl Acetate Copolymer "Portagen"	Ileostomy or colostomy conditions In compliance with authority procedures set out in sub- paragraph 14(d): Chyloascites Chylothorax Patients requiring a ketogenic diet for intractable child- hood epilepsy Proven malabsorption
Potassium Chloride B.P.	p
Potassium Chloride B.P. with Potassium Bicarbonate Efferverscent Tablets	=
Pralidoxime Iodide	_
Prazosin Hydrochloride	sultern
Prednisolone Acetate	In respect of the injection: For local intra-articular or peri-articular infiltration In respect of the eye drops:
Prednisolone Acetate with Phenylephrine Hydrochloride B.P.	Corneal grafts
	Uveitis

Column 1	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Prednisolone Sodium Phosphate B.P.	In respect of the retention enema: In compliance with authority procedures set out in sub- paragraph 14(d): Proctitis Ulcerative colitis In respect of the suppositories: Proctitis Ulcerative colitis In respect of the eye/ear drops:
Prednisolone Stearoylglycolate	
Prednisolone Tablets B.P.	
Prednisone Tablets B.P.	_
"Pregestimil"	In compliance with authority procedures set out in sub- paragraph 14(d): Biliary atresia Chyloascites Chylothorax Cystic fibrosis Enterokinase deficiency Intolerance to both milk protein and soya protein Severe diarrhoea of greater than two weeks duration in infants under the age of 4 months
Primaquine Tablets B.P.	-
Primidone Tablets B.P.	-
Probenecid Tablets B.P.	_
Probucol	In compliance with authority procedures set out in sub- paragraph 14(d): Familial hypercholesterolaemia inadequately responsive to diet and a cholesterol lowering resin
Procainamide Hydrochloride B.P.	
Procainamide Injection B.P.	~
Procaine Penicillin Injection B.P.	_
Procarbazine Hydrochloride	
Prochlorperazine	
Prochlorperazine Edisylate	
Prochlorperazine Injection B.P.	_
Prochlorperazine Tablets B.P.	
Procyclidine Tablets B.P.	_
Promethazine Hydrochloride Injection B.P.	_
Promethazine Theoclate Tablets B.P.	_
Propantheline Tablets B.P.	Chronic neurogenic incontinence of urine
Propranolol Injection B.P.	
Propranolol Tablets B.P.	_
Propylthiouracil Tablets B.P.	_
Protamine Sulphate Injection B.P.	
Pyrantel Embonate	***
Pyridostigmine Tablets B.P.	
Pyridoxine Hydrochloride B.P.	_
Pyridoxine Hydrochloride Tablets B.P.	Anaemia proved to be responsive to pyridoxine Convulsions responsive to pyridoxine Homocystinuria Primary hyperoxaluria Radiation sickness Sideroblastic (refractory) anaemia Treatment and prophylaxis of peripheral neuritis caused or likely to be caused by isoniazid therapy

Column I	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Pyrimethamine Tablets B.P.	_
Quinethazone	-
Quinidine Bisulphate B.P.	_
Quinidine Sulphate B.P.	_
Quinidine Sulphate Tablets B.P.	-
Quinine Bisulphate Tablets B.P.	_
Quinine Sulphate Tablets B.P. Ranitidine Hydrochloride	In respect of the tablet, 150 mg (base): In compliance with authority procedures set out in subparagraph 14(d): Duodenal ulcer (including pyloric and stomal ulcers) proven by current or prior x-ray, endoscopy or surgery Gastric ulcer, proven by x-ray, endoscopy or surgery within the previous 2 years Scleroderma oesophagus Severe ulcerating (erosive) oesophagitis, proven by endoscopy and unresponsive to other measures Zollinger-Ellison syndrome In respect of the tablet, 300 mg (base): In compliance with authority procedures set out in subparagraph 14(d): Duodenal ulcer (including pyloric and stomal ulcers) proven by current or prior x-ray, endoscopy or surgery Gastric ulcer, proven by x-ray, endoscopy or surgery within the previous 2 years Zollinger-Ellison syndrome
Red-back Spider Antivenom	
Rifampicin B.P.	Prophylactic treatment of contacts of patients with Hae mophilus influenzae type B Prophylaxis of meningococcal disease in close contacts and carriers In compliance with authority procedures set out in subparagraph 14(d): Leprosy in adults
Rolitetracycline	
Salbutamol B.P.	
Salbutamol Sulphate B.P.	_
Salbutamol Tablets B.P.	
Salcatonin Injection B.P. Semisodium Valproate	In compliance with authority procedures set out in sub- paragraph 14(d): Proven active Paget's disease of bone causing pain or disability Treatment initiated in a hospital (in-patient or out-patient) of hypercalcaemia For the continuation of treatment of patients already established on calcitonin under arrangements provided for by section 100 of the Act
Silver Sulphadiazine with Chlorhexidine Gluconate	In respect of the cream, 10 mg-2 mg per g, 50 g and the cream, 10 mg-2 mg per g, 100 g: For the prevention and treatment of infection in partia or full skin thickness loss due to burns or epidermo
Sodium Acid Citrate B.P. Sodium Acid Phosphate B.P.	lysis bullosa Stasis ulcers In respect of the cream, 10 mg-2 mg per g, 500 g: In compliance with authority procedures set out in sub- paragraph 14(d): Treatment, in a hospital, of burns In compliance with authority procedures set out in sub- paragraph 14(d): Familial hypophosphataemia Hypercalcaemia

Column I	Column 2
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act
Sodium Aurothiomalate Injection B.P.	_
Sodium Bicarbonate Intravenous Infusion B.P.	_
Sodium Chloride and Glucose Intravenous Infusion B.P.	
Sodium Chloride B.P. with Potassium Chloride B.P. and Calcium Chloride B.P. in Water for Injections	_
Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.	_
Sodium Chloride Intravenous Infusion B.P. Sodium Citro-Tartrate	-
Sodium Cromoglycate B.P.	In respect of the eye drops: In compliance with authority procedures set out in sub- paragraph 14(d): Vernal kerato-conjunctivitis In respect of the nebuliser solution: Use in a power-operated nebuliser for prophylaxis of asthma in persons who are unable to use the capsule or metered aerosol spray for oral inhalation
	In respect of the metered aerosol spray:
Sodium Cromoglycate Insufflation B.P.	-
Sodium Fusidate B.P.	In compliance with authority procedures set out in sub- paragraph 14(d): For use in combination with another antibiotic in the treatment of serious staphylococcal infections con- firmed by bacteriological evidence
Sodium Lactate Compound Intravenous Infusion B.P.	_
Sodium Lactate Compound Intravenous Infusion B.P. with Anhydrous Glucose B.P.	_
Sodium Nitroprusside B.P.	-
Sodium Valproate B.P.	_
Sodium Valproate Elixir B.P.	_
Sotalol Hydrochloride	In compliance with authority procedures set out in sub- paragraph 14(d): For the continuing treatment of severe refractory cardiac arrhythmias where treatment with sotalol hydrochlo- ride was initiated in a hospital (in-patient or out- patient)
Spectinomycin Hydrochloride B.P.	
Spironolactone Tablets B.P.	Female hirsutism
•	Hyperaldosteronism
Staphylococcus Toxoid B.P. 1968	-
Sterculia B.P.	Ileostomy or colostomy conditions
Sterculia B.P. with Frangula Bark B.P.	Any disease or condition in a paraplegic or quadriplegic patient For use by patients who are receiving long-term extensive nursing care in hospitals or nursing homes For use by patients for whose care a Commonwealth Domiciliary Nursing Care Benefit is approved
Streptokinase B.P.	
Sucralfate	_
Sulindac Tablets B.P.	_
Sulphacetamide Sodium B.P.	_
Sulphadimidine Tablets B.P.	_
Sulphafurazole B.P.	
Sulphamethizole Tablets B.P.	_
Sulphasalazine	Crohn's disease
oupmonazine .	Cronn's disease Rheumatoid arthritis Ulcerative colitis
Sulphinpyrazone Tablets B.P.	Glomerulonephritis Gout Renal transplantation

nn 2
nstances (if any) specified for the purposes of section f the Act
omy or colostomy conditions
disease or condition in a paraplegic or quadriplegicient
se with surgical appliances omy or colostomy conditions
disease or condition in a paraplegic or quadriplegic ient
se with surgical appliances omy or colostomy conditions
cancer in post-menopausal women with positive hor- ne receptor levels and local lymph node involvement n or without other evidence of metastases
rent or metastatic breast cancer
gonadism
felter's syndrome (seminiferous tubule dysgenesis)
tism
gonadism felter's syndrome (seminiferous tubule dysgenesis)
tism
gonadism elter's syndrome (seminiferous tubule dysgenesis)
gonadism felter's syndrome (seminiferous tubule dysgenesis)
a not adequately controlled by other drug therapy
mpliance with authority procedures set out in sub- agraph 14(d): osarrhythmia Itiple sclerosis, acute exacerbation atment of children who need long-term corticosteroic herapy but who are in danger of growth suppressior s a result atment of patients who are being withdrawn from ong-term corticosteroid therapy erative colitis, proven, not responding to parentera orticosteroids
ea
ractive states of organic or toxic delirium psychoses including: jor organic psychoses including arteriopathic demention depressive disorder—manic phase anoid states izophrenia ille dementia nant neoplasia (late stage) tion sickness

years

Thiotepa B.P.

Column 1	Column 2				
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of 88A of the Act				
Thioridazine Tablets R P	Chorea				

f section Thioridazine Tablets B.P. Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder-manic phase

Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness

Severe conduct disorders in patients under the age of 16 years

Thyroxine Tablets B.P. In compliance with authority procedures set out in sub-Ticarcillin Sodium paragraph 14(d): Infections where positive bacteriological evidence confirms that ticarcillin sodium is an appropriate therapeutic agent Septicaemia, suspected or proven

Timolol Maleate Eye Drops B.P. Timolol Maleate Tablets B.P.

Tinidazole Invasive ocular infection Tobramycin B.P. Perioperative use Suspected pseudomonal eve infection In compliance with authority procedures set out in sub-Tobramycin Injection B.P. paragraph 14(d):

Tolazamide Tablets B.P. Tolbutamide Tablets B.P. Tranexamic Acid Tablets B.P.

Tranyleypromine Tablets B.P. Triamcinolone Acetonide B.P.

Gramicidin and Nystatin B.P. Triamcinolone Acetonide Cream B.P. Triamterene B.P.

Triamcinolone Acetonide Ointment B.P. Trifluoperazine Hydrochloride B.P.

Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P.,

Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness years

Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder-manic phase

Severe conduct disorders in patients under the age of 16

Infections where positive bacteriological evidence confirms that tobramycin is an appropriate antibiotic

Depressive illness resistant to treatment with either tricyclic

antidepressants or electroconvulsive therapy

For local intra-articular or peri-articular infiltration

Septicaemia, suspected or proven

Hereditary angio-oedema

Phobic states

Keloid

Psoriasis

Chorea

Alopecia areata

Granulomata, dermal

Necrobiosis lipoidica

Lichen planus hypertrophic Lichen simplex chronicus

Lupus erythematosus, chronic discoid

Column 1 Pharmaceutical benefit Trifluoperazine Hydrochloride Tablets B.P.	Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness
	Chorea Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness
Trifluoperazine Hydrochloride Tablets B.P.	Hyperactive states of organic or toxic delirium Major psychoses including: Major organic psychoses including arteriopathic dementia Manic depressive disorder—manic phase Paranoid states Schizophrenia Senile dementia Malignant neoplasia (late stage) Radiation sickness
	Severe conduct disorders in patients under the age of 16 years
Triglycerides Oil, Medium Chain	In compliance with authority procedures set out in sub- paragraph 14(d): Abetalipoproteinaemia Chyloascites Chylothorax Intestinal lymphangiectasia Intractable childhood epilepsy requiring a ketogenic diet Short-term nutritional support in severe steatorrhoea Steatorrhoea due to distal small bowel resection
Trimethoprim Tablets B.P.	_
Trimipramine	_
Trimipramine Tablets B.P.	_
Trioxysalen	Vitiligo
Urea Cream B.P.	_
Vancomycin Hydrochloride B.P.	In respect of the capsules: In compliance with authority procedures set out in sub- paragraph 14(d): Antibiotic associated colitis In respect of the injection: In compliance with authority procedures set out in sub- paragraph 14(d): Any disease or condition in a patient receiving treat- ment in a hospital Endophthalmitis
Vasopressin Tannate	
Verapamil Hydrochloride B.P.	-
Verapamil Hydrochloride Injection B.P.	
Verapamil Hydrochloride Tablets B.P.	_
Vidarabine	Eye infections caused by herpes simplex virus
Vinblastine Sulphate B.P.	· · · ·
Vincristine Sulphate B.P.	_
Warfarin Tablets B.P.	
Water for Injections B.P.	_
Wool Alcohols Ointment B.P.	****
Zinc Oxide B.P.	
Zinc Sulphate B.P. with Phenylephrine Hydrochloride B.P.	

Allowable compounds of ready-prepared pharmaceutical benefits

Column 1	Column 2					
Pharmaceutical benefit	Allowable compounds					
Aluminium Hydroxide, Dried B.P.	Aluminium Hydroxide, Dried B.P. with Magnesium Hydroxide B.P. Aluminium Hydroxide, Dried B.P. with Magnesium Hydroxides B.P. with Magnesi					
Aluminium Hydroxide Mixture B.P.	droxide B.P. and Magnesium Trisilicate B.P. Aluminium Hydroxide Mixture B.P. with Light Kaolin B.P.					
Administrative Mixture D.F.	or Light Kaolin (Natural) B.P. Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. and Oxethazaine					
Amiloride Hydrochloride B.P.	Hydrochlorothiazide B.P. with Amiloride Hydrochloride B.P.					
Amoxycillin Trihydrate B.P.	Amoxycillin Trihydrate B.P. with Potassium Clavulanate Amoxycillin Trihydrate B.P. with Potassium Clavulanate and Purified Water B.P. Amoxycillin Trihydrate B.P. with Purified Water B.P.					
Ampicillin Trihydrate B.P.	Ampicillin Trihydrate B.P. with Purified Water B.P.					
Antazoline Phosphate	Antazoline Phosphate with Naphazoline Hydrochloride					
Antazoline Sulphate	Antazoline Sulphate with Naphazoline Nitrate B.P.					
Aspirin B.P.	Codeine Phosphate B.P. with Aspirin B.P.					
Atropine Sulphate B.P.	Difenoxin Hydrochloride with Atropine Sulphate B.P. Diphenoxylate Hydrochloride B.P. with Atropine Sulphate B.P.					
	Hyoscyamine Hydrobromide with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P. Hyoscyamine Sulphate B.P. with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P.					
Bacitracin B.P. 1968	Polymyxin B Sulphate B.P. with Bacitracin B.P. 1968 and Neomycin Sulphate B.P.					
Bacitracin Zinc B.P.	Neomycin Undecenoate with Bacitracin Zinc B.P. Polymyxin B Sulphate B.P. with Bacitracin Zinc B.P. an Neomycin Sulphate B.P.					
Benserazide	Levodopa B.P. with Benserazide					
Benzathine Penicillin B.P.	Benzathine Penicillin B.P. with Procaine Penicillin B.P. Benzylpenicillin Potassium B.P. and Water for Injections					
Benzylpenicillin Potassium B.P.	Benzathine Penicillin B.P. with Procaine Penicillin B.P. Benzylpenicillin Potassium B.P. and Water for Injections					
Betamethasone Acetate	Betamethasone Acetate with Betamethasone Sodium Phos- phate B.P.					
Betamethasone Sodium Phosphate B.P.	Betamethasone Acetate with Betamethasone Sodium Phos- phate B.P.					
Bisacodyl B.P.	Docusate Sodium B.P. with Bisacodyl B.P.					
Butyl Monoester Polymer	Butyl Monoester Polymer with Ethanol B.P. Butyl Monoester Polymer with Isopropyl Alcohol B.P.					
Caffeine B.P.	Ergotamine Tartrate B.P. with Caffeine B.P.					
Calcium Carbonate B.P. Calcium Chloride B.P.	Calcium Carbonate B.P. with Calcium Lactate-Gluconate Sodium Chloride B.P. with Potassium Chloride B.P. and Calcium Chloride B.P. in Water for Injections					
Calcium Lactate-Gluconate	Calcium Carbonate B.P. with Calcium Lactate-Gluconate					
Carmellose Sodium B.P.	Carmellose Sodium B.P. with Pectin and Gelatin B.P.					
Cephalexin B.P.	Cephalexin B.P. with Purified Water B.P.					
Chloramphenicol B.P.	Chloramphenicol B.P. with Polymyxin B Sulphate B.P.					
Chlorhexidine Gluconate	Silver Sulphadiazine with Chlorhexidine Gluconate					
Citric Acid B.P.	Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassiun Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P.					
Clindamycin Palmitate Hydrochloride	Clindamycin Palmitate Hydrochloride with Purified Water B.P.					
Codeine Phosphate B.P.	Codeine Phosphate B.P. with Aspirin B.P. Codeine Phosphate B.P. with Paracetamol B.P.					
Colistin Sulphate B.P.	Colistin Sulphate B.P. with Neomycin Sulphate B.P.					
Dexamethasone B.P.	Dexamethasone B.P. with Framycetin Sulphate B.P. and Gramicidin					

Gramicidin

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Column I Pharmaceutical benefit	Column 2 Allowable compounds				
Dexamethasone Sodium Metasulphobenzoate	Dexamethasone Sodium Metasulphobenzoate with Framy- cetin Sulphate B.P. and Gramicidin				
Dextran 40 Intravenous Infusion B.P.	Dextran 40 Intravenous Infusion B.P. with Glucose B.P. Dextran 40 Intravenous Infusion B.P. with Sodium Chloride B.P.				
Dextran 70 Dextran 70 Intravenous Infusion B.P.	Hypromellose 4500 B.P. with Dextran 70 Dextran 70 Intravenous Infusion B.P. with Glucose B.P. Dextran 70 Intravenous Infusion B.P. with Sodium Chloride B.P.				
Difenoxin Hydrochloride Diphenoxylate Hydrochloride B.P.	Difenoxin Hydrochloride with Atropine Sulphate B.P. Diphenoxylate Hydrochloride B.P. with Atropine Sulphate B.P.				
Docusate Sodium B.P.	Docusate Sodium B.P. with Bisacodyl B.P.				
Ergotamine Tartrate B.P. Erythromycin Ethyl Succinate B.P.	Ergotamine Tartrate B.P. with Caffeine B.P. Erythromycin Ethyl Succinate B.P. with Purified Water B.P.				
Ethanol B.P. Ethinyloestradiol B.P.	Butyl Monoester Polymer with Ethanol B.P. Ethynodiol Diacetate B.P. with Ethinyloestradiol B.P. Levonorgestrel B.P. with Ethinyloestradiol B.P. Norethisterone B.P. with Ethinyloestradiol B.P.				
Ethynodiol Diacetate B.P. Ferrous Sulphate, Dried B.P. Flucloxacillin Magnesium B.P. Flumethasone Pivalate	Ethynodiol Diacetate B.P. with Ethinyloestradiol B.P. Ferrous Sulphate, Dried B.P. with Folic Acid B.P. Flucloxacillin Magnesium B.P. with Purified Water B.P. Flumethasone Pivalate with Clioquinol B.P.				
Folic Acid B.P.	Ferrous Sulphate, Dried B.P. with Folic Acid B.P.				
Framycetin Sulphate B.P.	Dexamethasone B.P. with Framycetin Sulphate B.P. and Gramicidin Dexamethasone Sodium Metasulphobenzoate with Framycetin Sulphate B.P. and Gramicidin				
Frangula Bark B.P.	Sterculia B.P. with Frangula Bark B.P.				
Gelatin B.P. Glucose B.P.	Carmellose Sodium B.P. with Pectin and Gelatin B.P. Dextran 40 Intravenous Infusion B.P. with Glucose B.P. Dextran 70 Intravenous Infusion B.P. with Glucose B.P.				
Glucose, Anhydrous B.P.	Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P. Sodium Lactate Compound Intravenous Infusion B.P. with Anhydrous Glucose B.P.				
Gramicidin	Dexamethasone B.P. with Framycetin Sulphate B.P. and Gramicidin Dexamethasone Sodium Metasulphobenzoate with Framycetin Sulphate B.P. and Gramicidin Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Gramicidin Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P., Gramicidin and Nystatin B.P.				
Hydrochlorothiazide B.P.	Hydrochlorothiazide B.P. with Amiloride Hydrochloride B.P. Hydrochlorothiazide B.P. with Triamterene B.P.				
Hydrocortisone B.P.	Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Hydrocortisone B.P.				
Hydrocortisone Acetate B.P. Hyoscine Hydrobromide B.P.	Hydrocortisone Acetate B.P. with Neomycin Sulphate B.P. Hyoscyamine Hydrobromide with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P. Hyoscyamine Sulphate B.P. with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P. Papaveretum B.P.C. 1973 with Hyoscine Hydrobromide B.P.				
Hyoscyamine Hydrobromide	Hyoscyamine Hydrobromide with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P.				
Hyoscyamine Sulphate B.P.	Hyoscyamine Sulphate B.P. with Atropine Sulphate B.P. and Hyoscine Hydrobromide B.P.				
Hypromellose 4500 B.P.	Hypromellose 4500 B.P. with Dextran 70				
Insulin Injection, Isophane B.P.	Insulin Injection, Isophane B.P. and Insulin Injection, Neutral B.P.				
Insulin Injection, Neutral B.P.	Insulin Injection, Isophane B.P. and Insulin Injection, Neutral B.P.				

Column 1	Column 2				
Pharmaceutical benefit	Allowable compounds				
Insulin Isophane, Human (Synthetic)	Insulin Isophane, Human (Synthetic) and Insulin Neutral, Human (Synthetic)				
Insulin Neutral, Human (Synthetic)	Insulin Isophane, Human (Synthetic) and Insulin Neutra Human (Synthetic)				
Isopropyl Alcohol B.P.	Butyl Monoester Polymer with Isopropyl Alcohol B.P. Isopropyl Monoester Polymer with Isopropyl Alcohol B.P				
Isopropyl Monoester Polymer	Isopropyl Monoester Polymer with Isopropyl Alcohol B.I				
Kaolin, Light B.P. or Light Kaolin (Natural) B.P.	Aluminium Hydroxide Mixture B.P. with Light Kaolin B.P. or Light Kaolin (Natural) B.P. Kaolin, Light B.P. or Light Kaolin (Natural) B.P. with Pectin				
Lauramine Oxide	Lauramine Oxide with Octoxinol				
Levodopa B.P.	Levodopa B.P. with Benserazide				
Levonorgestrel B.P.	Levonorgestrel B.P. with Ethinyloestradiol B.P.				
Magnesium Chloride B.P.	Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.				
Magnesium Hydroxide B.P.	Aluminium Hydroxide, Dried B.P. with Magnesium Hydroxide B.P. Aluminium Hydroxide, Dried B.P. with Magnesium Trisilicate B.P. and Magnesium Hydroxide B.P. Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. and Oxethazaine				
Magnesium Trisilicate B.P.	Aluminium Hydroxide, Dried B.P. with Magnesium Trisilicate B.P. and Magnesium Hydroxide B.P.				
Mestranol B.P.	Norethisterone B.P. with Mestranol B.P.				
Morphine Sulphate B.P.	Morphine Sulphate B.P. with Tacrine Hydrochloride B.P.				
Naphazoline Hydrochloride	Antazoline Phosphate with Naphazoline Hydrochloride				
Naphazoline Nitrate B.P.	Antazoline Sulphate with Naphazoline Nitrate B.P.				
Neomycin Sulphate B.P.	Colistin Sulphate B.P. with Neomycin Sulphate B.P. Hydrocortisone Acetate B.P. with Neomycin Sulphate B.P. Polymyxin B Sulphate B.P. with Bacitracin B.P. 1968 and Neomycin Sulphate B.P. with Bacitracin Zinc B.P. and Neomycin Sulphate B.P. with Bacitracin Zinc B.P. and Neomycin Sulphate B.P. with Neomycin Sulphate B.P. and Gramicidin Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Hydrocortisone B.P. Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P., Gramicidin and Nystatin B.P.				
Neomycin Undecenoate	Neomycin Undecenoate with Bacitracin Zinc B.P.				
Norethisterone B.P.	Norethisterone B.P. with Ethinyloestradiol B.P. Norethisterone B.P. with Mestranol B.P.				
Nystatin B.P.	Tetracycline B.P. with a buffering agent and Nystatin B.P. Tetracycline Hydrochloride B.P. with Nystatin B.P. Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P., Gramicidin and Nystatin B.P.				
Octoxinol	Lauramine Oxide with Octoxinol				
Oxethazaine	Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. and Oxethazaine				
Papaveretum B.P.C. 1973	Papaveretum B.P.C. 1973 with Hyoscine Hydrobromide B.P.				
Paracetamol B.P.	Codeine Phosphate B.P. with Paracetamol B.P.				
Paraffin, Liquid B.P.	Paraffin, Soft White B.P. with Liquid Paraffin B.P.				
Paraffin, Soft White B.P. Pectin	Paraffin, Soft White B.P. with Liquid Paraffin B.P. Carmellose Sodium B.P. with Pectin and Gelatin B.P. Kaolin, Light B.P. or Light Kaolin (Natural) B.P. with Pectin				
Phenethicillin Potassium B.P.	Phenethicillin Potassium B.P. with Purified Water B.P.				
Phenylephrine Hydrochloride B.P.	Prednisolone Acetate with Phenylephrine Hydrochloride B.P. Zinc Sulphate B.P. with Phenylephrine Hydrochloride B.P.				

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Column 1	Column 2
Pharmaceutical benefit	Allowable compounds
Polymyxin B Sulphate B.P.	Chloramphenicol B.P. with Polymyxin B Sulphate B.P. Polymyxin B Sulphate B.P. with Bacitracin B.P. 1968 and Neomycin Sulphate B.P. Polymyxin B Sulphate B.P. with Bacitracin Zinc B.P. and Neomycin Sulphate B.P. Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Gramicidin
	Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Hydrocortisone B.P.
Polyvinyl Alcohol	Polyvinyl Alcohol with Povidone B.P.
Potassium Chloride B.P.	Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P. Sodium Chloride B.P. with Potassium Chloride B.P. and Calcium Chloride B.P. in Water for Injections Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.
Potassium Clavulanate	Amoxycillin Trihydrate B.P. with Potassium Clavulanate Amoxycillin Trihydrate B.P. with Potassium Clavulanate and Purified Water B.P.
Povidone B.P.	Polyvinyl Alcohol with Povidone B.P.
Prednisolone Acetate	Prednisolone Acetate with Phenylephrine Hydrochloride B.P.
Procaine Penicillin B.P.	Benzathine Penicillin B.P. with Procaine Penicillin B.P., Benzylpenicillin Potassium B.P. and Water for Injections
Silver Sulphadiazine	Silver Sulphadiazine with Chlorhexidine Gluconate
Sodium Acetate B.P.	Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.
Sodium Chloride B.P.	 Dextran 40 Intravenous Infusion B.P. with Sodium Chloride B.P. Dextran 70 Intravenous Infusion B.P. with Sodium Chloride B.P. Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P. Sodium Chloride B.P. with Potassium Chloride B.P. and Calcium Chloride B.P. in Water for Injections Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.
Sodium Citrate B.P.	Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P.
Sodium Gluconate	Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Magnesium Chloride B.P.
Sodium Lactate Compound Intravenous Infusion B.P.	Sodium Lactate Compound Intravenous Infusion B.P. with Anhydrous Glucose B.P.
Sterculia B.P.	Sterculia B.P. with Frangula Bark B.P.
Tacrine Hydrochloride	Morphine Sulphate B.P. with Tacrine Hydrochloride
Testosterone Decanoate B.P.	Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P., Testosterone Isocaproate B.P. and Testosterone Decanoate B.P.
Testosterone Isocaproate B.P.	Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P. and Testosterone Isocaproate B.P. Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P., Testosterone Isocaproate B.P. and Testosterone Decanoate B.P.
Testosterone Phenylpropionate B.P.	Testosterone Propionate B.P. with Testosterone Phenylpro-

Testosterone Propionate B.P.

Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P. and Testosterone Isocaproate B.P. Testosterone Propionate B.P. with Testosterone Phenylpro-

pionate B.P. and Testosterone Isocaproate B.P.
Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P., Testosterone Isocaproate B.P. and Testosterone

sterone Decanoate B.P.

estosterone Propionate B.P. with Testosterone Phenylpropionate B.P., Testosterone Isocaproate B.P. and Testosterone Decanoate B.P.

Column I	Column 2						
Pharmaceutical benefit	Allowable compounds						
Tetracycline B.P.	Tetracycline B.P. with a buffering agent Tetracycline B.P. with a buffering agent and Ny- statin B.P.						
Tetracycline Hydrochloride B.P.	Tetracycline Hydrochloride B.P. with Nystatin B.P.						
Triamcinolone Acetonide B.P.	Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P., Gramicidin and Nystatin B.P.						
Triamterene B.P.	Hydrochlorothiazide B.P. with Triamterene B.P.						
Water, Purified B.P.	 Amoxycillin Trihydrate B.P. with Potassium Clavulanate and Purified Water B.P. Amoxycillin Trihydrate B.P. with Purified Water B.P. Ampicillin Trihydrate B.P. with Purified Water B.P. Cephalexin B.P. with Purified Water B.P. Clindamycin Palmitate Hydrochloride with Purified Water B.P. 						
	Erythromycin Ethyl Succinate B.P. with Purified Water B.P. Flucloxacillin Magnesium B.P. with Purified Water B.P. Phenethicillin Potassium B.P. with Purified Water B.P.						
Zinc Sulphate B.P.	Zinc Sulphate B.P. with Phenylephrine Hydrochloride B.P.						

Drugs or medicinal preparations which may be used as ingredients of extemporaneously-prepared pharmaceutical benefits

Column 1	Column 2					
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act					
	ova y me zer					
Acacia B.P.	_					
Acetic Acid (33 per cent) B.P.	_					
Adrandina B.B.	— Managed to the constitution of the state o					
Adrenaline B.P. Alum B.P.	May only be prescribed in eye drops					
Aluminium Acetate Solution B.P.						
						
