HEALTH INSURANCE (PHARMACEUTICAL BENEFIT) (GENERAL PROVISIONS) (No. 2) (JERSEY) ORDER 2002

Revised Edition
26.500.22
Showing the law as at 1 January 2019
This is a revised edition of the law

HEALTH INSURANCE (PHARMACEUTICAL BENEFIT) (GENERAL PROVISIONS) (No. 2) (JERSEY) ORDER 2002

Arrangement

Article

Supporting Documents

Jersey

HEALTH INSURANCE (PHARMACEUTICAL BENEFIT) (GENERAL PROVISIONS) (No. 2) (JERSEY) ORDER 2002[1]

THE EMPLOYMENT AND SOCIAL SECURITY COMMITTEE, in pursuance of Articles 15, 17 and 36 of the Health Insurance (Jersey) Law 1967,[2] orders as follows –

Commencement [see endnotes]

1 Interpretation

(1) In this Order, unless the context otherwise requires –

"approved prescribing practitioner" means any medical practitioner, dentist, optician or other prescribing practitioner approved by the Minister under Article 26 of the Law;

"Drug Tariff" means the statement compiled and published by the Secretary of State for Health of the United Kingdom pursuant to Regulation 18(1) of the National Health Service (Pharmaceutical Services) Regulations 1992 of the United Kingdom, as that statement is for the time being in force;

"Health Card" means an Identity Card issued in accordance with the Health Insurance (Evidence) (Jersey) Order 2007[3];

"Law" means the Health Insurance (Jersey) Law 1967[4];

"prescription" has the same meaning as given to the expression "prescribed form" by Article 5.[5]

(2) In this Order, unless the context otherwise requires –

(a) a reference to an enactment (including an instrument made under any enactment of the Parliament of the United Kingdom) shall be taken to be a reference to –

(i) that enactment as from time to time amended or extended or applied by or under another enactment, and

(ii) any other enactment repealing and re-enacting that enactment with or without any further amendment;

(b) a reference in an Article or other division of this Order to a paragraph, sub-paragraph, clause or item by number or letter only is a reference to the paragraph, sub-paragraph, clause or item of that number or letter contained in the Article or other division of this Order in which it appears.

2 Approval of suppliers

(1) An application by a person conducting a retail pharmacy business for approval under Article 26(2) of the Law as a supplier of pharmaceutical benefit must be substantially in the form set out in Part 1 of Schedule 1.

(2) An application by any other person for that approval must be substantially in the form set out in Part 2 of Schedule 1.

3 Terms and conditions of supply of pharmaceutical benefit

For the purpose of Article 26(2) and (3) of the Law the terms and conditions a person in an application for approval as an approved supplier must undertake to supply pharmaceutical benefit on are those set out in Part 1 of Schedule 2.

4 Schemes for securing proper pharmaceutical services

(1) The Minister may act in accordance with paragraph (2) if at any time he or she is satisfied that there is an insufficient number of places of business of approved suppliers open outside normal business hours.

(2) The Minister may after consulting the Pharmaceutical Benefit Advisory Committee prepare a scheme designed to ensure that a sufficient number of places of business of approved suppliers are open at reasonable times outside normal business hours.

(3) The scheme must –

(a) specify the days and hours during which places of business of approved suppliers are to be open; and

(b) provide for payments to be made to those suppliers in respect of periods during which their premises are open under the scheme.

5 Prescribed form for the supply of pharmaceutical benefit

(1) For the purposes of Article 15(2) of the Law "prescribed form" means a form substantially in the form set out in Schedule 3 that –

(a) contains the particulars required to complete the form;

(b) is dated with the date on which it is to become effective;

(c) is signed by the approved prescribing practitioner that gave the form; and

(d) is printed and watermarked in such manner as may be designated by the Minister.[6]

(2) A form is not in the prescribed form if it authorizes the supply of pharmaceutical benefit for more than one person.

