



THE NATIONAL HEALTH SERVICE ACT 2001

**THE NATIONAL HEALTH SERVICE (GENERAL
PHARMACEUTICAL SERVICES) REGULATIONS 2004**

Laid before Tynwald

16th March 2004

Coming into operation

1st April 2004

In exercise of the powers conferred on the Department of Health and Social Security by section 8 of the National Health Service Act 2001¹, and of all other enabling powers, and after the consultations required by section 42(5) of that Act, the following Regulations are hereby made:—

1. Citation commencement and interpretation

(1) These Regulations may be cited as the National Health Service (General Pharmaceutical Services) Regulations 2004, and shall come into operation on the 1st April 2004.

(2) In these Regulations —

"the Act" means the National Health Service Act 2001;

"the Association" means the Isle of Man Pharmacy Contractors Association;

"contractor" means a person who has undertaken to provide general pharmaceutical services and whose name is included in the pharmaceutical list;

"dentist" means a dental practitioner (within the meaning of the Dental Act 1985²);

"doctor" means a registered medical practitioner who holds a licence to practise;

"drugs" includes medicines;

"Drug Tariff" has the meaning given by regulation 8;

"independent nurse prescriber" means —

(a) a person whose name is registered —

¹ 2001 c.14

² 1985 c.29

- (i) in Part 1 or 12 of the nurses and midwives' professional register and has a district nurse qualification additionally recorded in the nurses and midwives' professional register, or
- (ii) in Part 11 of the nurses and midwives' professional register as a health visitor,

and against whose name is recorded in the nurses and midwives' professional register an annotation signifying that he is qualified to order drugs, medicines and appliances from the Nurse Prescribers' Formulary for District Nurses and Health Visitors in Part XVIIIB(i) of the Drug Tariff; or

- (b) a person —
 - (i) whose name is registered in Parts 1, 3, 5, 8, 10, 11, 12, 13, 14 or 15 of the nurses and midwives' professional register, and
 - (ii) against whose name is recorded in the nurses and midwives' professional register an annotation signifying that he is qualified to order drugs, medicines and appliances from the Nurse Prescribers' Extended Formulary in Part XVIIIB(ii) of the Drug Tariff;

"nurses and midwives' professional register" means the register maintained by the Nursing and Midwifery Council pursuant to Article 2 of the Nursing and Midwifery Order 2002³;

"patient" means a person for whom a contractor has agreed to provide general pharmaceutical services;

"the pharmaceutical list" means a list of contractors maintained under section 9 of the Act;

"pharmacist" means a person registered in the register of pharmaceutical chemists maintained under section 2(1) of the Pharmacy Act 1954 (an Act of Parliament)⁴;

"pharmacy" means any premises registered under section 40 of the Medicines Act 2003⁵;

"prescription form" means a form provided by the Department, and issued by a doctor, dentist, supplementary prescriber or independent nurse prescriber to enable a person to obtain pharmaceutical services under the Act;

"scheduled drug" means a drug or other substance specified in the National Health Service (Scheduled Drugs) Regulations 2004⁶;

"supplementary prescriber" means a person whose name is registered in —

- (a) Part 1, 3, 5, 8, 10, 11, 12, 13, 14 or 15 of the nurses and midwives' professional register;

³ SI 2002/253

⁴ 1954 c.61

⁵ 2003 c.4

⁶ SD 21/04

- (b) the Register of Pharmaceutical Chemists maintained in pursuance of section 2(1) of the Pharmacy Act 1954 (an Act of Parliament); or
- (c) the register maintained in pursuance of Articles 6 and 9 of the Pharmacy (Northern Ireland) Order 1976⁷,

and against whose name is recorded in the relevant register an annotation signifying that he is qualified to order drugs, medicines and appliances as a supplementary prescriber;

"terms of service" means the terms set out in Schedule 1.

(3) Until the 1st January 2005 the definition of "doctor" in subsection (2) shall have effect with the omission of the words "who holds a licence to practice".

(4) Until the coming into operation of section 40 of the Medicines Act 2003, the reference to that section in the definition of "pharmacy" in subsection (2) shall have effect as a reference to section 74 of the Medicines Act 1976⁸.

2. Terms of service

The arrangements for the provision of general pharmaceutical services which it is the duty of the Department under section 8 of the Act to make shall incorporate the terms of service and the Drug Tariff.

