

Annex B

<u>Member State EU/EEA</u> ¹	<u>MRP/National² variation number</u>	<u>Marketing Authorisation number</u>	<u>Marketing Authorisation Holder</u> ³	<u>(Invented) name</u> ⁴	<u>Active substance(s)</u>	<u>Strength(s)</u> ⁵	<u>Pharmaceutical Form</u> ⁶	<u><Content (concentration)></u> ⁷
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¹ List all the EEA Countries where the medicinal product(s) included in the worksharing are authorised, in alphabetical order (i.e. all medicinal products authorised in Austria, followed by all medicinal products authorised in Belgium, etc.)

² If applicable

³ Name and address of the Marketing Authorisation Holder in the EEA Countries where the medicinal product is authorised

⁴ As registered in the respective official language of the EEA Country (no strength or pharmaceutical form should be mentioned unless it is an integral part of the authorised (invented) name)

⁵ It is possible to combine in the same row several strengths for the medicinal product in each Country. A separate row should however be used for each pharmaceutical form.

⁶ Information in English - use current standard terms from the Ph. Eur.

⁷ Complete only if applicable (only for liquids, creams and solid multidose forms, e.g. granules, powder, etc – not for tablets and capsules)