6 Period of supply

(1) An approved prescribing practitioner must not give a prescription that orders the supply of pharmaceutical benefit necessary to provide treatment for a person for a period exceeding 30 days.[7]

(2) Despite paragraph (1) an approved prescribing practitioner (other than a dentist) may give a prescription that orders the supply of pharmaceutical benefit necessary to provide treatment for a person for a period exceeding 30 days but not exceeding 90 days if the pharmaceutical benefit is of a type specified in Schedule 4.[8]

(3) An approved prescribing practitioner must not give more than one prescription in respect of the same pharmaceutical benefit at any one time.[9]

(4) Despite paragraph (3) an approved prescribing practitioner may give more than one but not more than 4 prescriptions in respect of the same pharmaceutical benefit at any one time if the prescriptions authorize consecutive periods of supply of pharmaceutical benefit to a person.[10]

(5) Where an approved prescribing practitioner has given a prescription or prescriptions in respect of a period of treatment of a person the practitioner must not during that period of treatment give a further prescription in respect of that person for the same pharmaceutical benefit unless the practitioner is satisfied that it is necessary or desirable to do so for the purpose of the treatment and –

(a) the pharmaceutical benefit is intended for the continuation of the treatment after the expiry of the present period of treatment and the prescription bears a date not earlier than 21 days before the end of that period;

(b) the prescription is for an increase in dosage; or

(c) the prescription is to replace pharmaceutical benefit previously supplied on prescription and accidentally lost or destroyed.[11]

(6) An approved supplier must not supply pharmaceutical benefit ordered on a prescription before the effective date of the prescription.

7 Supply

(1) This Article applies where in accordance with a prescription pharmaceutical benefit is supplied by an approved supplier in respect of an insured person or a dependant of an insured person.

(2) The person to whom the pharmaceutical benefit is supplied must produce and show to the approved supplier the insured person's Health Card.

(3) Subject to paragraph (4) –

(a) the person to whom the pharmaceutical benefit is supplied must surrender the prescription to the approved supplier; and

(b) the approved supplier must mark on the prescription the health insurance number specified on the insured person's Health Card.

(4) The person taking delivery of pharmaceutical benefit is not required to surrender the prescription if, by electronic means, the approved prescribing practitioner who signed the prescription –

(a) notifies the approved supplier of the particulars contained in the prescription and its effective date;

(b) confirms that the prescription is signed by the practitioner; and

(c) undertakes to surrender the prescription to the approved supplier within 72 hours of the notification,

and the approved supplier is satisfied that, by reason of an emergency, the practitioner has been unable to furnish the person taking delivery of the pharmaceutical benefit with the prescription so that it is available for surrender at the time of delivery.[12]

(5) [13]

(6) [14]

(7) [15]

(8) [16]

8 Offences

(1) Except as provided by paragraph (2), a person who fails to comply with a provision of this Order shall be guilty of an offence and liable to a fine not exceeding level 2 on the standard scale.

(2) A person who fails to surrender a prescription as required under Article 7(3)(a) is liable to a fine not exceeding level 1 on the standard scale.

9 Prescription costs payable by the Minister

(1) Except as otherwise provided by this Article, the amount to be paid by the Minister to an approved supplier for each item of pharmaceutical benefit supplied on a prescription in respect of an insured person or a dependant of an insured person is the sum of the following –

(a) the basic price of the ingredients; and

(b) the appropriate dispensing fee or fees as set out in Schedule 5.

(2) The aggregate amount payable to an approved supplier under paragraph (1)(a) in respect of pharmaceutical benefit supplied in any month shall be reduced in accordance with the scale set out in Schedule 6.

(3) Despite paragraph (2) a reduction shall not be made in respect of any item supplied by an approved supplier that is specified in the "ZD List" in the Drug Tariff if the approved supplier –

(a) purchased the item at a price not less than its basic price as calculated in accordance with the Drug Tariff; and

(b) has endorsed the prescription with the notation "ZD".

(4) In the circumstances specified in paragraph (5) the amount to be paid to the approved supplier under paragraph (1) shall be the amount that would be payable under that paragraph if each instalment referred to in paragraph (5) had been supplied on a separate prescription.

(5) The circumstances referred to in paragraph (4) are where an approved supplier supplies by way of pharmaceutical benefit a substance or product in accordance with a prescription signed by an approved medical practitioner that specifies that it is to be supplied in instalments at stipulated intervals or on given dates and, at the time of supply, the substance or product is specified in Part 1, 2 or 3 of Schedule 2 to the Misuse of Drugs (Jersey) Law 1978[17].[18]

(6) [19]

(7) For the purposes of this Article the basic price of an ingredient is its basic price calculated in accordance with the Drug Tariff as at the date of its supply.

