
STATUTORY INSTRUMENTS

2015 No. 570

NATIONAL HEALTH SERVICE, ENGLAND

**The National Health Service (Charges for
Drugs and Appliances) Regulations 2015**

<i>Made</i>	- - - -	<i>4th March 2015</i>
<i>Laid before Parliament</i>		<i>11th March 2015</i>
<i>Coming into force</i>	- -	<i>1st April 2015</i>

The Secretary of State for Health makes the following Regulations in exercise of the powers conferred by sections 172(1) and (2), 174, 178, 182, 184(1) and 272(7) and (8) of the National Health Service Act 2006(1).

Citation and commencement

1. These Regulations may be cited as the National Health Service (Charges for Drugs and Appliances) Regulations 2015 and come into force on 1st April 2015.

Interpretation

2.—(1) In these Regulations—

“the 2000 Act” means the Electronic Communications Act 2000(2);

“the 2006 Act” means the National Health Service Act 2006;

“accepted disablement” means physical or mental injury or disease which is accepted by the Secretary of State as attributable to, or aggravated by, service in the armed forces of the Crown or such other service as the Secretary of State may determine;

“advanced electronic signature” means an electronic signature which is—

- (a) uniquely linked to the signatory;
- (b) capable of identifying the signatory;
- (c) created using means that a signatory can maintain under their sole control; and

(1) 2006 c. 41. See section 275(1) of the National Health Service Act 2006 (“the 2006 Act”) which contains definitions of “prescribed” and “regulations” that are relevant to the powers being exercised. By virtue of section 271(1) of the 2006 Act, the functions of the Secretary of State being exercised in the making of these Regulations are exercisable only in relation to England.

(2) 2000 c. 7.

(d) linked to the data to which it relates in such a manner that any subsequent change of data is detectable;

“another primary care organisation” means—

- (a) as regards Wales, a Local Health Board;
- (b) as regards Scotland, a Health Board constituted under section 2 of the National Health (Scotland) Act 1978(3) (Health Boards);
- (c) as regards Northern Ireland, the Regional Health and Social Care Board established under section 7 of the Health and Social Care (Reform) Act (Northern Ireland) 2009(4) (Regional Health and Social Care Board); and
- (d) as regards England, in relation to any time prior to 1st April 2013, a Primary Care Trust;

“APMS contractor” means a person or a partnership that provides primary medical services under contractual arrangements with the Board under section 83(2) of the 2006 Act(5) (primary medical services), and “APMS contract” is to be construed accordingly;

“appliance”, unless the context otherwise requires, means an appliance included in a list approved by the Secretary of State for the purposes of section 126 of the 2006 Act(6) (arrangements for pharmaceutical services), but does not include a contraceptive appliance;

“approved”, in relation to forms, means approved by the Secretary of State;

“armed forces of the Crown” means the forces that are “regular forces” or “reserve forces” within the meanings given in section 374 of the Armed Forces Act 2006(7) (definitions applying for the purposes of the whole Act);

“arrangements for recharging” means arrangements under paragraph 3 of Schedule 12A to the 2006 Act(8) (pharmaceutical remuneration – other pharmaceutical remuneration) under which the Board requires a person to reimburse it for any pharmaceutical remuneration to which that paragraph applies;

“bank holiday” means any day that is, by virtue of section 1 of, or Schedule 1 to, the Banking and Financial Dealings Act 1971(9) (which relate to bank holidays), a bank holiday in England;

“batch issue” means an approved form, in the format required by the Board (or person exercising its functions), which—

- (a) is issued by a prescriber at the same time as a non-electronic repeatable prescription to enable a chemist or a dispensing doctor to receive payment for the provision of repeat dispensing services;
- (b) relates to a particular non-electronic repeatable prescription and contains the same date as that prescription;
- (c) is generated by a computer and not signed by a prescriber;

(3) 1978 c. 29. Section 2 has been amended by: the Health and Social Services and Social Security Adjudications Act 1983 (c. 41), Schedule 7, paragraph 1; the National Health Service and Community Care Act 1990 (c. 19), section 28, Schedule 9, paragraph 19(1), and Schedule 10; the National Health Service Reform (Scotland) Act 2004 (asp 7), Schedule 1, paragraph 1(2); the Smoking, Health and Social Care (Scotland) Act 2005 (asp 13), Schedule 2, paragraph 2(2); and the Health Boards (Membership and Elections) (Scotland) Act 2009 (asp 5), section 2(1).

(4) 2009 c. 1.

(5) Subsection (2) was substituted by the Health and Social Care Act 2012 (c. 7), Schedule 4, paragraph 30.

(6) Section 126 has been amended by the Health and Social Care Act 2012, sections 213(7)(k) and 220(7), and Schedule 4, paragraph 63. See section 126(9), which provides a definition of “listed” that includes the power for the Secretary of State to approve lists for the purposes of section 126.

(7) 2006 c. 52.

(8) Schedule 12A was inserted by the Health and Social Care Act 2012 (c. 7), Schedule 3.

(9) 1971 c. 80.

(d) is issued as one of a sequence of forms, the number of which is equal to the number of occasions on which the drugs or appliances ordered on the non-electronic repeatable prescription may be provided; and

(e) has included on it a number denoting its place in the sequence referred to in subparagraph (d);

“CCG” means a clinical commissioning group;

“chemist” means an LPS chemist or a person included in a pharmaceutical list of the type referred to in regulation 10(2)(a) or (b) of the Pharmaceutical and Local Pharmaceutical Services Regulations (pharmaceutical lists and EPS lists);

“chiropodist or podiatrist independent prescriber” means a chiropodist or podiatrist who is registered in Part 2 of the register maintained under article 5 of the Health and Social Work Professions Order 2001(10) (establishment and maintenance of register), and against whose name in that register is recorded an annotation signifying that the chiropodist or podiatrist is qualified to order drugs and appliances as a chiropodist or podiatrist independent prescriber;

“community treatment order” means an order under section 17A(1) of the Mental Health Act 1983(11) (community treatment orders);

“directed services” means additional pharmaceutical services provided in accordance with directions under section 127 of the 2006 Act(12) (arrangements for additional pharmaceutical services);

“dispensing doctor” is to be construed in accordance with regulation 46(1) of the Pharmaceutical and Local Pharmaceutical Services Regulations (dispensing doctor lists);

“dispensing services” means the provision of drugs or appliances that may be provided as pharmaceutical services by a medical practitioner in accordance with arrangements made under sections 126 and 132 of the 2006 Act(13) (arrangements for pharmaceutical services, and persons authorised to provide pharmaceutical services);

“doctor” means a registered medical practitioner;

“drugs” includes medicines, but does not include contraceptive substances;

“elastic hosiery” means anklet, legging, knee-cap, below-knee or thigh stocking;

“electronic communication” has the same meaning as in section 15(1) of the 2000 Act(14) (general interpretation);

“electronic prescription form” means data created in an electronic form for the purpose of ordering a drug or appliance, which—

(a) is compatible with the systems used by the Board (or a person exercising its functions) for—

(i) the remuneration of persons providing pharmaceutical services or local pharmaceutical services, and

(ii) any apportionment of, or any arrangements for recharging in respect of, that remuneration,

(10) S.I. 2002/254; article 5 has been amended by S.I. 2009/1182. This Order was so renamed by the Health and Social Care Act 2012, section 213(4) and (6).

(11) 1983 c. 20. Section 17A(1) was inserted by the Mental Health Act 2007 (c. 12), section 32(1) and (2).

(12) Section 127 has been amended by the Health and Social Care Act 2012, Schedule 4, paragraph 65.

(13) Section 132 has been amended by the Health and Social Care Act 2012, Schedule 4, paragraph 69, the Protection of Freedoms Act 2012 (c. 9), Schedule 9, paragraphs 120 and 122, and by S.I. 2007/289 and 2010/22 and 231.

(14) The definition of “electronic communication” has been amended by the Communications Act 2003 (c. 21), Schedule 17, paragraph 158.

unless the chemist or dispensing doctor dispensing the prescription is to receive no pharmaceutical remuneration of any kind in respect of any drug or appliance ordered on the form;

- (b) is signed with a prescriber's advanced electronic signature;
- (c) is transmitted as an electronic communication to a nominated dispensing contractor by the Electronic Prescription Service; and
- (d) does not indicate that the drug or appliance ordered may be provided more than once;

"electronic repeatable prescription" means data created in an electronic form, which—

- (a) is signed with a prescriber's advanced electronic signature;
- (b) is transmitted as an electronic communication to a nominated dispensing contractor by the Electronic Prescription Service;
- (c) indicates that the drug or appliance ordered may be provided more than once; and
- (d) specifies the number of occasions on which they may be provided;

"Electronic Prescription Service" means the service of that name which is managed by the Information Centre;

"electronic signature" has the same meaning as in section 7 of the 2000 Act (electronic signatures and related certificates);

"enhanced services" means the directed services which the Board is authorised (as opposed to required) to arrange by virtue of directions under section 127 of the 2006 Act;

"exemption certificate", unless the context otherwise requires, has the meaning assigned to it by regulation 15(1);

"GMS contract" means a general medical services contract, and "GMS contractor" is to be construed accordingly;

"independent nurse prescriber" means a person—

- (a) who is registered in the register maintained by the Nursing and Midwifery Council under article 5 of the Nursing and Midwifery Order 2001⁽¹⁵⁾ (establishment and maintenance of register); and
- (b) against whose name in that register is recorded an annotation signifying that they are qualified to order drugs and appliances as a community practitioner nurse prescriber, a nurse independent prescriber or a nurse independent/supplementary prescriber;

"Information Centre" means the Health and Social Care Information Centre established under section 252 of the Health and Social Care Act 2012⁽¹⁶⁾ (the Health and Social Care Information Centre);

"LPS chemist" means a party to an LPS scheme other than the commissioning body;

"mental disorder" has the same meaning as in section 1 of the Mental Health Act 1983⁽¹⁷⁾ (application of Act: "mental disorder");

"NHS services" means services provided as part of the health service (including services provided as part of the health service in pursuance of the public health functions of the Secretary of State or local authorities);

"nominated dispensing contractor" means a chemist or a dispensing doctor who has been nominated in a particular patient's PDS patient details to dispense the electronic prescriptions of that patient;

⁽¹⁵⁾ S.I. 2002/253. Article 5 was amended by S.I. 2009/1182.