Aminophylline B.P. Ammonia Spirit, Aromatic B.P.	_					
Ammoniated Mercury B.P. 1973						
Ascorbic Acid B.P.	Use on insteadient of Ferrous Culmbota Minture A.B.E. and					
Ascorbic Acid B.F.	Use as ingredient of Ferrous Sulphate Mixture A.P.F. a Ferrous Sulphate Mixture C.F. A.P.F. 13					
Aspirin B.P.	_					
Atropine Sulphate B.P.	_					
Beeswax, White B.P.	_					
Belladonna Tincture B.P.	•••					
Bentonite B.P.	_					
Benzocaine B.P.						
Benzoic Acid B.P.	_					
Benzoin Tincture, Compound B.P.	_					
Cade Oil B.P.C. 1973	_					
Calamine B.P.	_					
Calcium Hydroxide B.P.	_					
Camphor Spirit, Compound A.P.F.						
Cetomacrogol Emulsifying Wax B.P.	_					
Cetostearyl Alcohol B.P.	_					
Cetrimide B.P.	_					
Chlorbutol B.P.	_					
Chlorinated Lime B.P.	_					
Chlorocresol B.P.	_					
Citric Acid Monohydrate B.P.	_					
Clioquinol B.P.	_					
Coal Tar B.P.	_					
Coal Tar, Prepared B.P. 1973	_					
Coal Tar Solution B.P.	~					
Cocaine Hydrochloride B.P.	· -					
Coconut Oil B.P.	_					
Codeine Phosphate B.P.						
Collodion, Flexible B.P.	_					
Crystal Violet B.P.	_					
Dithranol B.P.	_					
Emulsifying Wax B.P.	-					
Ephedrine Hydrochloride B.P.	May only be prescribed in nasal instillations					
Ferrous Sulphate B.P.	_					
Ferrous Sulphate, Dried B.P.						
Formaldehyde Solution B.P.	_					
Gentian Infusion, Compound, Concentrated B.P.	_					
Glycerol B.P.	_					
Hydrochloric Acid B.P.	_					
Hyoscyamus Liquid Extract B.P.C. 1973	_					
Ichthammol B.P.	_					
Iodine B.P.	_					
Ipecacuanha Tincture B.P.	_					
Kaolin, Light B.P.	_					
Kaolin, Light (Natural) B.P.	-					

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Column 1	Column 2				
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act				
Lactic Acid B.P.					
Lactose B.P.	_				
Lavender Oil, Spike B.P.C. 1968	_				
Lemon Spirit B.P.	_				
Lignocaine Hydrochloride B.P.	_				
Liquorice Liquid Extract B.P.	_				
Lobelia Tincture, Ethereal B.P.C. 1973	_				
Magnesium Carbonate, Light B.P.	_				
Magnesium Sulphate B.P.	May only be prescribed for other than oral use				
Magnesium Trisilicate B.P.					
Menthol B.P.	_				
Methyl Salicylate B.P.	_				
Methyl Salicylate Ointment, Compound A.P.F. 1934	_				
Morphine Hydrochloride B.P.	_				
Oily Cream A.P.F.					
Oleic Acid B.P.					
Opium Tincture B.P.	_				
. ·	-				
Orange Tincture B.P.	-				
Paraffin, Hard B.P.	_				
Paraffin, Light Liquid B.P.	Manager to be access that for and a set of				
Paraffin, Liquid B.P.	May only be prescribed for other than oral use				
Paraffin, Soft White B.P.	_				
Paraffin, Soft Yellow B.P.					
Phenobarbitone B.P.	May only be prescribed for the treatment of epilepsy				
Phenobarbitone Sodium B.P.	May only be prescribed for the treatment of epilepsy				
Phenol B.P.	May not be prescribed in ear drops				
Phenoxyethanol B.P.					
Pilocarpine Hydrochloride B.P.					
Pilocarpine Nitrate B.P.					
Podophyllum Resin B.P.					
Potassium Citrate B.P.					
Potassium Iodide B.P.	- .				
Potassium Permanganate B.P.	_				
Propylene Glycol B.P.	_				
Pumilio Pine Oil B.P.					
Raspberry Syrup B.P.					
Red Syrup A.P.F.	_				
Resorcinol B.P.	_				
Salicylic Acid B.P.	_				
Siberian Fir Oil B.P.C. 1949					
Silver Nitrate B.P.					
Soap, Soft B.P.	_				
Sodium Acid Phosphate B.P.	_				
Sodium Bicarbonate B.P.	_				
Sodium Chloride B.P.	_				
Sodium Citrate B.P.	_				
Sodium Phosphate B.P.	_				
Starch B.P.	_				
Stramonium Tincture B.P.	-				
Sulphur, Precipitated B.P.					
Sulphurated Potash B.P.C. 1973	****				
Syrup B.P.	man.				
Talc, Purified B.P., sterilised	_				
Testosterone B.P.	May only be prescribed in creams				
Theophylline B.P.	—				
Thymol B.P.	_				
Thymol Mouth Wash, Compound A.P.F.	_				
mymor mouth mash, Compound A.F.F.					

Column 1	Column 2				
Pharmaceutical benefit	Circumstances (if any) specified for the purposes of section 88A of the Act				
Titanium Dioxide B.P.	***				
Tolu Syrup B.P.	_				
Tragacanth B.P.					
Tragacanth Powder, Compound B.P.	_				
Trichloroacetic Acid B.P.					
Triethanolamine B.P.	_				
Turpentine Liniment B.P.	_				
Water for Injections B.P.	May only be prescribed in eye drops and eye lotions				
Water, Purified B.P.	_				
Wool Alcohols B.P.	_				
Wool Fat B.P.	_				
Wool Fat, Hydrous B.P.	_				
Zinc Oxide B.P.	_				
Zinc Sulphate B.P.	_				

Additives

Acetone B.P.

Anise Water, Concentrated B.P.

Arachis Oil B.P.

Borax B.P.

Boric Acid B.P.

Castor Oil B.P.

Chlorhexidine Acetate B.P.

Chlorhexidine Gluconate Solution B.P.

Chloroform B.P.

Ethanol (96 per cent) B.P.

Ethanols, Dilute B.P.

Ether, Solvent B.P.

Eucalyptus Oil B.P.

Glucose for Oral Use B.P.

Honey, Purified B.P.

Industrial Methylated Spirit B.P.

Olive Oil B.P. 1973

Peppermint Oil B.P.

Peppermint Water, Concentrated A.P.F.

Pholcodine Citrate Syrup B.P.C. 1959

Sodium Thiosulphate B.P.

Turpentine Oil B.P.

Additional pharmaceutical benefits made available under arrangements provided for by section 100 of the Act

Anti-Haemophilic Factor of Animal Origin

Carmellose Sodium B.P.

Methadone Hydrochloride B.P.

Somatrem

Dated this 16th day of November 1988.

JOHN DEEBLE
Acting First Assistant Secretary
Health Benefits Division

Delegate of the Minister of State for Community Services and Health

N.N.-8852514

National Health Act 1953

PHARMACEUTICAL BENEFITS

DETERMINATION UNDER SECTIONS 85, 85A AND 88

- I, JOHN STEWART DEEBLE, Acting First Assistant Secretary, Health Benefits Division, Department of Community Services and Health and Delegate of the Minister of State for Community Services and Health, pursuant to sections 85, 85A and 88 of the National Health Act 1953, hereby make the following Determination:
- 1. This Determination shall come into operation on 1 December 1988.
- 2. The Determination under sections 85, 85A and 88 of the Act made with effect from 1 August 1988, as amended with effect from 1 October 1988, is hereby revoked.
 - 3. In the Schedules to this Determination:
 - 'amp.' means ampoule;
 - 'fl oz' means fluid ounce of 480 minims;
 - 'g' means gram;

 - 'I.M.' means intramuscular;
 'I.U.' means international unit;
- '1.V.' means intravenous;
- 'kg' means kilogram;
 'L' means litre;
- 'lb' means pound of 16 ounces;
- 'm' means metre;
- 'mg' means milligram;
- 'mL' means millilitre:
- 'mm' means millimetre;
- 'mmol' means millimole;
- 'Sch. 2' means the Second Schedule to this Determination.
- 4. Where the strength, type of unit, size of unit or other particulars of form are specified in the column headed 'Form' in the First Schedule to this Determination in relation to a drug or medicinal preparation referred to in subsection 85 (2) of the National Health Act 1953 the name of which is specified in that Schedule, those particulars refer to the form or forms of the drug or medicinal preparation that is or are allowable for the purposes of Part VII of that Act to the effect that the drug or medicinal preparation in that form or in each of those forms is a pharmaceutical benefit when prescribed by a medical practitioner.
- 5. Where the strength, type of unit, size of unit or other particulars of form are specified in the column headed 'Form' in the Third Schedule to this Determination in relation to a drug or medicinal preparation referred to in sub-section 85 (2) of the National Health Act 1953 the name of which is specified in that Schedule, those particulars refer to the form or forms of the drug or medicinal preparation that is or are allowable for the purposes of Part VII of that Act to the effect that the drug or medicinal preparation in that form or in each of those forms is a pharmaceutical benefit referred to in sub-section 88 (1A) of that Act for the supply of which a participating dental practitioner is authorized to write a prescription.
- 6. The purposes set out in the column headed 'Purposes' in Part 2 of the First Schedule to this Determination are the only purposes for which a medical practitioner may prescribe the maximum quantities and number of repeats specified in that Part in relation to those pharmaceutical benefits specified in that same Part.
- 7. The manner, if any, in which a pharmaceutical benefit specified in the First and Third Schedules to this Determination is to be administered in relation to the pharmaceutical benefit is set out in those Schedules in the column headed 'Manner of administration'.
- 8. The maximum quantity or number of units of a pharmaceutical benefit that may, in one prescription, be directed to be supplied on any one occasion is:

- (a) where the name of the pharmaceutical benefit is specified-
 - (i) in Part 1 of the First Schedule to this Determination—the quantity or number, if any, specified in that Part of the Schedule in the column headed 'Maximum quantity' in relation to the pharmaceutical benefit; or
 - (ii) in Part 2 of the First Schedule to this Determination and the pharmaceutical benefit is prescribed in accordance with the provisions of the column headed 'Purposes'-the quantity or number, if any, in that Part of the Schedule in the column headed 'Maximum quantity' in relation to the pharmaceutical benefit; or
 - (iii) if reference is made to the Second Schedule to this Determination in the column headed 'Maximum quantity' in the First Schedule to this Determination, the quantity or number, if any, specified in the column headed 'Maximum quantity' in the Second Schedule to this Determination in relation to the form of the pharmaceutical benefit; or
 - (iv) in the Third Schedule to this Determinationthe quantity or number, if any, specified in that Part of the Schedule in the column headed 'Maximum quantity' in relation to the pharmaceutical benefit; or
- (b) in any other case—the quantity or number, if any, specified in the column headed 'Maximum quantity' in the Second Schedule to this Determination in relation to the form of the pharmaceutical benefit.
- 9. The maximum number of occasions, if any, on which the supply of a pharmaceutical benefit may, in one prescription, be directed by a medical practitioner to be repeated
 - (a) where the name of the pharmaceutical benefit is specified-
 - (i) in Part 1 of the First Schedule to this Determination-the quantity or number, if any, specified in that Part of the Schedule in the column headed 'Number of repeats' in relation to the pharmaceutical benefit; or
 - (ii) in Part 2 of the First Schedule to this Determination and the pharmaceutical benefit is prescribed in accordance with the provisions of the column headed 'Purposes'—the quantity or number, if any, in that Part of the Schedule in the column headed 'Number of repeats' in relation to the pharmaceutical benefit; or
 - (iii) if reference is made to the Second Schedule to this Determination in the column headed 'Number of repeats' in the First Schedule to this Determination, the quantity or number, if any, specified in the column headed 'Number of repeats' in the Second Schedule to this Determination in relation to the form of the pharmaceutical benefit; or
 - (b) in any other case—the number, if any, specified in the column headed 'Number of repeats' in the Second Schedule to this Determination in relation to the form of the pharmaceutical benefit.
- 10. The following purposes are specified in relation to each pharmaceutical benefit the name of which is specified in Part 2 of the First Schedule to this Determination:
 - (a) Where a class of persons is specified in the column headed 'Purposes'-that the pharmaceutical benefit is to be supplied for the treatment of a person included in that class of persons;
 - (b) Where a disease or condition is specified in the column headed 'Purposes'-
 - (i) if sub-sub-paragraph (ii) does not apply—that the pharmaceutical benefit is to be supplied for the treatment of that disease or condition in relation to any person; or

- (ii) if the disease or condition is specified in relation to a specified class of persons—that the pharmaceutical benefit is to be supplied for the treatment of that disease or condition in a person included in that class of persons;
- (c) Where a purpose is specified in the column headed 'Purposes'—that the pharmaceutical benefit is to be supplied for that purpose;
- (d) Where it is specified in the column headed 'Purposes' that compliance with authority procedures set out in sub-paragraph 10 (d) is required—
 - (i) that an application for the written authority of the Secretary in relation to the supply of the pharmaceutical benefit has been made by a medical practitioner, on a form approved by the Secretary, and has been forwarded to the Secretary by that medical practitioner or on behalf of that medical practitioner by a person other than an approved pharmacist, and the Secretary has approved the application; or
 - (ii) where the medical practitioner believes that the supply of the pharmaceutical benefit is required urgently—that the medical practitioner has prepared an Authority Prescription on a form approved by the Secretary and has:
 - (A) issued the original and duplicate of the form to the person for whom the pharmaceutical benefit is to be supplied;
 - (B) on the departmental copy of the form declared that the pharmaceutical benefit is

- prescribed in accordance with the provisions of the column headed 'Purposes' and signed that copy in the medical practitioner's handwriting, and
- (C) forwarded the departmental copy to reach the Secretary within seven days from the date on which the original and duplicate were issued to the person for whom the pharmaceutical benefit is to be supplied.
- 11. Where a medical practitioner makes an application under sub-paragraph 10 (d) and the Secretary approves the application, the Secretary shall record the approval on a numbered authority and—
 - (a) where, in the approval, the Secretary requires the medical practitioner to vary the prescription for the supply of the pharmaceutical benefit—return the approval to the medical practitioner; or
 - (b) in any other case—return the approval to the medical practitioner or, if the medical practitioner so requests, forward the approval to the person to whom the pharmaceutical benefit is to be supplied.
- 12. The name of the manufacturer or the names of manufacturers denoted in accordance with the following table by letters appearing in the column headed 'Brand' in the First and Third Schedules to this Determination in relation to a drug or medicinal preparation the name of which is specified in those Schedules is the brand or brands under which the drug or medicinal preparation may be supplied under Part VII of the National Health Act 1953 as a pharmaceutical benefit:

<i>NO</i> . C	914 44, 25 INOVERIOEI 1900		Government departments 2000	
Letters Manufacturer's Name		Letters Manufacturer's Name		
AB	Abbott Australasia Pty Ltd	LY	Eli Lilly (Australia) and Company	
AD	Amrad Pharmaceuticals Pty Ltd	MB	May & Baker Pharmaceuticals	
AF	Alphapharm Pty Ltd	MG	J. McGloin Pty Ltd	
AG	Allergan Pharmaceuticals Pty Ltd	MJ	Mead Johnson	
AM	Ames Company, Division of Miles Laboratories	MK	Merck Sharp & Dohme (Australia) Pty Ltd	
7	Australia Pty Ltd	ML	Merrell Dow Pharmaceuticals Australia Pty Ltd	
ΑP	Astra Pharmaceuticals Pty Ltd	NN	Nelson Laboratories (Sales) Pty Ltd	
AQ	Alcon Laboratories (Australia) Pty Ltd	NR	Nordia, Denmark	
ΑŶ	Ayerst Laboratories Pty Ltd	NS	Nicholas Kiwi Pty Ltd	
BC	Bristol Laboratories Pty Ltd	NT	Nestlé Australia Ltd	
BF	Barnes-Hind Pty Limited	NW	Norwich Eaton Pharmaceuticals Pty Limited	
BH	Biopharm Pty Ltd	OL	Owen Laboratories	
BL	David Bull Laboratories Proprietary Limited	OR PD	Organon (Australia) Pty Ltd	
BN	Bayer Pharmaceutical Company	PF	Parke Davis Pty Ltd	
BO	Boehringer Mannheim, GmbH, Germany	PS	Pfizer Pty Ltd	
BR	Beecham Research Laboratories	PT	Pharmacia (Australia) Pty Ltd CP Protea	
BT	The Boots Company (Australia) Pty Ltd			
BW.	Wellcome Australia Limited	QE RC	Queensland Ethicals	
BX	Baxter Healthcare Pty Ltd	RG	Reckitts Pty Ltd	
BY	Boehringer Ingelheim Pty Ltd	RK	Rorer Australia Pty Ltd	
BZ	Boucher & Muir Pty Ltd	RL	Riker Laboratories Australia Pty Ltd	
CG	Ciba-Geigy Australia Limited	RO	Roussel Pharmaceuticals Pty Ltd	
CL	Cilag Pty Limited	RS	Roche Products Pty Ltd	
CN	CSL-Novo Pty Limited	RT	A. H. Robins Pty Limited	
CS	Commonwealth Serum Laboratories	SA	Rocke Tompsitt & Co. Ltd Sayco Pty Ltd	
DH	Drug Houses of Australia Pty Ltd	SB	Scientific Hospital Supplies (Australia) Pty Ltd	
DL	Dista Products (Australia) & Company	SC	Schering Pty Ltd, Australian subsidiary of Schering	
DY	Denyer Bros Pty Ltd	50	A. G., Berlin	
EG	Eagle Pharmaceuticals Pty Ltd	SD	Syntex Pharmaceuticals Ltd	
FA	F. H. Faulding and Co. Ltd	SE	Servier Laboratories (Aust.) Pty Ltd	
FC	Fisons Pty Ltd	SH	Essex Laboratories Pty Ltd, Australian subsidiary of	
FE	Farmitalia Carlo Erba		Schering Corporation, U.S.A.	
FM	Fawns and McAllan Pty Ltd	SI	Sigma Co. Ltd	
FR	Charles E. Frosst, Division of Merck Sharp &	SJ	Sharpe Laboratories Pty Ltd	
ГK		SK	Smith Kline & French Laboratories (Aust.) Ltd	
GL	Dohme (Australia) Pty Ltd Glaxo	SN	Smith & Nephew (Aust.) Pty Ltd	
GP	G.P. Laboratories	SQ	E. R. Squibb & Sons Pty Ltd	
HA		SR	Searle Laboratories	
НО	Hamilton Laboratories Pty Ltd	ST	A. E. Stansen & Co. Pty Ltd	
HP	Hollister Incorporated, U.S.A.	SU	Sauter Laboratories (Aust.) Pty Ltd	
	Hoechst Australia Ltd	SV	Stafford-Miller Limited	
IC	ICI Australia Operations Pty Ltd	SZ	Sandoz Australia Pty Ltd	
IQ	The Ioquin Company	TO	R. D. Toppin and Sons Pty Ltd	
ìC	Janssen-Cilag Pty Ltd	UP	Upjohn Pty Ltd	
JP	Janssen Pharmaceutica Pty Ltd	US	USV Pharmaceuticals, a Division of Rorer Australia	
KM	Kendall McGaw Laboratories, Division of Kendall Australia Pty Ltd	UW	Pty Ltd United Works of Pharmaceutical & Dietetic Prod	
KN	Knoll A. G., Germany		ucts, Hungary	
KY	Key Pharmaceuticals Pty Ltd	WH	H. W. Woods Pty Ltd	
LA	L.A. Chemicals Pty Ltd	WL	Winthrop Laboratories	
LE	Lederle Laboratories Division, Cyanamid Australia	ww	Wm R. Warner & Co. Pty Ltd	
	Pty Ltd	WY	Wyeth Pharmaceuticals Pty Ltd	
LH	Lipha Pharmaceuticals, London, U.K.	ZY	Zyma Pharmaceuticals	

FIRST SCHEDULE—PART 1

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Acetazolamide B.P.	Capsule, 500 mg (sustained release)	Oral	50	3	LE
Acetazolamide Sodium with any de- termined brand of Water for In- jections or other solvent	Injection, 500 mg vial (with required solvent)	Injection	1	••	LE
Acetazolamide Tablets B.P.	250 mg	Oral	100	3	LE
Acetylcysteine B.P.	Solution, 200 mg per mL,	Inhalation	5	3	AP
Acrylic Resin	10 mL Solution, 125 g aerosol spray pack	Application	1	••	AP
Acyclovir	Eye ointment, 30 mg per g, 4.5 g	Application to the eye	1	• •	BW
	Tablet, 200 mg	Oral	50		BW
	Tablet, 400 mg	Oral	70		BW
Adrenaline B.P.	Eye drops, 5 mg per mL, 7.5 mL	Application to the eye	2	3	BF
	Eye drops, 10 mg per mL, 7.5 mL	Application to the eye	2	3	BF
	Extemporaneously prepared eye drops	Application to the eye	Sch. 2	Sch. 2	
Adrenaline Hydrochloride	Eye drops, 5 mg (base) per mL, 10 mL	Application to the eye	2	3	AG
	Eye drops, 10 mg (base) per mL, 10 mL	Application to the eye	2	3	AG, AQ
	Eye drops, 20 mg (base) per mL, 10 mL	Application to the eye	2	3	AG, AQ
Adrenaline Injection B.P.	Adrenaline, 1 in 1,000, 1 mL	Injection	5	1	AP, SI
"Albumaid XP"	Powder, 200 g	Oral	10	5	SB
"Albumaid XPXT"	Powder, 200 g	Oral	10	5	SB
"Alfaré"	Powder, 400 g	Oral	2	20	NT
Allopurinol B.P.	Capsule, 100 mg	Oral	200	2	FM
	Capsule, 300 mg	Oral	60	2	FM
Allopurinol Tablets B.P.	100 mg 300 mg	Oral Oral	200 60	2 2	BW, DH, PT BW, DH, PT
Alprenolol Tablets B.P.	100 mg	Oral	100	5	AP, BR
Aluminium Hydroxide and Magne- sium Carbonate Co-dried Gel	Tablet, 375 mg	Orai	200	5	GL, WY
Aluminium Hydroxide, Dried B.P.	Tablet, 300 mg	Oral	200	5	WY
Aluminium Hydroxide, Dried B.P. with Magnesium Hydroxide B.P.	Tablet, 200 mg-200 mg Tablet, dispersible, 230 mg- 230 mg	Oral Oral	200 200	5 5	PD, WW BW
Aluminium Hydroxide, Dried B.P. with Magnesium Trisilicate B.P.	Suspension, 250 mg-120 mg- 120 mg per 5 mL, 500 mL	Oral	2	5	FM
and Magnesium Hydroxide B.P.	Tablet, 250 mg-120 mg-120 mg	Oral	200	5	FM
Aluminium Hydroxide Mixture B.P.	Suspension, 320 mg per 5 mL, 500 mL	Oral	2	5	DH, WY
Aluminium Hydroxide Mixture B.P. with Light Kaolin B.P. or Light Kaolin (Natural) B.P.	Suspension, 137 mg-1 g per 5 mL, 500 mL	Oral	1	2	WY
Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P.	Suspension, 200 mg-200 mg per 5 mL, 500 mL	Oral	2	5	DH, PD, SC, WW
	Suspension, 215 mg-80 mg per 5 mL, 500 mL	Oral	2	5	WY
	Suspension, 300 mg-100 mg per 5 mL, 500 mL	Oral	2	5	AY
Aluminium Hydroxide Mixture B.P. with Magnesium Hydroxide B.P. and Oxethazaine	Suspension, 306 mg-97.5 mg- 10 mg per 5 mL, 500 mL	Oral	2	5	AY
Amantadine Hydrochloride	Capsule, 100 mg	Oral	100	5	BT, CG
Ambenonium Chloride	Tablet, 10 mg	Oral	100	2	WL
Amiloride Hydrochloride Tablets B.P.	5 mg	Oral	100	1	AF, MK

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Aminacrine Hydrochloride B.P. 1968	Eye drops, 3 mg in 15 mL	Application to the eye	1	2	SI
Aminoglutethimide	Tablet, 250 mg	Oral	100	5	CG
"Aminogran Food Supplement"	Powder, 500 g	Oral	1	5	GL
"Aminogran Mineral Mixture"	Powder, 250 g	Oral	1	5	GL
Aminophylline B.P.	Mixture, 105 mg per 5 mL, 500 mL	Oral	1	5	FC
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Aminophylline Injection B.P.	Ampoule, 250 mg in 10 mL	I.V. injection	5	• •	BT, SI
Aminophylline Tablets B.P.	100 mg	Oral	100	5	HA
Amiodarone Hydrochloride	Tablet, 100 mg	Oral	30	5	RC
	Tablet, 200 mg	Oral	30	5	RC
Amitriptyline Tablets B.P.	10 mg	Oral	50	2	AF, MK, PT, RO
	25 mg	Oral	50	2	AF, MK, NS, PT, RO
	50 mg	Oral	50	2	AF
Ammonium Chloride B.P.	Tablet, 500 mg	Oral	100	3	FM
Amoxycillin Capsules B.P.	250 mg (base)	Oral	20	1	AF, BR, CS, PT, SI
	500 mg (base)	Oral	20	1	AF, BR, CS, PT, SI
Amoxycillin Sodium	Injection, 250 mg (base) vial with 3 mL solvent	Injection	5	1	PT
	Injection, 500 mg (base) vial with 3 mL solvent	Injection	5	1	PT
	Injection, l g (base) vial with 4 mL solvent	Injection	5	1	PT
Amoxycillin Trihydrate B.P.	Tablet, chewable, 250 mg (base)	Oral	20	1	BR, CS
	Tablet, dispersible, 3 g (base)	Oral	1	••	BR, CS
Amoxycillin Trihydrate B.P. with Potassium Clavulanate	Tablet, 250 mg (base)-125 mg (acid)	Oral	15	1	BR
	Tablet, 500 mg (base)-125 mg (acid)	Oral	15	1	BR
Amoxycillin Trihydrate B.P. with Potassium Clavulanate and Puri- fied Water B.P.	Powder for syrup, 125 mg (base)-31.25 mg (acid) per 5 mL, 75 mL	Oral	1	1	BR
	Powder for syrup, 250 mg (base)-62.5 mg (acid) per 5 mL, 75 mL	Oral	1	••	BR
Amoxycillin Trihydrate B.P. with Purified Water B.P.	Powder for paediatric drops, 100 mg (base) per mL, 20 mL	Oral	1	1	BR
	Powder for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	1	BR, CS, PT, SI
	Powder for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	BR, CS, PT, SI
Amphotericin B.P.	Cream, 30 mg per g, 15 g Ointment, 30 mg per g, 15 g	Application Application	1 1	1 1	SQ SQ
Amphotericin B.P. with any deter- mined brand of Water for Injec- tions or other solvent	Injection, 50 mg vial (with required solvent)	Injection	1		SQ
Amphotericin Lozenges B.P.	10 mg	Oral	20	1	SQ
Ampicillin Capsules B.P.	250 mg	Oral	24	1	BR, PT
•	500 mg	Oral	24		BR, PT
Ampicillin Sodium B.P. with any determined brand of Water for In-	Injection, 250 mg (base) vial (with required solvent)	Injection	5	1	CS, PT
jections or other solvent	Injection, 500 mg (base) vial (with required solvent)	Injection	5	1	BR, CS, PT
	Injection, I g (base) vial (with required solvent)	Injection	5	1	CS, PT

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Ampicillin Trihydrate B.P. with Pur- ified Water B.P.	Powder for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	1	BR
	Powder for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	BR
Amylobarbitone Sodium B.P. with any determined brand of Water for Injections or other solvent	Injection, 500 mg amp. (with required solvent)	Injection	2	••	LY
Antazoline Phosphate with Napha- zoline Hydrochloride	Eye drops, 5 mg-500 micro- grams per mL, 15 mL	Application to the eye	1	2	AG
Antazoline Sulphate with Naphazo- line Nitrate B.P.	Eye drops, 5 mg-250 micro- grams per mL, 10 mL	Application to the eye	1	2	ZY
Aspirin B.P.	Tablet, 325 mg (buffered)	Oral	100	1	AP
•	Tablet, 650 mg (enteric coated)	Oral	100	2	SK
	Tablet, 650 mg (sustained release)	Oral	100	2	BT
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Aspirin Tablets B.P.	300 mg	Oral	100	1	SI
Aspirin Tablets, Dispersible B.P.	300 mg	Oral	100	1	RC
Atenolol	Tablet, 50 mg	Oral	30	5	AF, IC
Atropine Eye Ointment B.P.	10 mg per g, 4 g	Application to the eye	1	••	PD
Atropine Sulphate B.P.	Eye drops, 5 mg per mL, 15 mL	Application to the eye	1	2	SI
	Eye drops, 10 mg per mL, 15 mL	Application to the eye	1	2	SI
	Forms specified in Sch. 2	·	Sch. 2	Sch. 2	
Atropine Sulphate Injection B.P.	Ampoule, 600 micrograms in 1 mL	Injection	5	1	AP, BT
	Ampoule, 1.2 mg in 1 mL	Injection	5	1	AP, BT
Atropine Sulphate Tablets B.P.	600 micrograms	Oral	100	2	FM
Auranofin	Tablet, 3 mg	Oral	60	5	SK
Aurothioglucose	Injection, 50 mg per mL, 10 mL	Injection	1	••	SH
Azathioprine Tablets B.P.	50 mg	Oral	100	2	AF, BW
Baclofen Tablets B.P.	10 mg	Oral	100	5	CG
	25 mg	Oral	100	5	CG
"Banish"	Liquid, 15 mL		1		DY
Beclomethasone Dipropionate B.P.	Capsule, 100 micrograms	Inhalation by mouth	100	5	GL
	Spray, metered aerosol, 50 micrograms per dose, 200 doses, 20.4 g	Inhalation by mouth	1	5	GL, SH
	Spray, metered aerosol, 100 micrograms per dose, 200 doses, 20.4 g	Inhalation by mouth	1	5	GL
	Spray, metered aerosol, 250 micrograms per dose, 200 doses, 20.4 g	Inhalation by mouth	1	5	GL
Bendrofluazide Tablets B.P.	2.5 mg 5 mg	Oral Oral	100 100	1	BT BT, PT
Benzathine Penicillin B.P.	Injection, 1.8 g in 4 mL disposable syringe	Injection	1		WY
Benzathine Penicillin B.P. with Pro- caine Penicillin B.P., Benzylpeni- cillin Potassium B.P. and Water for Injections	Injection, 450 mg-300 mg-187 mg vial with 2 mL water for injections	Injection	1		WY
Benzhexol Tablets B.P.	2 mg 5 mg	Oral Oral	200 200	2	LE, PT LE, PT
Benzoin Tincture, Compound B.P.	Spray, aerosol, 3.5 mL per	Application	1		EG.
someoni i moturo, compound B.F.	10 mL, 167 mL	Application	•	••	20
			Sch. 2	Sch. 2	
Benztropine Injection B.P.	Forms specified in Sch. 2 Ampoule, 2 mg in 2 mL	Injection	Sch. 2 5	Sch. 2	MK

	Form	Manner of adminis-	Maxi- mum	Number of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration		repeats	Brand
Benzyl Benzoate Application B.P.	50 g in 200 mL	Application	1	2	MG
Benzyl Benzoate B.P.	Application, 50 g in 200 mL	Application	1	2	MB
Benzylpenicillin Potassium B.P. with any determined brand of Water	Injection, 300 mg vial (with required solvent)	Injection	5	1	CS
for Injections or other solvent	Injection, 600 mg vial (with required solvent)	Injection	5	1	CS
	Injection, 3 g vial (with required solvent)	Injection	10	••	CS
	Injection, 6 g vial (with required solvent)	Injection	10	• •	CS
Betamethasone Acetate with Betamethasone Sodium Phosphate B.P.	Injection, 3 mg-3.9 mg in 1 mL amp.	Injection	5		SH
Betamethasone Dipropionate	Cream, 500 micrograms (base) per g, 15 g	Application	1	1	SH
	Ointment, 500 micrograms (base) per g, 15 g	Application	1	1	SH
	Scalp lotion, 500 micrograms (base) per mL, 30 mL	Application to the scalp	1	1	SH
Betamethasone Tablets B.P.	500 micrograms	Oral	30	4	SH
Betamethasone Valerate B.P.	Gel, 500 micrograms (base) per g, 15 g	Application	1	1	GL
Betamethasone Valerate Cream B.P.	200 micrograms (base) per g, 100 g	Application	2		GL, SH
·	500 micrograms (base) per g, 15 g	Application	1	1	GL, SH
Betamethasone Valerate Ointment B.P.	200 micrograms (base) per g, 100 g	Application	2	••	SH
	500 micrograms (base) per g, 15 g	Application	1	1	GL, SH
Bethanechol Chloride	Injection, 5 mg in 1 mL amp.	Injection	2	••	MK
	Tablet, 10 mg	Oral	100	2	HA, MK
Biperiden Hydrochloride	Tablet, 2 mg	Oral	200	2	KN
Bisacodyl B.P.	Enema, 10 mg in 5 mL, 25	Enema	1	2	PT
Bisacodyl Suppositories B.P.	10 mg, 10		1		BY
Bisacodyl Tablets B.P.	5 mg	Oral	200	1	PT
Bismuth Subcitrate	Tablet, 107.7 mg (as Bi)	Oral	112	2	PD
Bleomycin Sulphate with any deter- mined brand of Water for Injec- tions or other solvent	Injections, 15 mg (base) (with required solvent), 10	Injection	1		ВС
Bromocriptine Mesylate Capsules	5 mg (base)	Oral	60	5	SZ
B.P.	10 mg (base)	Oral	100	5	SZ
Bromocriptine Mesylate Tablets B.P.	• , ,	Oral	30	••	SZ
Bumetanide	Tablet, 1 mg	Oral	100	1	AP
Busulphan Tablets B.P.	2 mg	Oral	100	• •	BW
Butyl Monoester Polymer with Ethanol B.P.	Paste, 60 g	Application	1	••	SQ
Butyl Monoester Polymer with Iso- propyl Alcohol B.P.	Protective dressing aerosol, 120 g	Application	1	••	DY
	Protective dressing solution, 59 mL	Application	1	••	DY
	Protective dressing wipes, 50		1	••	DY, SA
Calciferol Tablets, High-Strength B.P.	250 micrograms	Oral	100	5	GL
Calcitonin (Human)—Synthetic	Injections, 0.5 mg with 2 mL amp. solvent, 5	Injection	3	5	CG
Calcitonin (Pork) B.P.	Injections, 160 I.U. vial with 2 mL vial gelatin solvent, 10	Inj e ction	2	5	RG
Calcitriol	Capsule, 0.25 micrograms	Oral	100	5	RO
Calcium Carbonate B.P.	Tablet, 1.5 g (equivalent to 600 mg calcium)	Oral	120	1	LE