(8) If the term "aqua" is used in a prescription without qualification the approved supplier shall –

(a) interpret it to mean wholesome drinking water; and

(b) not charge for it.

(9) An approved supplier shall not charge for distilled water unless –

(a) its use is specified on the prescription; or

(b) it is necessary to use distilled water to conform with standard dispensing practice.

10 Citation

This Order may be cited as the Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Jersey) Order 2002.

SCHEDULE 1

(Article 2)

APPROVAL OF SUPPLIERS OF PHARMACEUTICAL BENEFIT

PART 1

HEALTH INSURANCE (JERSEY) LAW 1967

Form of application for approval as supplier of pharmaceutical benefit of person(s) lawfully conducting a retail pharmacy business

To: The Minister for Social Security

I/We, ...

of .....

being a person or persons lawfully conducting a retail pharmacy business within the meaning of the Medicines (Jersey) Law 1995, apply for approval in accordance with Article 26(2) of the Health Insurance (Jersey) Law 1967. I/we undertake to dispense medicines and supply drugs at the prices fixed and in accordance with the terms and conditions prescribed under the Health Insurance (Jersey) Law 1967. I/We understand that those prices, terms and conditions are subject to variation in the manner provided by that Law.

The address(es) of my/our business premises registered in accordance with the Medicines (Jersey) Law 1995 and the pharmacist(s) in charge of those premises will be as follows –

Address(es) of premises	Full name(s) of pharmacist(s) in charge

Signed: .....

Date: .....

PART 2

HEALTH INSURANCE (JERSEY) LAW 1967

Form of application for approval as supplier of pharmaceutical benefit of person(s) other than person(s) lawfully conducting a retail pharmacy business

To: The Minister for Social Security

I/We .....

of .....

apply for approval in accordance with Article 26(3) of the Health Insurance (Jersey) Law 1967. I/we undertake to supply drugs (except poisons in Part 1 of the Poisons List set out in the Schedule to the Poisons List (Jersey) Order 1986) at the prices fixed and in accordance with the terms and conditions prescribed under the Health Insurance (Jersey) Law 1967. I/We understand that those prices, terms and conditions are subject to variation in the manner provided by that Law.

The address(es) of my/our business premises for this purpose will be .....

Signed: .....

Date: .....

SCHEDULE 2[20]

(Article 3)

PART 1

TERMS AND CONDITIONS TO BE OBSERVED BY AN APPROVED SUPPLIER

1 Supplier to supply pharmaceutical benefit

(1) The supplier must supply pharmaceutical benefit with reasonable promptness to a person who presents a prescription for them.

(2) Sub-paragraph (1) does not require a supplier to supply a pharmaceutical benefit that the supplier does not ordinarily supply.

(3) If under this paragraph a supplier is required to supply a medicine or drug the supplier must supply the medicine or drug in a suitable container being –

(a) in relation to capsules, tablets, pills or any other medicine or drug in solid form (other than those prepacked in foil or paper-board or strip card containers by the manufacturer) – an airtight container of glass, aluminium or rigid plastic;

(b) in relation to ointments, creams or pastes (other than those prepacked by the manufacturer) – a container of glass, aluminium or rigid plastic;

(c) in relation to eye, ear or nasal drops (other than those prepacked by the manufacturer) – a container of glass either incorporating or having a separate dropper attachment;

(d) in relation to liquid medicines (other than those prepacked by the manufacturer) – a container of glass or rigid plastic, including, in the case of an oral liquid medicine, a 5 ml. plastic measuring spoon (unless the patient already has one or the manufacturer's pack includes one).

(4) The supplier must not give, promise or offer a gift or a reward as an inducement to or in consideration of a person presenting a prescription to the supplier.

2 Place and hours of business

(1) The supplier must supply pharmaceutical benefit at the place or places of business specified in the supplier's application for approval under Article 26 of the Law.

(2) The supplier must keep that place or those places open for the supply of pharmaceutical benefit –

(a) during normal business hours; and

(b) on the days and during the hours specified in any scheme made by the Minister under Article 4.

(3) At each such place of business the supplier must display a notice to be provided by the Minister in the form set out in Part 2 or Part 3 of this Schedule.