⁽¹⁶⁾ 2012 c. 7.

⁽¹⁷⁾ Section 1 has been amended by the Mental Health Act 2007 (c. 12), sections 1, 2 and 3, and Schedule 11, Part 1.

“non-electronic prescription form” means an approved form for ordering a drug or appliance which—

(a) is compatible with the systems used by the Board (or person exercising its functions) for—

- (i) the remuneration of persons providing pharmaceutical services and local pharmaceutical services, and
- (ii) any apportionment of, or any arrangements for recharging in respect of, that remuneration,

unless the chemist or dispensing doctor dispensing the prescription is to receive no pharmaceutical remuneration of any kind in respect of any drug or appliance ordered on the form;

(b) has been provided for use by a prescriber by—

- (i) the Board,
- (ii) another primary care organisation,
- (iii) a local authority, under arrangements for providing NHS services which include, with the consent of the Board, the dispensing of prescriptions as part of pharmaceutical or local pharmaceutical services,
- (iv) the Secretary of State,
- (v) a CCG, under arrangements for providing NHS services which include, with the consent of the Board, the dispensing of prescriptions as part of pharmaceutical or local pharmaceutical services,
- (vi) an NHS trust, or
- (vii) an NHS foundation trust;

(c) has been issued by a prescriber; and

(d) does not indicate that the drug or appliance ordered may be provided more than once;

“non-electronic repeatable prescription” means a repeatable prescription which is not an electronic repeatable prescription;

“optometrist independent prescriber” means a person—

(a) who is registered in the register of optometrists maintained by the General Optical Council in pursuance of section 7 of the Opticians Act 1989⁽¹⁸⁾ (register of opticians); and

(b) against whose name in that register is recorded an annotation that the person is qualified to order drugs and appliances as an optometrist independent prescriber;

“out of hours period” means—

(a) the period beginning at 6.30pm on any day from Monday to Thursday and ending at 8am on the following day;

(b) the period beginning at 6.30pm on Friday and ending at 8am on the following Monday; and

(c) Good Friday, Christmas Day and bank holidays;

“out of hours services” means the services commissioned in respect of all, or part of, the out of hours period to enable the registered patients of a GMS, PMS or APMS contractor to receive primary medical services outside core hours, and in this definition—

(18) 1989 c. 44. Section 7 has been amended by S.I. 2005/484.

- (a) “core hours” means the period beginning at 8am and ending at 6.30pm on any day from Monday to Friday except Good Friday, Christmas Day and bank holidays; and
- (b) “part” of an out of hours period means any part of one or more of the periods described in paragraphs (a) to (c) of the definition of “out of hours period”;

“PDS patient details” means the information held about a patient in the Patient Demographics Service managed by the Information Centre;

“the Pharmaceutical and Local Pharmaceutical Services Regulations” means the National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013(19);

“pharmacist independent prescriber” means a registered pharmacist—

- (a) against whose name in Part 1 of the register maintained under article 19 of the Pharmacy Order 2010(20) (establishment, maintenance of and access to the register) or in the register maintained under articles 6 and 9 of the Pharmacy (Northern Ireland) Order 1976(21) (which relates to the registers and the registrar) is recorded an annotation signifying that the pharmacist is qualified to order drugs and appliances as a pharmacist independent prescriber; and
- (b) who is prescribing under arrangements for the provision of NHS services, but for these purposes those arrangements must not be arrangements for the provision of—
 - (i) pharmaceutical services, unless they are arrangements for the provision of enhanced services, or
 - (ii) local pharmaceutical services, unless they are arrangements that have been made by the Board with an LPS chemist for the provision of services that are equivalent to services provided as enhanced services;

“physiotherapist independent prescriber” means a physiotherapist who is registered in Part 9 of the register maintained under article 5 of the Health and Social Work Professions Order 2001, and against whose name in that register is recorded an annotation signifying that the physiotherapist is qualified to order drugs and appliances as a physiotherapist independent prescriber;

“PMS agreement” means an agreement made under section 92 of the 2006 Act(22) (arrangements by the Board for the provision of primary medical services);

“PMS contractor” means a contractor within the meaning of the National Health Service (Personal Medical Services Agreements) Regulations 2004(23);

“pre-payment certificate”, unless the context otherwise requires, is to be construed in accordance with regulation 16;

“prescriber” means a doctor, a dental practitioner, a pharmacist independent prescriber, a supplementary prescriber, a chiropodist or podiatrist independent prescriber, a physiotherapist independent prescriber, an independent nurse prescriber or an optometrist independent prescriber;

“prescription form”, except in the context of the expression “electronic prescription form” or “non-electronic prescription form”, means an electronic prescription form or a non-electronic prescription form;

“provider of NHS services” means a person or a body which provides services under the 2006 Act pursuant to arrangements made with—

(19) S.I. 2013/349; amended by S.I. 2014/417 and 2015/58.

(20) S.I. 2010/231.

(21) S.I. 1976/1213 (N.I. 22).

(22) Section 92 has been amended by the Health and Social Care Act 2012 (c. 7), Schedule 4, paragraph 36.

(23) S.I. 2004/627. The definition of “contractor” in regulation 2(1) of those Regulations was substituted by S.I. 2013/363.

- (a) the Board;
- (b) a CCG;
- (c) an NHS trust;
- (d) an NHS foundation trust;
- (e) a local authority in the exercise of its public health functions—
 - (i) under section 2B of the 2006 Act⁽²⁴⁾ (functions of local authorities and Secretary of State as to improvement of public health),
 - (ii) under section 111 of the 2006 Act⁽²⁵⁾ (dental public health),
 - (iii) under paragraphs 1 to 7B and 13 of Schedule 1 to the 2006 Act⁽²⁶⁾ (further provision about the Secretary of State and services under the Act), or
 - (iv) pursuant to regulations made under section 6C of the 2006 Act⁽²⁷⁾ (regulations as to the exercise by local authorities of certain public health functions); or
- (f) the Secretary of State when exercising the public health functions of the Secretary of State (as defined in section 1H(5)(a) of the 2006 Act⁽²⁸⁾ (the Board and its commissioning functions));

“provider of out of hours services” means—

- (a) an APMS contractor or a PMS contractor which provides out of hours services under its APMS contract or PMS agreement;
- (b) a GMS contractor which provides out of hours services under its GMS contract; or
- (c) an out of hours services sub-contractor of a GMS contractor, PMS contractor or APMS contractor;

“registered pharmacist” means a person who is registered in Part 1 or 4 of the register maintained under article 19 of the Pharmacy Order 2010 or in the register maintained under articles 6 and 9 of the Pharmacy (Northern Ireland) Order 1976;

“repeat dispensing services” means pharmaceutical services or local pharmaceutical services which involve the provision of drugs or appliances by a chemist in accordance with a repeatable prescription;

“repeatable prescription” means an electronic repeatable prescription or an approved form for ordering drugs or appliances which—

- (a) is compatible with the systems used by the Board (or a person exercising its functions) for—
 - (i) the remuneration of persons providing pharmaceutical services or local pharmaceutical services, and
 - (ii) any apportionment of, or any arrangements for recharging in respect of, that remuneration,

unless the chemist or dispensing doctor dispensing the prescription is to receive no pharmaceutical remuneration of any kind in respect of any drug or appliance ordered on the form;

- (b) has been provided for use by a prescriber by—

⁽²⁴⁾ Section 2B was inserted by the Health and Social Care Act 2012 (c. 7), section 12.

⁽²⁵⁾ Section 111 has been amended by the Health and Social Care Act 2012, section 29.

⁽²⁶⁾ Relevant amendments have been made to Schedule 1 by the Health and Social Care Act 2012, sections 17 and 143(1), and Schedule 14, paragraph 6, and by [S.I. 2010/1158](#).

⁽²⁷⁾ Section 6C was inserted by the Health and Social Care Act 2012, section 18(1).

⁽²⁸⁾ Section 1H was inserted by the Health and Social Care Act 2012, section 9(1).

