ANNEX

REGISTER FORMAT REFERRED TO IN ARTICLE 2(1)

Registration number	**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**	**Manufacturer**	**Type of product (generic) and subtype if applicable**	**Production phase conformity module[a]**	**Notified body undertaking production phase conformity assessment[a]**	**Additional information**

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.