(4) If the supplier is a person lawfully conducting a retail pharmacy business the supplier must also display a notice to be provided by the Minister in the form set out in Part 4 of this Schedule when the supplier's place of business is closed.

(5) The notice must indicate –

(a) the addresses of other people lawfully conducting a retail pharmacy business where medicines and drugs may be obtained; and

(b) the times when they may be obtained at those premises.

(6) Each notice must be displayed in a manner that makes it easily visible to members of the public.

3 Dispensing medicines

The supplier must ensure that the supply of medicines on prescriptions is performed by or under the direct supervision of a pharmacist.

4 Names of pharmaceutical chemists

Whenever required to do so by the Minister the supplier must furnish to the Minister the name of each pharmacist employed by the supplier in dispensing medicines on prescription.

5 Charges

Except for charges that are required or authorized to be made by this or any other Order made under the Law the supplier must supply a pharmaceutical benefit and any container free of charge.

6 Advertising

(1) The supplier must not advertise either directly or by implication that the supplier is an approved supplier or that the supplier provides or is authorized to provide pharmaceutical benefit.

(2) Despite sub-paragraph (1) the supplier may –

(a) display a notice required by paragraph 2;

(b) include in an advertisement a statement of the days and hours at which pharmaceutical benefit is supplied.

7 Information to be provided

(1) This paragraph does not apply except where the Minister requires information to determine the amount payable under Article 10 to the supplier for pharmaceutical benefit supplied by the supplier.

(2) The supplier must furnish to the Minister or to such person or body as the Minister directs information that the Minister requires concerning so much of the supplier's business that relates to the supply of pharmaceutical benefit.

(3) The supplier must permit a person authorized in writing to do so by the Minister to conduct surveys at each place of business at or from which the supplier supplies pharmaceutical benefit.

8 Payment

(1) On dates specified by the Minister the supplier must furnish to the Minister or to such person or body as the Minister directs the prescriptions on which pharmaceutical benefit has been supplied by the supplier.

(2) The prescriptions must be arranged in the manner the Minister directs and must be accompanied by a declaration.

(3) The declaration must contain such particulars relating to the supply by the supplier of pharmaceutical benefit as the Minister specifies.

9 Withdrawal

If the supplier wishes to cease to be an approved supplier the supplier must give at least 3 months written notice to the Minister (or such shorter notice as the Minister may agree) that the supplier no longer wishes to supply pharmaceutical benefit.

PART 2

Form of notice to be displayed by an approved supplier who is a person lawfully conducting a retail pharmacy business

Health Insurance Scheme

(Name of approved supplier)

Approved under the Health Insurance (Jersey) Law 1967 to dispense medicines and supply drugs.

These premises are open at the following times –

PART 3

Form of notice to be displayed by an approved supplier other than a person lawfully conducting a retail pharmacy business

Health Insurance Scheme

(Name of approved supplier)

Approved under the Health Insurance (Jersey) Law 1967 to supply drugs (except poisons in Part 1 of the Poisons List set out in the Schedule to the Poisons List (Jersey) Order 1986).

These premises are open at the following times –

PART 4

Form of notice to be displayed by an approved supplier who is a person lawfully conducting a retail pharmacy business at times when the person's premises are closed