- (i) the Board,
- (ii) another primary care organisation,
- (iii) a local authority, under arrangements for providing NHS services which include, with the consent of the Board, the dispensing of prescriptions as part of pharmaceutical or local pharmaceutical services,
- (iv) the Secretary of State,
- (v) a CCG, under arrangements for providing NHS services which include, with the consent of the Board, the dispensing of prescriptions as part of pharmaceutical or local pharmaceutical services,
- (vi) an NHS trust, or
- (vii) an NHS foundation trust;
- (c) has been issued by a prescriber;
- (d) indicates that the drugs or appliances ordered may be provided more than once; and
- (e) specifies the number of occasions on which they may be provided;

“the Standing Rules Regulations” means the National Health Service Commissioning Board and Clinical Commissioning Groups (Responsibilities and Standing Rules) Regulations 2012⁽²⁹⁾;

“supplementary prescriber” means—

- (a) a registered pharmacist against whose name in the register (“register” has the same meaning here as in the definition of “registered pharmacist”), is recorded an annotation signifying that they are qualified to order drugs and appliances as a supplementary prescriber;
- (b) a person—
 - (i) who is registered in a part of the register under article 5 of the Health and Social Work Professions Order 2001 which relates to chiropodists and podiatrists, physiotherapists or radiographers, and
 - (ii) against whose name in that register is recorded an annotation signifying that they are qualified to order drugs and appliances as a supplementary prescriber; or
- (c) an optometrist against whose name in the register of optometrists maintained under section 7 of the Opticians Act 1989 is recorded an annotation signifying that the optometrist is qualified to order drugs and appliances as a supplementary prescriber;

“the Board” means the National Health Service Commissioning Board⁽³⁰⁾;

“the Travel Expenses and Remission of Charges Regulations” means the National Health Service (Travel Expenses and Remission of Charges) Regulations 2003⁽³¹⁾;

“treatment” includes examination and diagnosis; and

“walk-in centre” means a centre at which information and treatment for minor conditions is provided to the public under arrangements made by the Board or a CCG.

(2) For the purposes of these Regulations, the supply against an order on one prescription form, or on one repeatable prescription (but only where the supply is against one batch issue relating to that repeatable prescription)—

⁽²⁹⁾ S.I. 2012/2996.

⁽³⁰⁾ The National Health Service Commissioning Board is established by section 1H of the National Health Service Act 2006. Section 1H is inserted by section 9 of the Health and Social Care Act 2012 (c. 7).

⁽³¹⁾ S.I. 2003/2382.

- (a) of quantities of the same drug in more than one container is to be treated as the supply of only one quantity of a drug;
- (b) of more than one appliance of the same type, except in the case of elastic hosiery and tights, or of two or more component parts of the same appliance, is to be treated as the supply of only one appliance.

Supply of drugs and appliances by chemists

3.—(1) Except as provided in paragraph (2), a chemist who provides pharmaceutical services or local pharmaceutical services to a patient must, subject to paragraphs (5) to (7), make and recover from that patient for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) each other appliance, a charge of £8.20;
- (c) each quantity of a drug, a charge of £8.20.

(2) A chemist who provides repeat dispensing services to a patient must, subject to paragraphs (5) to (7), make and recover from that patient in respect of each batch issue and each electronic prescription form for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) each other appliance, a charge of £8.20;
- (c) each quantity of a drug, a charge of £8.20.

(3) Where a charge is paid under paragraph (1), the person paying the charge must on doing so either—

- (a) where a non-electronic prescription form has been issued, sign a declaration in writing on the non-electronic prescription form that the relevant charge has been paid; or
- (b) where an electronic prescription form has been created, provide a declaration that the relevant charge has been paid on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a chemist.

(4) Where a charge is paid under paragraph (2), the person paying the charge must on doing so either—

- (a) in respect of a batch issue, sign a declaration in writing on the batch issue that the relevant charge has been paid; or
- (b) in respect of an electronic repeatable prescription, provide a declaration that the relevant charge has been paid on an approved form provided by the Board for recording patient declarations in respect of electronic repeatable prescriptions and issued by a chemist.

(5) No charge is to be made and recovered under paragraph (1) or paragraph (2) where—

- (a) there is an exemption by virtue of regulation 10(1) and, subject to regulation 10(5), a declaration of entitlement to an exemption is duly completed by or on behalf of the patient—
 - (i) in cases involving a batch issue referred to in paragraph (2), on the batch issue at the time that the drug or appliance is supplied,
 - (ii) in cases involving an electronic repeatable prescription referred to in paragraph (2) on an approved form provided by the Board for recording patient declarations in respect of electronic repeatable prescriptions and issued by a chemist,
 - (iii) in cases involving an electronic prescription form, on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a chemist, or

- (iv) in all other cases, on the non-electronic prescription form;
 - (b) there is entitlement to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽³²⁾ (entitlement to full remission and payment) and a declaration of entitlement to remission is duly completed by or on behalf of the patient either—
 - (i) in cases involving a batch issue referred to in paragraph (2), on the batch issue at the time that the drug or appliance is supplied,
 - (ii) in cases involving an electronic repeatable prescription referred to in paragraph (2), on an approved form provided by the Board for recording patient declarations in respect of electronic repeatable prescriptions and issued by a chemist,
 - (iii) in cases involving an electronic prescription form, on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a chemist, or
 - (iv) in all other cases, on the non-electronic prescription form; or
 - (c) the patient is resident in a school or institution, the name of which is inserted on the non-electronic prescription form by a prescriber under the term of an arrangement to provide primary medical services under section 83(2) of the 2006 Act⁽³³⁾ (primary medical services).
- (6) No charge is to be made and recovered under paragraph (1) or (2) where there is an exemption by virtue of section 173(1)(d) of the 2006 Act⁽³⁴⁾ (exemptions from general charging) or regulation 10(2), 11, 12, 13 or 14.
- (7) For the purposes of this regulation, where a drug ordered on a single prescription form is supplied by instalments, the charge of £8.20 payable for that drug is payable on the supply of the first instalment.
- (8) A chemist is under no obligation to supply drugs or appliances in the course of providing pharmaceutical services or local pharmaceutical services where a charge is required to be made and recovered under paragraphs (1) or (2) unless the patient first pays that charge (notwithstanding any provisions in the chemist's terms of service).
- (9) Where a patient requests a receipt for a charge made and recovered under paragraph (1) or (2), the chemist must give the patient a receipt for the amount received on the relevant approved form.
- (10) Any sum which would otherwise be payable by the Board to a chemist in respect of the provision by that chemist of pharmaceutical services or local pharmaceutical services is to be reduced by the amount of any charges which must be made and recovered under paragraph (1) or (2).
- (11) In paragraph (8), "terms of service" means the terms on which pharmaceutical services or local pharmaceutical services are provided under the 2006 Act.

Supply of drugs and appliances by doctors

- 4.—**(1) A doctor who provides dispensing services to a patient must, subject to paragraphs (3), (4) and (8), make and recover from that patient for the supply of—
- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
 - (b) each other appliance, a charge of £8.20;
 - (c) each quantity of a drug, a charge of £8.20.

⁽³²⁾ Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

⁽³³⁾ Subsection (2) was substituted by the Health and Social Care Act 2012 (c. 7), Schedule 4, paragraph 30.

⁽³⁴⁾ Subsection (1) has been amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

(2) Where a charge is paid under paragraph (1), the person paying the charge must on doing so either—

- (a) where a non-electronic prescription form has been issued, sign a declaration in writing on the non-electronic prescription form that the relevant charge has been paid; or
- (b) where an electronic prescription form has been created, provide a declaration that the relevant charge has been paid on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a doctor.

(3) No charge is to be made and recovered under paragraph (1) where—

- (a) there is an exemption by virtue of regulation 10(1) and, subject to regulation 10(5), a declaration of entitlement to an exemption is duly completed by or on behalf of the patient either—
 - (i) on the non-electronic prescription form, or
 - (ii) in cases involving an electronic prescription form, on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a doctor;
- (b) there is entitlement to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽³⁵⁾ (entitlement to full remission and payment) and a declaration of entitlement to remission is duly completed by or on behalf of the patient either—
 - (i) on the non-electronic prescription form, or
 - (ii) in cases involving an electronic prescription form, on an approved form provided by the Board for recording patient declarations in respect of electronic prescription forms and issued by a doctor;
- (c) the drugs or appliances are supplied in respect of two or more persons in a school or institution in which 20 persons are normally resident, of whom at least 10 are that doctor's patients; or
- (d) there is an exemption by virtue of section 173(1)(d) of the 2006 Act⁽³⁶⁾ (exemptions from general charging) or regulation 10(2), 11(1), 12, or 13.

(4) For the purposes of this regulation, where a drug ordered on a single prescription form is supplied by instalments, the charge of £8.20 payable for that drug is payable on the supply of the first instalment.

(5) A doctor is under no obligation to supply drugs or appliances in the course of providing dispensing services where a charge is required to be made and recovered under paragraph (1) unless the patient first pays that charge (notwithstanding any provisions in the relevant GMS contract, PMS agreement or APMS contract).

(6) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the doctor must give the patient a receipt for the amount received on the relevant approved form.

(7) Any sum which would otherwise be payable by the Board to a contractor under a GMS contract, a PMS agreement or an APMS contract that includes terms of service by virtue of regulation 47 of the Pharmaceutical Services and Local Pharmaceutical Services Regulations (terms of service of dispensing doctors: general) is to be reduced by the amount of any charges which must be made and recovered under paragraph (1).

(8) Nothing in this regulation authorises the payment of a charge where the drug or appliance supplied is either—

⁽³⁵⁾ Relevant amendments were made to regulation 5 by [S.I. 2004/663](#) and [936](#), [2006/562](#), [2008/1697](#), [2009/411](#), [2013/475](#) and [2014/2667](#).

⁽³⁶⁾ Subsection (1) was amended by the Health Act 2009 ([c. 21](#)), Schedule 1, paragraphs 6 and 7(c).

- (a) needed for immediate treatment and no order for the drug or appliance is made on a prescription form; or
- (b) administered or applied to the patient by the doctor personally.