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	Tablet (chewable), 1.25 g (equivalent to 500 mg calcium)	Oral	120	1	RK
Calcium Carbonate B.P. with Calcium Lactate-Gluconate	Tablet, compound efferves- cent, equivalent to 1 g calcium	Oral	60	1	SZ
Calcium Folinate	Injection, 3 mg in 1 mL amp.	Injection	5		BL
	Tablet, 15 mg	Oral	10		LE
Calcium Glubionate	Injection, 1.375 g in 10 mL amp.	Injection	5	1	SZ
Captopril	Tablet, 12.5 mg	Oral	90	5	SQ
	Tablet, 25 mg	Oral	90	5	SQ
	Tablet, 50 mg	Oral	90	5	SQ
Carbachol B.P. 1973	Eye drops, 15 mg per mL, 15 mL	to the eye	2	3	AQ
	Eye drops, 30 mg per mL, 15 mL	Application to the eye	2	3	AQ
Carbamazepine B.P.	Syrup, 100 mg per 5 mL, 300 mL	Oral	1	5	CG
Carbamazepine Tablets B.P.	100 mg	Oral	200	2	CG
	200 mg	Oral	200	2	AF, CG
Carbimazole Tablets B.P.	5 mg	Oral	200	2	NS
Carboplatin	Solution for I.V. injection, 50 mg in 5 mL vial	Injection	2	• •	BL
	Solution for I.V. injection, 150 mg in 15 mL vial	Injection	6	• •	BL
	Solution for I.V. injection, 450 mg in 45 mL vial	Injection	2	••	BL
Carmellose Sodium B.P. with Pectin	Paste, 167 mg-167 mg-167	Application	1		SQ
and Gelatin B.P.	mg per g, 15 g Powder, 333 mg-333 mg-333 mg per g, 15 g	Application	1		SQ
Cefotaxime Sodium with any deter- mined brand of Water for Injec-	Injection, l g (base) (with required solvent)	Injection	5		RL
tions or other solvent	Injection, 2 g (base) (with required solvent)	Injection	5	••	RL
Ceftriaxone Sodium with any deter- mined brand of Water for Injec-	Injection, 250 mg (base) (with required solvent)	Injection	2	••	RO
tions or other solvent	Injection, 500 mg (base) (with required solvent)	Injection	2		RO
	Injection, l g (base) (with required solvent)	Injection	2	••	RO
	Injection, 2 g (base) (with required solvent)	Injection	2	••	RO
Cephalexin B.P. with Purified Water B.P.	Granules for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	1	GL, LY
	Granules for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	GL, LY
Cephalexin Capsules B.P.	250 mg	Oral	20	1	GL, LY
	500 mg	Oral	20	1	GL, LY
Cephalothin Sodium B.P. with any determined brand of Water for	Injection, 1 g (base) (with required solvent)	Injection	5	••	GL, LY
Injections or other solvent	Injection, 2 g (base) (with required solvent)	Injection	1	••	LY
	Injection, 4 g (base) (with required solvent)	Injection	1	• •	LY
Cephazolin Sodium with any deter- mined brand of Water for Injec-	Injection, 500 mg (base) (with required solvent)	Injection	5		LY
tions or other solvent	Injection, 1 g (base) (with required solvent)	Injection	5	••	LY
Charcoal, Activated B.P.	Tablet, 300 mg	Oral	500	1	EG, QE, TO
Chloral Hydrate B.P.	Capsule, 500 mg	Oral	25		SQ
-	Mixture, 250 mg per 5 mL, 200 mL	Oral	1	••	DĤ

		Manner of	Maui	Nember	
	Form	Manner of adminis-	Maxi- mum	Number of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
Chlorambucil Tablets B.P.	2 mg 5 mg	Oral Oral	100 100	2 2	BW BW
Chloramphenicol B.P.	Ear drops (aqueous), 5 mg per mL, 5 mL	Application to the ear	1	2	PD
Chloramphenicol B.P. with Polymyxin B Sulphate B.P.	Eye drops, 5 mg-5,000 units per mL, 10 mL	Application to the eye	1	2	PD
, <u> </u>	Eye ointment, 10 mg-5,000 units per g, 4 g	Application to the eye	1	••	PD
Chloramphenicol Capsules B.P.	250 mg	Oral	16		PD
Chloramphenicol Eye Drops B.P.	5 mg per mL, 10 mL	Application to the eye	1	2	AG, PD, SI
Chloramphenicol Eye Ointment B.P.	10 mg per g, 4 g	Application to the eye	1	••	PD, SI
Chloramphenicol Palmitate Mixture B.P.	125 mg per 5 mL, 100 mL	Oral	1		PD
Chloramphenicol Sodium Succinate B.P. with any determined brand of Water for Injections or other solvent	Injection, I g vial (with required solvent)	Injection	3		PD
Chlorhexidine Gluconate	Solution, 50 mg per mL, 200 mL		1	1	IC
Chlormethiazole Capsules B.P.	192 mg	Oral	50		AP
Chlormethiazole Edisylate B.P.	Injection, 8 mg per mL, 500 mL	Injection	1	••	AP
Chloroquine Phosphate Tablets B.P.	250 mg	Oral	100		PT
Chloroquine Sulphate B.P.	Tablet, 200 mg	Oral	100		MB
Chlorothiazide Tablets B.P.	500 mg	Oral	100	1	FM, FR
Chlorpromazine Elixir B.P.	25 mg per 5 mL, 100 mL	Oral	2	2	MB
Chlorpromazine Injection B.P.	Ampoule, 50 mg in 2 mL	Injection	10		MB
Chlorpromazine Suppositories B.P.	100 mg, 5	·	1	2	MB
Chlorpromazine Tablets B.P.	10 mg	Oral	200		MB, PT
•	25 mg	Oral	200		MB, PT
	50 mg	Oral	200	:•	MB, PT
	100 mg	Oral	100	1	MB, PT
Chlorpropamide Tablets B.P.	250 mg	Oral	100	5	PF
Chlorthalidone Tablets B.P.	25 mg	Oral	100	1	CG
Cholestyramine	Sachets, containing 9 g pow- der (equivalent to 4 g cho- lestyramine), 50	Oral	2	5	AP
Choline Theophyllinate B.P.	Elixir, 50 mg per 5 mL, 500 mL	Oral	1	5	PD
	Syrup, 50 mg per 5 mL, 500 mL	Oral	1	5	ww
	Tablet, 100 mg	Oral Oral	100 100	5	WW
Chorionic Gonadotophin B.P.	Tablet, 200 mg Injection set containing 3 ampoules powder for injection 500 units and 3 ampoules solvent 1 mL	Injection	1	5	PD, WW CS
	Injection set containing 3 ampoules powder for injec- tion 1,000 units and 3 am- poules solvent 1 mL	Injection	1	5	CS
	Injection set containing 3 ampoules powder for injec- tion 2,000 units and 3 am- poules solvent 1 mL	Injection	1	5	CS
	Injection set containing 1 ampoule powder for injec- tion 5,000 units and 1 am- poule solvent 1 mL	Injection	2	••	CS
Cimetidine	Tablet, 200 mg	Oral	120	2	CS, SK
	Tablet, 400 mg	Oral	60	2	CS, SK
C	Tablet, 800 mg	Oral	30	1	CS, SK
Ciprofloxacin Hydrochloride	Tablet, 250 mg (base) Tablet, 500 mg (base)	Oral Oral	14 14		BN BN

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	Tablet, 750 mg (base)	Oral	14		BN
Clindamycin Capsules B.P.	75 mg	Oral	25		UP
	150 mg	Oral	25	• •	UP
Clindamycin Palmitate Hydrochlo- ride with Purified Water B.P.	Granules for syrup, 75 mg (base) per 5 mL, 100 mL	Oral	1	••	UP
Clioquinol B.P.	Cream, 10 mg per g, 30 g Forms specified in Sch. 2	Application	I Sch. 2	Sch. 2	CG
Clofibrate Capsules B.P.	500 mg	Oral	100	5	IC, PT
Clomiphene Tablets B.P.	50 mg, 5	Oral	2	5	ML
Clomipramine Hydrochloride B.P.	Tablet, 25 mg	Oral	50	1	CG
Clonazepam	Injection, 1 mg in 2 mL (set containing solution 1 mg in 1 mL and 1 mL diluent) Paediatric drops, 2.5 mg per	Injection Oral	5	••	RO RO
	mL, 10 mL	0.1	200		no.
	Tablet, 500 micrograms Tablet, 2 mg	Oral Oral	200 200	2 2	RO RO
Clonidine Hydrochloride Tablets	100 micrograms	Oral	100	5	BY
B.P.	150 micrograms	Oral	100	5	BY
Clopamide	Tablet, 5 mg	Oral	100	1	SZ
Clotrimazole B.P.	Cream, 10 mg per g, 20 g Lotion, 10 mg per mL, 20 mL	Application Application	1	1	BN, SH BN, SH
	Pessaries, 100 mg, 6		1		BN, SH
	Pessary, 500 mg		1		BN BN SH
	Vaginal cream, 50 mg per 5 g, 35 g Vaginal cream, 100 mg per		1		BN, SH BN
O	5 g, 20 g		•		
Cloxacillin Capsules B.P.	250 mg (base) 500 mg (base)	Oral Oral	24 24		AF, BR AF, BR
Cloxacillin Sodium B.P. with any determined brand of Water for	Injection, 250 mg (base) (with required solvent)	Injection	5		BR, CS
Injections or other solvent	Injection, 500 mg (base)	Injection	5		BR, CS
	(with required solvent) Injection, l g (base) (with required solvent)	Injection	5	••	BR, CS
Codeine Phosphate B.P. with Aspirin B.P.	Tablet, 30 mg-325 mg Forms specified in Sch. 2	Oral	20 Sch. 2	 Sch. 2	·BW
Codeine Phosphate B.P. with Paracetamol B.P.	Tablet, 30 mg-500 mg	Oral	20		WL
Codeine Phosphate Tablets B.P.	30 mg	Oral	20	••	DH, FA, FM, US
Colchicine Tablets B.P.	500 micrograms	Oral	100	2	DH, PT, RT
Colestipol Hydrochloride	Sachets, 5 g, 120	Oral	1	5	UP
Colistin Sulphate B.P. with Neomycin Sulphate B.P.	Ear drops, 3 mg (base)- 3.3 mg (base) per mL, 10 mL	Application to the ear	1	2	ww
Colistin Sulphomethate Sodium B.P. with any determined brand of Water for Injections or other solvent	Injection, 150 mg (with required solvent)	Injection	5	••	ww
Copper Sulphate B.P.C. 1973	Tablets, diagnostic com- pound, 36		2	3	AM
Cortisone Tablets B.P.	5 mg 25 mg	Oral Oral	50 60	4	PT PT
Co-trimoxazole Mixture, Paediatric B.P.	40 mg-200 mg per 5 mL, 100 mL	Oral	1	1	AF, BW, PT, RO
Co-trimoxazole Tablets B.P.	80 mg-400 mg	Oral	10		AF, BW, PT,
	160 mg-800 mg	Oral	10		RO AF, BW, PT, RO
Cyclopenthiazide Tablets B.P.	500 micrograms	Oral	100	1	CG

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Cyclophosphamide B.P. with any de-	Injection, 100 mg vial (with	Injection	6		FE
termined brand of Water for In- jections or other solvent	required solvent) Injection, 200 mg vial (with	Injection	6		FE
	required solvent) Injection, 500 mg vial (with	Injection	2		FE
	required solvent) Injection, 1 g vial (with required solvent)	Injection	1	••	FE
Cyclophosphamide Tablets B.P.	50 mg	Oral	50	2	FE
Cyproheptadine Tablets B.P.	4 mg	Oral	100	2	FR
Cytarabine B.P.	Injection, 40 mg in 2 mL amp.	Injection	10	1	BT
	Injection, 100 mg in 5 mL amp.	Injection	10	1	BT
	Injection, 500 mg in 25 mL amp.	Injection	2	1	BT
	Injection set containing 100 mg and 5 mL solvent	Injection	10	1	UP
Danazol	Capsule, 100 mg	Oral	100	5	WL
	Capsule, 200 mg	Oral	100	5	WL
Dantrolene Sodium	Capsule, 25 mg	Oral	100	2	NW
	Capsule, 50 mg	Oral	100	2	NW
Debrisoquine Tablets B.P.	10 mg (base)	Oral	100	5	RO
	20 mg (base)	Oral	100	5	RO
"De-Lact Infant"	Powder, 500 g	Oral	2		SJ
Demeclocycline Capsules B.P.	150 mg	Oral	100	3	LE
Desipramine Tablets B.P.	25 mg	Oral	50	2	CG
Desmopressin	Nasal solution, 100 micro- grams per mL, 2.5 mL dropper bottle	Nasal	5	2	FC
Dexamethasone B.P.	Eye drops, 1 mg per mL, 5 mL	Application to the eye	1	2	AQ
Dexamethasone B.P. with Framycetin Sulphate B.P. and Gramicidin	Ear ointment, 500 micro- grams-5 mg-50 micrograms per g, 5 g	Application to the ear	1	2	RL
Dexamethasone Sodium Metasul- phobenzoate with Framycetin Sul- phate B.P. and Gramicidin	Ear drops, 500 micrograms (base)-5 mg-50 micrograms per mL, 8 mL	Application to the car	1	2	RL
Dexamethasone Sodium Phosphate B.P.	Injection, 1 mL amp. con- taining equivalent of 4 mg Dexamethasone Phosphate	Injection	5	••	MK
	Injection, 1 mL vial containing equivalent of 5 mg Dexamethasone Phosphate	Injection	5		OR
	Injection, 2 mL vial contain- ing equivalent of 8 mg Dexamethasone Phosphate	Injection	5	••	MK
	Injection, 5 mL vial contain- ing equivalent of 120 mg Dexamethasone Phosphate	Injection	1		MK
Dexamethasone Tablets B.P.	500 micrograms 4 mg	Oral Oral	30 30	4	OR, PT PT
Dexamphetamine Tablets B.P.	5 mg	Oral	100	5	SI
Dextran 40 Intravenous Infusion B.P. with Glucose B.P.	100 mg per mL-139 mmol per 500 mL, 500 mL	Injection	3	••	PS
Dextran 40 Intravenous Infusion B.P. with Sodium Chloride B.P.	100 mg per mL-77 mmol per 500 mL, 500 mL	Injection	3	••	PS
Dextran 70 Intravenous Infusion B.P. with Glucose B.P.	60 mg per mL-139 mmol per 500 mL, 500 mL	Injection	3	••	PS
Dextran 70 Intravenous Infusion B.P. with Sodium Chloride B.P.	60 mg per mL-77 mmol per 500 mL, 500 mL	Injection	3	••	PS
Diazepam B.P.	Syrup, 2 mg per 5 mL, 100	Oral	1		RO
Diazepam Injection B.P.	mL 10 mg in 2 mL	Injection	5		BL, RO

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Diazepam Tablets B.P.	2 mg	Oral	50		AF, PT, RO,
	5 mg	Oral	50		SU AF, PT, RO, SU
Diazoxide Injection B.P.	300 mg in 20 mL	Injection	1		BL, SH
Dichlorphenamide Tablets B.P.	50 mg	Oral	50	6	MK
Diclofenac Sodium	Tablet, 25 mg (enteric coated)	Oral	50	3	CG
	Tablet, 50 mg (enteric coated)	Oral	50	3	CG
Dienoestrol B.P.	Cream, 500 micrograms per 5 g, 85 g		1	1	JC
Difenoxin Hydrochloride with Atropine Sulphate B.P.	Tablet, 500 micrograms-25 micrograms	Oral	20	••	PT
Diffunisal Tablets B.P.	250 mg	Oral	50	3	MK
"Digestelact"	Powder, 500 g	Oral	2	20	SJ
Digoxin Elixir, Paediatric B.P.	50 micrograms per mL, 100 mL	Oral	1	3	BW
Digoxin Injection B.P.	Ampoule, 500 micrograms in 2 mL	Injection	5	1	BW
Digoxin Tablets B.P.	62.5 micrograms	Oral	200	i	BW
Dihydroergotamine Mesylate B.P.	250 micrograms Injection, 1 mg in 1 mL amp.	Oral Injection	100 5	1	BW SZ
Dihydrotachysterol B.P.	Capsule, 125 micrograms	Oral	100	5	WL
Diloxanide Furoate Tablets B.P.	500 mg	Oral	30		ВТ
Diltiazem Hydrochloride	Tablet, 60 mg	Oral	100	5	IC
Dimercaprol Injection B.P.	Ampoule, 100 mg in 2 mL	Injection	12	••	BT
Dimethicones B.P.	Solution, 100 mg per mL, 180 mL aerosol spray pack	Application	1		EG
Diphenoxylate Hydrochloride B.P. with Atropine Sulphate B.P.	Tablet, 2.5 mg-25 micrograms	Oral	20	• •	SR
Diphtheria and Tetanus Vaccine, Adsorbed B.P.	Injection, 0.5 mL amp.	Injection	3	••	CS
Diphtheria and Tetanus Vaccine, Adsorbed B.P., Diluted	Injection, 0.5 mL amp. For immunization of adults and children over the age of 8 years	Injection	3		CS
Diphtheria Antitoxin B.P.	Injection, 10,000 units amp.	Injection	2	1	CS
Diphtheria, Tetanus and Pertussis Vaccine, Adsorbed B.P.	Injection, 0.5 mL amp.	Injection	3	••	CS, PD
Diphtheria Vaccine, Adsorbed B.P.	Injection, 0.5 mL amp.	Injection	2	1	CS
Diphtheria Vaccine, Adsorbed B.P., Diluted	Injection, 0.5 mL amp. For immunization of adults and children over the age of 8 years	Injection	2	••	CS
Dipivefrine Hydrochloride	Eye drops, 1 mg per mL, 10 mL	Application to the eye	1	6	AG
Disodium Etidronate	Tablet, 200 mg	Oral	60	5	NW
Disopyramide Capsules B.P.	100 mg 150 mg	Oral Oral	100 100	5 5	RL RL
Disopyramide Phosphate Capsules B.P.	100 mg (base) 150 mg (base)	Oral Oral	100 100	5 5	SR SR
Docusate Sodium B.P. with Bisacodyl B.P.	Suppositories, 100 mg-10 mg, 5		2	• •	FM
Domperidone	Tablet, 10 mg	Oral	25		JC
Dothiepin Capsules B.P.	25 mg	Oral	50	2	BT
Dothiepin Hydrochloride B.P.	Tablet, 75 mg	Oral	30	2	BT
Doxepin Capsules B.P.	10 mg (base) 25 mg (base)	Oral Oral	50 50	2 2	AF, PF AF, PF
Doxepin Hydrochloride B.P.	Tablet, 50 mg (base)	Oral	50	2	AF

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Doxorubicin Hydrochloride	Powder for I.V. injection or intravesical administration, 10 mg vial	Injection or intra- vesical admini- stration	4		FE
	Powder for I.V. injection or intravesical administration, 20 mg vial	Injection or intra- vesical admini- stration	4	••	FE
	Powder for I.V. injection or intravesical administration, 50 mg vial	Injection or intra- vesical admini- stration	3		FE
Doxycycline Capsules B.P.	100 mg	Oral	7	1	PT
Doxycycline Hydrochloride B.P.	Capsule, 50 mg (base) (containing enteric coated pellets)	Oral	25	5	FA
	Capsule, 100 mg (base) (containing enteric coated pellets)	Oral	7	1	FA
	Tablet, 50 mg (base)	Oral	25	5	AF, PF
	Tablet, 100 mg (base)	Oral	7	1	AF, PF
Dydrogesterone Tablets B.P.	10 mg	Oral	50	5	1C
Econazole Nitrate B.P.	Lotion, 10 mg per mL, 20 mL	Application	1	1	SK, SQ
Econazole Nitrate Cream B.P.	10 mg per g, 20 g	Application	1	1	SK, SQ
	75 mg per 5 g, 35 g	Application	1	• •	SQ
Econazole Nitrate Pessaries B.P.	150 mg, 3		1	• •	SK, SQ
Ecothiopate Iodide B.P.	Eye drops, 300 micrograms per mL (1.5 mg and 5 mL vial of solvent)	Application to the eye	1	6	AY
	Eye drops, 600 micrograms per mL (3 mg and 5 mL vial of solvent)	Application to the eye	1	6	AY
	Eye drops, 1.25 mg per mL (6.25 mg and 5 mL vial of solvent)	Application to the eye	1	6	AY
	Eye drops, 2.5 mg per mL (12.5 mg and 5 mL vial of solvent)	Application to the eye	1	6	AY
Enalapril Maleate	Tablet, 5 mg	Oral	30	5	AD, MK
	Tablet, 10 mg	Oral	30	5	AD, MK
	Tablet, 20 mg	Oral	30	5	AD, MK
Ergometrine and Oxytocin Injection B.P.	Ampoule, 500 micrograms-5 units in 1 mL	Injection	5	••	SZ
Ergometrine Injection B.P.	Ampoule, 250 micrograms in 1 mL	Injection	5	••	BL
Ergometrine Tablets B.P.	200 micrograms	Oral	24		LY
Ergotamine Tablets B.P.	1 mg	Oral	50	2	WL
Ergotamine Tartrate B.P. with Caf- feine B.P.	Suppositories, compound, 6 Tablet, 1 mg-100 mg	Oral	1 50	2 2	SZ SZ
Erythromycin B.P.	Capsule, 125 mg (containing enteric coated pellets)	Oral	25	1	FA
	Capsule, 175 mg (containing enteric coated pellets)	Oral	25	1	FA
Cauthannusia Fatalata B.B.	Capsule, 250 mg (containing enteric coated pellets)	Oral	25	1	FA, LY
Erythromycin Estolate B.P.	Orops, paediatric, 100 mg (base) per mL, 10 mL Suspension, 125 mg (base)	Oral Oral	1	1	LY LY
Erythromycin Ethyl Succinate B.P.	per 5 mL, 100 mL Injection, 100 mg (base) in 2	I.M.	5		AB
,	mL (case) in 2	injection	-	**	
	Tablet, chewable, 200 mg (base)	Oral	25	1	AB

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	Tablet, 400 mg (base)	Oral	25	1	AB
Erythromycin Ethyl Succinate B.P. with Purified Water B.P.	Granules for suspension, 200 mg (base) per 5 mL, 100 mL	Orai	1	i	AB
Erythromycin Lactobionate	Infusion, I.V., 300 mg (base) Infusion, I.V., 1 g (base)	Injection Injection	5 1		AB AB
Erythromycin Stearate B.P.	Capsule, 250 mg (base)	Oral	25	1	AB
•	Suspension, 125 mg (base) per 5 mL, 100 mL	Oral	1	i	AB
	Suspension, 250 mg (base) per 5 mL, 100 mL	Oral	t	• •	AB
Erythromycin Stearate Tablets B.P.	250 mg (base)	Oral	25	1	AB
Erythromycin Tablets B.P.	250 mg	Oral	25	1	AB, UP
Ethacrynic Acid Tablets B.P.	50 mg	Oral	50	3	MK
Ethinyloestradiol Tablets B.P.	10 micrograms	Oral	200	1	GL
	20 micrograms	Oral	200	1	GL
	50 micrograms	Oral	200	1	GL
Ethosuximide Capsules B.P.	250 mg	Oral	200	2	PD
Ethosuximide Elixir B.P.	250 mg per 5 mL, 250 mL	Oral	1	4	PD
Ethyloestrenol Tablets B.P.	2 mg	Oral	100	3	OR
Ethynodiol Diacetate B.P. with Ethinyloestradiol B.P.	Pack containing 21 tablets, 500 micrograms-50 micro- grams and 7 inert tablets	Oral	4	2	SR
	Pack containing 21 tablets, 1 mg-50 micrograms and 7 inert tablets	Oral	4	2	SR
	Tablets, 500 micrograms- 50 micrograms, 21	Oral	4	2	SR
	Tablets, 1 mg-50 micrograms, 21	Oral	4	2	SR
Etretinate	Capsule, 10 mg	Oral	100	5	RO
	Capsule, 25 mg	Oral	100	5	RO
Famotidine	Tablet, 20 mg	Oral	60	2	MK
	Tablet, 40 mg	Oral	30	1	MK
Felodipine	Tablet, 5 mg Tablet, 10 mg	Oral Oral	60 60	5 5	AP, HP AP, HP
Fenoterol Hydrobromide B.P.	Solution, 1 mg per mL, 20 mL Spray, metered aerosol, 200 micrograms per dose, 300 doses, 21 g		1	5	BY BY
Ferrous Aminoacetosulphate	Syrup, 100 mL	Oral	1	4	NN
Ferrous Gluconate B.P.	Elixir, 300 mg per 5 mL, 100 mL	Oral	1	4	WL
Ferrous Gluconate Tablets B.P.	300 mg	Oral	100	2	WL
Ferrous Sulphate, Dried B.P.	Capsule, 320 mg, equivalent to 96 mg Ferrous Iron (de-	Oral	30	2	SK
	layed release) Tablet, 320 mg, equivalent to 96 mg Ferrous Iron (de-	Oral	30	2	CG
	layed release) Tablet, 350 mg, equivalent to 105 mg Ferrous Iron (sustained release)	Oral	30	2	AB, PT
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Ferrous Sulphate, Dried B.P. with	•	Oral	30	2	SK
Folic Acid B.P.	80 mg Ferrous Iron-300 mi- crograms (delayed release)				
	Tablet, 270 mg, equivalent to 80 mg Ferrous Iron-300 mi- crograms (delayed release)	Orai	30	2	PT
	Tablet, 270 mg, equivalent to 80 mg Ferrous Iron-300 micrograms (sustained release)	Oral	30	2	AB
Flecainide Acetate	Tablet, 100 mg	Oral	60	5	RK
Fluctorolone Acetonide B.P.	Cream, 250 micrograms per g,	Application	1	1	SD
	15 g	• •			

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Flucioxacillin Capsules B.P.	250 mg (base)	Oral	24		AF, BR,
	500 mg (base)	Oral	24		CS AF, BR, CS
Flucioxacillin Magnesium B.P. with Purified Water B.P.	Powder for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	••	BR, CS
Turned Water D.I.	Powder for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	• •	BR, CS
Flucloxacillin Sodium B.P. with any determined brand of Water for In-	Injection, 250 mg (base) (with required solvent)	Injection	5	••	CS
jections or other solvent	Injection, 500 mg (base) (with required solvent)	·	5	••	BR, CS
	Injection, 1 g (base) (with required solvent)	•	5		BR, CS
Flucytosine Tablets B.P.	500 mg	Oral	100	• •	RO
Fludrocortisone Tablets B.P.	100 micrograms	Oral	200	1	SQ
Flumethasone Pivalate with Clioquinol B.P.	Ear drops, 200 micrograms-10 mg per mL, 7.5 mL	Application to the ear	1	••	CG
Fluocortolone Pivalate and Fluocortolone Hexanoate Cream B.P.	1 mg-1 mg per g, 15 g	Application	1	1	SC
Fluocortolone Pivalate and Fluocortolone Hexanoate Ointment B.P.	1 mg-1 mg per g, 15 g Anhydrous base, 1 mg-1 mg per g, 15 g	Application Application	1	1	SC SC
Fluorometholone	Eye drops, 1 mg per mL, 5 mL	Application to the eye	1	6	AG, AQ
Fluorouracil Injection B.P.	Ampoule, 250 mg in 10 mL Ampoule, 500 mg in 10 mL	Injection Injection	10 5	 1	RO FE
Fluoxymesterone Tablets B.P.	5 mg	Oral	100	3	UP
Fluphenazine Decanoate Injection	Ampoule, 12.5 mg in 0.5 mL	Injection	5		so
B.P.	Ampoule, 50 mg in 2 mL	Injection	5		šò
2	Syringe, disposable, 25 mg in 1 mL	Injection	5	•••	sõ
Fluphenazine Tablets B.P.	1 mg	Oral	200		SQ
•	2.5 mg	Oral	200		SQ
	5 mg	Oral	100	1	SQ
Folic Acid B.P.	Injection, 15 mg in 1 mL amp.	Injection	10	••	AB
Folic Acid Tablets B.P.	500 micrograms	Oral	200		DH, PT, SI
	5 mg	Oral	200	1	DH, FM, NN, PT, RT, SI, US
Fosfestrol Sodium B.P.	Injection, 250 mg in 5 mL	Injection	10	:-	BC
	Tablet, 120 mg	Oral	100	2	BC
Framycetin Sulphate B.P.	Eye/ear drops, 5 mg per mL, 8 mL	to the	1	2	RL
	Eye ointment, 5 mg per g, 5 g	eye/ear Application to the eye	1		RL
Frusemide B.P.	Solution, 10 mg per mL, 30 mL	Oral	1	3	HP
Frusemide Injection B.P.	Ampoule, 20 mg in 2 mL	Injection	5		AF, HP
Frusemide Tablets B.P.	20 mg	Oral	100	1	FM, HP
7.1000	40 mg	Oral	100	ī	FM, HP, PT
	500 mg	Oral	50	3	FM, HP
Fusidic Acid Mixture B.P.	50 mg per mL, 90 mL	Oral	1		SK
Gas-gangrene Antitoxin, Mixed B.P.	Injection, 1 amp. containing 10,000 units Perfringens; 5,000 units Septicum; 10,000 units Novyi	Injection	2	1	CS
Gentamicin Eye Drops B.P.	3 mg (base) per mL, 5 mL	Application to the eye	1	2	AG
Gentamicin Injection B.P.	Ampoule, 40 mg (base) in 1 mL	Injection	5	• •	BL, SH
	Ampoule, 60 mg (base) in 1.5 mL	Injection	5	••	BL, SH