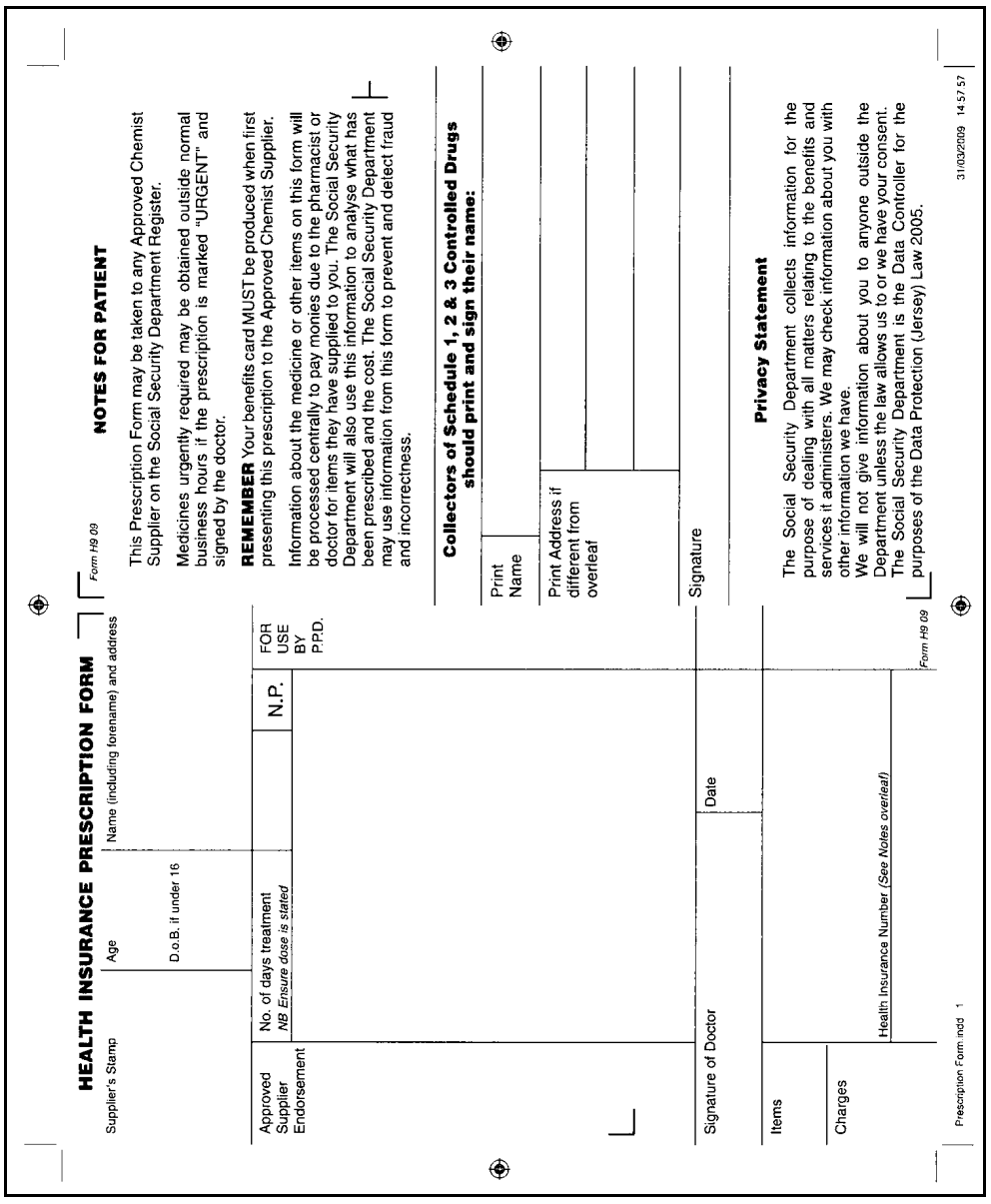
Health Insurance Scheme

When these premises are closed, medicines and drugs may be obtained at the addresses and times shown below –

SCHEDULE 3[21]

(Article 5)

FORM OF PRESCRIPTION



SCHEDULE 4[22]

(Article 6(2))

PHARMACEUTICAL BENEFIT WHICH MAY BE SUPPLIED FOR 90 DAYS ON ANY ONE PRESCRIPTION

PART 1

Name of drug and form	
A. Anticonvulsants	
Acetazolamide	Tablets 250 mg
Carbamazepine	Tablets 100 mg; 200 mg; 400 mg Tablets M.R. 200 mg; 400 mg Oral liquid 100 mg per 5 ml
Clobazam	Tablets 10 mg
Clonazepam	Tablets 0.5 mg; 2 mg
Ethosuximide	Capsules 250 mg Oral liquid 250 mg per 5 ml
Lamotrigine	Tablets 25 mg; 50 mg; 100 mg; 200 mg Dispersible Tablets 2 mg; 5 mg; 25 mg; 100 mg
Levetiracetam	Tablet 250 mg; 500 mg; 750 mg; Oral solution 500 mg/5 ml
Phenobarbitone	Tablets 15 mg; 30 mg; 60 mg Oral solution15 mg/5 ml
Phenytoin	Capsules 25 mg; 50 mg; 100 mg; 300 mg Infatabs 50 mg Suspension 30 mg per 5 ml
Primidone	Tablets 250 mg
Sodium Valproate	Syrup 200 mg per 5 ml; Tablets enteric coated 200 mg; 500 mg Tablets and Crushable 100 mg Tablets M.R. 200 mg; 300 mg; 500 mg Sugar free liquid 200 mg/5 ml Capsules M.R. 150 mg; 300 mg