Out of hours supply of drugs and appliances by providers of out of hours services

5.—(1) A provider of out of hours services which are not dispensing services who supplies drugs or appliances to a patient in the course of providing out of hours services must, subject to paragraphs (3), (4) and (8), make and recover from that patient for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) each other appliance, a charge of £8.20;
- (c) each quantity of a drug, a charge of £8.20.

(2) Any person paying a charge under paragraph (1) must on doing so sign a declaration in writing on the approved form that the relevant charge has been paid.

(3) No charge is to be made and recovered under paragraph (1) where—

- (a) there is an exemption by virtue of regulation 10(1) and a declaration of entitlement to an exemption is duly completed by or on behalf of the patient on the approved form;
- (b) there is entitlement to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽³⁷⁾ (entitlement to full remission and payment) and a declaration of entitlement to remission is duly completed by or on behalf of the patient on the approved form;
- (c) the drugs or appliances are supplied in respect of two or more persons in a school or institution in which at least 20 persons are normally resident provided that the name of the school or institution is inserted on the approved form; and
- (d) in the cases described in sub-paragraphs (a) and (b), such evidence of entitlement to an exemption or remission is provided as the provider of out of hours services may reasonably require.

(4) No charge is to be made and recovered under paragraph (1) where there is an exemption by virtue of section 173(1)(d) of the 2006 Act⁽³⁸⁾ (exemptions from general charging) or regulation 10(2), 11(1), 12 or 13.

(5) A provider of out of hours services is under no obligation to supply drugs or appliances in the course of providing out of hours services where a charge is required to be made and recovered under paragraph (1) unless the patient first pays that charge (notwithstanding any provisions in the relevant agreement to provide out of hours services).

(6) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the provider of out of hours services must give the patient a receipt for the amount received on the relevant approved form.

(7) Any sum which would otherwise be payable by the Board to a provider of out of hours services in respect of the provision of primary medical services is to be reduced by the amount of any charges which must be made and recovered under paragraph (1).

(8) Nothing in this regulation authorises the payment of a charge where the drug or appliance supplied is either—

- (a) needed for immediate treatment and no order for the drug or appliance is made on an approved form; or

⁽³⁷⁾ Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

⁽³⁸⁾ Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

- (b) administered or applied to the patient by the provider of out of hours services personally.

Supply of drugs and appliances by NHS trusts and NHS foundation trusts

6.—(1) Where an NHS trust or an NHS foundation trust supplies a drug or appliance to a patient for the purpose of treatment, the NHS trust or the NHS foundation trust (as the case may be) must, subject to paragraphs (3) to (6), make and recover from the patient for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) an item specified in column (1) of Schedule 1 (which has effect), the charge specified in relation to it in column (2) of that Schedule;
- (c) tights, a charge of £16.40;
- (d) each other appliance, a charge of £8.20;
- (e) each quantity of a drug, a charge of £8.20.

(2) Any person paying a charge under paragraph (1) must on doing so sign a declaration in writing that the relevant charge has been paid.

(3) No charge is to be made and recovered under this regulation for the supply of drugs administered or appliances fitted or put into service at the hospital or other establishment or facility which forms part of the NHS trust or NHS foundation trust which supplies the drugs or appliances.

(4) No charge is to be made and recovered under this regulation from a patient who is exempt—

- (a) by virtue of section 173(1)(a), (c) or (d) of the 2006 Act⁽³⁹⁾ (exemptions from general charging) or regulation 10(2), 11(1), 12 or 13;
- (b) by virtue of regulation 10(3); or
- (c) by reason of being entitled to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽⁴⁰⁾ (entitlement to full remission and payment),

and who, in the case described in section 173(1)(c) of the 2006 Act (aged under 16 or aged under 19 and in full-time education) and in the cases described in sub-paragraph (b) and (c), completes a declaration of entitlement to such exemption or remission and provides any other evidence of entitlement as the NHS trust or NHS foundation trust may reasonably require.

(5) No charge is to be made and recovered under this regulation from a patient who is accepted by the person supplying the drug as—

- (a) suffering from tuberculosis, in respect of any drug supplied to that patient for the treatment of tuberculosis; or
- (b) being subject to a community treatment order, in respect of any drug supplied to that patient for the treatment of a mental disorder.

(6) A partially remitted charge must be made and recovered under this regulation from a patient who is entitled to partial remission by virtue of regulation 10(6) if a declaration in writing is provided that the relevant part of the charge has been paid and a declaration of entitlement, and any other evidence of entitlement to partial remission, as may be required, is provided.

(7) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the NHS trust or an NHS foundation trust must give the patient a receipt for the amount received on the relevant approved form.

⁽³⁹⁾ Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

⁽⁴⁰⁾ Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

Supply of drugs and appliances at walk-in centres

7.—(1) Where drugs or appliances are supplied to a patient, including during the out of hours period, for the purpose of treating that patient, by a prescriber at a walk-in centre, the NHS trust, NHS foundation trust or other person responsible for the management of the centre, must, subject to paragraphs (3) to (5), make and recover from that patient for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) each other appliance, a charge of £8.20;
- (c) each quantity of a drug, a charge of £8.20.

(2) Any person paying a charge under paragraph (1), must on doing so, sign a declaration in writing that the relevant charge has been paid.

(3) No charge is to be made and recovered under this regulation for the supply of drugs administered or appliances fitted or put into service at the walk-in centre.

(4) No charge is to be made and recovered under this regulation from a patient who is exempt—

- (a) by virtue of section 173(1)(d) of the 2006 Act⁽⁴¹⁾ (exemptions from general charging) or regulation 10(2), 11(1), 12 or 13;
- (b) by virtue of regulation 10(1); or
- (c) by reason of being entitled to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽⁴²⁾ (entitlement to full remission and payment),

and in the cases described in sub-paragraphs (b) and (c), completes a declaration of entitlement to such exemption or remission and provides any other evidence of entitlement as the NHS trust, NHS foundation trust or other person responsible for the management of the walk-in centre supplying the drug or appliance may reasonably require.

(5) For the purpose of this regulation, where a drug ordered is supplied by instalments, the charge of £8.20 payable for that drug is payable on the supply of the first instalment.

(6) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the person making and recovering the charge must give the patient a receipt for the amount received on the relevant approved form.

Supply of drugs under Patient Group Directions

8.—(1) Where, in the course of providing NHS services, drugs are supplied to a patient in accordance with a Patient Group Direction, including during the out of hours period, the person supplying the drugs must, subject to paragraphs (3) to (5), make and recover from the patient for the supply of each quantity of a drug, a charge of £8.20.

(2) Any person paying a charge under paragraph (1) must on doing so sign a declaration in writing that the relevant charge has been paid.

(3) No charge is to be made and recovered under this regulation from a patient who is exempt—

- (a) by virtue of section 173(1)(a) of the 2006 Act⁽⁴³⁾ (exemptions from general charging) or regulation 10(2), 11(1), 12 or 13;
- (b) by virtue of regulation 10(1); or

(41) Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

(42) Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

(43) Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

- (c) by reason of being entitled to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations(44) (entitlement to full remission and payment),

and who, in the cases described in sub-paragraphs (b) and (c), completes a declaration of entitlement to such exemption or remission and provides any other evidence of entitlement as the person supplying the drug may reasonably require.

(4) No charge is to be made and recovered under this regulation from a patient who is accepted by the person supplying the drug as—

- (a) suffering from tuberculosis, in respect of any drug supplied to that patient for the treatment of tuberculosis; or
- (b) being subject to a community treatment order, in respect of any drug supplied to that patient for the treatment of a mental disorder.

(5) No charge is to be made and recovered under this regulation where the drug is supplied for personal administration by the person making the supply in accordance with the Patient Group Direction.

(6) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the person making that charge must give the patient a receipt for the amount received on the relevant approved form.

(7) For the purpose of this regulation, the reference to the supply of a drug in accordance with a Patient Group Direction is a reference to the supply of a drug for that purpose as provided for in the Human Medicines Regulations 2012(45).

Supply of drugs and appliances by other providers of NHS services

9.—(1) A provider of NHS services who, otherwise than as mentioned in regulations 3 to 8, supplies drugs or appliances to a patient for the purpose of treatment must, subject to paragraphs (3) to (8), make and recover from the patient for the supply of—

- (a) an item of elastic hosiery, a charge of £8.20 or £16.40 per pair;
- (b) tights, a charge of £16.40;
- (c) an item specified in column (1) of Schedule 1, the charge specified in relation to it in column (2) of that Schedule;
- (d) each other appliance, a charge of £8.20;
- (e) each quantity of a drug, a charge of £8.20.

(2) Any person paying a charge under paragraph (1) must on doing so sign a declaration in writing that the relevant charge has been paid.

(3) No charge is to be made and recovered under this regulation from a patient who is exempt—

- (a) by virtue of section 173(1)(a), (c) or (d) of the 2006 Act(46) (exemptions from general charging) or regulation 10(2), 11(1), 12 or 13;
- (b) by virtue of regulation 10(3); or

(44) Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

(45) S.I. 2012/1916. See regulations 229 to 235 of those Regulations, which provide for the exemptions relating to Patient Group Directions from the general restrictions on the sale and supply of prescription only and pharmacy medicines.

(46) Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

- (c) by reason of being entitled to remission of the charge by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽⁴⁷⁾ (entitlement to full remission and payment),

and who, in the cases described in section 173(1)(c) of the 2006 Act and in the cases described in subparagraphs (b) and (c), completes a declaration of entitlement to such exemption or remission and provides any other evidence of entitlement as the provider of NHS services may reasonably require.

