



04 June 2026
EMA/102302/2005 v. 7.1
Human Medicines Division

Member states contact points for translations review

The Eudralink message should include at least the following information:

Eudralink subject: Post-opinion review – Day <...> – <Product name> – <EMEA procedure number>
– Deadline for comments: <...>

Eudralink text: Please find attached the product information for <Product name> in your national language.

<For PSUR: Period covered by the PSUR: <dd.mm.yyyy> - <dd.mm.yyyy> >

<Please note that due to the existence of usage patent(s) the Annexes reflect differences in SmPC/PL compared to the reference medicinal product> [only for generics/hybrids/biosimilars – delete if not applicable]

Comments in Word must be sent directly to <MAH translation contact point e-mail> with copy to the Agency <Product Shared Mailbox>.

Deadline for the linguistic review is: <...>

Initial and line extension applications

Translations sent by Applicant/MAH to QRD@ema.europa.eu with a copy to the product mailbox

Renewals, variations, referrals, annual re-assessments

Translations sent by MAH to be below NCA mailing list with a copy to the procedure assistant/vet e-mail address and product mailbox

| Member states | Human | Veterinary |
|-----------------|--|--|
| Austria | Translations reviewed by Germany | Translations reviewed by Germany |
| Belgium | Translations reviewed by France, Germany and the Netherlands | Translations reviewed by France, Germany and the Netherlands |
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| Member states | Human | Veterinary |
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*German translations to be sent to both BfArM and PEI (Human) and to PEI and BVL (Veterinary)

** If applicable