3. Pharmaceutical list

- (1) The pharmaceutical list shall contain —
 - (a) the names of persons who are entitled to be included in it;
 - (b) the addresses of any places at which they have undertaken to provide general pharmaceutical services;
 - (c) particulars of the days on which and hours between which general pharmaceutical services will normally be available at those addresses;
 - (d) the names of every other pharmacist who is regularly engaged as a deputy, director or employee in the provision of pharmaceutical services at any of those addresses.

(2) The Department shall send a copy of the pharmaceutical list to the Isle of Man Medical Society and the Association and shall notify them of any change made in that list.

4. Application for inclusion in list

(1) A pharmacist who wishes to be included in the pharmaceutical list shall send to the Department an application to that effect which shall indicate whether

⁷ SI 1976/1213

⁸ 1976 c.22

the applicant is a pharmacist or a body corporate carrying on business as pharmacists, and shall include —

- (a) an undertaking to provide general pharmaceutical services and to comply with the terms of service; and,
- (b) the information, as respects the matters mentioned in regulation 3, which it is proposed shall be contained in the pharmaceutical list.

(2) The application shall be determined by an officer nominated by the Department for the purpose, who shall inform the applicant of their decision and, where the application is refused, the reasons for refusal and the right to appeal under regulation 5.

(3) A contractor shall within 14 days of any change or addition affecting the entries which the pharmaceutical list is required to contain in relation to him notify the Department accordingly.

5. Appeal

(1) Where an application to be included in the pharmaceutical list has been refused, the applicant may appeal to the Department by giving notice in writing to the Department, stating the grounds of appeal, within 21 days after refusal is notified to him, or within such further time as the Department may allow.

(2) The Department shall appoint a person (being a member or officer of the Department) to determine the appeal, who, if he considers that it can properly be determined without an oral hearing, shall determine the appeal summarily and notify the applicant of his decision.

(3) If the appointed person does not determine the appeal summarily under paragraph (2), he shall, not less than 7 days before the date fixed for the hearing, notify the appellant and the nominated officer of the time and place at which the appeal will be heard.

(4) The applicant and the nominated officer may attend the hearing and be heard either in person or by an advocate or other representative. The procedure at the hearing shall be such as the appointed person may determine.

(5) The appointed person shall as soon as may be thereafter notify the applicant and the nominated person in writing of his decision on the appeal, with the grounds therefor, and if he allows the appeal shall direct the nominated officer to grant the application.

6. Withdrawal from list

(1) Subject to paragraphs (2) and (3), where a contractor gives notice in writing to the Department that he wishes to withdraw from the pharmaceutical list, his name shall be removed from the list at the expiration of 3 months from the date of that notice or of such shorter period as the contractor and the Department may agree.

(2) If an enquiry is instituted as to whether the continued inclusion of a contractor would be prejudicial to the efficiency of the general pharmaceutical services, he shall not, except with the consent of the Department and subject to such conditions as it may impose, be entitled to have his name removed from the list pending the termination of the inquiry.

(3) The Department shall not agree to a contractor's withdrawal from the pharmaceutical list unless and until it is satisfied that satisfactory arrangements have been made for the completion of any general pharmaceutical services which he has undertaken to provide.

7. Removal from pharmaceutical list

(1) Where the Department has determined that a contractor —

- (a) has died, or
- (b) has otherwise ceased to be a pharmacist,

it shall remove his name from the pharmaceutical list

(2) If it appears to the Department that a contractor whose name has been included for the preceding 6 months in the pharmaceutical list has not during that period provided general pharmaceutical services for persons on the Island, it shall refer the matter to an officer nominated by it, who shall —

- (a) give to the contractor 28 days notice of the proposal to remove his name from the list;
- (b) afford the contractor an opportunity to make representations to him in writing or, if he so wishes, orally; and
- (c) consult the Association.

(3) No determination under this regulation shall be made in respect of any contractor who is called into —

- (a) whole-time service in the Armed Forces of the Crown in a national emergency as a volunteer or otherwise; or,
- (b) compulsory whole-time service in those forces, including service resulting from an reserve liability or any equivalent service by a person liable for whole-time service in those forces,

until 6 months after completion of that service.

(4) Nothing in these regulations prejudices the right of a person to have his name included again in the pharmaceutical list.

8. Drug Tariff

For the purpose of enabling arrangements to be made for the provision of general pharmaceutical services, the Drug Tariff, or any variation thereof, prepared by the Secretary of State under the National Health Act 1977 (an Act of Parliament) shall apply in the Island in respect of those services provided by the Department.