(4) No charge is to be made and recovered under this regulation from a patient where the drugs supplied to the patient are administered, or the appliances supplied to the patient are fitted or put into service, at the hospital or other establishment or facility from which the provider of NHS services provides such services.

(5) No charge is to be made and recovered under this regulation where the drug or appliance supplied is administered or applied to the patient personally by a person employed by, or contracted to provide services for, a provider of NHS services.

(6) No charge is to be made and recovered under this regulation from a patient who is accepted by the person supplying the drug as—

- (a) suffering from tuberculosis, in respect of any drug supplied to that patient for the treatment of tuberculosis; or
- (b) being subject to a community treatment order, in respect of any drug supplied to that patient for the treatment of a mental disorder.

(7) A partially remitted charge must be made and recovered under this regulation from a patient who is entitled to partial remission by virtue of regulation 10(6) if a declaration in writing is provided that the relevant part of the charge has been paid and a declaration of entitlement, and any other evidence of entitlement to partial remission, as may be required, is provided.

(8) For the purpose of this regulation, where a drug ordered on a single written direction is supplied by instalments, the charge of £8.20 is payable for that drug on the supply of the first instalment.

(9) Where a patient requests a receipt for a charge made and recovered under paragraph (1), the provider of NHS services must give the patient a receipt for the amount received on the relevant approved form.

Exemptions: general

10.—(1) Subject to paragraph (4), no charge is payable under regulations 3, 4, 5, 7 or 8 by a person who—

- (a) is under 16 years of age;
- (b) is under 19 years of age and is receiving qualifying full-time education within the meaning of section 173(2) and (3) of the 2006 Act (exemptions from general charging);
- (c) is 60 years of age or older;
- (d) has a valid exemption certificate on the grounds that they—
 - (i) are expecting a child, or
 - (ii) have within the last twelve months given birth to a child (including a child whose death was registrable under the special provisions as to the registration of still-births in the Births and Deaths Registration Act 1953⁽⁴⁸⁾),

⁽⁴⁷⁾ Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

⁽⁴⁸⁾ 1953 c. 20. See in particular section 11 of that Act.

- or has a valid exemption certificate issued under equivalent arrangements (to regulation 15) having effect in Scotland, Wales or Northern Ireland;
- (e) has a valid exemption certificate on the grounds that they are suffering from one or more of the following conditions—
- (i) permanent fistula (including caecostomy, colostomy, laryngostomy or ileostomy) requiring continuous surgical dressing or an appliance,
 - (ii) the following disorders—
 - forms of hypoadrenalism (including Addison’s disease) for which specific substitution therapy is essential,
 - diabetes insipidus and other forms of hypopituitarism,
 - diabetes mellitus – except where treatment is by diet alone,
 - hypoparathyroidism,
 - myasthenia gravis, or
 - myxoedema,
 - (iii) epilepsy requiring continuous anti-convulsive therapy, or
 - (iv) a continuing physical disability which prevents the person from leaving the person’s residence without the help of another person;
- (f) has a valid exemption certificate on the grounds that the person is undergoing treatment for—
- (i) cancer,
 - (ii) the effects of cancer, or
 - (iii) the effects of cancer treatment;
- (g) has a valid exemption certificate issued on any of the grounds in sub-paragraph (e) or (f) under equivalent arrangements which have effect in Scotland, Wales or Northern Ireland;
- (h) has a valid exemption certificate in respect of the supply of drugs and appliances for the treatment of accepted disablement, but in either case only in respect of those supplies to which the certificate relates; or
- (i) has a valid pre-payment certificate granted under regulation 16 or a valid pre-payment certificate granted under equivalent arrangements (to regulation 16) having effect in Scotland, Wales or Northern Ireland.
- (2) No charge is payable under these Regulations in respect of the supply of any drug for the treatment of a sexually transmitted disease within the meaning of section 173(1)(b) of the 2006 Act⁽⁴⁹⁾.
- (3) Subject to paragraph (4), no charge is payable under regulation 6 or 9—
- (a) by a person of a description specified in paragraph (1)(h) in respect of the supply of an item specified in column (1) of Schedule 1; or
 - (b) by a person of any description specified in paragraph (1) in respect of the supply of an appliance not specified in column (1) of Schedule 1, or of tights or of drugs.
- (4) Subject to paragraph (5), a person who wishes to claim entitlement to exemption by virtue of paragraph (1) or (3) must provide any declaration of entitlement required under regulation 3(5) or 4(3), or any declaration and evidence of entitlement required under regulation 5(3), 6(4), 7(4), 8(3) or 9(3).

⁽⁴⁹⁾ Subsection (1) was amended by the Health Act 2009 (c. 21), Schedule 1, paragraphs 6 and 7(c).

(5) A person of a description specified in paragraph (1)(a) or (c) is not required to provide any declaration of entitlement required under regulations 3(5) and 4(3) where—

- (a) an electronic prescription form or an electronic repeatable prescription is created and the person's date of birth is recorded on the person's PDS patient details and is set out on the electronic prescription form or electronic repeatable prescription; or
- (b) a non-electronic prescription form or a non-electronic repeatable prescription is issued and the person's date of birth is printed by means of a computer on the non-electronic prescription form or non-electronic repeatable prescription.

(6) A charge referred to in column (2) of Schedule (1) must, in the case of a person referred to in regulation 6 of the Travel Expenses and Remission of Charges Regulations⁽⁵⁰⁾ (entitlement to partial remission and payment), be remitted to the extent specified in that regulation.

(7) Where a charge is remitted in part under paragraph (6), the person making the part payment must, on doing so, sign a declaration in writing that the relevant part of the charge has been paid, complete a declaration of entitlement and provide such other evidence of entitlement to partial remission as may be required by the person supplying the drug or appliance.

(8) An exemption by reference to age or the validity of an exemption certificate must be determined by reference to the age or validity on the day on which—

- (a) in the case of pharmaceutical services or local pharmaceutical services provided by a chemist, the order for drugs or appliances is presented for dispensing; and
- (b) in any other case, the drugs or appliances are supplied.

(9) Where a claim to an exemption has been made but is not substantiated, and in consequence of the claim a charge has not been recovered, if—

- (a) the drugs or appliances were supplied by a chemist as mentioned in regulation 3 or by a doctor as mentioned in regulation 4, then the Board must recover that charge from the person concerned;
- (b) the drugs or appliances were supplied by an NHS trust or an NHS foundation trust as mentioned in regulation 6, then that NHS trust or NHS foundation trust must recover that charge from the person concerned; or
- (c) the drugs or appliances were supplied as mentioned in regulations 5, 7, 8 or 9, then the body which made the arrangements for the provision of NHS services with the person or body that supplied the drugs or appliances must recover that charge from the person concerned.

Exemption from charges for prisoners and persons detained in other secure accommodation

11.—(1) A prisoner or person detained in other secure accommodation is not liable to pay any charge under these Regulations.

(2) No charge is payable under regulation 3(1), 3(2), or 4 by a person where the charge is for drugs or appliances ordered on a non-electronic prescription form, and—

- (a) that person was a prisoner or a person detained in other secure accommodation at the time that the non-electronic prescription form was issued; and
- (b) that non-electronic prescription form has printed upon it—
 - (i) the letters "HMP", and
 - (ii) the name and address of the prison or other secure accommodation in which the person was detained at the time the non-electronic prescription form was issued.

(3) In this regulation—

⁽⁵⁰⁾ Regulation 6 has been amended by [S.I. 2006/562](#) and [675](#).

“other secure accommodation” means—

- (a) a court in which criminal proceedings against a person are heard;
- (b) secure training centre accommodation in which offenders in respect of whom detention and training orders have been made under section 100 of the Powers of Criminal Courts (Sentencing) Act 2000(51) (offenders under 18: detentions and training orders) may be detained and given training and education and prepared for their release and in which children who have been remanded to youth detention accommodation under section 91(4) of the Legal Aid, Sentencing and Punishment of Offenders Act 2012(52) (remands of children otherwise than on bail) may be detained; or
- (c) a secure children’s home in which accommodation is provided within the meaning of the Care Standards Act 2000(53), which provides accommodation for the purposes of restricting liberty and in respect of which a person is registered under Part 2 of that Act (establishments and agencies);

“person detained in other secure accommodation” means a person who is detained in other secure accommodation in which medical, dental, ophthalmic, pharmaceutical or nursing services are provided under the 2006 Act under arrangements made by the Board pursuant to Parts 4 to 7 of the 2006 Act or regulation 10 of the Standing Rules Regulations(54) (services for prisoners and other detainees);

“prison” includes a young offenders institution but not a naval, military or air force prison; and

“prisoner” means a person who is detained in a prison in which medical, dental, ophthalmic, pharmaceutical or nursing services are provided under the 2006 Act under arrangements made by the Board pursuant to Parts 4 to 7 of the 2006 Act or regulation 10 of the Standing Rules Regulations.

Exemption from charges for detainees

12.—(1) A detainee is not liable to pay any charge under these Regulations.

(2) In this regulation—

“detainee” means a person who is detained—

- (a) under the Immigration Act 1971(55) or section 62 of the Nationality, Immigration and Asylum Act 2002(56)(detention by the Secretary of State); and
- (b) in a removal centre in which medical, dental, ophthalmic, pharmaceutical or nursing services are provided under the 2006 Act under arrangements made by the Board pursuant to Parts 4 to 7 of the 2006 Act or regulation 10 of the Standing Rules Regulations(57) (services for prisoners and other detainees); and

“removal centre” has the same meaning as in section 147 of the Immigration and Asylum Act 1999(58) (interpretation of Part 8).