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	Ampoule, 80 mg (base) in 2 mL	Injection	5		BL, RL, SH
	Syringe, disposable, 80 mg (base)	Injection	5		SH
Glibenclamide Tablets B.P.	5 mg	Oral	100	5	AF, HP, RC
Gliclazide	Tablet, 80 mg	Oral	100	5	SE
Glipizide	Tablet, 5 mg	Oral	100	5	FE
Glucagon Hydrochloride	Injection, 1 I.U. with diluent	Injection	1	1	CN, LY
Glucose and Ketone Indicator— Urine	100 reagent strips	·	1	2	AM
Glucose, Anhydrous B.P., Sodium Chloride B.P., Potassium Chloride B.P., Citric Acid B.P. and Sodium Citrate B.P.	Powder, 36.354 g-354 mg-903 mg-606 mg-1.781 g per sachet, 6	Oral	i	••	DH
Glucose Indicator—Blood	50 reagent strips (Ames-BG) 50 reagent strips (BM-Test-		2 2	5 5	AM BO
	BG) 50 reagent strips (BM-Test-		2	5	ВО
	Glycemie 20-800)				
Character 112	50 reagent strips (Glucostix)		2	5	AM
Glucose Indicator—Urine	4 m dispenser 100 reagent strips (Clinistix)		1	2 2	LY AM
	100 reagent strips (Diastix)		i	2	AM
Glucose Intravenous Infusion B.P.	278 mmol per L, 1 L	Injection	5	1	AB, BX, KM
	555 mmol per L, 1 L	Injection	5	1	AB, BX
	1,110 mmol per L, 1 L	Injection	2	1	AB
	1,387 mmol per L, 1 L 1,387 mmol per 500 mL, 500 mL	Injection Injection	2	1	AB, BX AB
	Ampoule, 5 g in 10 mL	Injection	5		AP
Glyceryl Trinitrate	Ointment, 20 mg per g, 60 g	Application	1	5	FC
	Transdermal disc, 16 mg	Application	30	2	SR
	Transdermal disc, 32 mg Transdermal pad, 25 mg	Application	30 30	2 2	SR CG
	Transdermal pad, 25 mg	Application Application	30	2	CG
Glyceryl Trinitrate Tablets B.P.	600 micrograms, 100	Oral	1	5	BW
Griscofulvin Tablets B.P.	125 mg	Oral	100	2	GL, IC
	330 mg	Oral	28	2	SH
	500 mg	Oral	28	2	GL, IC
Haloperidol B.P.	Injection, 5 mg in 1 mL amp.	Injection	10		SR
Haloperidol Solution B.P.	2 mg per mL, 15 mL	Oral	1	2	SR
	2 mg per mL, 100 mL	Oral	1	1	SR
Haloperidol Tablets B.P.	1.5 mg	Oral	200		SR
Heparin Injection B.P. (Calcium	5 mg 5,000 I.U. in 0.2 mL	Oral Injection	50 5	5	SR BL, CS,
Salt)	5,000 I.U. in 0.5 mL	Injection	5	5	FC CS
	12,500 I.U. in 0.5 mL	Injection	2	5	BL, CS
	25,000 I.U. in 1 mL	Injection	2	5	BL, CS
Heparin Injection B.P. (Sodium Salt)	Ampoule, 5,000 units in 0.2 mL	Injection	5	5	BL, CS, FC
	Ampoule, 5,000 units in 1 mL	Injection	5	5	BL, CS, FC
	Ampoule, 20,000 units in 20 mL	Injection	12	5	BL SS FS
	Ampoule, 25,000 units in 5 mL	Injection	2	5	CS, FC
**************************************	Vial, 35,000 units in 35 mL	Injection	12	5	CS, FC
Hexamine Hippurate	Tablet, 1 g	Oral	100	5	RK
Hexamine Mandelate	Tablet, 250 mg	Oral	200	2	ww
	Tablet, 500 mg Tablet, 1 g	Oral Oral	200 100	2 5	ww ww

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Homatropine Hydrobromide B.P.	Eye drops, 20 mg per mL, 15 mL	Application to the eye	1	2	AG, AQ
	Eye drops, 50 mg per mL, 15 mL		1	2	AG, AQ
Hydralazine Hydrochloride Tablets B.P.	25 mg	Oral	200	2	AF, CG, PT
	50 mg	Oral	200	2	AF, CG, PT
Hydrochlorothiazide B.P. with Amiloride Hydrochloride B.P.	Tablet, 50 mg-5 mg	Oral	100	1	MK, PT
Hydrochlorothiazide B.P. with Triamterene B.P.	Tablet, 25 mg-50 mg	Oral	100	1	SK
Hydrochlorothiazide Tablets B.P.	25 mg 50 mg	Oral Oral	100 100	1	CG, MK CG, MK
Hydrocortisone B.P.	Tablet, 4 mg Tablet, 20 mg	Oral Oral	50 60	4	PT PT
Hydrocortisone Acetate B.P.	Eye drops, 5 mg per mL, 10	Application	1	2	SI
	mL Eye drops, 10 mg per mL, 10 mL	to the eye Application to the eye	1	2	SI
	Eye drops, 25 mg per mL, 5	Application	1	2	SI
	mL Eye drops, 25 mg per mL, 10 mL	to the eye Application to the eye	1	2	SI
	Eye ointment, 5 mg per g,	Application to the eye	i	••	UP
	Eye ointment, 5 mg per g, 5 g	Application to the eye	1	••	SI
	Eye ointment, 10 mg per g, 5 g	Application to the eye	1	• •	SI
	Rectal foam, 100 mg per g, aerosol, 25 g	to the eye	2	3	sv
Hydrocortisone Acetate B.P. with	Ear drops, 15 mg-3.5 mg	Application to the ear	1	••	UP
Neomycin Sulphate B.P.	(base) per mL, 5 mL Ear ointment, 15 mg-3.5 mg (base) per g, 4 g	Application to the ear	1	2	UP
Hydrocortisone Acetate Cream B.P.	10 mg per g, 30 g	Application	1	1	NN, PD, SI, SQ, UP, US
	10 mg per g, 50 g	Application	1	1	NN, PD, SI, SQ, UP
Hydrocortisone Acetate Injection B.P.	Ampoule, 25 mg in 1 mL	Intra-articu- lar injection	5		PT
Hydrocortisone Acetate Ointment B.P.	10 mg per g, 30 g	Application	1	1	NN, PD, SI, SQ, UP
	10 mg per g, 50 g	Application	1	1	NN, PD, SI, SQ, UP
Hydrocortisone Sodium Succinate B.P.	Injection set containing equivalent of 100 mg Hy- drocortisone and 2 mL solvent	Injection	2	••	NR, UP
	Injection set containing equivalent of 250 mg Hy- drocortisone and 2 mL solvent	Injection	1	••	UP
	Injection set containing equivalent of 500 mg Hy- drocortisone and 4 mL solvent	Injection	2	••	UP
Hydroxocobalamin Injection B.P.	1 mg in 1 mL amp.	Injection	3		GL
Hydroxychloroquine Tablets B.P.	200 mg	Oral	100	1	WL
Hydroxyurea Capsules B.P.	500 mg	Oral	100	••	SQ

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Hyoscyamine Hydrobromide with Atropine Sulphate B.P. and Hyos- cine Hydrobromide B.P.	Tablet, 101.1 micrograms- 14.8 micrograms-10.7 micrograms	Oral	100	2	FM
	Tablet, 151.6 micrograms- 22.2 micrograms-16.0 micrograms	Oral	100	2	FM
Hyoscyamine Sulphate B.P. with Atropine Sulphate B.P. and Hyos- cine Hydrobromide B.P.	Tablet, 103.7 micrograms- 19.4 micrograms-6.5 micrograms	Oral	100	2	RS
Hypromellose 4500 B.P.	Eye drops, 5 mg per mL, 15 mL	Application to the eye	2	3	AG, AQ, SI
	Eye drops, 10 mg per mL, 15 mL		2	3	SI
Hypromellose 4500 B.P. with Dex- tran 70	Eye drops, 3 mg-1 mg per mL, 15 mL	Application to the eye	2	3	AQ
Ibuprofen Tablets B.P.	200 mg 400 mg	Oral Oral	50 50	3	AF, BT, PT BT, PT
Idoxuridine B.P.	Eye ointment, 5 mg per g,	Application	1	••	SK
	5 g Ointment, 5 mg per g, 5 g	to the eye Topical application	1		SK
Idoxuridine Eye Drops B.P.	1 mg per mL, 15 mL	Application to the eye	1	2	AG, SK
Imipramine Hydrochloride B.P.	Injection, 25 mg in 2 mL amp.	Injection	10		CG
Imipramine Tablets B.P.	10 mg	Oral	50	2	CG, PT
	25 mg	Oral	50	2	CG, PT, UW
Indapamide	Tablet, 2.5 mg	Oral	60	I	SE
Indomethacin Capsules B.P.	25 mg	Oral	50	3	AF, MK, PT
Indomethacin Suppositories B.P.	100 mg		40	3	MK
Influenza Vaccine (Split Virion), Inactivated B.P.	Injection containing antigens representative of the following types: A/Singapore/6/86 (H1N1)-like strain 15 micrograms haemagglutinin; A/Leningrad/360/86 (H3N2)-like strain 15 micrograms haemagglutinin; B/Victoria/2/87-like strain 15 micrograms haemagglutinin; 15 micrograms haemagglutinin; 0.5 mL pre-filled syringe	Injection	1		CS, MB
Insect Allergen Extract—Honey Bee Venom	Injection set containing 550 micrograms vial with 9 mL vial solvent and 3 vials di- luent 1.8 mL	Injection	1	••	BN
Insect Allergen Extract—Paper Wasp Venom	Injection set containing 550 micrograms vial with 9 mL vial solvent and 3 vials diluent 1.8 mL	Injection	1	••	BN
Insulin Injection, Acid B.P.	Vial, (bovine) 100 units per mL, 10 mL	Injection	5	2	CN
	Vial, (bovine) 300 units per mL, 5 mL	Injection	10	2	CN
Insulin Injection, Biphasic B.P.	Vial, (mixed porcine/bovine) 100 units per mL, 10 mL	Injection	5	2	CN
Insulin Injection, Isophane B.P.	Vial, (bovine) 100 units per mL, 10 mL	Injection	5	2	CN, FC
	Vial, (porcine) 100 units per mL, 10 mL	Injection	5	2	BW, CN
Insulin Injection, Isophane B.P. and Insulin Injection, Neutral B.P.	Vial, (porcine) 50 units-50 units per mL, 10 mL	Injection	5	2	BW
	Vial, (porcine) 70 units-30 units per mL, 10 mL	Injection	5	2	BW, CN

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Insulin Injection, Neutral B.P.	Cartridges, (porcine) 100	Injection	7	2	CN
	units per mL, 1.5 mL, 5 Cartridges, (porcine) 100	Injection	5	2	BW
	units per mL, 2 mL, 5 Vial, (bovine) 100 units per	Injection	5	2	FC
	mL, 10 mL Vial, (porcine) 100 units per mL, 10 mL	Injection	5	2	BW, CN
Insulin Injection, Protamine Zinc B.P.	Vial, (bovine) 100 units per mL, 10 mL	Injection	5	2	CN
Insulin Isophane, Human (Synthetic)	Injection, 100 units per mL, 10 mL vial	Injection	5	2	BW, CN, LY
	Injection, cartridges, 100 units per mL, 1.5 mL, 5	Injection	7	2	CN
	Injection, cartridges, 100 units per mL, 2.5 mL, 5	Injection	4	2	BW
Insulin Isophane, Human (Syn- thetic) and Insulin Neutral,	Injection, 50 units-50 units per mL, 10 mL vial	Injection	5	2	BW
Human (Synthetic)	Injection, 70 units-30 units per mL, 10 mL vial	Injection	5	2	BW, CN
	Injection, cartridges, 70 units-30 units per mL, 1.5	Injection	7	2	CN
	mL, 5 Injection, cartridges, 70 units-30 units per mL, 2.5 mL, 5	Injection	4	2	BW
Insulin Neutral, Human (Synthetic)	Injection, 100 units per mL, 10 mL vial	Injection	5	2	BW, CN, LY
	Injection, cartridges, 100 units per mL, 1.5 mL, 5	Injection	7	2	CN
	Injection, cartridges, 100 units per mL, 2.5 mL, 5	Injection	4	2	BW
Insulin Zinc Suspension B.P.	Injection, (bovine) 100 units per mL, 10 mL vial	Injection	5	2	CN
	Injection, (porcine) 100 units per mL, 10 mL vial	Injection	5	2	CN
Insulin Zinc Suspension (Amorphous) B.P.	Injection, (porcine) 100 units per mL, 10 mL vial	Injection	5	2	CN
Insulin Zinc Suspension (Crystal- line) B.P.	Injection, (bovine) 100 units per mL, 10 mL vial	Injection	5	2	CN
Insulin Zinc Suspension (Crystal- line), Human (Synthetic)	Injection, 100 units per mL, 10 mL vial	Injection	5	2	CN
Insulin Zinc Suspension, Human (Synthetic)	Injection, 100 units per mL, 10 mL vial	Injection	5	2	CN
Intraperitoneal Dialysis Solutions B.P.C. 1973	1 L (Glucose 76 mmol per L), 12	Injection	2	1	BX
	2 L (Glucose 76 mmol per L), 6	Injection	2	1	BX
	1 L (Glucose 126 mmol per L), 12	Injection	2	1	BX
	2 L (Glucose 126 mmol per L), 6	Injection	2	1	BX
	1 L (Glucose 214 mmol per L), 10	Injection	2	1	BX
	2 L (Glucose 214 mmol per L), 6	Injection	2	1	BX
Ipratropium Bromide	Nebuliser solution, 250 mi- crograms per mL, 20 mL	Inhalation	2	2	BY
	Spray, metered aerosol, 20 micrograms per dose, 200 doses, 10 mL	Inhalation by mouth	1	5	BY
Iron Dextran Injection B.P.	Ampoule, 2 mL	Injection	5		FC
Iron Polymaltose Complex	Injection, 100 mg (Iron) in 2 mL amp.	Injection	5	••	SI
Incompanie Nitrat-	Tablet, 40 mg (Iron)	Oral	100	2	SI
Isoconazole Nitrate	Cream, 10 mg per g, 20 g Lotion, 10 mg per mL, 20 mL	Application Application	1	1	SC SC

	Form	Manner of adminis-	Maxi- mum	Number of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
	Pessaries, 300 mg, 2		1		SC
Isoniazid Tablets B.P.	100 mg	Oral	100	2	FM
Isopropyl Monoester Polymer with	Gel, adhesive protective,	Application	1		НО
Isopropyl Alcohol B.P.	28.35 g				
Isosorbide Dinitrate Tablets B.P.	5 mg (sublingual)	Oral	200	2	AY, PD
	10 mg	Oral	200	2	AY, PD
Isotretinoin	Capsule, 10 mg	Oral	60	3	RO
	Capsule, 20 mg	Oral	60	3	RO
Kaolin, Light B.P. or Light Kaolin (Natural) B.P. with Pectin	Suspension, 5.91 g-132 mg per 30 mL, 375 mL	Oral	1	2	UP
Ketoconazole	Tablet, 200 mg	Oral	30	5	JC
Ketoprofen B.P.	Capsule, 100 mg (sustained	Oral	50	3	MB
	release)				
	Suppository, 100 mg		40	3	MB
Ketoprofen Capsules B.P.	50 mg	Oral	50	3	MB
•	100 mg	Oral	50	3	MB
Labetalol Hydrochloride Tablets	100 mg	Oral	100	5	AF, GL
B.P.	200 mg	Oral	100	5	AF, GL
Lactulose Solution B.P.	3.34 g per 5 mL, 500 mL	Oral	1	5	JC
Lauramine Oxide with Octoxinol	Cleansing solution, com-		1		DY
	pound, 80 mg-10 mg per g, 240 mL				
Leuprorelin Acetate	Injection, 5 mg per mL, 2.8 mL vial	Injection	2	2	AB
Levodopa and Carbidopa Tablets	100 mg-10 mg	Oral	100	5	MK
B.P.	100 mg-25 mg	Oral	100	5	MK
	250 mg-25 mg	Oral	100	5	MK
Levodopa B.P. with Benserazide	Capsule, 50 mg-12.5 mg	Oral	100	5	RO
	Capsule, 100 mg-25 mg	Oral	100	5	RO
	Capsule, 200 mg-50 mg	Oral	100	5	RO
	Tablet, 200 mg-50 mg	Oral	100	5	RO
Levodopa Tablets B.P.	100 mg	Oral	50	5	RO
	250 mg	Oral	100	5	RO
	500 mg	Oral	250	5	RO
Levonorgestrel B.P.	Tablets, 30 micrograms, 28	Oral	4	2	SC, WY
Levonorgestrel B.P. with Ethinyloes- tradiol B.P.	Pack containing 21 tablets, 125 micrograms-50 micro-	Oral	4	2	SC, WY
	grams and 7 inert tablets	Oral	4	2	SC WV
	Pack containing 21 tablets, 150 micrograms-30 micro- grams and 7 inert tablets	Olai	•	۷	SC, WY
	Pack containing 21 tablets,	Oral	4	2	WY
	250 micrograms-50 micro-			_	
	grams and 7 inert tablets				
	Pack containing 11 tablets,	Oral	4	2	WY
	50 micrograms-50 micro-				
	grams and 10 tablets, 125				
	micrograms-50 micrograms	Orol	4	2	CC WY
	Pack containing 11 tablets,	Oral	4	2	SC, WY
	50 micrograms-50 micrograms, 10 tablets, 125 mi-				
	crograms-50 micrograms				
	and 7 inert tablets				
	Pack containing 6 tablets, 50	Oral	4	2	SC, WY
	micrograms-30 micrograms,				
	5 tablets, 75 micrograms-				
	40 micrograms and 10 tab-				
	lets, 125 micrograms-30				
	micrograms	Orol	4	2	SC WV
	Pack containing 6 tablets, 50	Oral	4	2	SC, WY
	micrograms-30 micrograms, 5 tablets, 75 micrograms-				
	40 micrograms, 10 tablets,				
	125 micrograms-30 micrograms and 7 inert tablets				
	125 micrograms-30 micro-	Oral	4	2	SC

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	Tablets, 150 micrograms-30 micrograms, 21	Oral	4	2	SC, WY
	Tablets, 250 micrograms-50 micrograms, 21	Oral	4	2	WY
Lignocaine Hydrochloride B.P.	Injection, 100 mg in 5 mL Forms specified in Sch. 2	Injection	2 Sch. 2	Sch. 2	AP
Lignocaine Hydrochloride Injection B.P.	Syringe, disposable, 300 mg in 3 mL	I.M. injection	1	••	AP
	500 mg in 5 mL	Infusion	5	• •	AP
Lincomycin Injection B.P.	Ampoule, 300 mg (base) in 1 mL Ampoule, 600 mg (base) in	Injection Injection	5	••	UP UP
L'alam D.D.	2 mL			••	
Lindane B.P.	Head lotion, 2 mg per mL, 200 mL Lotion concentrate, 10 mg	Application Application	1		IC IC
	per mL, 200 mL				
Lindane Cream B.P.	10 mg per g, 200 g	Application	1	2	IC
Liothyronine Tablets B.P.	20 micrograms	Oral	100	2	GL
Lithium Carbonate B.P.	Tablet, 400 mg (delayed release)	Oral	200	1	PT
Lithium Carbonate Tablets B.P.	250 mg	Oral	200	2	DH, PT
"Locasol New Formula"	Powder, 450 g	Oral	2	20	KY
"Lofenalac"	Powder, 1 lb	Oral	5	5	MJ
Loperamide Hydrochloride	Capsule, 2 mg	Oral	12		1C
Lypressin	Spray, nasal, 50 units per mL, 5 mL	Nasal spray	10	2	SZ
"Maxamaid RVHB"	Powder, 200 g	Oral	5	5	SB
"Maxamaid XP"	Powder, 200 g	Oral	5	5	SB
"Maxamum XP"	Powder, 200 g	Oral	5	5	SB
Medroxyprogesterone Acetate B.P.	Injection, 50 mg in 1 mL vial Injection, 150 mg in 1 mL vial	Injection Injection	1 1		UP UP
	Injection, 500 mg in 2.5 mL vial	Injection	1	••	FE
	Oral suspension, 100 mg per mL, 100 mL	Oral	1	2	UP
	Tablet, 100 mg	Oral Oral	30 100	5 2	UP FE, UP
	Tablet, 100 mg Tablet, 200 mg	Oral	60	2	FE, UP
	Tablet, 250 mg	Oral	60	2	UP
	Tablet, 500 mg	Oral	30	2	FE, UP
Medrysone	Eye drops, 10 mg per mL, 5 mL	Application to the eye	1	6	AG
Mefenamic Acid Capsules B.P.	250 mg	Oral	50	2	AF, PD
Mefruside	Tablet, 25 mg	Oral	100	1	BN
Megestrol Acetate B.P.	Tablet, 40 mg	Oral	100	2	BC
Melphalan Tablets B.P.	2 mg 5 mg	Oral Oral	100 100		BW BW
Menopausal Gonadotrophin, Human	Injection set containing 10 ampoules powder for injection providing 75 units follicle stimulating hormone and 75 units luteinizing hormone and 10 ampoules solvent 1 mL	Injection	1	5	CS, OR
Mercaptopurine Tablets B.P.	50 mg	Orai	100	2	BW
Metformin Tablets B.P.	500 mg	Oral	100	5	AF, FC, LH
Methacycline Hydrochloride B.P.	Capsule, 150 mg	Oral	21	1	ww
1973	Capsule, 300 mg	Oral	10	1	WW
Methadone Injection B.P.	Ampoule, 10 mg in 1 mL	Injection	5	• •	BW
Methadone Tablets B.P.	5 mg 10 mg	Oral Oral	20 20		BW BW

					
Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Methdilazine Hydrochloride	Tablet, 4 mg	Oral	100	2	GL
Methenolone Acetate	Tablet, 8 mg Tablet, 5 mg	Oral Oral	100 100	2	GL SC
Methotrexate B.P. with any deter-	Injection, 50 mg vial (with		5		LE
mined brand of Water for Injec- tions or other solvent	required solvent)	Injection	3	••	LE
Methotrexate Injection B.P.	Vial, 5 mg in 2 mL	Injection	5		FE, LE
	Vial, 50 mg in 2 mL	Injection	5	••	BL, FE, LE
Methotrexate Tablets B.P.	2.5 mg	Oral	100	2	FE, LE
Mathaurimida	10 mg	Oral	100	2	FE
Methyalothianida	Capsule, 300 mg	Oral	200	2	PD
Methyclothiazide	Tablet, 2.5 mg Tablet, 5 mg	Oral Oral	100 100	1	AB AB
Methyldopa Tablets B.P.	125 mg	Oral	100	5	MK
Weenyloopa Tablets D.I.	250 mg	Oral	100	5	AF, MK, US
Methylphenobarbitone B.P.	Tablet, 60 mg	Oral	200	2	WL
• •	Tablet, 200 mg	Oral	200	2	WL
Methylprednisolone Acetate Injection B.P.	Vial, 40 mg in 1 mL	Injection	5	••	UP
Methylprednisolone Sodium Succinate	Injection, 40 mg (base) in 1 mL amp.	Injection	5		UP
Methyl Salicylate Liniment A.P.F.	100 mL	Application	1	1	DH, MG, NN, QE,
	Forms specified in Sch. 2		Sch. 2	Sch. 2	SI, TO
Methyltestosterone Tablets B.P.	5 mg	Oral	100	2	PT
Methylestosterone rablets b.r.	25 mg	Oral	100	2	PT
	50 mg	Oral	100	2	PT
Methysergide Tablets B.P.	l mg (base)	Oral	100	2	SZ
Metoclopramide Hydrochloride B.P.	Syrup, 5 mg per 5 mL, 100 mL	Oral	1	• •	BR
Metoclopramide Injection B.P.	Ampoule, 10 mg in 2 mL	Injection	10		BR
Metoclopramide Tablets B.P.	10 mg	Oral	25		AF, BR, PT
Metolazone	Tablet, 2.5 mg	Oral	100	1	SR
Metoprolol Tartrate	Tablet, 50 mg	Oral	100	5	AP, CG
	Tablet, 100 mg	Oral	60	5	AP, CG
Metronidazole Benzoate	Suspension, 320 mg per 5 mL, 100 mL	Oral	1	••	MB
Metronidazole B.P.	Intravenous infusion, 500 mg in 100 mL	Injection	1	••	BL, MB
Metronidazole Suppositories B.P.	500 mg, 10		ı	• •	MB, PT
	1 g, 10		1	• •	MB
Metronidazole Tablets B.P.	200 mg	Oral	21	1	AF, MB, PT, SR
	250 mg 400 mg	Oral Oral	21 5	1	JC AF, MB
	500 mg	Oral	4	i	JC
Mexiletine Hydrochloride Capsules	50 mg	Oral	100	5	BY
B.P.	200 mg	Oral	100	5	BY
Mianserin Hydrochloride Tablets B.P.	10 mg 20 mg	Oral Oral	50 50	5 5	OR OR
Miconazole	Oral gel, 20 mg per mL, 20 g	Oral	1	1	JC
	Tincture, 20 mg per mL, 20 mL	Application	1	i	JC
Miconazole Nitrate B.P.	Lotion, 20 mg per g, 20 g Pessaries, 100 mg, 7	Application	1 1	t	JC CL, JP
	Pessaries, 200 mg, 3		1	• •	JC
Miconazole Nitrate Cream B.P.	20 mg per g, 20 g	Application	1	ï	CL, JP
	20 mg per g, 40 g	Application	1		CL, JP
Minocycline Hydrochloride	Capsule, 100 mg	Oral	11		LE
Minoxidil	Tablet, 10 mg	Oral	100	5	UP
Missesses	Tablet, 25 mg	Oral	100	5	UP
Misoprostol	Tablet, 200 micrograms	Oral	120	1	SR

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Mithramycin	Injection, 2.5 mg (with required solvent)	Injection	5		PF
Mitozantrone Hydrochloride	Injection, 20 mg (base) in 10 mL vial	Injection	1	• •	LE
	Injection, 25 mg (base) in 12.5 mL vial	Injection	1	••	LE
	Injection, 30 mg (base) in 15 mL vial	Injection	1	i i	LE
Morphine Sulphate B.P. with Tac- rine Hydrochloride	Tablet, 30 mg-15 mg	Oral	20	••	WH
Morphine Sulphate Injection B.P.	Ampoule, 10 mg in 1 mL	Injection	5	••	BL, BT, SI
	Ampoule, 15 mg in 1 mL	Injection	5	••	BL, BT, SI
	Ampoule, 30 mg in 1 mL	Injection	5		BL
Morphine Sulphate Tablets B.P.	30 mg	Oral	20		FM
"M.S.U.D. AID"	Powder, 200 g	Oral	5	5	SB
Mustine Hydrochloride B.P. with any determined brand of Water for Injections or other solvent	Injection, 10 mg amp. (with required solvent)	Injection	4	••	BT
Nalidixic Acid Tablets B.P.	500 mg	Oral	56	2	WL
Naloxone Hydrochloride	Injection, 40 micrograms in 2 mL amp.	Injection	5	••	ВТ
	Injection, 400 micrograms in 1 mL disposable injection set	Injection	1	••	CS
	Injection, 800 micrograms in 2 mL disposable injection set	Injection	1	••	CS
	Injection, 2 mg in 5 mL disposable injection set	Injection	1	••	CS
Nandrolone Decanoate Injection B.P.	Disposable syringe, 50 mg in 1 mL	Injection	1	3	OR
Nandrolone Phenylpropionate Injection B.P.	Ampoule, 25 mg in 1 mL	Injection	3	3	OR
Naphazoline Hydrochloride	Eye drops, 1 mg per mL, 15 mL	Application to the eye	1	2	AG
Naproxen B.P.	Suppository, 500 mg		40	3	SD
Naproxen Tablets B.P.	250 mg 500 mg	Oral Oral	50 50	3	AF, SD SD
Neomycin Sulphate Eye Drops B.P.	5 mg per mL, 10 mL	Application to the eye	1	2	SI
Neomycin Tablets B.P.	500 mg	Oral	25	1	PT
Neomycin Undecenoate with Bacitracin Zinc B.P.	Ear ointment, 12 mg (3.5 mg base)-400 units per g, 10 g	Application to the ear	1	••	НА
Neostigmine Injection B.P.	Ampoule, 500 micrograms in 1 mL	Injection	5	3	RO
	Ampoule, 2.5 mg in 1 mL	Injection	5	3	RO
Neostigmine Tablets B.P.	15 mg	Oral	200	1	RO
Niclosamide Tablets B.P.	500 mg	Oral	4		BN
Nicotinic Acid Tablets B.P.	25 mg	Oral	100	2	DH, US
	50 mg	Oral	100	2	DH, RT, SI, US
	100 mg	Oral	100	2	DH, US
	250 mg	Oral	100	2	DH
Nifedipine	Capsule, 5 mg	Oral	100	5	BN
	Capsule, 10 mg Tablet, 20 mg	Oral Oral	100 60	5	AF, BN BN
Nitrazepam Tablets B.P.	5 mg	Oral	25	• •	AF, RO
Nitrofurantoin B.P.	Capsule, 50 mg	Oral	30	1	NW
	Capsule, 100 mg	Oral	30	1	NW
Nitrofurantoin Mixture B.P.	25 mg per 5 mL, 200 mL	Oral	1	• •	NW
Nitrofurantoin Tablets B.P.	50 mg	Oral	25	1	NW
	100 mg	Oral	25	1	NW