	Granules 500 mg/sachet, 1 g/sachet
Topiramate	Tablets 25 mg; 50 mg; 100 mg; 200 mg Capsules 50 mg Sprinkle capsules 15 mg; 25 mg; 50 mg
B. Preparations for the management of diabetes	
Acarbose	Tablets 50 mg; 100 mg
Canagliflozin	Tablets 100 mg; 300 mg
Clinitest	Tablets
Dapagliflozin	Tablets 5 mg; 10 mg
Empagliflozin	Tablets 10 mg; 25 mg
Exenatide	Injection 250 mcg/ml; 2 mg
Glibenclamide	Tablets 2.5 mg; 5 mg
Gliclazide	Tablets 80 mg
Insulin preparations	All insulins
Linagliptin	Tablets 5 mg
Liraglutide	Injection 6 mg/ml
Liraglutide/degludec	Injection 3.6 mg/100 IU
Metformin	Tablets 500 mg; 850 mg Tablets M.R. 500 mg; 750 mg; 1g Oral powder 500 mg/sachet; 1 g/sachet
Metformin/canagliflozin	Tablets 850 mg/50 mg; 1 g/50 mg
Metformin/dapagliflozin	Tablets 850 mg/5 mg/1g/5 mg
Metformin/linagliptin	Tablets 850 mg/2.5 mg; 1 g/2.5 mg
Metformin/pioglitazone	Tablets 850 mg/15 mg
Metformin/empagliflozin	Tablets 1g/5 mg, 1g/12.5 mg; 85 0mg/5 mg; 850 mg/12.5 mg
Pioglitazone	Tablets 15 mg; 30 mg; 45 mg
Saxagliptin	Tablets 2.5 mg; 5 mg

Sitagliptin	Tablets 25 mg; 50 mg; 100 mg
Sitagliptin/Metformin	Tablets 50 mg/1000 mg
Tolbutamide	Tablets 500 mg
Vildagliptin	Tablets 50 mg
Vildagliptin/Metformin	Tablets 50 mg/850 mg; 50 mg/1000 mg
C. Thyroid and antithyroid drugs	
Carbimazole	Tablets 5 mg; 20 mg
Liothyronine Sodium	Tablets 20 mcg
Levothyroxine Sodium	Tablets 25 mcg; 50 mcg; 75 mcg; 100 mcg

PART 2

Oral or transdermal medicine used for contraception and containing any of the following –
Co-cyprindiol desogestrel dienogest drospirenone estradiol hemihydrate estradiol valerate ethinylestradiol gestodene levonorgestel mestranol nomegestrol norelgestromin norgestimate norethisterone
Oral, vaginal or transdermal medicines containing the following female sex hormones when used as hormone replacement therapy –

Conjugated oestrogens
Estradiol
Ethinylestradiol
Tibolone

SCHEDULE 5[23]

(Article 9(1)(b))

DISPENSING FEES

		Fee in pence for each item of pharmaceutical benefit supplied on a prescription
1.	Basic dispensing fee	
	For the period of 12 months beginning with 1st October 2014 and for each ensuing period of 12 months –	
	(a) for each of the first 50,000 items of pharmaceutical benefit supplied as described in Article 9(1) by an approved supplier	351
	(b) for each subsequent item of pharmaceutical benefit as described in Article 9(1) by that approved supplier	313
2.	Additional dispensing fees	
	(a) Preparations when dispensed extemporaneously and endorsed "Extemporaneously dispensed" –	
	(i) unit dosage forms, e.g. cachets, capsules, pills, lozenges, pastilles, pessaries, powders	256
	(ii) liquids, being "special formula preparations", e.g. mixtures, lotions, nasal drops (not including dilutions)	155
	(iii) liquid preparations, prepared by straightforward dilution (not including reconstitution)	85
	(iv) special formula powders	155
	(v) ointments, creams, pastes, being "special formula preparations" (not including dilutions)	310
	(vi) ointments, creams, pastes, prepared by dilution or admixture of standard or	155