9. Payment for services

(1) The Department shall make payments to contractors in accordance with provisions of the Drug Tariff.

(2) Where the Department considers that it has made a payment to a contractor owing to an error or in circumstances where it was not due, it shall, draw the overpayment to the attention of the contractor and —

- (a) where he admits overpayment; or
- (b) where he does not admit the overpayment but, the matter having been referred for investigation, the Department decides there has been an overpayment,

the amount overpaid shall be recoverable either by deduction from the contractor's remuneration or in some other manner.

(3) Recovery of an overpayment under this regulation shall be without prejudice to the investigation of an alleged breach of the terms of service.

10. Exercise of choice of contractor in certain cases

(1) Subject to paragraph (2), an application for general pharmaceutical services required by these Regulations may be made and a signature required by these Regulations may be given —

- (a) on behalf of a person under 16 years of age by either parent, or in the absence of both parents, the guardian or other person who has the care of him;
- (b) on behalf of any other person who is incapable of making the application or, giving the signature, by a relative or any other adult who has the care of that person; or
- (c) on behalf of any person under 18 years of age in the care of the Department, by any person duly authorised by the Department.

(2) A signature on an application may not be given by the contractor to whom the application is made.

11. Supplemental services

(1) A contractor may, in addition, undertake to provide additional professional services.

(2) In these Regulations, "additional professional services" means-

- (a) publishing a leaflet (a "practice leaflet") which shall include —
 - (i) a list of the pharmaceutical services which the contractor has undertaken to provide and in respect of which his name is included in the pharmaceutical list,
 - (ii) the name, address and telephone number of the pharmacy from which he provides those services and the hours in each day of

the week during which he provides those services from those premises,

- (iii) the arrangements made by the contractor to provide, or such arrangements as the contractor has made with other contractors to provide, pharmaceutical services to any person who needs those services in an emergency or outside of the normal hours during which the contractor provides pharmaceutical services
 - (iv) the procedure by which any person may comment upon the provision of pharmaceutical services undertaken by the contractor;
- (b) displaying such health promotion leaflets as the Department may approve;
 - (c) keeping records in connection with drugs supplied to any person —
 - (i) who claims exemption under regulation 4(1) of the National Health Service (Charges for Drugs and Appliances) Regulations 2004⁹, or
 - (ii) who, in the opinion of the pharmacist providing the drug, is likely to have difficulty understanding the nature and dosage of the drug provided and the times at which it is to be taken,

in circumstances where the nature of the drug is such that, in the opinion of the pharmacist providing it, the same or a similar drug is likely to be prescribed for that person regularly on future occasions.

- (3) In paragraph (2)(c), "records" includes a record of —
 - (a) the name and address of the person to whom the drug is supplied;
 - (b) the name, quantity and dosage of the drug provided; and
 - (c) the date on which it is provided.

12. Revocations

The Regulations specified in column 1 of Schedule 2 are revoked to the extent specified in column 3 of that Schedule.

⁹ SD 22/04

Regulation 2

SCHEDULE 1 TERMS OF SERVICE

Incorporation of provisions

1. Any provisions of the following affecting the rights and obligations of contractors shall be deemed to form part of the terms of service for contractors —

- (a) these Regulations; and
- (b) the Drug Tariff.

Provision of pharmaceutical services

2. (1) Where any person presents on a prescription form —
- (a) an order for drugs, not being scheduled drugs, or appliances, signed by a doctor or a supplementary prescriber; or
 - (b) an order for a drug specified in Schedule 2 to the National Health Service (Scheduled Drugs) Regulations 2004, signed by, and endorsed on its face with the reference "SLS" by, a doctor or a supplementary prescriber; or
 - (c) an order for listed drugs or medicines, signed by a dentist or his deputy or assistant, or
 - (d) an order for listed drugs or medicines or listed appliances, signed by an independent nurse prescriber,

a contractor shall, with reasonable promptness, provide the drugs or medicines so ordered, and such of the appliances so ordered as he supplies in the normal course of his business.

(2) If the person presenting the prescription form asks the contractor to do so he shall give an estimate of the time when the drugs, medicines or appliances will be ready; and if they are not ready by then, he shall give a revised estimate of the time when they will be ready (and so on).

(3) Where a contractor reasonably believes that a form presented to him as a prescription form in accordance with sub-paragraph (1) is not a genuine order for the person named on the form (for example because he reasonably believes the form has been stolen or forged), he may refuse to provide the drugs or medicines or listed appliances specified on the form presented.