		Manner of	Maxi-	Number	
	Form	adminis-	mum	of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
Norethisterone Acetate B.P.	Tablet, 10 mg	Oral	100	2	SC
Norethisterone B.P. with Ethinyloes-	Pack containing 21 tablets,	Oral	4	2	SD
tradiol B.P.	500 micrograms-35 micro-				
	grams and 7 inert tablets Pack containing 21 tablets,	Oral	4	2	SD
	1 mg-35 micrograms and				
	7 inert tablets Pack containing 12 tablets,	Oral	4	2	SD
	500 micrograms-35 micro-	· · · ·	•	_	32
	grams and 9 tablets, 1 mg- 35 micrograms				
	Pack containing 12 tablets,	Oral	4	2	SD
	500 micrograms-35 micro-				
	grams, 9 tablets, 1 mg-35 micrograms and				
	7 inert tablets		4		
	Tablets, 500 micrograms-35 micrograms, 21	Oral	4	2	JC, SD
	Tablets, 1 mg-35 micrograms,	Oral	4	2	SD
	21			_	
Norethisterone B.P. with Mestranol B.P.	Pack containing 21 tablets, 1 mg-50 micrograms and	Oral	4	2	SD
2	7 inert tablets				
	Tablets, 1 mg-50 micrograms, 21	Oral	4	2	JC, SD
Norethisterone Tablets B.P.	350 micrograms, 28	Oral	4	2	JC, SD
	5 mg	Oral	30	5	SC, SE
Norfloxacin	Tablet, 400 mg	Oral	14	1	MK
Nortriptyline Hydrochloride B.P.	Elixir, 10 mg per 5 mL, 100 mL	Oral	1	4	DL
Nortriptyline Tablets B.P.	10 mg	Oral	50	2	DL, SQ
Whiteman in a 21	25 mg	Oral	50	2	DL, SQ
"Nutramigen" Nystatin B.P.	Powder, 1 kg Capsule, 500,000 units	Oral Oral	1 50	20	MJ
Nystatiii D.F.	Cream, 100,000 units per g,	Application	1	i	LE, SQ LE, SQ
	Gel, 100,000 units per g,	Application	1	1	SQ
	Lozenge, 100,000 units	Oral	20	1	LE
	Tablet, 500,000 units Vaginal cream, 100,000 units	Oral	50 1	 1	LE SQ
	per 4 g, 75 g			-	-
	Vaginal cream, 100,000 units per 5 g, 75 g		I	1	LE
Nystatin Mixture B.P.	100,000 units per mL, 24 mL	Oral	1	1	LE, SQ
Nystatin Ointment B.P.	100,000 units per g, 15 g	Application	1	1	LE, SQ
Nystatin Pessaries B.P.	100,000 units, 15		1	1	LE, SQ
	100,000 units (cream base), 15		1	1	LE
Nystatin Tablets B.P.	500,000 units	Oral	50		SQ
"Odorgon"	Liquid, 15 mL		1		LA
Oestradiol Valerate	Injection, 10 mg in 1 mL	Injection	3		SC
	amp.	Oral	2	2	SC
	Tablets, 1 mg, 28 Tablets, 2 mg, 28	Oral Oral	2 2	2	SC
Oestrogens—Conjugated	Tablets, 300 micrograms, 28	Oral	2	2	AY
	Tablets, 625 micrograms, 28	Oral	2	2	AY
Oestrone B.P.C. 1954	Pessaries, 100 micrograms, 12		1	1	OR
Ossilono B.I.C. 1934	Pessaries, 1 mg, 12		1	i	OR
Olsalazine Sodium	Capsule, 250 mg	Oral	100	5	PS
Orphenadrine Hydrochloride Tablets	50 mg	Oral	200	2	RK ·
B.P.					
Oxazepam Tablets B.P.	15 mg 30 mg	Oral Oral	25 25	• •	AF, AY, WY AF, AY, WY
				• *	, ,

v (n)	Form	Manner of adminis-	Maxi- mum	Number of	ъ.	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand	
Oxprenolol Hydrochloride B.P. with any determined brand of Water for Injections or other solvent	Injection, 2 mg (with required solvent)	Injection	5	••	CG	
Oxprenolol Tablets B.P.	20 mg 40 mg	Oral Oral	100 100	5 5	AF, CG AF, CG	
Oxycodone Hydrochloride	Tablet, 5 mg	Oral	20		BT	
Oxycodone Pectinate	Suppositories, 30 mg (base),		1	••	ВТ	
Oxymetholone Tablets B.P.	50 mg 100 mg	Oral Oral	100 100	5	SD PD	
Oxytocin Injection B.P.	Ampoule, 2 units in 2 mL	Injection	5		SZ	
,	Ampoule, 5 units in 1 mL	Injection	5		SZ	
	Ampoule, 10 units in 1 mL	Injection	5		SZ	
Pancreatin B.P.	Capsule, providing not less than 6,500 B.P. units of	Oral	500	10	RS	
	lipase activity Tablet, providing not less than 6,500 B.P. units of lipase activity	Oral	500	10	RS	
Pancrelipase	Capsule, providing not less than 5,000 B.P. units of	Oral	500	10	JC	
	lipase activity Capsule, providing not less than 10,000 B.P. units of lipase activity	Oral	500	10	OR	
Papaveretum B.P.C. 1973	Injection, 20 mg in 1 mL amp.	Injection	5	••	RO	
Papaveretum B.P.C. 1973 with Hyoscine Hydrobromide B.P.	Injection, 20 mg-400 micro- grams in 1 mL amp.	Injection	5	••	RO	
Paracetamol B.P.	Mixture, 120 mg per 5 mL, 100 mL	Oral	1	2	BW, WL	
Paracetamol Tablets B.P.	500 mg	Oral	100	1	BW, DH, FM, WI	
Paraffin, Soft White B.P.	Cream, compound, 85 g Ointment, compound, 70 g	Application Application	1 1	• •	DY DY	
	Forms specified in Sch. 2		Sch. 2	Sch. 2		
Paraffin, Soft White B.P. with Liquid Paraffin B.P.	Eye ointment, compound, 3.5 g	Application to the eye	2	3	AQ	
Equit Futurini D.F.	Eye ointment, compound,	Application to the eye	2	3	AG	
	Forms specified in Sch. 2	·	Sch. 2	Sch. 2		
Penicillamine Tablets B.P.	125 mg	Oral	100	1	DL	
	250 mg	Oral	100	1	DL	
Perhexiline Maleate	Tablet, 100 mg	Oral	100	5	ML	
Pericyazine	Tablet, 2.5 mg	Oral	200	;	MB	
Pethidine Injection B.P.	Tablet, 10 mg Ampoule, 50 mg in 1 mL	Oral Injection	100 5	1	MB BL, BT,	
• • • • • • • • • • • • • • • • • • •	Ampoule, 100 mg in 2 mL	Injection	5		SI BL, BT,	
Dathidina Tahlata B.D	50 ma	Oral	20		SI	
Pethidine Tablets B.P.	50 mg	Oral	20		SI	
Phenelzine Tablets B.P.	15 mg	Oral	50	2	WW	
henethicillin Capsules B.P.	250 mg	Oral	25	1	SI	
Phenethicillin Potassium B.P. with Purified Water B.P.	500 mg Powder for syrup, 125 mg (Phenethicillin) per 5 mL, 100 mL	Oral Oral	25 1	1	SI SI	
	Powder for syrup, 250 mg (Phenethicillin) per 5 mL, 100 mL	Oral	1		SI	
Phenethicillin Tablets B.P.	250 mg	Oral	25	1	SI	
Phenindione Tablets B.P.	10 mg	Oral	100	2	GL	
	50 mg	Oral	100	2	GL	
Phenobarbitone Injection B.P.	Ampoule, 200 mg in 1 mL	Injection	5		FM	

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Phenobarbitone Tablets B.P.	30 mg	Oral	200	4	DH, FM, SI
Phenoxybenzamine Capsules B.P.	10 mg	Oral	100	5	SK
Phenoxymethylpenicillin (Benzathine Salt)	Suspension, 125 mg per 5 mL, 100 mL	Oral	1	1	CS, SI
	Suspension, 250 mg per 5 mL, 100 mL	Oral	l	1	CS, SI
Phenoxymethylpenicillin (Hydraba- mine Salt)	Suspension, 125 mg per 5 mL, 100 mL	Oral	1	1	AB
	Suspension, 250 mg per 5 mL, 100 mL Tablet, 125 mg	Oral Oral	1 25	1	AB AB
Phenoxymethylpenicillin Potassium	250 mg	Oral	25	1	AB, CS,
Capsules B.P.	500 mg	Oral	25	1	LY, SI CS, SI
Phenoxymethylpenicillin Potassium	250 mg	Oral	25	1	AB, CS,
Tablets B.P.	500 mg	Oral	25	1	LY, SI AB, CS, FM, LY
Phensuximide B.P.C. 1973	Capsule, 500 mg	Oral	200	2	PD
Phenylephrine Hydrochloride B.P.	Eye drops, 1.2 mg per mL, 15 mL	Application to the eye	1	2	AG, AQ
	Ointment, compound, 60 g Suppositories, compound, 12		1 1	1 1	RL RL
Phenytoin B.P.	Tablet, 50 mg	Oral	200	2	PD
Phenytoin Capsules B.P.	30 mg 100 mg	Oral Oral	200 200	2 2	PD PD
Phenytoin Mixture B.P.	30 mg per 5 mL, 500 mL	Oral	1	3	PD
Phytomenadione Injection B.P.	Ampoule, 1 mg Ampoule, 10 mg	Injection Injection	5 5		RO RO
Pilocarpine	Eye disc, 5 mg (releasing 20 micrograms per hour)	Application to the eye	8	5	PT
	Eye disc, 11 mg (releasing 40 micrograms per hour)	Application to the eye	8	5	PT
Pilocarpine Hydrochloride B.P.	Eye drops, 5 mg per mL, 15 mL	Application to the eye	2	3	AG, AQ, IQ, SI
	Eye drops, 10 mg per mL, 15 mL	Application to the eye	2	3	AG, AQ, IQ, SI
	Eye drops, 20 mg per mL, 15 mL		2	3	AG, AQ, IQ, SI
	Eye drops, 30 mg per mL, 15 mL		2	3	AG, AQ, IQ, SI
	Eye drops, 40 mg per mL, 15 mL		2	3	AG, AQ, IQ, SI
	Eye drops, 60 mg per mL, 15 mL		2	3	AG, AQ, SI
P: 1115 P	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Pindolol B.P.	Injection, 400 micrograms in 2 mL amp.	Injection	5		SZ
Pindolol Tablets B.P.	5 mg 15 mg	Oral Oral	100 50	5 5	AF, SZ AF, SZ
Piperazine Oestrone Sulphate	Tablets, 730 micrograms, 28 Tablets, 1.46 mg, 28	Oral Oral	2 2	2 2	AB AB
Piroxicam	Capsule, 10 mg Capsule, 20 mg	Oral Oral	50 25	3	PF PF
Pizotifen Malate	Tablet, 500 micrograms (base)	Oral	100	2	SZ
"PK AID I"	Powder, 250 g	Oral	2	5	SB
"PK AID II"	Powder, 250 g	Oral	2	5	SB
Pneumococcal Vaccine, Polyvalent	Injection, 0.5 mL in disposa- ble syringe (17 valent)	Injection	1	• •	SK
	Injection, 0.5 mL vial (23 valent)	Injection	1	• •	CS

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Polygeline	Intravenous infusion, 17.5 g per 500 mL with Na + 145 mmol per L, K + 5.1 mmol per L, Ca ²⁺ 6.25 mmol per L and Cl = 145 mmol per L; 500 mL	Injection	3		НР
Polyisobutylene	Wafers, compound adhesive, pack of 5	Topical	1	5	DY, HO
	Wafers, compound adhesive with discs, pack of 5	Topical	1	5	BH, SQ
Polymyxin B Sulphate B.P. with Bacitracin B.P. 1968 and Neomy- cin Sulphate B.P.	Eye ointment, 5,000 units- 400 units-5 mg per g, 4 g	400 units-5 mg per g, 4 g to the eye		• •	UP
Polymyxin B Sulphate B.P. with Bacitracin Zinc B.P. and Neomy- cin Sulphate B.P.	Ear drops, 10,000 units-500 units-3.5 mg (base) per mL, 15 mL	Application to the ear	1	••	JC
	Eye ointment, 5,000 units- 400 units-5 mg per g, 4 g	Application to the eye	1	• •	BW
Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Gramicidin	Eye drops, 5,000 units-2.5 mg-25 micrograms per mL, 10 mL	Application to the eye	1	2	BW
Polymyxin B Sulphate B.P. with Neomycin Sulphate B.P. and Hy- drocortisone B.P.	Ear drops, 10,000 units-5 mg-10 mg per mL, 7 mL	Application to the ear	1	2	BW
Polyvinyl Alcohol	Eye drops, 14 mg per mL, 15		2	3	AG
	mL Eye drops, 30 mg per mL, 15 mL	to the eye Application to the eye	2	3	AG
Polyvinyl Alcohol with Povidone B.P.	Eye drops, 14 mg-6 mg per mL, 15 mL	Application to the eye	2	3	AG
Polyvinylpyrrolidone and Vinyl Acetate Copolymer	Spray, aerosol, 60 mg per mL, 121 mL	Application	1	••	EG
"Portagen" Potassium Chloride B.P.	Powder, 450 g Elixir, 1.5 g per 15 mL, 500 mL	Oral Oral	2 2	20 1	MJ SC
	Tablet, 600 mg (sustained release)	Oral	200	1	AF, CG, PT
Potassium Chloride B.P. with Potassium Bicarbonate Effervescent Tablets	14 mmol K ⁺ and 8 mmol Cl ⁻	Oral	60	1	PT
Pralidoxime Iodide	Injection, 500 mg	Injection	5		AB
Prazosin Hydrochloride	Tablet, 1 mg (base) Tablet, 2 mg (base)	Oral Oral	100 100	5 5	PF PF
	Tablet, 5 mg (base)	Oral	100	5	PF
Prednisolone Acetate	Eye drops, 5 mg per mL, 5 mL	Application to the eye	1	6	AQ
	Injection, 10 mg in 1 mL amp.	Intra-articu- lar injection	5		SC
Prednisolone Acetate with Phenyl- ephrine Hydrochloride B.P.	Eye drops, 10 mg-1.2 mg per mL, 10 mL	Application to the eye	l	2	AG
Prednisolone Sodium Phosphate B.P.	Enema, retention, 20 mg in 100 mL	Enema	28	3	GL
	Eye/ear drops, 5 mg per mL, 5 mL	Application to the eye/ear	1	6	GL
	Suppositories, 5 mg (base), 10	- /	3	3	GL
Prednisolone Stearoylglycolate	Tablet, 6.65 mg	Oral	60	1	FE
Prednisolone Tablets B.P.	1 mg 5 mg	Oral Oral	100 60	4	PT FM, NN, PT, UP, US
	25 mg	Oral	30	4	FM, PT
Prednisone Tablets B.P.	1 mg 5 mg	Oral Oral	100 60	4	PT DH, FM, NN, PT, US

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	25 mg	Oral	30	4	FM, PT
"Pregestimil"	Powder, 454 g	Oral	2	20	MJ
Primaquine Tablets B.P.	Tablet, equivalent to 7.5 mg Primaquine base	Oral	42		IC
Primidone Tablets B.P.	250 mg	Oral	200	2	IC
Probenecid Tablets B.P.	500 mg	Oral	100	5	FR
Probucol	Tablet, 250 mg	Oral	112	5	ML
Procainamide Hydrochloride B.P.	Capsule, 250 mg	Oral	200	1	SQ.
n	Capsule, 375 mg	Oral	100	2	SQ
Procainamide Injection B.P.	Vial, 100 mg per mL, 10 mL	Injection	2	••	SQ
Procaine Penicillin Injection B.P.	Syringe, disposable, 1 g Syringe, disposable, 1.5 g	Injection Injection	5 5		SI SI
Procarbazine Hydrochloride	Capsule, 50 mg	Oral	100	2	RO
Prochlorperazine	Suppositories, 3 mg, equiva- lent to 5 mg Prochlorpera- zine Maleate, 5 Suppositories, 15 mg, equiva-	Olui	1	2	MB MB
	lent to 25 mg Prochlorper- azine Maleate, 5				
Prochlorperazine Edisylate	Injection, 12.5 mg in 1 mL	Injection	10	• •	SK
Prochlorperazine Injection B.P.	Ampoule, 12.5 mg in 1 mL	Injection	10	••	MB
Prochlorperazine Tablets B.P.	5 mg	Oral	25		MB, PT, SK
Procyclidine Tablets B.P.	5 mg	Oral	200	2	BW
Promethazine Hydrochloride Injection B.P.	Ampoule, 50 mg in 2 mL	Injection	10		BL
Promethazine Theoclate Tablets B.P.	25 mg	Oral	30	1	MB
Propantheline Tablets B.P.	15 mg	Oral	200	2	PT
Propranolol Injection B.P.	1 mg in 1 mL	Injection	5		IC
Propranolol Tablets B.P.	10 mg	Oral	100	5	AF, IC
•	40 mg	Oral	100	5	AF, IC, PT
	160 mg	Oral	50	5	AF, IC, PT
Propylthiouracil Tablets B.P. Protamine Sulphate Injection B.P.	50 mg Ampoule, 10 mg per mL, 10 mL	Oral Injection	200 6	2	JC BT
Pyrantel Embonate	Tablet, 125 mg (base)	Oral Oral	6		AF AF
Duridantiamina Taklata D D	Tablet, 250 mg (base)		100	2	
Pyridostigmine Tablets B.P.	10 mg 60 mg	Oral Oral	100	2	RO RO
	180 mg (sustained release)	Oral	100	ī	RO
Pyridoxine Hydrochloride B.P.	Injection, 50 mg in 1 mL amp.	Injection	5	1	FM
Pyridoxine Hydrochloride Tablets B.P.	25 mg	Oral	100	••	DH, FM, PT
Pyrimethamine Tablets B.P.	25 mg	Oral	50		BW
Quinethazone	Tablet, 50 mg	Oral	100	1	LE
Quinidine Bisulphate B.P.	Tablet, 250 mg (sustained release)	Oral	100	5	AP
Quinidine Sulphate B.P.	Tablet, 300 mg (sustained release)	Oral	100	5	RS
Quinidine Sulphate Tablets B.P.	200 mg	Oral	100	5	BW, DH, NN, PT, QE
Quinine Bisulphate Tablets B.P.	300 mg	Oral	50	2	AF, DH, FM, NN, PT, SI, US
Quinine Sulphate Tablets B.P.	300 mg	Oral	50	2	AF, DH, FM, NN, PT, SI, US
Ranitidine Hydrochloride	Tablet, 150 mg (base)	Oral	60	2	GL
n a	Tablet, 300 mg (base)	Oral	30	1	GL
Red-back Spider Antivenom	Injection, 500 units amp.	Injection	2	• •	CS
Rifampicin B.P.	Capsule, 150 mg Capsule, 300 mg	Oral Oral	10 10	• •	AF AF

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of reneats	Brand
	strength, type, size, etc.)	17411071	quantity	repears	
Rolitetracycline	Injection, 275 mg amp. and 10 mL solvent	I.V. injection		••	НР
	Injection, 350 mg amp. and 2 mL solvent	I.M. injection	5	• •	НР
Salbutamol B.P.	Spray, metered aerosol, 100 micrograms per dose, 200 doses, 17 g	Inhalation by mouth	2	5	GL
Salbutamol Sulphate B.P.	Capsule, 200 micrograms Inhalation by mouth		100	5	GL
	Solution, 2.5 mg (base) in 2.5 mL ampoules, 30	Inhalation	I	5	GL
	Solution, 5 mg (base) in 2.5 mL ampoules, 30	Inhalation	1	5	GL
	Solution, 5 mg (base) per mL, 30 mL	Inhalation	2	2	GL, RK
	Spray, metered aerosol, 100 micrograms (base) per dose, 200 doses, 13.5 g	Inhalation by mouth	2	5	RK
	Spray, metered aerosol, 100	Inhalation	1	5	RK
	micrograms (base) per	by mouth			
	dose, 400 doses, 13.5 g Syrup, 2 mg (base) per 5 mL, 300 mL	Oral	1	5	GL
Salbutamol Tablets B.P.	4 mg	Oral	100	5	GL
Salcatonin Injection B.P.	50 I.U. in 1 mL amp., 5	Injection	6	5	SZ
	80 I.U. in 0.8 mL amp., 5	Injection	3	5	SZ
	100 I.U. in 1 mL amp., 5 400 I.U. in 2 mL vial	Injection Injection	3 4	5	RG, SZ RG
Semisodium Valproate	Tablet, 135 mg (enteric	Oral	200	2	AB
	coated) Tablet, 269 mg (enteric	Oral	200	2	AB
	coated) Tablet, 538 mg (enteric coated)	Oral	200	2	AB
Silver Sulphadiazine with Chlorhexidine Gluconate	Cream, 10 mg-2 mg per g, 50 g	Application	1	••	SN
	Cream, 10 mg-2 mg per g, 100 g	Application	1	• •	SN
	Cream, 10 mg-2 mg per g, 500 g	Application	1	••	SN
Sodium Acid Citrate B.P.	Mixture, 5.5 g per 20 mL, 500 mL	Oral	1	4	PD
Sodium Acid Phosphate B.P.	Tablet, compound efferves- cent, equivalent to 500 mg phosphorus	Oral	100	5	SZ
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Sodium Aurothiomalate Injection B.P.	Ampoule, 10 mg Ampoule, 20 mg	Injection Injection	5 10	 1	MB MB
D.1 .	Ampoule, 50 mg	Injection	10	1	MB
Sodium Bicarbonate Intravenous Infusion B.P.	100 mmol in 100 mL	Injection	5		AB
Sodium Chloride and Glucose Intra- venous Infusion B.P.	31 mmol-222 mmol per L,	Injection	5	1	AB, BX, KM
	19 mmol-104 mmol per 500 mL, 500 mL	Injection	5	1	AB, BX
	39 mmol-69 mmol per 500 mL, 500 mL	Injection	5	1	AB, BX
	154 mmol-278 mmol per L, 1 L	Injection	2	1	AB, BX
Sodium Chloride B.P. with Potas- sium Chloride B.P. and Calcium Chloride B.P. in Water for Injections	Sodium Chloride Compound Injection—1 L	Injection	4	1	AB, BX
Sodium Chloride B.P. with Sodium Acetate B.P., Sodium Gluconate, Potassium Chloride B.P. and Mag- nesium Chloride B.P.	Electrolyte Replacement Solution, 5.26 g-3.68 g-5.02 g-370 mg-300 mg per L, 1 L	Injection	2	1	AB, BX

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Sodium Chloride Intravenous Infu-	Ampoule, 9 mg per mL,	Injection	5	1	AP, BT
sion B.P.	2 mL Ampoule, 9 mg per mL, 5 mL	Injection	5	1	AP, BT
	Ampoule, 9 mg per mL, 10 mL	Injection	5	1	AP, BT
	154 mmol per L, 1 L	Injection	5	1	AB, BX, KM
S. J. City T. A.	513 mmol per L, 1 L	Injection	2	1	BX
Sodium Citro-Tartrate	Granules, effervescent, 100 g	Oral	1	4	AB, PT
Sodium Cromoglycate B.P.	Eye drops, 20 mg per mL, 10 mL	Application to the eye	1	5	FC
	Nebuliser solution, 20 mg per 2 mL, ampoule	Inhalation	120	3	FC
	Spray, metered aerosol, 1 mg per dose, 200 doses, 13.87 g	Inhalation by mouth	1	5	FC
Sodium Cromoglycate Insufflation B.P.	Capsule, 20 mg	Inhalation by mouth	100	5	FC
Sodium Fusidate B.P.	Tablet, 250 mg (enteric coated)	Oral	36	• •	SK
Sodium Lactate Compound Intravenous Infusion B.P.	1 L	Injection		1	AB, BX, KM
Sodium Lactate Compound Intrave- nous Infusion B.P. with Anhy- drous Glucose B.P.	278 mmol per L, 1 L	Injection	2	1	AB, BX
Sodium Nitroprusside B.P.	Infusion, 50 mg	Injection	10		RO
Sodium Valproate B.P.	Oral liquid, 200 mg per 5	Oral	3	2	RC
	mL, 200 mL Tablet, 200 mg (enteric	Oral	200	2	RC
	coated) Tablet, 500 mg (enteric coated)	Oral	200	2	RC
	Tablet, crushable, 100 mg	Oral	200	2	RC
Sodium Valproate Elixir B.P.	200 mg per 5 mL, 200 mL	Oral	3	2	RC
Sotalol Hydrochloride	Tablet, 160 mg	Oral	60	5	AP
Spectinomycin Hydrochloride B.P.	Injection, 2 g (base) with 3.2 mL diluent	Injection	1		UP
	Injection, 4 g (base) with 6.5 mL diluent	Injection	1	• •	UP
Spironolactone Tablets B.P.	25 mg	Oral	100	5	SR
Staphylococcus Toxoid B.P. 1968	Injection, 5 mL (diluted 1 in 10)	Injection	1	••	CS
	Injection, 5 mL (undiluted)	Injection	I		CS
Sterculia B.P.	Discs, 12		1	5	вн
	Discs, compound, 10		1	5	DY
	Paste, 127.6 g Powder, 71 g		1 1		HO DY
	Squares, 6		i	5	BH
Sterculia B.P. with Frangula Bark B.P.	Granules, 473 mg-83 mg per g, 250 g	Oral	2	1	SC
Streptokinase B.P. with any deter- mined brand of Water for Injec-	Injection, 100,000 I.U. (with required solvent)	Injection	2		PS
tions or other solvent	Injection, 250,000 I.U. (with required solvent)	Injection	2	• •	HP, PS
	Injection, 600,000 I.U. (with required solvent)	Injection	5	••	PS up
Suggeste	Injection, 750,000 I.U. (with required solvent)	Injection	5		HP
Sucralfate	Tablet, 1 g	Oral	120	2	BT
Sulindac Tablets B.P.	100 mg	Oral	50	3	FR
Sulphacetamide Sodium B.P.	Eye drops, 100 mg per mL, 15 mL	Application to the eye	1	2	AG, SI
Culabadia: dia T. U. a. D. D.	Eye drops, 200 mg per mL, 15 mL	Application to the eye	1	2	SI
Sulphadimidine Tablets B.P.	500 mg	Oral	40	2	PT

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Sulphafurazole B.P.	Eye drops, 40 mg per mL, 10 mL	Application	1	2	RO
Sulphamethizole Tablets B.P.	500 mg	to the eye Oral Oral	40 20	2 2	ww ww
Sulphasalazine	Tablet, 500 mg Tablet, 500 mg (enteric coated)	Oral Oral	200 200	5 5	AF, PS PS
Sulphinpyrazone Tablets B.P. Sulthiame Tablets B.P.	100 mg 50 mg	Oral Oral Oral	100 200 200	5 2 2	CG BN
"Super Banish"	200 mg Liquid, 7.5 mL	Orai	1		BN DY
Surgical Cement	Skin Bond Adhesive, 118 mL	Application	1	2	DY
Surgical Cement Solvent	227 mL 240 mL 250 mL	Application Application Application	1 1 1	2 2 2	DY EG EG
Tamoxifen Citrate Tablets B.P.	10 mg (base) 20 mg (base)	Oral Oral	60 60	5 5	IC IC
Temazepam	Capsule, 10 mg	Oral	25		AF, SI, WY
Terbutaline Sulphate B.P.	Elixir, 300 micrograms per mL, 300 mL	Oral	1	5	AP
	Injection, 100 micrograms in 1 mL amp. Injection, 500 micrograms in	Injection Injection	5		AP AP
	1 mL amp. Nebuliser solution, 10 mg per	Inhalation	1	5	AP
	mL, 50 mL Spray, metered aerosol, 250 micrograms per dose, 400 doses, 14 g	Inhalation by mouth	1	5	AP
Terbutaline Sulphate Tablets B.P.	5 mg	Oral	100	5	AP
Testosterone Enanthate B.P.	Injection, 250 mg in 1 mL amp.	Injection	3	3	SC
Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P. and Testosterone Isocaproate B.P.	Injection, 20 mg-40 mg-40 mg amp.	Injection	3	3	OR
Testosterone Propionate B.P. with Testosterone Phenylpropionate B.P., Testosterone Isocaproate B.P. and Testosterone Decanoate B.P.	Injection, 30 mg-60 mg-60 mg-100 mg amp.	Injection	3	3	OR
Testosterone Propionate Injection B.P.	Ampoule, 50 mg in 1 mL	Injection	9	••	SC
Tetanus Vaccine, Adsorbed B.P.	Injection, 0.5 mL amp.	Injection	3		CS
Tetrabenazine	Tablet, 25 mg	Oral	100	2	RO
Tetracosactrin Zinc Injection B.P.	Ampoule, 500 micrograms in 1 mL Ampoule, 1 mg in 1 mL	Injection Injection	5	5	CG CG
Tetracycline B.P. with a buffering agent	Capsule, 250 mg	Oral	25	1	AP, CS, FM, HP,
Tetracycline B.P. with a buffering agent and Nystatin B.P.	Capsule, 250 mg-250,000 units	Oral	25	1	LE, SQ SQ
Tetracycline Capsules B.P.	250 mg	Oral	25	1	BZ, LE, UP
Tetracycline Hydrochloride B.P.	Eye ointment, 10 mg per g, 5 g	Application to the eye	1	••	BZ
Tetracycline Hydrochloride B.P. with Nystatin B.P.	Capsule, 250 mg-250,000 units	Oral	25	1	AP, LE
Theophylline B.P.	Capsule, 50 mg (containing sustained release beads)	Oral	100	5	RG, RK
	Capsule, 100 mg (containing sustained release beads)	Oral	100	5	RG, RK
	Capsule, 100 mg (controlled release)	Oral	100	5	FC

Name of Pharmacoutical Panels	Form	Manner of adminis-	Maxi- mum	Number of	Praud
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
	Capsule, 250 mg (controlled release)	Oral	100	5	FC
	Elixir, 80 mg per 15 mL, 500 mL	Oral	1	5	SC
	Syrup, 80 mg per 15 mL, 500 mL	Oral	1	5	RK
	Tablet, 50 mg	Oral	100	5	RK
	Tablet, 125 mg	Oral	100	5	RK
	Tablet, 200 mg	Oral	100	5	RK
	Tablet, 200 mg (sustained release)	Oral	100	5	AP
	Tablet, 250 mg (sustained release) Tablet, 300 mg (sustained	Oral Oral	100	5	RK AP
	release) Forms specified in Sch. 2	Olai	Sch. 2	Sch. 2	Al
Thinks denote Tables D.B.	-	01			1414
Thiabendazole Tablets B.P.	500 mg	Oral	16	• •	MK
Thiamine Hydrochloride Injection B.P.	Ampoule, 100 mg in 1 mL	Injection	5	1	FM
Thiamine Hydrochloride Tablets B.P.	100 mg	Oral	100	2	FM, NN, US
Thiethylperazine Malate	Injection, 6.5 mg (base) in 1 mL amp.	Injection	10	••	SZ
Thiethylperazine Maleate	Tablet, 6.5 mg (base)	Oral	50		SZ
	Suppositories, 6.5 mg (base),		1	2	SZ
Thioguanine Tablets B.P.	40 mg	Oral	25	1	BW
Thiopropazate Hydrochloride Tablets B.P.	5 mg	Oral	200	••	SR
Thioridazine Hydrochloride B.P.	Solution, 30 mg per mL, 30 mL	Oral	2	2	SZ
Thioridazine Tablets B.P.	10 mg	Oral	200		AF, SZ
	25 mg	Oral	200		AF, SZ
	50 mg	Oral	200		AF, SZ
	100 mg	Oral	100	1	AF, SZ
Thiotepa B.P. with any determined	Eye drops, set, 15 mg with	Application	1	5	LE
brand of Water for Injections or other solvent	required solvent Injection, 15 mg (solvent required)	to the eye Injection	2	1	LE
Thurswine Tobleto D.D.	• •	01	200	i	вw
Thyroxine Tablets B.P.	50 micrograms 100 micrograms	Oral Oral	200 200	1	BW BW
	200 micrograms	Oral	200	1	BW
Tinggaillia Cadinas mish ann datas					
Ticarcillin Sodium with any deter- mined brand of Water for Injec- tions or other solvent	Injection, vial containing equivalent 1 g Ticarcillin (with required solvent)	Injection	10	••	BR, CS
	Injection, vial containing equivalent 3 g Ticarcillin (with required solvent)	Injection	10	••	BR, CS
Timolol Maleate Eye Drops B.P.	2.5 mg per mL, 5 mL	Application to the eye	1	6	FR, SI
	5 mg per mL, 5 mL	Application to the eye	1	6	FR, SI
Timolol Maleate Tablets B.P.	5 mg	Oral	100	5	FR
Tinidazole	Tablet, 500 mg	Oral	4		PF
Tobramycin B.P.	Eye drops, 3 mg per mL,	Application	1	2	AQ
	5 mL Eye ointment, 3 mg per g,	to the eye	1		AQ
Tobramycin Injection B.P.	3.5 g 40 mg	to the eye	5	• •	LY
yyy	80 mg	Injection	5		ĹŸ
Tolazamide Tablets B.P.	250 mg	Oral	100	5	UP
Tolbutamide Tablets B.P.	500 mg	Oral Oral	200 100	2 5	HP HP
Tranexamic Acid Tablets B.P.	l g 500 mg	Oral	100	2	PS
Tranylcypromine Tablets B.P.	10 mg	Oral	50	2	SK