(51) 2000 c. 6. Section 100 has been amended by: the Criminal Justice and Court Services Act 2000 (c. 6), Schedule 7, paragraphs 160 and 184; the Criminal Justice Act 2003 (c. 44), Schedule 32, paragraphs 90 and 111, and Schedule 37, Part 7; and the Legal Aid, Sentencing and Punishment of Offenders Act 2012 (c. 10), Schedule 21, paragraphs 7 and 13, and Schedule 26, paragraphs 9 and 11.

(52) 2012 c. 10.

(53) 2000 c. 14.

(54) S.I. 2012/2996. Regulation 10 has been amended by S.I. 2013/261 and 2014/452.

(55) 1971 c. 77.

(56) 2002 c. 41. Section 62 has been amended by the Prevention of Terrorism Act 2005 (c. 2), section 16(2)(c), and the Immigration Act 2014 (c. 22), Schedule 9, paragraphs 3 and 13.

(57) S.I. 2012/2996. Regulation 10 has been amended by S.I. 2013/261 and 2014/452.

(58) 1999 c. 33. Section 147 has been amended by section 66 of the Nationality, Immigration and Asylum Act 2002 (c. 41).

Exemption from charges: risks to public health

13.—(1) In the circumstances set out in paragraph (2), the following drugs are exempt from any charge under these Regulations—

- (a) drugs for the prevention or treatment of a condition or disease arising from an emergency that threatens, is causing or has caused serious damage or risk to public health in England or any part of England; and
- (b) drugs, the supply of which is for, or is in anticipation of, a pandemic disease where the supply is in accordance with protocols relating to that disease as provided for in regulation 247 of the Human Medicines Regulations 2012⁽⁵⁹⁾ (exemption for supply in the event or anticipation of pandemic disease).

(2) Those circumstances are where—

- (a) the Secretary of State has made arrangements for supplying the drug to patients free of charge; or
- (b) the NHS body⁽⁶⁰⁾ that is responsible for the arrangements under which the drug is supplied has made arrangements, with the approval of the Secretary of State, for supplying the drug to patients free of charge,

and the patient is supplied with the drug under those arrangements.

Exemption for prescriptions provided or issued in Northern Ireland

14. No charge is payable under regulation 3 in respect of any pharmaceutical services or local pharmaceutical services which are obtained on the presentation of a prescription form which was provided or issued in Northern Ireland.

Certificates of exemption: application and issue

15.—(1) A person who wishes to claim exemption from charges payable under these Regulations by virtue of regulation 10(1)(d), (e), (f), or (h) must apply for a certificate conferring exemption (in these Regulations referred to as an “exemption certificate”) as follows—

- (a) a claim for an exemption by virtue of regulation 10(1)(d) must be made on an approved form provided by the Secretary of State;
- (b) a claim for an exemption by virtue of regulation 10(1)(e) or (f) must be made on the approved form, “FP92A—1 January 2009”, provided by the Secretary of State; and
- (c) a claim for an exemption by virtue of regulation 10(1)(h) must be made to an office of the Ministry of Defence on a form provided by the Secretary of State.

(2) The Secretary of State, on being satisfied that an applicant is entitled to exemption by virtue of regulation 10(1)(d), must issue an exemption certificate which is to be valid from the date which is one month before the date on which the Secretary of State received the application made under paragraph (1) until, in the case of an expectant mother, the end of that mother’s pregnancy, and in the case of a mother who gives birth—

- (a) to a child whose death is registrable under the special provisions as to the registration of still-births in the Births and Deaths Registration Act 1953⁽⁶¹⁾, thereafter until the end of the period of twelve months beginning with the expected date of confinement; or
- (b) to a live child (who is not registrable as still-born), thereafter until the end of the period of twelve months beginning with the date of birth of that child.

⁽⁵⁹⁾ S.I. 2012/1916. Regulation 247 has been amended by S.I. 2013/235.

⁽⁶⁰⁾ “NHS body” is defined in section 275(1) of the National Health Service Act 2006 (c. 41).

⁽⁶¹⁾ 1953 c. 20. See in particular section 11 of that Act.

(3) The Secretary of State, on being satisfied that an applicant, not being a person entitled to exemption by virtue of regulation 10(1)(a), (b) or (c), is entitled to exemption by virtue of regulation 10(1)(e), must issue to the applicant an exemption certificate which is to be valid—

- (a) from the date one month prior to the date on which the Secretary of State received the application made under paragraph (1); and
- (b) for such period as the Secretary of State may determine.

(4) The Secretary of State, on being satisfied that an applicant, not being a person entitled to exemption by virtue of regulation 10(1)(a), (b), (c), or (e) is entitled to exemption by virtue of regulation 10(1)(f), must issue to the applicant an exemption certificate which is to be valid—

- (a) from the date one month prior to the date on which the Secretary of State received the application under paragraph (1); and
- (b) for such period as the Secretary of State may determine.

(5) The Secretary of State, on being satisfied that an applicant is entitled to exemption by virtue of regulation 10(1)(h), must issue to the applicant an exemption certificate which is to be valid for such period as the Secretary of State may determine.

Pre-payment certificates: application and grant

16.—(1) A person applying to the Secretary of State for a pre-payment certificate (referred to in this regulation as “the applicant”) must—

- (a) pay, or undertake to pay by means of a direct debit in ten monthly instalments, to the Secretary of State, the sum payable for the pre-payment certificate as set out in paragraph (3); and
- (b) provide the Secretary of State with the information required to determine that application by—
 - (i) duly completing and submitting an application for the pre-payment certificate on an approved form provided for the purpose, or
 - (ii) providing the Secretary of State with that information in a manner that is acceptable to the Secretary of State.

(2) A pre-payment certificate is valid for a period of either 3 months or 12 months and an application for such a certificate must indicate the period for which it is required to be valid.

(3) The applicant must pay to the Secretary of State—

- (a) in the case of a 3 month pre-payment certificate, £29.10;
- (b) in the case of a 12 month pre-payment certificate payable by means of a single instalment, £104.00; or
- (c) in the case of a 12 month pre-payment certificate payable by means of a direct debit in ten monthly instalments, instalments amounting to £10.40,

and the payment amount referred to for each type of pre-payment certificate is referred to in this regulation and regulation 17 as “the issuing amount”.

(4) Subject to paragraph (6), the Secretary of State must grant a pre-payment certificate to the applicant as soon as reasonably practicable after the applicant has fulfilled the requirements set out in paragraph (1).

(5) The Secretary of State may refuse to accept payment for a pre-payment certificate valid for 12 months by means of direct debit in ten monthly instalments, and refuse to grant such a certificate to an applicant, where the applicant has—

- (a) previously been granted a pre-payment certificate valid for 12 months payable by direct debit in ten monthly instalments;

- (b) failed to pay in full the amount payable for that previously granted prepayment certificate without good reason; and
 - (c) failed to return without good reason that previously granted pre-payment certificate to the Secretary of State within 28 days of the date on which it was requested to be returned.
- (6) A pre-payment certificate is valid—
- (a) from a date before the application is made, where that date—
 - (i) was specified by the applicant when making the application, and
 - (ii) is no more than one month prior to the date when the application was made under paragraph (1); or
 - (b) from a date after the application is made, where that date—
 - (i) was specified by the applicant when making the application, and
 - (ii) is not more than one month after the date when the application was made under paragraph (1); or
 - (c) if the applicant does not specify a date in accordance with sub-paragraphs (a)(i) or (b)(i), from the date when the application is received.
- (7) Pre-payment certificates granted under this regulation are not valid in respect of the items specified in column (1) of Schedule 1.

Pre-payment certificates: repayment

17.—(1) Where the issuing amount in respect of a pre-payment certificate has been paid and, not more than one month after the date when the pre-payment certificate became valid, the relevant person—

- (a) dies; or
- (b) becomes resident in a hospital and thereafter dies while resident in hospital before the expiry of the pre-payment certificate,

an application for repayment of the entirety of the issuing amount which has already been paid may be made on behalf of that person's estate in accordance with paragraph (6) and (7).

(2) Where the issuing amount in respect of a pre-payment certificate valid for 12 months has been paid and during the period beginning on the date when the pre-payment certificate became valid and ending on the date of its expiry, the relevant person becomes—

- (a) entitled to exemption by virtue of regulation 10(1)(b) to (h);
- (b) entitled to remission by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations⁽⁶²⁾ (entitlement to full remission and payment); or
- (c) resident in hospital and thereafter remains in hospital until the expiry of the pre-payment certificate,

an application for a refund may be made, by or on behalf of that person or that person's estate, in accordance with paragraph (6) and (7), and is to be calculated in accordance with paragraph (4), in respect of each complete month following the date on which one of the events listed in sub-paragraph (a) to (c) occurred.

(3) Where the issuing amount in respect of a pre-payment certificate has been paid and during the period beginning one month after the date when the pre-payment certificate became valid and ending with the date of its expiry, the relevant person—

⁽⁶²⁾ Relevant amendments were made to regulation 5 by S.I. 2004/663 and 936, 2006/562, 2008/1697, 2009/411, 2013/475 and 2014/2667.

- (a) dies; or
- (b) becomes resident in hospital and thereafter dies while resident in hospital before the expiry of the pre-payment certificate,

an application for a refund may be made on behalf of that person's estate in accordance with paragraph (6) and (7), and is to be calculated in accordance with paragraph (4), in respect of each complete month following the date when the person died or became resident in hospital.