	proprietary ointments, creams and pastes	
(b)	Preparations when aseptically dispensed and endorsed "Aseptically dispensed" (excluding proprietary preparations) –	
	(i) unit dosage forms, e.g. injections	1277 per ten or part thereof
	(ii) non-unit dosage forms, e.g. eye drops	767
(c)	Liquid preparations dispensed extemporaneously other than items (a)(iii) and (b) above which are ordered by the approved medical practitioner or registered dentist to be supplied in more than one container, each extra quantity ordered	128
(d)	A preparation which requires the addition of a vehicle/diluent by the approved supplier, and results in a liquid of stability of less than 14 days, and for pharmaceutical reasons necessitates supply in more than one container and the prescription is endorsed with the number of extra quantities supplied, for each extra supply	155
(e)	the following drugs if the prescription is endorsed "C.D." by the approved supplier –	
	(i) the drugs listed in Schedules 2 and 3 to the Misuse of Drugs (General Provisions) (Jersey) Order 1989[24]	128
	(ii) chlordiazepoxide, diazepam, flurazepam, lorazepam, nitrazepam, oxazepam, zolpidem and zopiclone	43
(f)	When the prescription has been dispensed at a time when the premises are not open for dispensing on the day or (in the case of a prescription dispensed after midnight) the day following that on which it was written and –	
	(i) is endorsed "URGENT" by the approved medical practitioner or	713

		registered dentist, and dispensed between the time the premises close for dispensing and the time the premises open for dispensing on a day other than a Sunday or a public holiday	(1756 non-resident rate)
	or		
	(ii)	is endorsed "URGENT" by the approved medical practitioner or registered dentist, or "DISPENSED URGENTLY" by the approved supplier and is signed by the patient (or the patient's representative) and dispensed on a Sunday or a public holiday URGENT FEES are not payable for prescriptions dispensed by an approved supplier when the supplier's premises are open for dispensing in accordance with a scheme prepared by the Minister under Article 4.	920 (2118)
(g)	items of pharmaceutical benefit the cost of which exceeds –		
	(i)	£75 but does not exceed £99.99	300
	(ii)	£100 but does not exceed £199.99	500
	(iii)	£200 but does not exceed £499.99	1000
	(iv)	£500 or over	2500

NOTES:

(1) All "URGENT" prescriptions must be endorsed by the approved supplier with the date and time of dispensing.

(2) In order to qualify for the non-resident rates contained in sub-paragraph (f) an approved supplier shall –

(a) normally live elsewhere than on the supplier's business premises;

(b) have left those premises; and

(c) have returned to open them to dispense an "URGENT" prescription.

In the absence of an endorsement "NON-RESIDENT" on an "URGENT" prescription, payment will automatically be made at

the "RESIDENT" rate.

SCHEDULE 6[25]

(Article 9(2))

REDUCTION OF AMOUNT PAYABLE TO APPROVED SUPPLIERS

Aggregate basic ingredient price of pharmaceutical benefit supplied during month		Percentage by which basic ingredient price reduced %
exceeding £	not exceeding £	
0	1500	0.37
1500	1750	0.40
1750	2000	0.43
2000	2250	0.45
2250	2500	0.46
2500	2750	0.50
2750	3000	0.52
3000	3250	0.55
3250	3500	0.57
3500	3750	0.59
3750	4000	0.61
4000	4250	0.64
4250	4500	0.66
4500	4750	0.68
4750	5000	0.70
5000	5250	0.73
5250	5500	0.75
5500	5750	0.77
5750	6000	0.78
6000	6250	0.79
6250	6500	0.81

6500	7000	0.83
7000	8900	0.91
8900	10300	0.95
10300	11700	0.99
11700	13100	1.01
13100	14400	1.01
14400	15800	1.02
15800	17200	1.02
17200	18600	1.03
18600	20000	1.03
20000	21400	1.04
21400	22800	1.04
22800	24200	1.05
24200	25800	1.05
25800	27400	1.06
27400	29000	1.06
29000	30600	1.07
30600	32200	1.07
32200	33800	1.08
33800	35400	1.08
35400	37000	1.09
37000	38600	1.10
38600	40200	1.10
40200	41800	1.11
41800	43400	1.11
43400	45000	1.12
45000	46600	1.12
46600	48200	1.13
48200	49800	1.13
49800	51400	1.14
51400	53000	1.14