		Manner of	Maxi-	Number	
	Form	Manner of adminis-	mum	of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
Triamcinolone Acetonide B.P.	Injection, 10 mg in 1 mL amp.	Injection	5	••	SQ
Triamcinolone Acetonide B.P. with Neomycin Sulphate B.P., Gramici- din and Nystatin B.P.	Ear cream, 1 mg-2.5 mg (base)-250 micrograms- 100,000 units per g, 5 g	Application to the ear	1	2	SQ
dir and rystatii D.F.	Ear drops, 1 mg-2.5 mg (base)-250 micrograms- 100,000 units per g, 7.5 mL	Application to the ear	1	2	SQ
	Ear ointment, 1 mg-2.5 mg (base)-250 micrograms- 100,000 units per g, 5 g	Application to the ear	1	2	SQ
Triamcinolone Acetonide Cream B.P.	200 micrograms per g, 45 g 200 micrograms per g, 100 g 500 micrograms per g, 15 g	Application Application Application	1 2 1		SQ LE, SQ LE, SQ
Triamcinolone Acetonide Ointment	200 micrograms per g, 45 g	Application	1	•	SO SO
B.P.	200 micrograms per g, 100 g 500 micrograms per g, 15 g	200 micrograms per g, 100 g Application 2		i i	LE, SQ LE, SQ
Triamterene B.P.	Tablet, 100 mg	Oral	100	1	SK
Trifluoperazine Hydrochloride B.P.	Injection, 1 mg (base) in 1 mL amp.	Injection	10		SK
Trifluoperazine Hydrochloride	l mg	Oral	200		PT, SK
Tablets B.P.	2 mg	Oral Oral	200 100	;	PT, SK
Triglycerides Oil, Medium Chain	5 mg 1 L	Oral	100	1	PT, SK KY
Trimethoprim Tablets B.P.	300 mg	Oral	7	1	AF, BW
Trimipramine	Capsule, 50 mg	Oral	50	2	MB
Trimipramine Tablets B.P.	25 mg (base)	Oral	50	2	MB
Trioxysalen	•	Oral	100	2	PT
Urea Cream B.P.	Tablet, 5 mg		100	2	
Orea Cream B.P.	100 mg per g, 100 g	Application	1	2	AG, HA, NW, OL, PS
Vancomycin Hydrochloride B.P.	Capsule, 125 mg (125,000 I.U.) vancomycin activity	Oral	40	••	LY
	Capsule, 250 mg (250,000 I.U.) vancomycin activity	Oral	40	••	LY
Vancomycin Hydrochloride B.P. with any determined brand of Water for Injections or other solvent	Injection, 500 mg (500,000 I.U.) vancomycin activity (with required solvent)	Injection	5	••	LY
Vasopressin Tannate	Injection, (oily) 5 units in 1 mL amp.	Injection	10	1	PD
Verapamil Hydrochloride B.P.	Tablet, 40 mg Tablet, 80 mg	Oral Oral	100 100	5 5	KN, SC KN, SC
Verapamil Hydrochloride Injection B.P.	Ampoule, 5 mg in 2 mL	Injection	5		KN, SC
Verapamil Hydrochloride Tablets B.P.	40 mg	Oral	100	5	AF, PT, US
	80 mg	Oral	100	5	AF, PT, US
	120 mg	Oral	100	5	KN, PT, SC, US
	160 mg	Oral	60	5	KN, SC
Vidarabine	Eye ointment, 30 mg per g, 3.5 g	Application to the eye	1	••	PD
Vinblastine Sulphate B.P.	Injection, 10 mg and 10 mL solvent	Injection	2	••	BL
Vinblastine Sulphate B.P. with any determined brand of Water for Injections or other solvent	Injection, 10 mg (with required solvent)	Injection	2	••	LY
Vincristine Sulphate B.P.	Injection, 1 mg and 10 mL solvent	Injection	5	••	BL, LY
	Injection, 5 mg and 10 mL solvent	Injection	1	••	BL
Warfarin Tablets B.P.	1 mg 2 mg	Oral Oral	50 50	2 2	BT, GL BT

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
	2.5 mg	Oral	50	2	BT
	3 mg	Oral	50	2	GL
	5 mg	Oral	50	2	BT, GL
	7.5 mg	Oral	50	2	BT [']
	10 mg	Oral	50	2	BT
Water for Injections B.P.	Ampoule, 2 mL	Injection	5	3	AP, BT, BZ, SI
	Ampoule, 5 mL	Injection	5	3	AP, BT, BZ, SI
	Ampoule, 10 mL	Injection	5	3	AP, BT, BZ, SI
Wool Alcohols Ointment B.P.	100 g		1	1	SN
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Zinc Oxide B.P.	Ointment, compound, 50 g		1	1	ww
	Suppositories, compound, 12		i	i	ww
	Forms specified in Sch. 2		Sch. 2	Sch. 2	
Zinc Sulphate B.P. with Phenyle- phrine Hydrochloride B.P.	Eye drops, 2.5 mg-1.2 mg per mL, 15 mL	Application to the eye	1	2	AG, AQ

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Purposes	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Acyclovir	Tablets, 200 mg, 90	In compliance with authority procedures set out in sub-paragraph 10 (d): Moderate to severe recurrent (more than ten at- tacks per year) genital herpes, confirmed by ap- propriate microbiological technique Suppression of genital herpes in severely immuno- compromised patients	Oral	ì	2	BW
Bromocriptine Mesylate Tablets B.P.	2.5 mg (base)	In compliance with authority procedures set out in sub-paragraph 10 (d): Acromegaly, prior to surgery or radiotherapy or where surgery or radiotherapy is inappropriate Parkinson's disease Pathological hyperprolactinaemia where appropriate surgery or radiotherapy is not indicated or has already been used with incomplete resolution	Oral	60	5	SZ
Codeine Phosphate B.P. with Aspirin B.P.	Tablet, 30 mg- 325 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain not responding to non-narcotic analgesics	Oral	40		BW
Codeine Phosphate B.P. with Paracetamol B.P.	Tablet, 30 mg- 500 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain not responding to non-narcotic analgesics	Oral	40	••	WL
Diazepam B.P.	Syrup, 2 mg per 5 mL, 100 mL	In compliance with authority procedures set out in sub-paragraph 10 (d): Disabling spasticity For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal Malignant neoplasia (late stage)	Oral	2		RO
Diazepam Injection B.P.	10 mg in 2 mL	In compliance with authority procedures set out in sub-paragraph 10 (d): Disabling spasticity For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal Malignant neoplasia (late stage)	Injection	10		BL, RO

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Purposes	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand	2618 C
Diazepam Tablets B.P.	2 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Disabling spasticity For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal Malignant neoplasia (late stage)	Oral	100		AF, PT, RO, SU	Government departments
	5 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Disabling spasticity For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal Malignant neoplasia (late stage)	Oral	100		AF, PT, RO, SU	
Doxycyline Capsules B.P.	100 mg	Urethritis	Oral	21		Dar	
Doxycycline Hydrochloride B.P.	Capsule, 100 mg (base) (con- taining enteric coated pellets)	Urethritis		21		PT FA	
	Tablet, 100 mg (base)	Urethritis	Oral	21		AF, PF	
Hydrocortisone Sodium Succinate B.P.	Injection set containing equivalent of 100 mg Hy- drocortisone and 2 mL solvent	For use in a hospital	Injection	6		NR, UP	ν̈́
	Injection set containing equivalent of 250 mg Hy- drocortisone and 2 mL solvent	For use in a hospital	Injection	6		UP	No. GN 44, 23 November 198
Medroxyprogesterone Acetate B.P.	Tablet, 10 mg	Endometriosis	Oral	100	2	UP	Sm.
Methadone Injection B.P.	Ampoule, 10 mg in 1 mL	In compliance with authority procedures set out in sub-paragraph 10 (d):		10		BW	ner 190

ame of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Purposes	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
		Severe disabling pain associated with proven malig- nant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)		-	•	
1ethadone Tablets B.P.	5 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain associated with proven malig- nant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)	Oral	40		BW ,
	10 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain associated with proven malig- nant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)	Oral	40		BW
letronidazole Tablets B.P.	400 mg	Treatment of anaerobic infections	Oral	21	1	AF, MB
lorphine Sulphate Tablets B.P.	30 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain associated with proven malig- nant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)	Oral	40		FM
iclosamide Tablets B.P.	500 mg	Hymenolepiasis nana	Oral	16		BN
itrazepam Tablets B.P.	5 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal Myoclonic epilepsy	Oral	50	5	AF, RO
xazepam Tablets B.P.	15 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal	Oral	50	5	AF, AY, WY
	30 mg	In compliance with authority procedures set out in sub-paragraph 10 (d):	Oral	50	5	AF, AY, WY

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Purposes	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
		For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal				
Oxycodone Hydrochloride	Tablet, 5 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain associated with proven malignant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)	Oral	40	••	вт
Oxycodone Pectinate	Suppositories, 30 mg (base), 12	In compliance with authority procedures set out in sub-paragraph 10 (d): Severe disabling pain associated with proven malignant neoplasia Severe disabling pain where treatment is initiated in a hospital (in-patient or out-patient)		2		ВТ
Pethidine Injection B.P.	Ampoule, 50 mg in 1 mL	In compliance with authority procedures set out in sub-paragraph 10 (d): Late stage malignant neoplasia	Injection	10	••	BL, BT, SI
	Ampoule, 100 mg in 2 mL	In compliance with authority procedures set out in sub-paragraph 10 (d): Late stage malignant neoplasia	Injection	10	••	BL, BT, SI
Phenethicillin Capsules B.P.	250 mg	Prophylaxis of recurrent streptococcal infections (in- cluding rheumatic fever)	Oral	50	5	SI
Phenethicillin Tablets B.P.	250 mg	Prophylaxis of recurrent streptococcal infections (in- cluding rheumatic fever)	Oral	50	5	SI
Phenoxymethylpenicillin (Hydrabamine Salt)	Tablet, 125 mg	Prophylaxis of recurrent streptococcal infections (in- cluding rheumatic fever)	Oral	50	5	AB
Phenoxymethylpenicillin Potassium Capsules B.P.	250 mg	Prophylaxis of recurrent streptococcal infections (in- cluding rheumatic fever)	Oral	50	5	AB, CS, LY, SI
Phenoxymethylpenicillin Potassium Tablets B.P.	250 mg	Prophylaxis of recurrent streptococcal infections (in- cluding rheumatic fever)	Oral	50	5	AB, CS, LY, SI
Prochlorperazine Tablets B.P.	5 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Emesis associated with malignant disease Rotational vertigo	Oral	100	5	MB, PT, SK
Rifampicin B.P.	Capsule, 150 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Leprosy in adults	Oral	100	••	AF ,
	Capsule, 300 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): Leprosy in adults	Oral	100	••	AF

2621

Name of Pharmaceutical Benefit	Form (strength, type, size, etc.)	Purposes	Manner of adminis- tration	Maxi- mum quantity	Number of repeats	Brand
Temazepam	Capsule, 10 mg	In compliance with authority procedures set out in sub-paragraph 10 (d): For use by patients receiving long-term nursing care in hospitals and nursing homes who have been demonstrated, within the past six months, to be benzodiazepine dependent by an unsuccessful attempt at gradual withdrawal	Oral	50	5	AF, SI, WY
Tetracycline B.P with a buffering agent	Capsule, 250 mg	Bronchiectasis in patients over the age of 8 years Chronic bronchitis in patients over the age of 8 years Severe acne	Oral	50	5	AP, CS, FM, HP, LE, SQ
Tetracycline B.P with a buffering agent and Nystatin B.P.	Capsule, 250 mg-250,000 units	Bronchiectasis in patients over the age of 8 years Chronic bronchitis in patients over the age of 8 years Severe acne	Oral	50	5	SQ .
Tetracycline Capsules B.P.	250 mg	Bronchiectasis in patients over the age of 8 years Chronic bronchitis in patients over the age of 8 years Severe acne	Oral	50	5	BZ, LE, UP
Tetracycline Hydrochloride B.P. with Nystatin B.P.	Capsule, 250 mg-250,000 units	Bronchiectasis in patients over the age of 8 years Chronic bronchitis in patients over the age of 8 years Severe acne	Oral	50	5	AP, LE

SECOND SCHEDULE

Form of Pharmaceutical Benefit	Maximum quantity	Number of repeats
Creams	100 g	1
Ear Drops	15 mL	2 4
Elixirs	100 mL	4
Eye Drops		
Cocaine	15 mL	
Pilocarpine	15 mL	6
Others	15 mL	6
Eye Lotions	200 mL	1
Gargles	200 mL	4
Glycerins	100 mL	1
Inhalations	50 mL	1
Inhalations, Solid	4 g	1
Linctuses	100 mL	2
Liniments	100 mL	2 1 2 4
Lotions	200 mL	2
Mixtures	200 mL	4
Mixtures for Children	100 mL	4
Mouth Washes	200 mL	1
Nasal Instillations	15 mL	2
Ointments and Waxes	100 g	1
Paints	25 mL	1
Pastes		
Cocaine	25 g	
Others	100 g	1
Powders, Dusting	100 g	1
Powders for Internal Use	100 g	2
Powders, Irrigation	100 g	2 1
Soaps	200 mL	1
Solutions	200 mL	2
Syrups	100 mL	4
Tinctures	25 mL	1

THIRD SCHEDULE

Name of Pharmacon at 1 D Co	Form	Manner of adminis-	Maxi- mum	Number of	n .
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
Amoxycillin Capsules B.P.	250 mg (base)	Oral	20	••	AF, BR, CS, PT, SI
	500 mg (base) .	Oral	20	• •	AF, BR, CS, PT, SI
Amoxycillin Trihydrate B.P.	Tablet, chewable, 250 mg (base)	Oral	20	••	BR, CS
	Tablet, dispersible, 3 g (base)	Oral	1	••	BR, CS
Amoxycillin Trihydrate B.P. with Purified Water B.P.	Powder for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	••	BR, CS, PT, SI
	Powder for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	BR, CS, PT, SI
Amphotericin B.P.	Lozenge, 10 mg	Oral	20		SQ
	Ointment, 30 mg per g, 15 g	Application	1	• •	SQ
Ampicillin Sodium B.P. with any determined brand of Water for	Injection, 250 mg (base) vial (with required solvent)	Injection	5	••	CS, PT
Injections or other solvent	Injection, 500 mg (base) vial (with required solvent)	Injection	5	••	BR, CS, PT
	Injection, 1 g (base) vial (with required solvent)	Injection	5	••	CS, PT
Aspirin Mixture A.P.F. 13 with Codeine Phosphate B.P.	Mixture, 500 mg-30 mg per 10 mL	Oral	200 mL	• •	
Benzylpenicillin Potassium B.P. with any determined brand of Water	Injection, 300 mg vial (with required solvent)	Injection	5	••	CS
for Injections or other solvent	Injection, 600 mg vial (with required solvent)	Injection	5	••	CS
	Injection, 3 g vial (with required solvent)	Injection	5	••	CS
Cephalexin B.P. with Purified Water B.P.	Granules for syrup, 125 mg (base) per 5 mL, 100 mL	Oral	1	• •	GL, LY
	Granules for syrup, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	GL, LY
Cephalexin Capsules B.P.	250 mg	Oral	20		GL, LY
0 1 1 2 1 0 E DD 12	500 mg	Oral	20	• •	GL, LY
Cephalothin Sodium B.P. with any determined brand of Water for Injections or other solvent	Injection, I g (base) (with required solvent) Injection, 2 g (base) (with	Injection Injection	5 1	••	GL, LY LY
	required solvent) Injection, 4 g (base) (with	Injection	1		LY
Codeine Phosphate B.P. with	required solvent) Tablet, 30 mg-325 mg	Oral	20		BW
Aspirin B.P. Codeine Phosphate B.P. with Paracetamol B.P.	Tablet, 30 mg-500 mg	Oral	20	••	WL
Co-trimoxazole Mixture, Paediatric B.P.	40 mg-200 mg per 5 mL, 100 mL	Oral	1		AF, BW, PT, RO
Co-trimoxazole Tablets B.P.	80 mg-400 mg	Oral	10	••	AF, BW, PT, RO
	160 mg-800 mg	Oral	10	••	AF, BW, PT, RO
Doxycycline Capsules B.P.	100 mg	Oral	7		PT
Doxycycline Hydrochloride B.P.	Capsule, 100 mg (base) (containing enteric coated pellets)	Oral	7	••	FA
	Tablet, 100 mg (base)	Oral	7		AF, PF
Erythromycin B.P.	Capsule, 125 mg (containing enteric coated pellets)	Oral	25	• •	FA
	Capsule, 250 mg (containing enteric coated pellets)	Oral	25	••	FA, LY
Erythromycin Estolate B.P.	Drops, paediatric, 100 mg (base) per mL, 10 mL	Oral	1	••	LY
	Suspension, 125 mg (base) per 5 mL, 100 mL	Oral	1	••	LY
Erythromycin Ethyl Succinate B.P.	Tablet, chewable, 200 mg (base)	Отаі	25	••	AB
	Tablet, 400 mg (base)	Oral	25	• •	AB
Erythromycin Ethyl Succinate B.P.	Granules for suspension, 200	Oral	1	• •	AB
with Purified Water B.P.	mg (base) per 5 mL, 100 mL				

	Form	Manner of adminis-	Maxi- mum	Number of	
Name of Pharmaceutical Benefit	(strength, type, size, etc.)	tration	quantity	repeats	Brand
Erythromycin Stearate B.P.	Capsule, 250 mg (base)	Oral	25		AB
•	Suspension, 125 mg (base) per 5 mL, 100 mL	Oral	1		AB
	Suspension, 250 mg (base) per 5 mL, 100 mL	Oral	1	••	AB
Erythromycin Stearate Tablets B.P.	250 mg (base)	Oral	25		AB
Erythromycin Tablets B.P.	250 mg	Oral	25		AB, UP
Metronidazole Tablets B.P.	200 mg	Oral	21	• •	AF, MB, PT, SR
N. C. J. BB	250 mg	Oral	21		JC
Nystatin B.P.	Lozenge, 100,000 units	Oral	20		LE
Montanto Millia D.D.	Tablet, 500,000 units	Oral	25	• •	LE
Nystatin Mixture B.P.	100,000 units per mL, 24 mL		1	• •	LE, SQ
Nystatin Tablets B.P.	500,000 units	Oral	25	• •	SQ
Paracetamol B.P.	Mixture, 120 mg per 5 mL, 100 mL	Oral	1	••	BW, US, WL
Phenethicillin Capsules B.P.	250 mg	Oral	25		SI
	500 mg	Oral	25		SI
Phenethicillin Potassium B.P. with Purified Water B.P.	Powder for syrup, 125 mg (Phenethicillin) per 5 mL, 100 mL	Oral	1	••	SI
	Powder for syrup, 250 mg (Phenethicillin) per 5 mL, 100 mL	Oral	1		SI
Phenethicillin Tablets B.P.	250 mg	Oral	25		SI
Phenoxymethylpenicillin (Benzathine Salt)	Suspension, 125 mg per 5 mL, 100 mL	Oral	1	••	CS, SI
	Suspension, 250 mg per 5 mL, 100 mL	Oral	1	••	CS, SI
Phenoxymethylpenicillin (Hydraba- mine Salt)	Suspension, 125 mg per 5 mL, 100 mL	Oral	1	••	AB
	Suspension, 250 mg per 5 mL, 100 mL	Oral	1	• •	AB
	Tablet, 125 mg	Oral	25	• •	AB
Phenoxymethylpenicillin Potassium Capsules B.P.	250 mg	Oral	25	••	AB, CS, LY, SI
	500 mg	Oral	25		CS, SI
Phenoxymethylpenicillin Potassium Tablets B.P.	250 mg	Oral	25	• •	AB, CS, LY, SI
	500 mg	Oral	25	• •	AB, CS, FM, LY
Procaine Penicillin Injection B.P.	Syringe, disposable, 1 g	Injection	5		SI
	Syringe, disposable, 1.5 g	Injection	5		SI
Sulphadimidine Tablets B.P.	500 mg	Oral	40		PT
•	U		25		
Tetracycline B.P. with a buffering agent	Capsule, 250 mg	Oral		• •	AP, CS, FM, HP, LE, SQ
Tetracycline Capsules B.P.	250 mg	Oral	25		BZ, LE, UP
Water for Injections B.P.	Ampoule, 2 mL	Injection	5	• •	AP, BT, BZ, SI
	Ampoule, 5 mL	Injection	5	••	AP, BT, BZ, SI
	Ampoule, 10 mL	Injection	5	••	AP, BT, BZ, SI

Dated this 16th day of November 1988.

JOHN DEEBLE

Acting First Assistant Secretary Health Benefits Division

Delegate of the Minister of State for Community Services and Health

Industrial Relations

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE OF APPLICATION DECLARE THE ACTORS (THEATRICAL) AWARD 1981 AS VARIED TO DATE, A COMMON RULE IN THE NORTHERN TERRITORY

(C No. 22679 of 1988)

Notice is hereby given that at 2.15 p.m. on 5 December 1988 at CML Building, University Ave, Canberra, ACT the Australian Conciliation and Arbitration Commission will hear an application by Actors Equity of Australia to declare the award known as the Actors (Theatrical) Award 1981 as varied to date, a Common Rule of the industry covered by the said award namely actors in the live entertainment industry in the Northern Territory.

A copy of the said award as varied to date may be inspected free of charge at the Office of the Industrial Registrar, 80 Collins St, Melbourne Vic.; the Office of the Deputy Industrial Registrar, 50 Grenfell St, Adelaide, South Australia; the Office of the Deputy Industrial Registrar, University Ave, Canberra, Australian Capital Territory; the Office of the Deputy Industrial Registrar, 1 Briggs St Darwin.

Compliance with Regulation 35 is not required.

Persons and organisations not appearing or represented will be bound by any order or declaration made by the Commission in the matter.

Dated this sixteenth day of November 1988.

L. J. SOETENS Deputy Industrial Registrar

N.N.-8878095

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE OF APPLICATION DECLARE THE ACTORS ETC. (TELEVISION) AWARD 1979 AS VARIED TO DATE, A COMMON RULE IN THE NORTHERN TERRITORY

(C No. 22680 of 1988)

Notice is hereby given that at 2.15 p.m. on 5 December 1988 at CML Building, University Ave, Canberra, ACT the Australian Conciliation and Arbitration Commission will hear an application by Actors Equity of Australia to declare the award known as the Actors Etc. (Television) Award 1979 as varied to date, a Common Rule of the industry covered by the said award namely actors in the live entertainment industry in the Northern Territory.

A copy of the said award as varied to date may be inspected free of charge at the Office of the Industrial Registrar, 80 Collins St, Melbourne Vic.; the Office of the Deputy Industrial Registrar, 50 Grenfell St, Adelaide, South Australia, the Office of the Deputy Industrial Registrar, University Ave, Canberra, Australian Capital Territory, the Office of the Deputy Industrial Registrar, 1 Briggs St Darwin.

Compliance with Regulation 35 is not required.

Persons and organisations not appearing or represented will be bound by any order or declaration made by the Commission in the matter.

Dated this sixteenth day of November 1988.

L. J. SOETENS Deputy Industrial Registrar

N.N.--8878096

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE UNDER SUB-SECTION 49A (3) IN RELATION TO VARIATION OF A COMMON RULE In the matter of the LIOUOR TRADES HOTELS

(AUSTRALIAN CAPITAL TERRITORY) CONSOLIDATED AWARD 1977

C. No. 32885 of 1988

And in the matter of the variation of the award dated 25 July 1977 in the above matter.

Notice is hereby given:

- (a) that on 3 November 1988 the Commission varied the terms of the above mentioned award referred to in the Schedule as set out in the Schedule:
- (b) that the variation will be a common rule in the Australian Capital Territory in the industry in respect of which the dispute arose with effect from the first pay period to commence on or after 12 September 1988 and 12 March 1989; and
- (c) that any person or organisation having an objection to the variation binding that person or organisation and desiring to be heard in relation to that objection is invited to lodge with the Commission a notice of that objection is invited to lodge with the Commission a notice of that objection within 28 days after the date specified in paragraph (a).

A copy of the award may be inspected at the office of the Registrar. Objections should be lodged with the Registrar at 4th Floor, CML Building, University Ave, Canberra, by 4.30 p.m. on 1 December 1988.

SCHEDULE TERMS VARIED

Clause No.	Subject	Substance of variation
	PRINT NO. H46	660
6	Minimum Wage	National Wage Case August 1988
7	Wages	National Wage Case August 1988
7 A	No Extra Claims	National Wage Case August 1988
10	Overtime and Penalty Rates	National Wage Case August 1988
11	Casual Employees	
15	Board and Residence	National Wage Case August 1988

Dated 16 November 1988.

CHRISTOPHER BRENDON Deputy Industrial Registrar

N.N.-8878097

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE UNDER SUB-SECTION 49A (3) IN RELATION TO VARIATION OF A COMMON RULE In the matter of the AUSTRALIAN WORKERS' UNION CONSTRUCTION-ON-SITE AND CIVIL

ENGINEERING (ACT) AWARD 1981

C No. 90220 of 1988

And in the matter of the variation of the award dated 13 October 1982 in the above matter.

Notice is hereby given:

- (a) that on 3 November 1988 the Commission varied the terms of the above mentioned award referred to in the Schedule as set out in the Schedule:
- (b) that the variation will be a common rule in the Australian Capital Territory in the industry in respect of which the dispute arose with effect from the first pay period to commence on or after 1 September 1988; and
- (c) that any person or organisation having an objection to the variation binding that person or organisation and desiring to be heard in relation to that objection is invited to lodge with the Commission a notice of that objection within twenty-eight days after the date specified in paragraph (a).

A copy of the award may be inspected at the office of the Registrar. Objections should be lodged with the Registrar at 4th Floor, CML Building, University Ave, Canberra, by 4.30 p.m. on 1 December 1988.

SCHEDULE TERMS VARIED

Clause No.	Subject	Substance of variation
	PRINT NO. H46	65
7	Classifications and	0
	Wage Rates	August 1988
8	Leading hands	National Wage Case August 1988
9	Industry Allowance	National Wage Case August 1988
10	Extra Rates	National Wage Case August 1988
16	Meal Breaks and Meal Allowance	National Wage Case August 1988
17	Compensation for Travel Patterns etc	National Wage Case August 1988
27	First Aid	National Wage Case
33	Ambulance Chest Fares and Travelling Expenses	August 1988 National Wage Case August 1988

Dated 16 November 1988

CHRISTOPHER BRENDON
Deputy Industrial Registrar

N.N.-8878098

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE UNDER SUB-SECTION 49A (3) IN RELATION TO VARIATION OF A COMMON RULE

In the matter of the PARKING STATIONS ETC. EMPLOYEES (ACT) AWARD 1983

C No. 21171 of 1988

And in the matter of the variation of the award dated 10 October 1983 in the above matter.

Notice is hereby given:

- (a) that on 7 November 1988 the Commission varied the terms of the above mentioned award referred to in the Schedule as set out in the Schedule;
- (b) that the variation will be a common rule in the Australian Capital Territory in the industry in respect of which the dispute arose with effect from the first pay period to commence on or after 27 September 1988; and

(c) that any person or organisation having an objection to the variation binding that person or organisation and desiring to be heard in relation to that objection is invited to lodge with the Commission a notice of that objection within twenty-eight days after the date specified in paragraph (a).

A copy of the award may be inspected at the office of the Registrar. Objections should be lodged with the Registrar at 4th Floor, CML Building, University Ave, Canberra, by 4.30 p.m. on 5 December 1988.

SCHEDULE TERMS VARIED

Clause No.	Subject	Substance of variation
	PRINT NO. H5	020
4	Wages	Second Tier Increase
19	Uniforms	Second Tier Increase
21	Wages etc. Time and payment of	Second Tier Increase
22	Terms of engagement	Second Tier Increase
27	Holidays	Second Tier Increase
28	Sick Leave	Second Tier Increase

Dated 16 November 1988

CHRISTOPHER BRENDON

Deputy Industrial Registrar

N.N.-8878099

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

COMMON RULE

- 1. Application has been made that the following Award shall be declared common rule in the Australian Capital Territory in the industry in connection with which the dispute arose which led to the making of the Award. The said industry is described in the said Award.
- 2. That Award is: Actors etc. (Television) Award 1979
- 3. A copy of the Award may be inspected free of charge at the office of the Registrar at the address shown in item 4.
- 4. The application will be heard at 2.15 p.m. Monday, 5 December 1988 at 4th Floor, CML Building, University Ave. Canberra ACT.
- 5. All persons and organisations interested and desiring to be heard may appear or be represented before the Commission and shall be heard.
- 6. Persons and organisations not so appearing or represented will be bound by any declaration made by the Commission in the matter which is applicable to them.
- 7. Compliance with Regulation 35 is not required.

Dated 15 November 1988

CHRISTOPHER BRENDON
Registrar

N.N.-8878100

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

COMMON RULE

1. Application has been made that the following Award shall be declared common rule in the Australian Capital Territory in the industry in connection with which the

dispute arose which led to the making of the Award. The said industry is described in the said Award.

- 2. That Award is: Actors (Theatrical) Award 1981
- 3. A copy of the Award may be inspected free of charge at the office of the Registrar at the address shown in item 4.
- 4. The application will be heard at 2.15 p.m. Monday, 5 December 1988 at 4th Floor, CML Building, University Ave, Canberra ACT.
- 5. All persons and organisations interested and desiring to be heard may appear or be represented before the Commission and shall be heard.
- 6. Persons and organisations not so appearing or represented will be bound by any declaration made by the Commission in the matter which is applicable to them.
- 7. Compliance with Regulation 35 is not required.

Dated 15 November 1988.

CHRISTOPHER BRENDON

Registrar

N.N.-8878101

IN THE AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

In the matter of an application by the Federated Miscellaneous Workers Union of Australia for the CHILD CARE INDUSTRY (AUSTRALIAN CAPITAL TERRITORY) AWARD 1985 to be declared a common rule in the Australian Capital Territory

(C No. 21686 of 1988)

Mr Commissioner Laing Canberra, 9 November 1988

DECLARATION

- 1. That the whole of the terms of the Child Care Industry (Australian Capital Territory) Award including variations thereto, be a common rule of the relevant industry in the Australian Capital Territory and shall be binding on all employers in the said industry in respect of the employment by them of employees in the classifications for which provision is made in the said award and shall be binding on all such employees.
- 2. That this declaration shall not apply to the ACT Health Authority in respect of employees under the Capital Territory Health Commission Ordinance 1975 (as amended) and Calvary Hospital ACT Incorporated.
- 3. That the foregoing declaration shall operate from 9 November 1988.

By the Commission,

COMMISSIONER

Appearances:

- D. Cseti for the Federated Miscellaneous Workers Union of Australia.
- B. G. Cooke for Confederation of ACT Industry on behalf of respondents.

Date and place of hearing: 1988; Canberra;

(1) Print G0220

N.N.-8878102

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE UNDER SUB-SECTION 49A (3) IN RELATION TO VARIATION OF A COMMON RULE

In the matter of the NURSES PRIVATE **EMPLOYMENT (ACT) AWARD 1972**

C No. 90275 of 1988

And in the matter of the variation of the award dated 16 August 1972 in the above matter.

Notice is hereby given:

- (a) that on 8 November 1988 the Commission varied the terms of the above mentioned award referred to in the Schedule as set out in the Schedule;
- (b) that the variation will be a common rule in the Australian Capital Territory in the industry in respect of which the dispute arose with effect on and from 13 October 1988; and
- (c) that any person or organisation having an objection to the variation binding that person or organisation and desiring to be heard in relation to that objection is invited to lodge with the Commission a notice of that objection within twenty-eight days after the date specified in paragraph (a).

A copy of the award may be inspected at the office of the Registrar. Objections should be lodged with the Registrar at 4th Floor, CML Building, University Ave, Canberra, by 4.30 p.m. on 6 December 1988.

SCHEDULE

TERMS VARIED

Clause No.	Subject	Substance of variation
	PRINT NO. H52	76
4	Wages	National Wage Case August 1988
4A	No Extra Claims	National Wage Case August 1988
5 (a) (b) (c) (d) (i) (ii) (iii)	Special Allowance	National Wage Case August 1988

Dated 11 November 1988.