(4) A contractor shall provide any drug which he is required to provide under this paragraph in a suitable container.

(5) A contractor shall not give, promise or offer to any person any gift or reward (whether by way of a share of or dividend on the profits of the business or by way of discount or rebate or otherwise) as an inducement to or in consideration of his presenting an order for drugs or appliances on a prescription form.

(6) In this paragraph "listed", in relation to drugs, medicines or appliances, means specified in the Drug Tariff as being drugs, medicines or appliances which may be prescribed by a dentist or an independent nurse prescriber, as the case may be.

Provision of drugs: supplementary

3. (1) Any drug which is provided as part of pharmaceutical services and included in the Drug Tariff, the British National Formulary (including any Appendix published as part of that Formulary), the Dental Practitioner's Formulary, the European Pharmacopoeia or the British Pharmaceutical Codex, shall comply with the standard or formula specified therein.

(2) Subject to any regulations in force under the Weights and Measures Act 1989¹⁰ and subject to sub-paragraphs (3) to (11) a contractor shall provide pharmaceutical services only in response to and in accordance with an order on a prescription form, signed as specified in sub-paragraph (1).

¹⁰ 1989 c.1

(3) Where an order, not being an order to which the Poisons Rules 1982¹¹ (as they have effect in the Island¹²) or the Misuse of Drugs Regulations 2001¹³ (as they have effect in the Island¹⁴) apply, which is issued by a doctor, dentist, supplementary prescriber or independent nurse prescriber on a prescription form for drugs does not prescribe their quantity, strength or dosage, a contractor may provide the drugs in such strength and dosage as in the exercise of his professional skill, knowledge and care he considers to be appropriate and, subject to sub-paragraph (5), in such quantity as he considers to be appropriate for a course of treatment, for the patient to whom the order relates, for a period not exceeding 5 days.

(4) Where an order to which sub-paragraph (3) applies is for —

- (a) an oral contraceptive substance;
- (b) a drug, which is available for supply as part of pharmaceutical services only together with one or more drugs; or
- (c) an antibiotic in a liquid form for oral administration in respect of which pharmaceutical considerations require its provision in an unopened package,

which is not available for provision as part of pharmaceutical services except in such packages that the minimum available package contains a quantity appropriate to a course of treatment for a patient for a period of more than 5 days, the contractor may provide that minimum available package.

(5) Where any drug, not being one to which the Misuse of Drugs Regulations 2001 (as they have effect in the Island) apply, ordered by a doctor, dentist, a supplementary prescriber or independent nurse prescriber on a prescription form, is available for provision by a contractor in a pack in a quantity which is different from the quantity which has been so ordered, and that drug is —

- (a) sterile;
- (b) effervescent or hygroscopic;
- (c) a liquid preparation for addition to bath water;
- (d) a coal tar preparation;
- (e) a viscous preparation; or
- (f) packed at the time of its manufacture in a calendar pack or special container,

the contractor shall, subject to sub-paragraph (6), provide the drug in the pack whose quantity is nearest to the quantity which has been so ordered.

(6) A contractor shall not provide pursuant to sub-paragraph (5) a drug in a calendar pack where, in his opinion, it was the intention of the doctor, dentist, a supplementary prescriber or independent nurse prescriber who ordered the drug that it should be provided only in the exact quantity ordered.

(7) In this paragraph —

- (a) "calendar pack" means a blister or strip pack showing the days of the week or month against each of the several units in the pack; and
- (b) "special container" means any container with an integral means of application or from which it is not practicable to dispense an exact quantity.

(8) Where, in a case of urgency, a doctor, a supplementary prescriber or independent nurse prescriber personally known to a pharmacist requests him to provide a drug, the pharmacist may provide that drug before receiving a prescription form, provided that —

- (a) that drug is not a scheduled drug;