- (4) The refund referred to in paragraph (2) and (3) is to be calculated as follows—
 - (a) in the case of a pre-payment certificate valid for 3 months, one third of £29.10 for each complete month during which the pre-payment certificate is or was valid following the relevant date;
 - (b) in the case of a pre-payment certificate valid for 12 months where a person has paid for the pre-payment certificate by means of a single payment, one twelfth of £104.00 for each complete month during which the pre-payment certificate is or was valid following the relevant date; or
 - (c) in the case of a pre-payment certificate valid for 12 months where a person is or was paying for the pre-payment certificate in ten monthly instalments by means of direct debit either—
 - (i) where the sum payable for the pre-payment certificate has been paid in full, one twelfth of £104.00 for each complete month during which the pre-payment certificate is or was valid following the relevant date, or
 - (ii) where the sum payable for the pre-payment certificate has not been paid in full, the amount paid for the pre-payment certificate at the point of refund, minus £104.00, plus one twelfth of £104.00 for each complete month during which the pre-payment certificate is or was valid following the relevant date,

and for the purposes of these calculations, a “complete month” is a month beginning on the monthly anniversary of the date on which the pre-payment certificate became valid and ending on the date immediately preceding that date in the following month, “the relevant date” is the date on which one of the events listed in paragraph (2) or (3) occurred, and if a calculation produces a minus figure for the refund, no refund is payable.

(5) Where the issuing amount in respect of a pre-payment certificate valid for 3 months has been paid and not more than one month after the date on which the pre-payment certificate became valid, the relevant person becomes—

- (a) entitled to exemption by virtue of regulation 10(1)(b) to (h);
- (b) entitled to remission by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations; or
- (c) resident in a hospital and remains there until expiry of the pre-payment certificate,

an application for a refund of two thirds of £29.10 may be made by or on behalf of that person in accordance with paragraph (6) and (7).

(6) An application for a refund under this regulation is to be made to the Secretary of State and must be accompanied by the pre-payment certificate (where granted) and a declaration in support of the claim.

- (7) An application for a refund under this regulation is to be made—
 - (a) where the relevant person dies or becomes resident in hospital and thereafter dies, within 24 months of the date of that death;
 - (b) where the relevant person becomes—
 - (i) entitled to exemption by virtue of regulation 10(1)(b) to (h), or

- (ii) entitled to remission by virtue of regulation 5 of the Travel Expenses and Remission of Charges Regulations,
 - within 3 months of the date on which the event specified in paragraph (i) or (ii) occurred; or
 - (c) where the relevant person becomes resident in hospital and remains there until the expiry of the pre-payment certificate, within 3 months of the date of expiry.
- (8) Any repayment is to be made in such manner and subject to such conditions as the Secretary of State may determine.
- (9) Where the calculation of an amount under this regulation produces a sum which is not a whole number, the amount must be rounded up to the nearest penny.
- (10) In this regulation, “the relevant person” means the person in respect of whom an issuing amount was paid.

Repayment of charges

18.—(1) Where a charge has been paid under these Regulations by or on behalf of a person who was at the time of payment exempt from the requirement to pay that charge, an application for repayment may be made in accordance with paragraph (2) by or on behalf of that person.

- (2) An application for repayment must be—
- (a) made to the person or body specified in the receipt which is given under regulation 3(9), 4(6), 5(6), 6(7), 7(6), 8(6) or 9(9) as being the person or body to whom application for repayment of charges may be made;
 - (b) made in such form and manner as the Secretary of State may determine for the applicant, any class of applicant or applicants generally;
 - (c) made within 3 months from the date on which the drug or appliance was supplied to the applicant or within such longer period as the Secretary of State may allow; and
 - (d) accompanied by the receipt which is given pursuant to regulation 3(9), 4(6), 5(6), 6(7), 7(6), 8(6) or 9(9) for the charge paid and a declaration as to the grounds of exemption.
- (3) In the case of a charge under regulation 6 or 9 in respect of an item specified in column (1) of Schedule 1, the application must be accompanied by the exemption certificate referred to in regulation 10(1)(h) and, if the patient was referred by a doctor to the NHS trust, NHS foundation trust or other provider of NHS services for treatment—
- (a) a certificate from the doctor certifying that the treatment was for an accepted disablement; or
 - (b) a statement that such a certificate was surrendered to the NHS trust, NHS foundation trust or other provider of NHS services before the supply of the item.
- (4) The Secretary of State must make arrangements for the repayment of any charge paid under these Regulations by a person who is entitled to exemption.

Transitional provision

19. Where, on or after 1st April 2015, an item specified in column (1) of the table in Schedule 1 is supplied pursuant to an order made before that date, the National Health Service (Charges for Drugs and Appliances) Regulations 2000(**63**) are to have effect in relation to the supply of that appliance as if these Regulations had not come into force (and so those Regulations had not been revoked).

(63) *S.I. 2000/620.* The relevant amendments to this instrument are listed in Schedule 3 as a consequence of being revoked by these Regulations.

Consequential amendments

20. The amendments set out in Schedule 2 have effect.

Revocations

21. The Regulations and Orders specified in column (1) of Schedule 3 are hereby revoked in relation to England to the extent specified in column (3) of that Schedule.

Signed by authority of the Secretary of State for Health.

4th March 2015

Earl Howe
Parliamentary Under-Secretary of State,
Department of Health

Status: This is the original version (as it was originally made).

SCHEDULE 1

Regulation 6(1)

Charges for fabric supports and wigs

<i>(1) Specified item</i>	<i>(2) Specified charge</i>
Surgical brassiere	£27.45
Abdominal or spinal support	£41.50
Stock modacrylic wig	£67.75
Partial human hair wig	£179.45
Full bespoke human hair wig	£262.45

SCHEDULE 2

Regulation 20

Consequential amendments

Amendments to the National Health Service (Travel Expenses and Remission of Charges) Regulations 2003

1.—(1) The National Health Service (Travel Expenses and Remission of Charges) Regulations 2003(64) are amended as follows.

(2) In regulation 2 (interpretation), for the definition of “the Charges Regulations” substitute the following definition—

““the Charges Regulations” means the National Health Service (Charges for Drugs and Appliances) Regulations 2015;”.

(3) In regulation 5(2)(e) (entitlement to full remission and payment), for “regulation 3(1)(b)” substitute “regulation 3(1)(b) or (c)”.

(4) In regulation 12(1)(a)(ia) (repayments), for “regulation 6B” substitute “regulation 9”.

Amendment to the National Health Service (General Medical Services Contracts) Regulations 2004

2. In regulation 2(1) of the National Health Service (General Medical Services Contracts) Regulations 2004(65) (interpretation), for the definition of “listed medicines” substitute the following definition—

““listed medicines” means the drugs mentioned in regulation 13(1) of the National Health Service (Charges for Drugs and Appliances) Regulations 2015;”.

Amendment to the National Health Service (Personal Medical Services Agreements) Regulations 2004

3. In regulation 2(1) of the National Health Service (Personal Medical Services Agreements) Regulations 2004(66) (interpretation), for the definition of “listed medicines” substitute the following definition—

(64) S.I. 2003/2382. Relevant amendments have been made by S.I. 2004/663, 696 and 936, 2005/578, 2006/562, 2008/571 and 1697 and 2013/475.

(65) S.I. 2004/291. The original definition of “listed medicines” was inserted by S.I. 2009/2230.

(66) S.I. 2004/627. The original definition of “listed medicines” was inserted by S.I. 2009/2230.

““listed medicines” means the drugs mentioned in regulation 13(1) of the National Health Service (Charges for Drugs and Appliances) Regulations 2015;”.

Amendment to the Primary Ophthalmic Services Regulations 2008

4. In regulation 3(2)(c) of the Primary Ophthalmic Services Regulations 2008⁽⁶⁷⁾ (sight tests –eligibility), for “National Health Service (Charges for Drugs and Appliances) Regulations 2000” substitute “National Health Service (Charges for Drugs and Appliances) Regulations 2015”.

Amendments to the National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013

5.—(1) The National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013⁽⁶⁸⁾ are amended as follows.

(2) In regulation 2 (interpretation), for the definition of “Charges Regulations” substitute the following definition—

““Charges Regulations” means the National Health Service (Charges for Drugs and Appliances) Regulations 2015;”.

(3) In regulation 96(2)(a) (refunds of prescription charges), for “regulation 10(2)(b)” substitute “regulation 18(2)(b)”.

(4) In Schedule 4 (terms of service of NHS pharmacists), in paragraph 7 (preliminary matters before providing ordered drugs or appliances)—

(a) in sub-paragraph (3), for “or (1A)” substitute “or (2)”; and

(b) for “regulation 7(1)”, at each place where it occurs, substitute “regulation 10(1)”.

(5) In Schedule 5 (terms of service of NHS appliance contractors), in paragraph 6 (preliminary matters before providing appliances)—

(a) in sub-paragraph (3), for “or (1A)” substitute “or (2)”; and

(b) for “regulation 7(1)”, at each place where it occurs, substitute “regulation 10(1)”.

(6) In Schedule 6 (terms of service of dispensing doctors), in paragraph 4 (preliminary matters before providing ordered drugs or appliances)—

(a) in sub-paragraph (a), for “regulation 7(1)” substitute “regulation 10(1)”, and for “regulation 7” substitute “regulation 10”; and

(b) in sub-paragraph (c)(i)(aa), for “regulation 7(1)” substitute “regulation 10(1)”.