DAVID SINCLAIR Deputy Industrial Registrar

N.N.-8878103

AUSTRALIAN CONCILIATION AND ARBITRATION COMMISSION

Conciliation and Arbitration Act 1904

NOTICE UNDER SUB-SECTION 49A (3) IN RELATION TO VARIATION OF A COMMON RULE

In the matter of the NURSES PRIVATE **EMPLOYMENT (ACT) AWARD 1972**

C No. 90278 of 1988

And in the matter of the variation of the award dated 16 August 1972 in the above matter.

Notice is hereby given:

- (a) that on 4 November 1988 the Commission varied the terms of the abovementioned award referred to in the Schedule as set out in the Schedule;
- (b) that the variation will be a common rule in the Australian Capital Territory in the industry in respect of which the dispute arose with effect on and from 23 August 1988; and
- (c) that any person or organisation having an objection to the variation binding that person or organisation and desiring to be heard in relation to that objection is invited to lodge with the Commission a notice of that objection within twenty-eight days after the date specified in paragraph (a).

A copy of the award may be inspected at the office of the Registrar. Objections should be lodged with the Registrar at 4th Floor, CML Building, University Ave, Canberra, by 4.30 p.m. on 1 December 1988.

SCHEDULE TERMS VARIED

Clause No.	Subject	Substance of variation
	PRINT NO. H52	275
4	Wages	Second Tier Increase
5 (d)	Special Allowance	Second Tier Increase

Dated 11 November 1988.

DAVID SINCLAIR
Deputy Industrial Registrar

CA03I048 CRACTOIR\$

N.N.-8878104

COMMONWEALTH OF AUSTRALIA

Commonwealth Teaching Service Act 1972

DECLARATION UNDER SUBSECTION 50A (1)

WHEREAS on 13 October 1988, Allan Clyde Holding, Minister of State for the Arts and Territories, certified in accordance with subsection 50A (1) of the Commonwealth Teaching Service Act 1972 ('the Act') that the performance of teaching and associated duties, being a function that has been performed by persons appointed or engaged as Commonwealth Teaching Service staff under the Act in the Canberra School of Art and the Canberra School of Music is to be performed by the Canberra Institute of the Arts, a Commonwealth authority within the meaning of subsection 4 (1) of the Act:

Now, therefore I, RAE MARTIN TAYLOR, Secretary to the Department of Industrial Relations, declare, pursuant to subsection 50A (1) of the Act that the officers and employees specified in the Schedule attached to this instrument are in the employment of the Canberra Institute of the Arts and that 16 November 1988 is the specified date for the purposes of subsection 50A (2).

Dated 16 November 1988.

R. M. TAYLOR

Secretary

Department of Industrial Relations

SCHEDULE

Permanent Officers

Boynes, Robert John
Dabros, Ante
Hansen, Finn Ragnar
Harry, Patricia Joyce
Herel, Peter
Hill, Vernon Kingsley
Huff-Johnston, William
Langridge, Anne Balcome
Kleinert, Ingo Dietmar
Mann-Hay, Gillian
Martin, Amanda Laura De Rose
Meadham, Christopher William
Reid, John Colin
Schmeisser, Jorg Giselher
Swen, Hiroe

Temporary Fixed-Term Employees

Butler, Michael Anthony Crisp, Deborah Gay Crisp, Leanne Christine Croft, Christopher (LWOP) De Borah, Anna Bommett, Leonard Bertram Douglas, Anne Eaton, Janenne Pearl Ewington, Julie Mary Gadd, Charmian Jane Geier, Helen Maria Greenwood, Garry Grey-Smith, Mark Fancis Hawkey, William Haynes, Peter John Ingham, George Richard Johnson, Donald Duane Kain, Timothy Patrick Kean, Roselyn Annette Kleinert, Sylvia Joyce Legrand, Michael Martin Leibbrandt, Ingrid Olga Lewis, Barbara Dianne Luxton, John Fraser Maconaghie, Ronald Dereck McClure, Elizabeth McDonald, Hector Harcourt McElhinny, Christopher Michael Mulcahy, Michael Noone, Michael John Nuttall, David Christopher Palmgvist, Bengt-Olov Plunkett, Paul Pratt, Daryl Loring Robertson-Swann, Ronald Charles Taylor, Viginia Teakel, Wendy Evelyn Tremblay, Theodore Paul Trew, Dennis Michael Van Hove, Marie Celeste Vivian, Alan Edward Watt, Alan Hugh Douglas West, Susan Dawn Worrall, David Robert

N.N.-8878112

Industry, Technology and Commerce

COMMONWEALTH OF AUSTRALIA
DECLARATION BY THE MINISTER FOR
INDUSTRY, TECHNOLOGY AND COMMERCE,
PURSUANT TO SUB-SECTION 6 (7) OF THE
BOUNTY (METAL WORKING MACHINES AND
ROBOTS) ACT 1985

I, JOHN NORMAN BUTTON, the Minister of State for Industry, Technology and Commerce, pursuant to subsection 6 (7) of the Bounty (Metal Working Machines and Robots) Act 1985, hereby declare that this Act is to have effect as if there were substituted for the definition of 'modification' in subsection 4 (1) the definition set out below:

'modification', in relation to bountiable equipment B,

- (a) a conversion of the equipment that, in the opinion of the Minister, will substantially improve the designed capability of the equipment; or
- (b) the conversion of the equipment from equipment that is not designed to form part of a flexible manufacturing cell or system into equipment intended to form part of such a cell or system,

but does not include:

- (c) the conversion of the equipment in the course of the normal manufacturing of the equipment; or
- (d) retrofit manufacture of, or involving, the equipment.

This declaration shall take effect on and from the first day of July 1985.

Dated this 28th day of October 1988.

JOHN NORMAN BUTTON

Minister of State for Industry, Technology and Commerce

Bounty (Computers) Act 1984

NOTICE OF DECLARATION OF EQUIPMENT

I, BARRY OWEN JONES, Minister for Science Customs and Small Business hereby declare, pursuant to subsection 5 (1) of the Bounty (Computers) Act 1984, that each class of computer equipment appearing in Column 1 of the Schedule hereunder shall, as from the date specified opposite it in Column 2 of the said Schedule, be a class of computer equipment to which paragraph (e) to the definition of 'bountiable equipment' in subsection 3 (1) of the Bounty (Computers) Act 1984 applies.

Dated this 11th day of November 1988.

BARRY O. JONES

Minister for Science, Customs and Small Business

Reference in the Schedule hereunder to the word 'subheading' followed by a set of numerals shall be a reference to the relevant subheading, as indicated by those numerals, in Schedule 3 to the Customs Tariff Act 1987 as in force on the date of this Notice and as amended from time to time thereafter.

SCHEDULE

Column 1 Class of computer equipment	Column 2 Date of effect
Automatic Monitoring and Control Systems for in-motion weighing ma- chines to which if imported, subhead- ing 9032.89.90 of Schedule 3 of the Customs Tariff Act 1987 would apply.	6 July 1984

N.N.-8878106

Social Security

COMMONWEALTH OF AUSTRALIA

Social Security Act 1947

APPOINTMENT OF SENIOR MEMBER

I, SIR NINIAN MARTIN STEPHEN, Governor-General of the Commonwealth of Australia, acting with the advice of the Federal Executive Council, pursuant to subsection 218 (1) of the Social Security Act 1947, hereby appoint Anne Coghlan as National Convener, Social Security Appeals Tribunal for five years.

Dated 8 November 1988.

N. M. STEPHEN Governor-General

By His Excellency's Command, **BRIAN HOWE** Minister of State for Social Security (Ex. Min. No. 12)

N.N.-8878107

Transport and Communications

COMMONWEALTH OF AUSTRALIA

Telecommunications Act 1975

APPOINTMENT OF PART-TIME COMMISSIONER OF THE AUSTRALIAN TELECOMMUNICATIONS COMMISSION

I, SIR NINIAN MARTIN STEPHEN, Governor-General of the Commonwealth of Australia acting with the advice of the Federal Executive Council pursuant to subsection 22 (2) of the Telecommunications Act 1975, hereby appoint Graham Charles Evans to be a part-time Commissioner of the Australian Telecommunications Commission referred to in paragraph 22 (1) (b) of that Act.

Dated 27 October 1988.

N. M. STEPHEN Governor-General

By His Excellency's Command, **RALPH WILLIS** Minister of State for Transport and Communications

N.N.-8878108

COMMONWEALTH OF AUSTRALIA

TELECOMMUNICATIONS ACT 1975

DETERMINATION OF RENTAL AND CHARGES

I, Mario Dominic Zilko, the General Manager, Operator Assisted Services of the Australian Telecommunications Commission, being a person to whom the Commission has, by instrument in writing under Section 33 of the Telecommunications Act 1975, delegated its powers and functions under the Act in pursuance of Section 11 of the Act, hereby make the following determination.

 The determination of rentals and charges made by the Commission on 21 August 1975*, as varied to date Ø, is further varied by including at the end of Item 1L of Schedule 1 the following item:

"ITEM 1M CHARGES FOR TELEPHONE ANSWERING SERVICE

The fee payable by a subscriber to a personalised telephone answering service (TAS) provided by the Commission, not being a service as described in Item IF of this Schedule shall be in accordance with the following:

(a) An annual fee in respect of a personalised telephone answering service as follows -

(i) For each standard message telephone answering service 600.00

(ii) For each premium message telephone answering service 720.00

with a minimum charge equal to one-twelfth of the annual rate prescribed in paragraph (a)

(b) a fee in respect of each printed report of messages processed by a telephone answering service requested by the subscriber of

10.00"

This determination will take effect on and from 1 November 1988.

DATED this thirty-first day of October 1988.

M.D. Zilko
Delegate of the Australian
Telecommunications Commission

- * Notified in Gazette No. S170 of 29.8.75.
- Ø For previous amendments see footnote Ø appearing on Pages 2223 - 2229 of Gazette No. GN36 of 28.9.88.

AUSTRALIAN BROADCASTING TRIBUNAL

BROADCASTS BY MR RON CASEY AND MR LES THOMPSON ON 2KY—NOTICE OF INQUIRY

IP/88/196 November 1988

PUBLIC SUBMISSIONS INVITED

Following a Federal Court decision, the Australian Broadcasting Tribunal is commencing a new inquiry into certain broadcasts by Mr Ron Casey and Mr Les Thompson on radio station 2KY ('2KY'). The inquiry is also being held in consequence of an application dated 23 March 1988 to the Tribunal by the Anti-Discrimination Board, the Ethnic Affairs Commission of New South Wales, the Human Rights and Equal Opportunity Commission, the Ethnic Communities Council of New South Wales and the Federation of Ethnic Communities Councils of Australia ('the joint applicants') requesting the exercise by the Tribunal of its powers under sections 85, 99, 101 and 119 of the Broadcasting Act 1942 ('the Act') in respect of a broadcast by Mr Ron Casey on 2KY at 10.50 a.m. on 18 February 1988. The joint applicants alleged this broadcast was in breach of Radio Program Standard 3 ('RPS3'). Further particulars of the application may be found in Appendix A to the Tribunal's information paper relating to the inquiry.

The Tribunal is interested in your views on whether any of the broadcasts (listed at Issue 1, below) breached RPS3 and the Act. It would also like to know what action, if any, you believe the Tribunal should take in accordance with sections 85, 99, 101 and 119 of the Act if it is found that any of these broadcasts breached RPS3 and the Act.

The inquiry is to be conducted in accordance with section 17C of the Act and the issues to be considered in the inquiry are:

1. Whether any of the programs respectively presented on radio station 2KY ('2KY') by Ron Casey on 23 April 1987, 24 April 1987, 19 October 1987 and 18 February 1988 and by Les Thompson on 22 October 1987 failed to comply with Radio Program Standard 3 ('RPS3').

If any of the programs did fail to comply with RPS3:

- 2. Whether the licensee of 2KY contravened subsection 99 (1A) of the Broadcasting and Television Act 1942 by failing to supervise the broadcasting of the programs in such a manner as to ensure that RPS3 was complied
- 3. Whether the Tribunal should give any direction to the licensee of 2KY pursuant to subsection 99 (2) of the Act for the purpose of ensuring that program standards are complied with in relation to the broadcasting of programs by 2KY and in relation to programs to be broadcast by 2KY and, if so, the form of any such direction.
- 4. Whether the Tribunal should impose further conditions in respect of the licence held by the licensee of 2KY pursuant to subsection 85 (1) of the Act requiring the licensee to adopt management procedures relating to talk-back radio programs broadcast by 2KY for the purpose of preventing breaches of program standards and the Act.
- 5. Whether the Tribunal should issue a written reprimand or admonishment to the licensee of 2KY pursuant to subsection 101 (1) of the Act and, if so, the form of any such reprimand or admonishment.

- 6. Whether the Tribunal should also direct the licensee of 2KY pursuant to subsection 101 (2) of the Act to publish the reprimand or admonishment and, if so, in what form and manner.
- 7. (a) Whether the Tribunal should give a direction pursuant to subsection 119 (1) of the Act:
 - (i) prohibiting Ron Casey and/or Les Thompson from presenting programs in live broadcasts;
 - (ii) imposing restrictions on the presenting by Ron Casey and/or Les Thompson of programs in live broadcasts;
 - (iii) prohibiting any person who passed or selected for broadcasting the whole or a part of the programs from passing or selecting programs for broadcasting;
 - (iv) imposing restrictions on the passing or selecting by that person of programs for broadcasting.
- 7. (b) If the Tribunal decides to give any such direction, whether the direction should have effect indefinitely, or for a specified period.

You are invited to make submissions in relation to the inquiry. Your submissions should reach the Tribunal's office at Tandem House, 76 Berry St, North Sydney 2060 or be sent to the General Manager, Australian Broadcasting Tribunal, PO Box 1308, North Sydney 2059, before 5.00 p.m. on 20 January 1989. Each submission will become a public document and will be attached to the inquiry file unless access is restricted by the Tribunal for the whole or part of it. The inquiry files which contain copies of the application for the inquiry and other relevant information can be inspected at the Tribunal's North Sydney office.

A guide and information package is available by contacting the Tribunal on (02) 959 7941 or 959 7946.

N.N.-8878109

Treasurer

Insurance Act 1973

NOTIFICATION OF GRANT OF AUTHORITY IN **ACCORDANCE WITH SECTION 28**

In pursuance of section 28 of the Insurance Act 1973, I, RICHARD MATTHEW BEETHAM, Insurance and Superannuation Commissioner, hereby give notice that I did on 9th day of November 1988, pursuant to section 23 of the Insurance Act 1973, grant to Continental Pacific Insurance Company (Australia) Limited, the Registered Office of which is situated at Level 32, 100 Miller St, North Sydney, an authority to carry on insurance business.

In accordance with section 122 of the Insurance Act 1973, copies of grants of authority are contained in the Register of Authorised Insurers which can be inspected at my Office.

Dated this 9th day of November 1988.

RICHARD M. BEETHAM

Commissioner

N.N.-8878110

Commonwealth of Australia Gazette No. GN 44, 23 November 1988

Australian Banking Statistics

Savings Banks

STATEMENTS PREPARED FROM RETURNS DELIVERED TO THE STATISTICIAN UNDER SECTION 53 OF THE BANKING ACT 1959 OR UNDER SPECIAL ARRANGEMENTS WITH BANKS NOT SUBJECT TO THAT ACT

LIABILITIES AND ASSETS WITHIN AUSTRALIA OF EACH SAVINGS BANK AT THE END OF SEPTEMBER 1988(a) (\$'000)

	Liabilities	·		Assets		-												
		Balances due to					Australia	n public s	ecurities			Loans	Cheques and bills of other					
	Demonisons	other banks, bills payable and all other		Coin, bullion and Aus-	Deposits with	Deposits in Australia with		wealth and	States	Local and semi- govern- mental	Other	to authorised dealers in the	banks and balances with and due from other	dealers in short-tern market), i	uthorised the money	Bank premises, furniture	Bills receiv- able and all	
Savings Banks (b)	Depositors' balances	(c)	Total	tralian notes	Reserve Bank	Trading Banks	Treasury Bills	Treasury Notes	Other	author- ities	securi- ties	money market	banks (d)	Housing	Other	and sites	other assets	Tota
Commonwealth	. 15 227 007	705 650	15 932 657	131 859		19 441		130 000	1 708 925	2 653 440	208 653	17 800	104 605	9 501 844	1 428 948	846 622	126 936	16 879 07
State Banks (e)— S.B. of Vic S.B. of S.A	. 10 322 956 . 1 596 110		14 327 527 2 566 941	68 197	725	398		498 112	996 648 543 642	394 561	103 859		125 269	5 479 488	3 199 542	405 093	3 683 075	14 954 96
R. & I. of W.A. Tasmania Bank	1 303 113 690 531	17 275	1 303 113 707 806	596 5 807	51	31 282		19 525 13 625	153 295 62 079	40 020 106 811	13	16 800	2 499	2 023 299 933 843 216 813	255 661	23 009	222 13 126	
Total, State	. 13 912 710	4 992 677	18 905 387	74 600	776	31 680		531 262	1 755 664	541 392	103 872	16 800	127 768	8 653 443	3 455 203	428 102	3 696 423	
Trustee Banks— S.B. of Tas	. 411 772	29 705	441 477	3 128	104	10 011		3 000	35 000	95 619	37	8 900	22 072	116 061	152 306	7 985	10 557	
Private Banks— A.N.Z. S.B.	. 5 968 138	375 012		197		59 011		590 000	335 555	240 485	47 610		500	4 370 841	875 036	.,,03	10 337	6 519 23
National Aust. S.B. Westpac S.B Advance	. 7 885 404 . 10 698 554 . 3 827 316	526 778 117 738 384 052	10 816 292			69 485		341 500 1 428 050	733 493 382 322	992 280 1 170 070	500 195		500	4 710 441 7 082 282	1 762 453 814 300	131 464		8 543 51 11 413 17
B.N.Z. S.B	. 78 346 . 2 051 391	6 038 434 015	84 384 2 485 406	13 928 3 929	327	64 219 28 325		27 287 136 724	526 951 11 617 155 723	108 040 16 632 67 440	98 119 49 910	46 800	225 295 251	2 132 515 59 271	910 813	205 209	467	4 535 94 88 25
Citibank		34 942 28 999	1 355 091 529 517	1 460	 7		••	72 000	109 000 51 416	3 980		43 550 25 145	63 641 27 53 550	1 741 857 1 067 253 297 020	329 565 168 061 105 750	19 626	-1 294 6 728	1 423 06
Metway Nat. Mutual Old S.B	. 1 129 265 . 2 021 583 . 262 234	43 126 151 287 21 554	1 172 391 2 172 870 283 788	4 638 13 517	7	30 351 1 000		19 844 4 979	133 531 266 452	9 872 46 020	90 16 697	12 030	1 257 61 358	925 591 1 467 237	86 379 441 696	13 313 17 834 27 996	11 423 7 456 307	
Total, Private				27.660	241	660	• •		35 808				••	262 880	4 273		167	303 78
Total, All Banks					341	253 051			2 741 868			127 525		24 117 188	5 498 342	415 442	539 046	39 625 17
I UMI, All Banks	03 294 38 /	/ 831 3/3	13 143 960	24 / 256	1 221	314 183		3 284 646	6 241 457	5 945 270	525 683	171 025	660 824	42 388 536	10 534 700	1 698 151	4 372 962	76 206 OI

⁽a) Particulars for The State Bank of Victoria, The Rural and Industries Bank of Western Australia (Savings Bank Division) and The Hobart Savings Bank relate to the end of the month, those for Tasmania Bank to the last Monday in the month, and those for the other Savings Banks to the last Wednesday in the month. Excludes capital and reserves and inter-branch accounts. (b) For legend on abbreviations of Bank names see footnote on page 2633. (c) Includes provisions for contingencies. (d) Includes deposits with and loans to specified banks other than Trading Banks. (e) The information for the State Savings Banks is supplied and published under a special arrangement with the banks and not under the Banking Act.

Note: Any discrepancies between totals and sums of components are due to rounding.

DEPOSITORS' BALANCES IN EACH STATE AND THE TERRITORIES FOR EACH SAVINGS BANK AT THE END OF SEPTEMBER 1988

(Excludes inter-branch accounts)

(\$'000)

Savings Banks†	New South Wales	Victoria	Queensland	South Australia	Western Australia	Tasmania	Northern Territory	Australian Capital Territory	Other Territories
Commonwealth	6 702 585	3 170 314	2 743 849	913 898	1 106 567	245 524	79 864	254 321	10 085
State Banks (e)— S.B. of Vic	5 677	10 322 956 	236	1 586 471	1 297 436	690 531	9 403		
Total, State	5 677	10 322 956	236	1 586 471	1 297 436	690 531	9 403		
Trustee Banks— S.B. of Tas						411 772			
Private Banks— A.N.Z.S.B	1 339 251 2 512 646 4 519 238	2 454 058 3 138 819 2 294 226	958 334 1 349 321 2 072 554	567 373 336 254 465 668	400 872 407 501 943 306	134 957 34 783 185 292	60 695 27 882 84 213	52 598 78 198 126 890	7 167
Advance	3 459 841 59 693 90 430	270 648 5 160 432 484	96 679 11 362 83 044	148 838 1 776	1 293 1 443 657				
Citibank	935 840 15 138	117 553	181 992 1 129 265	46 263 	16 586 	6 082	••	15 833 485 380	
Nat. Mutual	1 626 342	380 658	4 971 262 234	6 081	2 448		•••	1 083	
Total, Private	14 558 419	9 093 606	6 149 756	1 424 401	3 215 663	361 114	172 790	759 982	7 167
Total, All Banks	21 266 681	22 586 876	8 893 841	3 924 770	5 619 666	1 708 941	262 057	1 014 303	17 252

[†] Commonwealth-Commonwealth Savings Bank of Australia.

S.B. of Vic.—The State Bank of Victoria.
S.B. of S.A.—The State Bank of South Australia.

R. & I. of W.A.—The Rural and Industries Bank of Western Australia (Savings Bank Division).

Tasmania Bank-Tasmania Bank.

S.B. of Tas.—The Hobart Savings Bank (trading as The Savings Bank of Tasmania).

A.N.Z.S.B.—Australia and New Zealand Savings Bank Limited.

National Aust. S.B.—National Australia Savings Bank Limited.

Westpac S.B.—Westpac Savings Bank Limited.

Advance-Advance Bank Australia Limited.

B.N.Z.S.B.-Bank of New Zealand Savings Bank Limited.

Challenge-Challenge Bank Limited.

Citibank-Citibank Savings Limited.

Civic Advance-Civic Advance Bank Limited.

Metway-Metway Bank Limited.

Nat. Mutual-National Mutual Royal Savings Bank Limited.

Qld S.B.-Bank of Queensland Savings Bank Limited.

Australian Banking Statistics

Trading Banks

STATEMENTS PREPARED FROM RETURNS DELIVERED TO THE STATISTICIAN UNDER SECTION 52 OF THE BANKING ACT 1959 OR UNDER SPECIAL ARRANGEMENTS WITH BANKS NOT SUBJECT TO THAT ACT

LIABILITIES

AVERAGES OF THE LIABILITIES WITHIN AUSTRALIA OF EACH BANK AT THE CLOSE OF BUSINESS ON WEDNESDAYS, 7, 14, 21 AND 28 SEPTEMBER 1988 (a) (\$'000)

	Deposits re	payable in	Australia													Bills	
	Fixed			-	Current								_			payable and all	
					Bearing intere	st			Not bearing	interest						other liabilities	
ink(b)	Common- wealth and State Govern- ments	Other banks	Other	Total fixed deposits	Common- wealth and State Govern- ments	Other banks	Other	Total	Common- wealth and State Govern- ments	Other banks	Other	Total	Total current deposits	Total deposits	Balances due to other banks	to the public (excluding share- holders' funds) (c)	To liabili
ajor Banks— A.N.Z. Commonwealth National Westpac	19 709 14 725 119 363 3 321	15 1 969 5 981	8 805 399 7 123 328 8 021 060 5 591 310	8 825 122 7 140 022 8 146 403 5 594 630	4 861 29 600 ~ 14 088 25 796	165 11 876 1 329 39 565	730 123 527 463 1 327 686 2 174 329	735 149 568 939 1 343 103 2 239 689	4 558 13 676 2 588 52 534	437 504 63 299 274 958 441 091	2 993 641 2 910 194	3 536 873 3 070 615 3 187 739 4 765 039	3 639 554 4 530 842	10 779 576 12 677 245	630 505 74 875 931 629 558 806	16 160 965 10 403 874 13 251 387 12 390 121	21 258 26 860
Total, Major	157 117	7 964	29 541 096	29 706 177	74 345		4 759 600		73 355			14 560 267				52 206 347	
her Banks-											13 270 001	14 300 207	17 447 147	49 133 324	2 193 614	32 200 347	103 333
America Aust. Bank Bankers Trust Barclays Chase A.M.P.	 43	49 378	91 312 3 351 67 107 212 389 328 340	91 312 3 351 67 107 212 389 377 761		7 340	73 400 52 832 264 471	73 400 52 832 271 811			5 419 5 380 2 437 7 674 6 846	5 419 5 380 2 437 7 674	5 419 78 781 2 437 60 505	96 732 82 131 69 544 272 895	33 156 126 328 198 494 162 668	318 746 84 874 893 124 1 008 400	293 1 161 1 443
China	3 534	29 550 3 000	8 574 49 205 91 109 44 967	8 574 78 755 97 643 44 967	• •		5 615 21 432	5 615 21 432			21 248 3 311	6 846 21 248 3 311	278 657 5 615 42 680 3 311	656 418 14 189 121 435 100 953	90 985 20 358 24 231 16 557	943 677 4 642 1 512 213 265 062	1 65° 382
I.B.J	363	188	78 130 47 975 101 574	78 130 48 525 101 574	•••		6 005 21 192 106 257	6 005 21 192 106 257	••		8 289 7 597 44 977	8 289 7 597 44 977	117 575 6 295 28 790 151 233	162 542 84 424 77 315 252 807	47 667 22 750 262 864 126 225	1 253 422 326 955 365 187 287 473	434 70:
Nat. Mutual		••	60 662 82 476 330 046 217 103	60 662 82 476 330 046 217 103	•••		4 473 74 161 8 600 296 756	4 473 74 161 8 600 296 756			99 14 012 72 834	99 14 012 72 834	4 572 74 161 22 612 369 590	65 233 156 636 352 658 586 693	30 462 164 531 134 871 159 920	194 863 1 089 928 955 571 1 062 478	1 41 1 44
Paris P.I.B.A. Queensland Queensland Queensland		2 220 500	204 206 155 845 72 653 134 294	206 426 156 345 72 653 134 294		65	6 133 53 738 9 985	6 133 53 803 9 985	119	4 916 	29 794	34 710 119	40 843 53 922	247 269 156 345 126 574	148 451 45 668 10 343	727 446 476 032 50 116	1 12 67 18
tandard Okyo tate Bank N.S.W. (d) tate Bank S.A. (d)		• • • • • • • • • • • • • • • • • • • •	162 746 95 659 3 295 613	162 746 95 659 3 295 613	91 268		45 483 2 486 301 164	45 483 2 486 392 432	10 508	73 	3 966 8 886 549 736	3 966 8 959 560 245	9 985 49 450 11 445 952 676		46 851 63 263 39 761 168 771	249 781 674 388 250 078 4 055 967	94 39
tate bank S.A. (a)	45 151		1 608 566 1 942 944	1 608 566 1 988 095	1 470		200 833 107 718	200 833 109 187	98		73 198 301 983	73 198 302 081	274 031 411 268	1 882 596 2 399 363	68 230 88 453	2 942 830 649 265	4 89
Total, Other	49 090	84 835	9 486 843	9 620 769	92 738	7 405	1 780 299	1 880 441	10 725	4 989	1 159 694	1 175 408	3 055 849	12 676 617	2 301 855	20 642 513	35 62
Total, All Banks	206 207	92 800	39 027 939	39 326 946	167 082	60 340	6 539 899	6 767 321	84 080	1 221 840	14 429 755	15 735 675			4 497 669	72 848 860	
D.B	1 100	14 750	283 260 1 507 115	283 260 1 522 965	::			•••						283 260	29 709 127 918	87 516 33 608	40

⁽a) Excludes shareholders funds, inter-branch accounts and contingencies. (b) For legend on abbreviations of Bank names see footnote on page 2640. (c) Includes the Australian dollar equivalent of foreign currency balances. (d) The information for these banks is supplied and published under a special arrangement with the banks and not under the Banking Act.

Note: Any discrepancies between totals and sums of components are due to rounding.

