

*Status: EU Directives are published on this site to aid cross referencing from UK legislation. Since IP completion day (31 December 2020 11.00 p.m.) no amendments have been applied to this version.*

ANNEX

REGISTER FORMAT REFERRED TO IN ARTICLE 2(1)

<b>Registration number</b>	<b>Date of issue of EC type-examination certificate (Module B), certificate of conformity (Module G) or quality system approval (Module H) and date of expiry where applicable</b>	<b>Manufacturer</b>	<b>Type of product (generic) and subtype if applicable</b>	<b>Production phase conformity module<sup>a</sup></b>	<b>Notified body undertaking production phase conformity assessment<sup>†</sup></b>	<b>Additional information</b>

**a** Always has to be filled in if under responsibility of the notified body carrying out conformity assessment procedure referred to in Article 9(a) of Directive 2007/23/EC (Module B). Not required for conformity assessment procedures referred to in Article 9(b) and (c) (Modules G and H). Information shall be given (where known) if another notified body is involved.