53000		1.15

ENDNOTES

Table of Legislation History

Legislation	Year and No	Commencement
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Jersey) Order 2002	R&O.48/2002	1 October 2002
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment) (Jersey) Order 2003	R&O.85/2003	4 October 2003
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 2)) (Jersey) Order 2004	R&O.90/2004	1 October 2004
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 3) (Jersey) Order 2004	R&O.99/2004	1 October 2004
States of Jersey (Amendments and Construction Provisions No. 8) (Jersey) Regulations 2005	R&O.48/2005	9 December 2005
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 4) (Jersey) Order 2007	R&O.185/2007	21 January 2008 (applies to benefit supplied on or after 1 October 2007)
Health Insurance (Consequential Amendments) (Jersey) Order 2008	R&O.15/2008	28 January 2008
Health Insurance (Pharmaceutical Benefit) (No. 2) (Amendment No. 5) (Jersey) Order 2008	R&O.19/2008	1 February 2008
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 6) (Jersey) Order 2008	R&O.122/2008	1 October 2008
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 7) (Jersey) Order 2009	R&O.35/2009	1 May 2009
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 8) (Jersey) Order 2009	R&O.98/2009	1 November 2009
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 9) (Jersey) Order 2010	R&O.99/2010	1 November 2010

Legislation	Year and No	Commencement
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 10) (Jersey) Order 2013	R&O.44/2013	1 May 2013
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 11) (Jersey) Order 2013	R&O.119/2013	1 October 2013
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 12) (Jersey) Order 2014	R&O.141/2014	1 October 2014
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 13) (Jersey) Order 2014	R&O.174/2014	1 December 2014
Health Insurance (Pharmaceutical Benefit) (General Provisions) (No. 2) (Amendment No. 14) (Jersey) Order 2014	R&O.102/2017	13 October 2017
Health Insurance (Approved Prescribing Practitioners – Midwives and Nurses) (Jersey) Order 2018	R&O.55/2018	30 April 2018

Table of Renumbered Provisions

Original	Current
1(2)(b)	spent, omitted from this revised edition
1(2)(c)	1(2)(b)
10	spent, omitted from this revised edition
11	10

Table of Endnote References

[1] *This Order has been amended by the States of Jersey (Amendments and Construction Provisions No. 8) (Jersey) Regulations 2005. The amendments replace all references to a Committee of the States of Jersey with a reference to a Minister of the States of Jersey, and remove and add defined terms appropriately, consequentially upon the move from a committee system of government to a ministerial system of government*

[2] *chapter 26.500*

[3] *chapter 26.500.08*

[4] *chapter 26.500*

[5] *Article 1(1)* *amended by R&O.15/2008, R&O.55/2018*

[6] *Article 5(1)* *amended by R&O.55/2018*

[7] *Article 6(1)* *amended by R&O.55/2018*

[8] *Article 6(2)*	*amended by R&O.55/2018*
[9] *Article 6(3)*	*amended by R&O.55/2018*
[10] *Article 6(4)*	*amended by R&O.55/2018*
[11] *Article 6(5)*	*amended by R&O.55/2018*
[12] *Article 7(4)*	*amended by R&O.55/2018*
[13] *Article 7(5)*	*repealed by R&O.19/2008*
[14] *Article 7(6)*	*repealed by R&O.19/20084*
[15] *Article 7(7)*	*deleted by R&O.15/2008; repealed by R&O.19/2008*
[16] *Article 7(8)*	*deleted by R&O.15/2008; repealed by R&O.19/2008*
[17]	*chapter 08.680*
[18] *Article 9(5)*	*substituted by R&O.102/2017*
[19] *Article 9(6)*	*repealed by R&O.19/2008*
[20] *Schedule 2*	*amended by R&O.19/2008*
[21] *Schedule 3*	*substituted by R&O.35/2009*
[22] *Schedule 4*	*substituted by R&O.102/2017, amended by R&O.55/2018*
[23] *Schedule 5*	*amended by R&O.85/2003, R&O.90/2004, R&O.185/2007, R&O.122/2008, R&O.98/2009, R&O.99/2010, R&O.44/2013, R&O.119/2013, R&O.141/2014*
[24]	*chapter 08.680.60*
[25] *Schedule 6*	*substituted by R&O.141/2014*