(7) In Schedule 7 (mandatory terms for LPS schemes)—

(a) in paragraph 5 (preliminary matters before providing ordered drugs or appliances)—

(i) in sub-paragraph (3), for “or (1A)” substitute “or (2)”, and

(ii) for “regulation 7(1)”, at each place where it occurs, substitute “regulation 10(1)”; and

(b) in paragraph 18(2)(a) (refunds of prescription charges), for “regulation 10(2)(b)” substitute “regulation 18(2)(b)”.

⁽⁶⁷⁾ S.I. 2008/1186. Relevant amendments have been made by S.I. 2008/2449, 2009/409, 2013/365 and 2014/418 and 2667.

⁽⁶⁸⁾ S.I. 2013/349; amended by S.I. 2014/417 and 2015/58.

Status: This is the original version (as it was originally made).

Amendment to the National Health Service (Optical Charges and Payments) Regulations 2013

6. In regulation 8(3)(b) of the National Health Service (Optical Charges and Payments) Regulations 2013(69) (eligibility for a voucher – supply of optical appliances), for “the National Health Service (Charges for Drugs and Appliances) Regulations 2000” substitute “the National Health Service (Charges for Drugs and Appliances) Regulations 2015”.

Amendment to the Social Security (Information-sharing) (NHS Payments and Remission of Charges etc.) (England) Regulations 2015

7. In regulation 2 of the Social Security (Information-sharing) (NHS Payments and Remission of Charges etc.) (England) Regulations 2015(70) (interpretation), for the definition of “the CDA Regulations” substitute the following definition—

““the CDA Regulations” means the National Health Service (Charges for Drugs and Appliances) Regulations 2015;”.

Amendment to the Delegation of Additional Functions to the NHS Business Services Authority (Awdurdod Gwasanaethau Busnes y GIG) Regulations 2015

8. In regulation 2 of the Delegation of Additional Functions to the NHS Business Services Authority (Awdurdod Gwasanaethau Busnes y GIG) Regulations 2015(71) (functions of the National Health Service Commissioning Board to be exercised by the NHS Business Services Authority), for “regulation 7(7)(a) of the National Health Service (Charges for Drugs and Appliances) Regulations 2000” substitute “regulation 10(9)(a) of the National Health Service (Charges for Drugs and Appliances) Regulations 2015”.

SCHEDULE 3

Regulation 21

Revocations

<i>(1) Instruments revoked</i>	<i>(2) Reference</i>	<i>(3) Extent of revocation</i>
National Health Service (Charges for Drugs and Appliances) Regulations 2000	S.I. 2000/620	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2000	S.I. 2000/2393	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment (No. 2) Regulations 2000	S.I. 2000/3189	The whole Regulations
National Health Service (Charges for Drugs	S.I. 2001/746	The whole Regulations

(69) [S.I. 2013/461](#). Regulation 8 has been amended by [S.I. 2014/2667](#).

(70) [S.I. 2015/124](#).

(71) [S.I. 2015/127](#).

<i>(1) Instruments revoked</i>	<i>(2) Reference</i>	<i>(3) Extent of revocation</i>
and Appliances) Amendment Regulations 2001		
National Health Service (Charges for Drugs and Appliances) (Electronic Communications) Order 2001	S.I. 2001/2887	The whole Order
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2002	S.I. 2002/548	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment (No. 3) Regulations 2002	S.I. 2002/2352	The whole Regulations
National Health Service (Amendments Relating to Prescribing by Nurses and Pharmacists etc.) (England) Regulations 2003	S.I. 2003/699	The whole Regulations
National Health Service (Pharmaceutical Services) (General Medical Services) and (Charges for Drugs and Appliances) Amendment Regulations 2003	S.I. 2003/1084	The whole Regulations
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2004	S.I. 2004/663	Regulations 2, 3 and 4
Health and Social Care (Community Health and Standards) Act 2003 (Supplementary and Consequential Provision) (NHS Foundation Trusts) Order 2004	S.I. 2004/696	In Schedules 2, 3 and 4, the references to the National Health Service (Charges for Drugs and Appliances) Regulations 2000
General Medical Services and Personal Medical Services Transitional and Consequential Provisions Order 2004	S.I. 2004/865	Schedule 1, paragraph 26
Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004	S.I. 2004/1771	Paragraph 34 of Part 2 of the Schedule
National Health Service (Charges for Drugs and	S.I. 2005/578	Regulations 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13 and 14

Status: This is the original version (as it was originally made).

<i>(1) Instruments revoked</i>	<i>(2) Reference</i>	<i>(3) Extent of revocation</i>
Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2005		
General Dental Services, Personal Dental Services and Abolition of the Dental Practice Board Transitional and Consequential Provisions Order 2006	S.I. 2006/562	Schedule 1, paragraph 9
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2006	S.I. 2006/675	Regulations 2, 3 and 4
National Health Service (Miscellaneous Amendments Relating to Independent Prescribing) Regulations 2006	S.I. 2006/913	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment (No. 2) Regulations 2007	S.I. 2007/1510	The whole Regulations
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2007	S.I. 2007/1975	Regulations 2 and 3
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2008	S.I. 2008/571	Regulations 3, 4, 5, 6, 7, and 8
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment (No. 2) Regulations 2008	S.I. 2008/1697	Regulation 2
Primary Ophthalmic Services Amendment, Transitional and Consequential Provisions Regulations 2008	S.I. 2008/1700	Schedule 1, paragraph 7

<i>(1) Instruments revoked</i>	<i>(2) Reference</i>	<i>(3) Extent of revocation</i>
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2008	S.I. 2008/2593	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2009	S.I. 2009/29	The whole Regulations
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2009	S.I. 2009/411	Regulations 2 and 3
National Health Service (Charges) (Amendments Relating to Pandemic Influenza) Regulations 2009	S.I. 2009/1166	The whole Regulations
National Health Service (Prescribing and Charging Amendments Relating to Pandemic Influenza) Regulations 2009	S.I. 2009/2230	Regulation 2
National Health Service (Charges for Drugs and Appliances) and (Travel Expenses and Remission of Charges) Amendment Regulations 2010	S.I. 2010/1727	Regulation 2
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2011	S.I. 2011/518	The whole Regulations
National Health Service (Charges for Drugs and Appliances) Amendment Regulations 2012	S.I. 2012/470	The whole Regulations
Health and Social Care Act 2012 (Consequential Provision-Social Workers) Order 2012	S.I. 2012/1479	Paragraph 18 of the Schedule
Human Medicines Regulations 2012	S.I. 2012/1916	Schedule 34, paragraph 73(a)
National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013	S.I. 2013/349	Schedule 10, paragraph 1

Status: This is the original version (as it was originally made).

<i>(1) Instruments revoked</i>	<i>(2) Reference</i>	<i>(3) Extent of revocation</i>
National Health Service (Charges for Drugs and Appliances) (Dental Charges) and (Travel Expenses and Remission of Charges) (Amendment) Regulations 2013	S.I. 2013/475	Regulations 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 and 15
National Health Service (Charges, Payments and Remission of Charges) (Uprating, Miscellaneous Amendments and Transitional Provision) Regulations 2014	S.I. 2014/545	Regulations 2 and 3

EXPLANATORY NOTE

(This note is not part of the Regulations)

These Regulations consolidate with amendments the National Health Service (Charges for Drugs and Appliances) Regulations 2000. Those Regulations provided for the making and recovery of charges for drugs and appliances supplied under the National Health Service Act 2006.

These Regulations also increase the NHS charges that are levied for certain wigs and fabric supports by approximately 1.6% and the charges for drugs and other appliances by approximately 1.9%.

Regulations 3 and 4 provide that chemists and doctors who provide pharmaceutical services to a patient must make and recover from that patient the charges set out in those regulations.

Regulation 5 provides that providers of out of hours primary medical services who supply drugs and appliances to a patient in the course of providing those services must make and recover the charges set out in that regulation.

Regulation 6 provides that where drugs and appliances are supplied by NHS trusts and NHS foundation trusts, charges are to be made and recovered in the circumstances set out in that regulation.

Regulation 7 provides for charges to be made and recovered for the supply of drugs and appliances at walk-in-centres by a prescriber at a walk-in-centre.

Regulation 8 provides for charges to be made and recovered for the supply of drugs in accordance with a Patient Group Direction.

Regulation 9 provides for charges to be made and recovered from a patient for the supply of drugs or appliances by other providers of NHS services, not covered by regulations 3 to 8.

Regulations 10 to 14 set out the exemptions to any charges that may be made and recovered under the preceding regulations.

Regulation 15 sets out the mechanism for applying for an exemption certificate required in order to claim a medical or maternity related exemption, or an exemption related to service in the armed forces, by virtue of regulation 10.

Regulation 16 provides for patients to receive pre-payment certificates.

Regulation 17 provides for a system of granting repayments for amounts paid under regulation 16 for pre-payment certificates in the circumstances set out in regulation 17.

Regulation 18 provides for repayment of charges to persons who, at the time of payment, were exempt from the requirement to pay the charge.

Regulation 19 provides for a transitional provision in respect of the prescription charge to be paid for the appliances specified in Schedule 1.

Regulation 20 and Schedule 2 provide for consequential amendments to other secondary legislation to take account of this consolidation.

Regulation 21 and Schedule 3 provide for revocations of secondary legislation in relation to England to the extent specified in that Schedule.