ASSETS
AVERAGES OF THE ASSETS WITHIN AUSTRALIA OF EACH BANK AT THE CLOSE OF BUSINESS ON WEDNESDAYS, 7, 14, 21 AND 28 SEPTEMBER 1988 (a) (\$'000)

	Coi	in and bulli	on			Australian Pt	ublic Securiti	es			Loans to authorised	Statutory Reserve	Cheques and bills of other banks and balances	Loans (other than loans to authorised dealers in the short-term money		Bills	
						Commonweal	th and State	5	Local and semi-		dealers in the	Deposit account	with and due from	market), advances	Bank	receivable and	
Bank(b)		Gold	Coin	Australian notes	Cash with Reserve Bank	Treasury bills	Treasury notes	Other securities	govern- mental authorities	Other securities (c)	short-term money market	with Reserve Bank	other banks (c)	and bills discounted (c)	premises, furniture and sites	all other assets (c)	Tota asse
Major Banks—	_									2.254.142	41 746	664 918	251 145	12 182 341	304 895	14 051 510	33 324 66
A.N.Z		9	14 411	155 800	218 428		1 309 275	1 876 048	3 773	2 254 142 1 104 587	41 745 31 875	564 931	399 281	10 590 076	801 453	7 500 375	23 516 30
Commonwealth .			8 973	98 807	184 645	• •	461 100	1 766 426 1 643 775	137 989	2 510 002	128 161	654 354	289 108	13 236 434	400 269	10 986 246	31 454 54
National .		• • •	13 863	167 351	280 318 219 576		1 006 675 924 000	1 406 789	137 969	2 279 876	82 301	665 478	136 600	14 734 523	541 899	7 745 529	28 938 9
Westpac			20 948	181 395	902 967	···	3 701 050	6 693 038	141 767	8 148 608	284 082	2 549 680	1 076 133	50 743 374	2 048 515	40 283 660	117 234 4
Total, Major	٠.	9	58 193	603 353	902 907		3 701 030	0 073 038	141 707	0 140 000	204 002	2377 000	10,0100				
other Banks—				3	1.053		23 414	34 414	25	26 431	13 456	3 084	22 719	258 002	3 590	133 630	5198
America		• •	4	82	1 053 1 446		41 336	3 116	4 033	16 446	11 250	4 188	9 419	179 436	2 641	84 647	358 (
Aust. Bank	•	• •		02	82	• •	41 330	145 883	4 033	122 335	36 850	4 007	202 041	617 275		181 723	1 310 1
Bankers Trust		• •	18	490	4 165		81 958	7 889	5 442	83 300	105 955	12 305	19 235	674 426	11 612	556 059	1 562 8
Barclays	•	• • •	62	972	10 970	• • • • • • • • • • • • • • • • • • • •	150 600		98 111	63 019	14 775	31 713	147 098	808 555	29 208	382 683	1 737 7
China	•			188	260		6 250			2 200	8 773	756	495	131 694	4 787	82	155 4
Citibank	•		2	1 089	5 512		58 500	133 750		217 268	89 970	5 369	136 716	501 796	: :	1 136 435	2 286 4
Deutsche				78	1 850		18 750	13 202		16 971	37 320	5 644	2 219	512 289	3 805	79 812	691 9
HongKong			23	699	2 795		137 143	27 215		77 425	11 450	7 903	22 737	659 833	9 733	736 803 57 427	1 693 7 618 1
I.B.J				39	1 388			45 922	11 708	30 796	3 325	3 915	22 906	438 418	2 276 3 789	61 243	824 9
Lloyds N.Z.A				68	1 548		39 000	:	5 458	17 102	36 569	4 531	40 246 46 421	615 367 476 206	6 920	119 434	840
Macquarie			500	245	2 831		50 230	25 156	35 824	44 111	24 450 1 083	8 649 5 005	1 026	301 192	1 782	57 326	443
Mitsubishi				123	23		20 000	30 300	26 000 3 031	147 590	1 003	6 227	16 481	349 789	1 702	966 670	1 654
Nat. Mutual				126	2 488		3 248 29 835	159 239 21 315		149 135	183 534	17 407	32 390	1 171 384	15 294	545 075	2 171
Nat West .		• •	36	135	5 823 210		63 400	84 374	• •	149 133	118 248	27 334	33 913	883 513	10 029	984 323	2 206 5
New Zealand			35 20	1 141 517	3 517		100 683	20 000		36 587	31 575	10 192	109 341	521 798	4 832	483 653	1 322
Paris			20	317	2 911	• • • • • • • • • • • • • • • • • • • •	9 480	36 525	• • • • • • • • • • • • • • • • • • • •	10 002	9 765	8 529	365 246	280 181	559	451	723 (
P.I.B.A			340	3 491	1 839		3 250	15 100	4 842	40 423	2 730	5 082	10 586	109 447	9 166	45 730	252 (
Singapore			4	313	3 048		6 131	52 959	6 580		1 269	9 506	2 511	315 000	3 571	80 272	481
Standard		• • • • • • • • • • • • • • • • • • • •	17	529	4 199		39 514	55 924		85 701	8 625	13 232	2 165	315 081	5 774	563 337	1 094 (
Tokyo			3	150	1 586		27 818			11 372	19 263	4 672	17 834	186 042	3 625	159 185	431 : 10 542 (
State Bank N.S.W. (4	d)		2 700	40 242	610		437 500	553 688	8 952	488 206	161 556		76 984	6 443 934 2 949 987	243 677 99 087	2 084 609 871 592	5 329
State Bank S.A. (d)			1 957	26 653	1 330		128 241	252.000	174 850	226 471	28 117	• •	820 925 84 261	2 451 085	191 832	559 957	3 957 2
R. & I. of W.A. (d)	•		1 544	21 360	862		100 163	253 909	80 042	48 898	163 325	100.240				10 932 152	
Total, Other .			7 228	98 606	62 342	• •	1 576 442	1 719 877	464 897	1 961 788	1 123 230	199 248	2 245 912	22 151 727	667 589		160 445
Total, All Banks		9	65 422	701 958	965 309	••	5 277 492	8 412 915	606 664	10 110 396	1 407 312	2 748 928	3 322 045	72 895 101	2 716 104	51 215 811	
A.R.D.B					16			00.206		4 086			155 069 7 310	278 549 1 845 204	191 2 485	-3 744 -43 324	434 1 900
C.D.B								88 296		775			, 310	1 040 204	4 703	73 324	. ,00

⁽a) Excludes inter-branch accounts and contingencies. (b) For legend on abbreviations of Bank names see footnote on page 2640. (c) Includes the Australian dollar equivalent of foreign currency balances. (d) The ir for these banks is supplied and published under a special arrangement with the banks and not under the Banking Act. Note: Any discrepancies between totals and sums of components are due to rounding.

DEPOSITS AND ADVANCES

AVERAGES OF DEPOSITS AND ADVANCES IN EACH STATE AND THE TERRITORIES FOR EACH BANK AT THE CLOSE OF BUSINESS ON WEDNESDAYS, 7, 14, 21 and 28 SEPTEMBER 1988 (\$'000)

	D		44					Loans (other than loans to authorised
		repayable in wealth and ents	<u> </u>	Other that State Gov	n Commonw ernments	ealth and		dealers in the short-term money
	<u> </u>	Current			Current			market),
Bank(a)	Fixed	Bearing Interest	Not Bearing Interest	Fixed	Bearing Interest	Not Bearing Interest	Total Deposits	advances and bills discounted (b)
NEW SOUTH WALES	<u> </u>						·	
Major Banks— A.N.Z	9 989 5 750	9 492	13 696	3 320 481 3 466 421	176 332 259 844	866 016 1 534 848	4 372 831 5 277 050	2 958 719 4 637 308
National	54 900	3 223 1 610	1 174 10 103	2 628 610 2 878 395	476 248 937 193	930 814 2 308 409	4 094 967 6 135 710	3 540 471 7 730 974
Total, Major	70 639	14 325	11 985	12 293 907	1 849 616	5 640 086	19 880 557	18 867 471
Other Banks-			····					
America	• •		• •	34 129 2 983 51 663	73 400	2 856 5 380 2 437	36 984 81 763 54 100	104 067 179 310 617 275
Barclays	43	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	212 389 195 195 8 574	22 301 167 377 5 615	7 674 6 846	242 364 369 461 14 189	585 865 744 907 131 694
Citibank	• •	• • • • • • • • • • • • • • • • • • • •	• •	70 100 44 201 27 236	14 345 54 438	1 850 2 317	86 295 46 517 81 674	446 246 225 532 308 978
I.B.J	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	• •	57 684 28 302 38 866	2 261 17 229 57 442	221 7 597 23 976	60 166 53 128 120 284	199 982 481 840 321 035
Mitsubishi Nat. Mutual NatWest New Zealand	• •	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	13 156 31 796 155 370 124 298	3 380 34 921 3 175 123 990	99 10 910	16 634 66 717 169 455 282 487	233 004 123 908 624 857 565 379
Paris	••		• •	124 298 152 889 77 070 14 135	4 399 6 967	34 200 21 254	178 542 77 070 21 101	271 983 136 548 183 548
Standard	••	91 268	10 508	26 557 65 765 3 248 896	20 570 1 807 298 798	3 003 7 056 541 190	50 130 74 627 4 190 660	136 560 142 507 6 230 147
R. & I. of W.A.(c)				183 992	1 901	344	186 237	165 205
Total, Other	43	91 268	10 508	4 865 242	914 313	679 208	6 560 583	13 160 375
Total, All Banks .	70 682	105 593	22 494	17 159 149	2 763 929	6 319 294	26 441 140	32 027 846
A.R.D.B	1 100			125 513 1 521 865			125 513 1 522 965	104 162 514 824
VICTORIA								
Major Banks— A.N.Z	1 343	4 861	4 116	3 020 283	290 040	1 589 232	4 909 874	5 490 112
Commonwealth	4 468 50 689 2 828	5 856 8 366 23 300	47 1 078 41 899	1 311 296 2 547 642 941 733	94 706 469 768 410 587	554 842 1 311 888 919 078	1 971 215 4 389 430 2 339 425	2 276 911 6 031 371 2 502 445
Total, Major	59 328	42 383	47 140	7 820 953	1 265 100	4 375 040	13 609 943	16 300 838
Other Banks— America Aust. Bank				53 391 368		384	53 775 368	153 625 126
Bankers Trust	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	• •	2 979 89 612	30 531 46 458		2 979 30 531 136 070	17 864 43 535
Citibank			• •	1 949	3 318	19 398	24 664	32 414

⁽a) For legend on abbreviations of Bank names see footnote on page 2640. (b) See note (c) on Statement of Assets. (c) See note (d) on Statement of Assets.

Note. Any discrepancies between totals and sums of components are due to rounding.

Bank(a) Deutsche HongKong		repayable in wealth and i ents Current Bearing Interest		Other than State Gove	Commonw	ealth and		dealers in the short-term
Deutsche		Bearing	Not					money
Deutsche			Not		Current			market), advances
	3 534	interest	Bearing Interest	Fixed	Bearing Interest	Not Bearing Interest	Total Deposits	and bills discounted (b)
	2 22 .			49 909		994	54 436	286 757
Hongrong				11 613	45 751	8	57 372	241 049
I.B.J				1 290	755		2 045	185 261
Lloyds N.Z.A				4 505	772		5 277	45 708
Macquarie				60 359	43 463	13 673	117 496	122 842
Mitsubishi				47 506	1 093		48 599	68 188
Nat. Mutual				28 158	35 165		63 323	212 965
NatWest				21 891	4 193	2 004	28 088	265 765
New Zealand				27 870	73 927	7 437	109 233	94 022
Paris				18 478	729	5 914	25 120	170 188
P.I.B.A				38 855			38 855	21 619
Singapore				120 159	3 018		123 178	124 001
Standard				27 186	2 534	84	29 804	28 178
Tokyo				29 895	679	1 903	32 477	43 536
Total, Other	3 534			635 972	292 385	51 798	983 689	2 157 640
Total, All Banks .	62 861	42 383	47 140	8 456 925	1 557 485	4 426 838	14 593 632	18 458 478
A.R.D.B		, .		77 379			77 379	154 027
C.D.B		···	···		···			318 970
QUEENSLAND								
Major Banks-								
A.N.Z	2 650		272	1 295 906	103 574	492 372	1 894 775	1 540 785
Commonwealth	1 228	3 343	10 988	1 181 562	89 397	495 677	1 782 196	1 528 441
National	2 327		302	1 717 437	205 045	570 329	2 495 440	1 829 853
Westpac				917 782	425 811	798 633	2 142 226	2 242 679
Total, Major	6 205	3 343	11 562	5 112 687	823 828	2 357 012	8 314 636	7 141 757
Other Banks—								
America				1 631		1 354	2 985	
Bankers Trust				3 420			3 420	
Barclays				• •				48 27€
Chase A.M.P				49 960	30 514		80 474	10 029
Citibank		• •		6 707	3 397		10 104	12 424
HongKong	- : :			1 965	9 356		11 321	41 785
Lloyds N.Z.A	350			13 046	3 191		16 586	83 719
Macquarie				2 349	5 352	7 327	15 027	32 329
Nat. Mutual				12 624	2 126		14 750	8 348
Nat West		• •	• •	66 012	54.225	15 550	66 012	62 147
New Zealand				24 037	54 235	15 558	93 830	120 970
Paris	• •	• •	• •	17 462 12 135	566	2 755	20 782 12 135	29 410 32 760
P.I.B.A	• •	• •	110		53 803	• •	12 135	
Queensland Standard			119	72 653 36 442	4 098	104	40 644	109 447 44 200
Total, Other.	350	•••	119	320 439	166 638	27 098	514 643	635 842
Total, All Banks	6 555	3 343	11 681	5 433 126	990 465	2 384 109	8 829 279	7 777 599
A.R.D.B				20 851			20 851	892
C.D.B			• •	20 851		• •	20 831	357 562

⁽a) For legend on abbreviations of Bank names see footnote on page 2640. (b) See note (c) on Statement of Assets. (c) See note (d) on Statement of Assets.

Note. Any discrepancies between totals and sums of components are due to rounding.

	Deposits	repayable ii	n Australia					Loans (other than loans to authorised dealers
	Common Governme	wealth and . ents	State	Other than State Gove	Commonw rnments	ealth and		in the short-term money
		Current			Current			market),
Bank(a)	Fixed	Bearing Interest	Not Bearing Interest	Fixed	Bearing Interest	Not Bearing Interest	Total Deposits	advances and bills discounted (b)
SOUTH AUSTRALIA Major Banks—								
A.N.Z			68	468 336	94 056	224 510	786 970	1 047 163
Commonwealth	398	2 805	343	427 965	20 857	122 950	575 318	712 650
National	1 106	• •	33	490 850	59 991	106 359	658 338	583 960
Westpac				173 948	68 977	156 906	399 831	639 703
Total, Major	1 504	2 805	443	1 561 098	243 881	610 725	2 420 456	2 983 476
Other Banks—								100
Barclays	• •	• •	• •	25 143	14 364	• •	39 506	400 5 005
Citibank				23 143	310	• • • • • • • • • • • • • • • • • • • •	310	5 454
Lloyds N.Z.A	13			2 311			2 324	
Nat. Mutual				4 972	1 691		6 663	3 081
Nat West	• •	,	••	36 011	0.540	726	36 011	52 363
Paris	• •		•••	1 441 2 955	9 548 22	726 282	11 714 3 259	16 662 26 997
P.I.B.A				18 859			18 859	20 660
Standard				15 817	8 450	515	24 781	48 697
State Bank S.A. (c)				1 606 588	168 290	73 198	1 848 076	2 906 906
Total, Other	13			1 714 097	202 674	74 720	1 991 502	3 086 224
Total, All Banks .	1 517	2 805	443	3 275 195	446 555	685 444	4 411 958	6 069 700
A.R.D.B				21 854		•••	21 854	11 667
C.D.B		••						241 645
WESTERN AUSTRALIA								
Major Banks—	1.260		26	400.005	21 120	017.500	720.000	(20.000
A.N.Z	1 369 427	1 489	35 115	488 905 498 695	31 420 18 047	217 508 216 921	739 238 735 692	628 932 916 053
National	4 511	1 407		518 999	72 335	196 557	792 402	895 932
Westpac			• • •	469 955	221 220	338 873	1 030 047	1 015 070
Total, Major	6 307	1 489	150	1 976 554	343 022	969 859	3 297 379	3 455 987
Other Banks—								
America				2 163		826	2 988	310
Bankers Trust				9 046			9 046	
Barclays	• •	• •	• •	9 027	4 794	• •	13 821	22 022 3 413
Citibank	• •			3021	51		51	4 055
HongKong		• • •	• • • • • • • • • • • • • • • • • • • •	4 153	8 022	• •	12 175	68 022
1.B.J				19 156	2 990	69	22 214	53 176
Lloyds N.Z.A	• •	• •	• •	4 926	• •	• •	4 926	3 600 86
NatWest			• •	42 864	1 232	1 099	45 195	124 145
New Zealand			,,	39 457	35 057	14 915	89 428	86 481
Paris				13 593	272	3 602	17 468	20 436
P.I.B.A	• •	••	• •	1 760	• •	• •	1 760	68 594 7 451
Standard				49 529	6 884	97	56 509	7 451 43 947
R.& I. of W.A. (c)	45 151	1 470	98	1 758 952	105 817	301 639	2 213 126	2 285 881
Total, Other	45 151	1 470	98	1 954 624	165 118	322 245	2 488 706	2 791 617
Total, All Banks .	51 458	2 958	248	3 931 177	508 140	1 292 104	5 786 084	6 247 604
A.R.D.B				19 451			19 451	
C.D.B	• • •							292 727

 ⁽a) For legend on abbreviations of Bank names see footnote on page 2640.
 (b) Se on Statement of Assets.
 Note. Any discrepancies between totals and sums of components are due to rounding. (b) See note (c) on Statement of Assets. (c) See note (d)

(\$'000)

	Deposits	repayable ir	n Australia					Loans (other than loans to authorised dealers
	Common Governme	wealth and . ents	State	Other than State Gove	Commonwo rnments	ealth and		in the short-term money
		Current			Current			market), advances
Bank(a)	Fixed	Bearing Interest	Not Bearing Interest	Fixed	Bearing Interest	Not Bearing Interest	Total Deposits	and bills
TASMANIA							- D op our	
Major Banks—								
A.N.Z	4		26	129 970	25 083	79 040	234 123	332 489
Commonwealth	517	364	1 472	91 591	14 101	44 295	152 339	173 011
National	1 641	• •		40 527	7 301	17 173	66 643 205 972	108 925
Westpac	493	264	1.400	91 118	36 833	77 529		253 645
Total, Major Other Banks	2 655	364	1 499	353 206	83 318	218 036	659 077	868 069
Chase A.M.P				8 563	7 762		16 325	882
Citibank					• • •			252
Lloyds N.Z.A				2016		• •	2046	500
NatWest				2 846 6 723			2 846 6 723	20 484
Total, Other				18 132	7 762		25 894	22 118
Total, All Banks .	2 655	364	1 499	371 338	91 080	218 036	684 971	890 187
A.R.D.B				15 787		• •	15 787	7 801
C.D.B		•••	···			••		84 249
NORTHERN TERRITO	RY							
Major Banks-			•	45.460		22.620	00.000	
A.N.Z	4 354 98	60	28 16	45 168 31 608	6 053 2 017	32 620 17 964	88 222 51 764	111 895 56 980
National	2 1 1 5			35 147	10 061	15 097	62 420	57 787
Westpac		885	532	21 705	13 924	38 244	75 289	137 486
Total, Major	6 567	945	576	133 628	32 055	103 925	277 695	364 148
Other Banks-							C 42	200
Chase A.M.P		• •	• •	100	447	• •	547	290 4 398
P.I.B.A.		• •		12		• •	12	4370
Standard				7 216	2 948	164	10 328	13 500
State Bank S.A. (c).				1 977	32 542		34 520	43 080
Total, Other		··-		9 305	35 938	164	45 406	61 267
Total, All Banks .	6 567	945	576	142 933	67 993	104 088	323 101	425 415
C.D.B			···		•••	•••	•••	22 563
AUSTRALIAN CAPITA	L TERRI	TORY						
Major Banks—				26 266	2 720	21.017	71 112	72 240
A.N.Z	1 839	6 176	• •	36 366 98 040	3 729 39 334	31 017 68 093	71 112 213 482	72 248 281 539
National	2 075	2 500	2	47 829	28 267	36 936	117 607	188 137
Westpac				48 409	98 492	72 079	218 981	202 825
Total, Major	3 9 1 4	8 676	2	230 643	169 822	208 124	621 180	744 748
Other Banks—				110	0.6		215	404
Chase A.M.P			• • •	119	96 12		215 12	496 951
Nat. Mutual		• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •		258	• • •	258	1 402
NatWest				5 052			5 052	17 225
Paris	• •	• •	• •	1 049	145	905	2 099	2 784

⁽a) For legend on abbreviations of Bank names see footnote on page 2640. (b) So on Statement of Assets.

Note. Any discrepancies between totals and sums of components are due to rounding. (c) See note (d) (b) See note (c) on Statement of Assets.

		repayable in wealth and . ents		Other than State Gove	a Commonwernments	ealth and		Loans (other than loans to authorised dealers in the short-term money	
		Current			Current			market), advances	
Bank(a)	Fixed_	Bearing Interest	Not Bearing Interest	Fixed	Bearing Interest	Not Bearing Interest	Total Deposits	and bills discounted (b)	
P.I.B.A State Bank N.S.W. (c)			• •	931 46 717	2 366	8 546	931 57 630	213 787	
Total, Other				53 868	2 876	9 451	66 195	236 645	
Total, All Banks .	3 914	8 676	2	284 511	172 698	217 575	687 375	981 393	
A.R.D.B		••		2 425		• •	2 425	12 665	
OTHER TERRITORIES									
Major Banks— Commonwealth Westpac		16		18 120 48 264	1 036 859	1 351 2 757	20 523 51 879	7 185 9 696	
Total, Major		16		66 385	1 895	· 4 107	72 403	16 881	
Total, All Banks .		, 16		66 385	1 895	4 107	72 403	16 881	

(a) A.N.Z.—Australian and New Zealand Banking Group Limited.

Commonwealth—Commonwealth Bank of Australia.

National-National Australia Bank Limited.

Westpac-Westpac Banking Corporation.

America—Bank of America Australia Limited.

Aust. Bank-Australian Bank Limited.

Bankers Trust-Bankers Trust Australia Limited.

Barclays-Barclays Bank Australia Limited.

Chase A.M.P.—Chase A.M.P. Bank Limited.

China—Bank of China.

Citibank—Citibank Limited.

Deutsche-Deutsche Bank Australia Limited.

HongKong-HongKong Bank of Australia Limited.

I.B.J.—I.B.J. Australia Bank Limited.

Lloyds N.Z.A.—Lloyds Bank N.Z.A. Limited.

Macquarie—Macquarie Bank Limited.
Mitsubishi—Mitsubishi Bank of Australia Limited.

Nat. Mutual-National Mutual Royal Bank Limited.

NatWest-NatWest Australia Bank Limited.

New Zealand-Bank of New Zealand.

Paris-Banque Nationale de Paris.

P.I.B.A.—Primary Industry Bank of Australia.

Queensland-Bank of Queensland Limited.

Singapore—Bank of Singapore (Australia) Limited.

Standard—Standard Chartered Bank Australia Limited.

Tokyo-Bank of Tokyo Australia Limited.

State Bank N.S.W.—State Bank of New South Wales (General Bank Department).

State Bank S.A.—The State Bank of South Australia.

R. & I. of W.A.—The Rural and Industries Bank of Western Australia (General Banking Department).

A.R.D.B.—Australian Resources Development Bank.

C.D.B.—Commonwealth Development Bank of Australia.

(b) See note (c) on Statement of Assets.

(c) See note (d) on Statement of Assets.

Note: Any discrepancies between totals and sums of components are due to rounding.

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SPECIAL

COMMONWEALTH OF AUSTRALIA

FISHERIES ACT 1952

FISHERIES NOTICE NO. ECC1

EAST COAST DEEP-WATER CRUSTACEAN TRAWL FISHERY: PROHIBITION ON TRAWLING FOR CRUSTACEANS WITHOUT ENDORSEMENT

- I, JOHN CHARLES KERIN, Minister of State for Primary Industries and Energy, hereby:
- (a) provide that this Notice shall come into force on 15 November 1988;
- (b) provide that, in this Notice, words have the same meaning as in the Act and -

"licence" means a licence granted to a person, in respect of a boat, under subsection 9(2) of the Act; and

"the Act" means the Fisheries Act 1952;

- (c) pursuant to paragraph 8(1)(c) of the Act, prohibit the taking of fish of the Class <u>Crustacea</u>, in the area of proclaimed waters specified in <u>Schedule 1</u>, by the method known as trawling;
- (d) pursuant to paragraph 8(1)(da) of the Act, prohibit a person from having in the person's possession or in the person's charge in a boat, in the area of proclaimed waters specified in Schedule 1, a trawl net the mesh of which is less than, or equal to, 60 millimetres when measured in accordance with either Schedule 2 or Schedule 3;
- (e) pursuant to paragraph 8(4D)(b) of the Act, declare that the activities to which the prohibitions contained in paragraphs (c) and (d) of this Notice relate are activities in respect of which an endorsement may be made under subsection 9(4) of the Act; and
- (f) revoke Fisheries Notice No. 44, published in Gazette 57 E on 15 July 1974, in its application to the taking of fish by the method known as trawling with the use of a boat, the licence in respect of which is endorsed under subsection 9(4) of the Act pursuant to this Notice.

SCHEDULE 1

AREA OF THE EAST COAST DEEP-WATER CRUSTACEAN TRAWL FISHERY

The area of proclaimed waters contained within the area bounded by a line:

- (1) commencing at the point of Latitude 33°35' South, Longitude 153°20'East;
- (2) thence northerly along the geodesic to the point of Latitude 30°South, Longitude 154°45'East:
- (3) thence northerly along the geodesic to the point of Latitude 27°48'South, Longitude 154°40'East;
- (4) thence north along that meridian to its intersection by the parallel of Latitude 22°15'South;
- (5) thence north-westerly along the geodesic to the point of Latitude 18⁰45'South, Longitude 152⁰35'East;
- (6) thence westerly along the geodesic to the point of Latitude 18°20'South, Longitude 148°20'East;
- (7) thence north-westerly along the geodesic to the point of Latitude 15°South, Longitude 146°East;
- (8) thence north-westerly along the geodesic to the point of Latitude 13°South, Longitude 145°East;
- (9) thence north along that meridian to its intersection by the outer limit of the Australian fishing zone;
- (10) thence generally southerly along that outer limit to its intersection by the parallel of Latitude 33^o35'South; and
- (11) thence west along that parallel to the point of commencement.

SCHEDULE 2

Manner of measuring mesh of net

 Immediately before a measurement is taken, the part of the net to be measured shall be soaked in fresh or salt water for not less than 5 minutes.

- The part of the net to be measured shall then be suspended vertically and a weight of 3170 grams shall be suspended from one knot of that part of the net.
- The distance between the surface of the knot from which the weight is suspended and the surface of the knot in the diagonally opposite corner of the mesh immediately above the first-mentioned knot shall be measured with a metal rule to the nearest millimetre.
- 4. The measurement of the mesh shall be taken to be the average measurement of 10 adjacent meshes at any place in the net.

SCHEDULE 3

Manner of measuring mesh of net

- Immediately before a measurement is taken the part of the net to be measured shall be soaked in fresh or salt water for not less than 5 minutes.
- The part of the net to be measured shall be suspended vertically. The distance between the surface of a knot and the surface of the knot diagonally opposite the first-mentioned knot in the same mesh shall be measured by means of a device designed for the measuring of the mesh of nets by means of two arms, the upper of which, when the device is in use, shall be fixed to the body of the device and the lower of which when the device is in use shall:
 - (a) be directly below the upper arm;
 - (b) move freely in relation to the upper arm; and
 - (c) have a weight of 3170 grams acting on it.
- The device shall be suspended in such a manner as to ensure that the upper edge shall remain in a fixed position and that the lower edge may move freely in relation to the upper edge, within the confines of the mesh to be measured.

4 Fisheries Act 1952

4. The device shall be equipped with a scale from which the distance between the upper edge of the upper arm and the lower edge of the lower arm can be ascertained at any time to the nearest millimetre.

Dated this Tenth

day of November 1988.

JOHN KERIN

COMMONWEALTH OF AUSTRALIA

FISHERIES ACT 1952

FISHERIES NOTICE NO. 117D

PROHIBITION ON THE TAKING OF PRAWNS OFF THE EAST COAST OF NEW SOUTH WALES WITHOUT ENDORSEMENT: VARIATION OF FISHERIES NOTICE NO. 117

I, JOHN CHARLES KERIN, Minister of State for Primary Industries and Energy, pursuant to section 8 of the Fisheries Act 1952, hereby vary Fisheries Notice No. 117, published in Gazette No. S 272 on 4 November 1983 by omitting all the words after "hereby" and substituting the following:

- " (a) provide that this Notice shall come into force on 15 November 1988;
 - (b) provide that, in this Notice, words have the same meaning as in the Act and "the Act" means the Fisheries Act 1952:
 - (c) pursuant to paragraph 8(1)(a) of the Act, prohibit the taking, processing or carrying of fish of the family <u>Penaeidae</u> (commonly known as prawns or shrimps) in the area of proclaimed waters specified in the Schedule: and
 - (d) pursuant to paragraph 8(4D)(b) of the Act, declare that the activities to which the prohibition contained in this Notice relates are activities in respect of which an endorsement may be made under subsection 9(4) of the Act.

SCHEDULE

AREA OF THE IN-SHORE PRAWN FISHERY OFF THE EAST COAST OF NEW SOUTH WALES

The area of proclaimed waters contained within the area bounded by a line:

- (1) commencing at the point of intersection of the eastern shore of Australia by the parallel of Latitude 28010'South;
- (2) thence easterly along the geodesic to the point of Latitude 27°58'South, Longitude 154°East;

6 Fisheries Act 1952

- (3) thence north-easterly along the geodesic to the point of Latitude 27°48'South, Longitude 154°22'East:
- (4) thence east along that parallel to its intersection by the meridian of Longitude 154°40'East;
- (5) thence southerly along the geodesic to the point of Latitude 30°South, Longitude 154°45'East;
- (6) thence south-westerly along the geodesic to the point of Latitude 33⁰35'South, Longitude 153⁰20'East;
- (7) thence east along that parallel to its intersection by the eastern shore of Australia; and
- (8) thence generally northerly along that shore to the point of commencement."

Dated this Touth

day of November 1988. John Kerin

JOHN KERIN

Fisheries Act 1952

Fisheries Notice No. 216.

REVOCATION OF FISHERIES NOTICES

In pursuance of section 8 of the Fisheries Act 1952, I, JOHN CHARLES KERIN, Minister of State for Primary Industries and Energy, hereby revoke the Fisheries Notices specified in the Schedule

	SCHEDULE	
Notice No.	Gazette No.	Gazette date
Notice No. 17 40 40A 46 51 52 53 54 54A 555 55A 56 57 58 59 69 75 75A 75B 87 90 99 107 109 112 129 136 146 149	Gazette No. 97 129 S 1 57 E 103 E 103 E 103 E 103 E 103 E 103 E S 133 G 13 G 13 G 13 G 13 G 13 S 191 S 79 S 133 S 340 S 144 S 142 S 209 S 229 S 133 S 1470 S 340 S 371 S 413	Gazette date
153 164	S 435 S 154	24 October 1985 08 April 1986

Dated this Tenth day of November 1988

Mahn Van

JOHN CHARLES KERIN



Commonwealth of Australia

Gazette

No. S 352, Monday, 14 November 1988

Published by the Australian Government Publishing Service, Canberra

SPECIAL

PROCLAMATION

Commonwealth of Australia N. M. STEPHEN Governor-General By His Excellency the Governor-General of the Commonwealth of Australia

I, SIR NINIAN MARTIN STEPHEN, Governor-General of the Commonwealth of Australia, acting with the advice of the Federal Executive Council and pursuant to subsection 2 (9) of the Statute Law (Miscellaneous Provisions) Act 1987, hereby fix 1 December 1988 as the day on which amendments to the Commonwealth Prisoners Act 1967 made by the first-mentioned Act shall come into operation.

(L.S.) GIVEN under my hand and the Great Seal of Australia on 8 November 1988

By His Excellency's Command,

LIONEL BOWEN Attorney-General

GOD SAVE THE QUEEN!

AN ARRANGEMENT BETWEEN HIS EXCELLENCY THE GOVERNOR-GENERAL OF THE COMMON-WEALTH OF AUSTRALIA, acting with the advice of the Federal Executive Council, and HIS HONOUR THE ADMINISTRATOR OF THE NORTHERN TERRITORY OF AUSTRALIA, acting with the advice of the Executive Council of the Northern Territory.

IT IS HEREBY ARRANGED, pursuant to section 21 of the Commonwealth Prisoners Act 1967 of the Commonwealth and all powers thereunto enabling, as follows:

- Persons who hold office as Stipendiary Magistrates in the Northern Territory of Australia may perform the functions of a prescribed authority under the Commonwealth Prisoners Act 1967 or that Act as amended from time to time.
- Officers of the Northern Territory who perform the functions of parole officer or probation officer in that Territory may perform the functions of a parole officer under the Commonwealth Prisoners Act 1967 or that Act as amended from time to time.

Dated 8 November 1988

N. M. STEPHEN Governor-General

By His Excellency's Command LIONEL BOWEN Attorney-General

E. E. JOHNSTON Administrator

By His Honour's Command DONALD F. DALE Minister for Health and Community Services



Commonwealth of Australia

Gazette

No. S 353, Tuesday, 15 November 1988

Published by the Australian Government Publishing Service, Canberra

SPECIAL

NOTIFICATION OF THE MAKING OF STATUTORY RULES

Notice is hereby given that the undermentioned Statutory Rules have been made. Copies of the Statutory Rules may be purchased at the Commonwealth Government Bookshop, 70 Alinga St, Canberra City, ACT.

Act under which the Statutory Rules were made	Description of the Statutory Rules	Year and number of the Statutory Rules
Superannuation Act 1976	Superannuation (Approved Authorities) Regulations (Amendment)	1988 No. 273
Superannuation Act 1976	Superannuation (Interest) Regulations (Amendment)	1988 No. 274
Superannuation Act 1976	Superannuation (Existing Invalidity Pensioners) Regulations	1988 No. 275
Marriage Act 1961	Marriage Regulations (Amendment)	1988 No. 276
Air Navigation Act 1920	Air Navigation (Aircraft Noise) Regulations (Amendment)	1988 No. 277



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