SCHEDULE 7

STANDARD PROVISIONS FOR MANUFACTURING AUTHORISATIONS

PART 3

PROVISIONS WHICH MAY BE INCORPORATED IN AN AUTHORISATION RELATING TO THE IMPORTATION OF INVESTIGATIONAL MEDICINAL PRODUCTS

[F1 9A. The holder of the authorisation shall only import EAMS medicinal products if and to the extent that such products are required for the Early Access to Medicines Scheme.**]**

Textual Amendments

F1 Sch. 7 Pt. 3 para. 9A inserted (15.4.2022) by The Human Medicines (Amendments Relating to the Early Access to Medicines Scheme) Regulations 2022 (S.I. 2022/352), regs. 1(2), **18(3)(f)** (with reg. 19)

Status:
Point in time view as at 15/04/2022.

Changes to legislation:
There are currently no known outstanding effects for the The Medicines for Human Use (Clinical Trials) Regulations 2004, Paragraph 9A.