¹¹ SI 1982/218

¹² GC 137/82

¹³ SI 2001/3998

¹⁴ SD 72/02

- (b) that drug is not a controlled drug within the meaning of the Misuse of Drugs Act 1976¹⁵, other than a drug which is for the time being specified in Schedule 4 or 5 to the Misuse of Drugs Regulations 2001 (as they have effect in the Island); and
 - (c) the doctor, supplementary prescriber or independent nurse prescriber undertakes to give the contractor such a prescription form within 72 hours.
- (9) Except as provided in sub-paragraph (10), a contractor shall not provide a scheduled drug, by way of pharmaceutical services or otherwise, in response to an order by name, formula or other description on a prescription form.
- (10) Where a drug has is ordered on a prescription form either by a name or by a formula which is not included within Schedule 1 of the National Health Service (Scheduled Drugs) Regulations 2004, a contractor may provide a drug which has the same specification notwithstanding that it is a Scheduled drug, provided that where a Scheduled drug is a pack which consists of a drug in more than one strength, such provision does not involve the supply of part only of the pack.
- (11) Where a drug which is ordered as specified in sub-paragraph (10) combines more than one drug, that sub-paragraph shall apply only if the combination has an appropriate non-proprietary name, whether the individual drugs which it combines do so or not.

Premises and hours

4. (1) General pharmaceutical services shall be provided at each of the premises from which the contractor has undertaken to provide general pharmaceutical services at such times as, following an application in writing by the contractor, shall have been approved in his case by the Department.
- (2) The Department shall determine an application within 30 days of receiving it.
- (3) The Department shall notify the contractor in writing of his determination, and, where he refuses an application, it shall send the contractor a statement in writing of the reasons for the determination and of the contractor's right of appeal under regulation 5.
- (4) At each of the premises at which a contractor provides pharmaceutical services he shall exhibit —
- (a) a notice provided by the Department specifying the times at which the premises are open for the provision of drugs and appliances; and
 - (b) at times when the premises are not open, a notice, where practicable legible from outside the premises, specifying the addresses of other contractors included in the pharmaceutical list and the times at which drugs and appliances may be obtained from those addresses.
- (5) The Department shall notify the contractor in writing of the names and addresses of other contractors included in the pharmaceutical list and of the times at which they are required to provide general pharmaceutical services.
- (6) Where a contractor is prevented by illness or other reasonable cause from complying with his obligations under this paragraph, he shall, where practicable, make arrangements with one or more contractors whose premises are situated in the neighbourhood for the provision of pharmaceutical services during that time.

Provision of drugs and fitting of appliances

5. (1) Drugs shall be provided either by or under the direct supervision of a pharmacist.
- (2) Subject to paragraph 3(1), a contractor shall make all necessary arrangements —
- (a) for measuring a person who presents a prescription for a truss or other appliance of a type requiring measurement and fitting by the contractor; and
 - (b) for fitting the appliance.

¹⁵ 1976 c.21

Particulars of contractors

6. A contractor shall give the Department, if it so requires, the name of any pharmacist employed by him for the provision of drugs for persons from whom he has accepted an order for the provision of pharmaceutical services under paragraph 3.

Charges for drugs

7. (1) Subject to regulations made under section 32 of the Act, all drugs, containers and appliances provided under these terms of service shall be provided free of charge.

(2) Where a contractor supplies a container in response to an order for drugs signed by a doctor, a supplementary prescriber or independent nurse prescriber, other than equipment specified in the Drug Tariff as not returnable to the contractor, the container and equipment shall remain the property of the contractor.

Remuneration of contractors

8. (1) A contractor who has undertaken to provide additional professional services within the meaning of regulation 11 shall, on request, make available to the Department all records kept in accordance with regulation 11(2)(c); and permit the Department at any reasonable time to inspect the premises from which those services are provided for the purpose of satisfying itself that those services are being provided in accordance with the undertaking.

(2) The Department shall make payments, calculated in the manner provided by the Drug Tariff to contractors in respect of drugs and appliances, containers, medicines measures and dispensing fees and additional professional services within the meaning of regulation 11.

(3) Where a contractor so requires, the Department shall afford him reasonable facilities for examining all or any of the forms on which the drugs or appliances provided by him were ordered, together with particulars of the amounts calculated to be payable in respect of such drugs and appliances and the Department shall take into consideration any objections made by the contractor in relation to those amounts.

(4) Where so required by the Association the Department shall give them similar facilities for examining such forms and particulars relating to all or any of the contractors which it represents.

Professional standards

9. (1) A pharmacist whose name is on the pharmaceutical list shall provide pharmaceutical services and exercise any professional judgement in connection with the provision of such services in conformity with the standards generally accepted in the pharmaceutical profession.

(2) A contractor who employs a pharmacist in connection with the provision of pharmaceutical services shall secure that the pharmacist complies with the requirements set out in sub-paragraph (1).

Complaints

10. (1) Subject to sub-paragraph (2) and (3), a contractor shall establish, and operate in accordance with this paragraph, a procedure (a "complaints procedure") to deal with any complaints made by or on behalf of any person to whom he has provided pharmaceutical services.

