

SCHEDULE

Article 2

Amendments of Schedule to Patents (Isle of Man) Order 2013

1. After paragraph 24 insert—

“**24A.** In section 92(5) (obtaining evidence for proceedings under the European Patent Convention), for “Perjury Act 1911(1)” substitute “Perjury Act 1952 (an Act of Tynwald(2))” and omit “and Article 3(4) of the Perjury (Northern Ireland) Order 1979(3)”.

2. Omit paragraph 30.

3. In paragraph 34, in the inserted section 128B(2)(a), for “[Council Regulation \(EEC\) No 1768/92](#) of 18th June 1992 concerning the creation of a supplementary protection certificate for medicinal products(4)” substitute “[Regulation \(EC\) No 469/2009](#) of the European Parliament and of the Council of 6th May 2009 concerning the supplementary protection certificate for medicinal products(5)”.

4. In paragraph 37, in the inserted Schedule 4A—

(a) in paragraph 7(a), for “[Council Regulation \(EEC\) No 1768/92](#) of 18th June 1992 concerning the creation of a supplementary protection certificate for medicinal products” substitute “[Regulation \(EC\) No 469/2009](#) of the European Parliament and of the Council of 6th May 2009 concerning the supplementary protection certificate for medicinal products”;

(b) after paragraph 7 insert—

“Transitional provision

8.—(1) A reference (express or implied) in this Act to the Medicinal Products Regulation, or a provision of it, is to be read as being or (subject to context) including a reference to the old Regulation, or the corresponding provision of the old Regulation, in relation to times, circumstances or purposes in relation to which the old Regulation, or that provision, had effect.

(2) Other than in relation to times, circumstances or purposes referred to in subparagraph (1), anything done, or having effect as if done, under (or for the purposes of or in reliance on) the old Regulation or a provision of the old Regulation and in force or effective immediately before 1st June 2016 (the day on which the Patents (Isle of Man) (Amendment) Order 2016 came into force) has effect on or after that date for the purposes of this Act as if done under (or for the purpose of or in reliance on) the Medicinal Products Regulation or the corresponding provision of it.

(3) In this paragraph “the old Regulation” means [Council Regulation \(EEC\) No 1768/92](#) of 18th June 1992 concerning the creation of a supplementary protection certificate for medicinal products.”.

(1) [1911 c.6](#) (1 and 2 Geo 5).
(2) [1952 c.8](#) (Isle of Man).
(3) [S.I. 1979/1714 \(N.I. 19\)](#).
(4) [OJ No L182, 2.7.1992, p.1](#).
(5) [OJ No L152, 16.6.2009, p.1](#).