
STATUTORY INSTRUMENTS

2013 No. 3134

**The Consumer Contracts (Information, Cancellation
and Additional Charges) Regulations 2013**

PART 2

Information requirements

CHAPTER 1

Provision of information

Application of Part 2

7.—(1) This Part applies to on-premises, off-premises and distance contracts, subject to paragraphs (2), (3) and (4) and regulation 6.

(2) This Part does not apply to contracts to the extent that they are—

- (a) for the supply of a medicinal product by administration by a prescriber, or under a prescription or directions given by a prescriber;
- (b) for the supply of a product by a health care professional or a person included in a relevant list, under arrangements for the supply of services as part of the health service, where the product is one that, at least in some circumstances is available under such arrangements free or on prescription.

(3) This Part, except for regulation 14(1) to (5), does not apply to contracts to the extent that they are for passenger transport services.

(4) This Part does not apply to off-premises contracts under which the payment to be made by the consumer is not more than £42.

(5) In paragraph (2)—

“health care professional” and “prescriber” have the meaning given by regulation 2(1) of the National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013⁽¹⁾;

“health service” means—

- (a) the health service as defined by section 275(1) of the National Health Service Act 2006⁽²⁾ or section 206(1) of the National Health Service (Wales) Act 2006⁽³⁾,
- (b) the health service as defined by section 108(1) of the National Health Service (Scotland) Act 1978⁽⁴⁾, or

(1) S.I. 2013/349.

(2) 2006 c.41.

(3) 2006 c.42.

(4) 1978 c.29.

Status: This is the original version (as it was originally made). This item of legislation is currently only available in its original format.

- (c) any of the health services under section 2(1)(a) of the Health and Social Care (Reform) Act (Northern Ireland) 2009⁽⁵⁾;
“medicinal product” has the meaning given by regulation 2(1) of the Human Medicines Regulations 2012⁽⁶⁾;
“relevant list” means—
- (d) a relevant list for the purposes of the National Health Service (Pharmaceutical and Local Pharmaceutical Services) Regulations 2013⁽⁷⁾, or
- (e) a list maintained under those Regulations.

⁽⁵⁾ 2006 c.41 (N.I): section 1 was substituted by section 1 of the Health and Social Care Act 2012 (c.7).
⁽⁶⁾ S.I. 2012/1916.
⁽⁷⁾ S.I. 2013/349.