(2) The complaints procedure to be established by a contractor may be such that it also deals with complaints made in relation to one or more other contractors.

(3) The complaints procedure to be established by a contractor who provides general pharmaceutical services from more than one set of premises may be such that it relates to all those premises together.

(4) A complaints procedure shall apply to complaints made in relation to any matter reasonably connected with the contractor's provision of general pharmaceutical services and within the responsibility or control of—

- (a) the contractor;
- (b) where the contractor is a body corporate, any of its directors or former directors;
- (c) a former partner of the contractor;

- (d) any employee of the contractor .
- (5) A complaint may be made on behalf of any patient or former patient with his consent,
or —
 - (a) where he is under 16 years of age —
 - (i) by either parent, or in the absence of both parents, the guardian or other adult person who has care of the child, or
 - (ii) by the Department, where he is the care of the Department.
 - (b) where the patient is incapable of making a complaint, by a relative or other adult person who has an interest in his welfare.
- (6) Where a patient has died, a complaint may be made by a relative or other adult person who had an interest in his welfare or, where he was as described in paragraph (5)(a)(ii), by the Department.
- (7) A complaints procedure shall comply with the following requirements —
 - (a) the contractor must specify a person (who need not be connected with the contractor and who, in the case of an individual, may be specified by his job title) to be responsible for receiving and investigating all complaints.
 - (b) all complaints must be —
 - (i) recorded in writing.
 - (ii) acknowledged, either orally or in writing, within the period of 3 working days beginning with the day on which the complaint was received by the person specified under paragraph (a), or where that is not possible as soon as reasonably practicable, and
 - (iii) properly investigated;
 - (c) within the period of 10 working days beginning with the day on which the complaint was received by the person specified under paragraph (a) above, or where that is not possible as soon as reasonably practicable, the complainant must be given a written summary of the investigation and its conclusions;
 - (d) where the investigation of the complaint requires consideration of the patient's records, the person specified under (a) above must inform the patient or person acting on his behalf if the investigation will involve disclosure of information contained in those records to a person other than the contractor, or a director, partner, deputy or employee of the contractor; and
 - (e) the contractor must keep a record of all complaints and copies of all correspondence relating to complaints, but such records must be kept separate from any records relating to the person by whom the complaint was made.
- (8) At each of the premises at which the contractor provides general pharmaceutical services he must provide information about the complaints procedure and give the name (or title) and address of the person specified under paragraph (7)(a).

Complaints investigated by the Department

- 11. (1) A contractor shall co-operate with any investigation of a complaint by the Department, whether the investigation follows one under the contractor's complaints procedure or not.
 - (1) The co-operation required by sub-paragraph (1) includes —
 - (a) answering questions reasonably put to the contractor by the Department;
 - (b) providing any information relating to the complaint reasonably required by the Department; and
 - (c) attending any meeting to consider the complaint (if held at a reasonably accessible place and at a reasonable hour, and due notice has been given), if the contractor's presence at the meeting is reasonably required by the Department.

Regulation 12.

SCHEDULE 2
REGULATIONS REVOKED

<i>Reference</i>	<i>Title</i>	<i>Extent of revocation</i>
GC 226/78	National Health Service (Isle of Man) General Medical and Pharmaceutical Services Regulations 1978.	The whole Regulations
GC 2/80	National Health Service(Pharmaceutical List Committee) Regulations 1980	The whole Regulations
GC 153/85	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) Regulations 1985	The whole Regulations
GC 191/85	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) (No.2) Regulations 1985	The whole Regulations
GC 190/88	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) Regulations 1988	The whole Regulations
	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) (No.2) Regulations 1990	The whole Regulations
GC 76/91	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) (No.2) Regulations 1991	The whole Regulations
GC 472/92	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) (No.2) Regulations 1992	The whole Regulations
SD 254/93	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) Regulations 1993	The whole Regulations
SD 504/94	National Health Service (Isle of Man) General Medical and Pharmaceutical Services (Amendment) (No.2) Regulations 1995	The whole Regulations

MADE

27/2/04.

2004



Minister for Health and Social Security.

EXPLANATORY NOTE

(This note is not part of the Regulations.)

These Regulations consolidate, with amendments, those provisions of the National Health Service (Isle of Man) (General Medical and Pharmaceutical Services) Regulations 1978 which relate to general pharmaceutical services.