



25 March 2026
EMA/315185/2010 rev. 4#
Human Medicines Division

List of official languages per country

Austria	German	Italy	Italian, German****
Belgium	German, French, Dutch	Latvia	Latvian
Bulgaria	Bulgarian	Lithuania	Lithuanian
Croatia	Croatian	Luxemburg	Luxemburgish**, German, French
Cyprus	Greek	Malta	Maltese, English ***
Czechia	Czech	Netherlands	Dutch
Denmark	Danish	Poland	Polish
Estonia	Estonian	Portugal	Portuguese
Finland	Finnish, Swedish*	Romania	Romanian
France	French	Slovakia	Slovak
Germany	German	Slovenia	Slovenian
Greece	Greek	Spain	Spanish
Hungary	Hungarian	Sweden	Swedish
Ireland	English, Irish**	United Kingdom (Northern Ireland)	English

#Rev. 4 Changes since last revision: Change of country name for Czechia

* Summary of Product Characteristics in Finnish; Labelling and Package Leaflet in Finnish and Swedish.

** Product Information is not required to be submitted in Irish and Luxemburgish.

*** Labelling and Package Leaflet can be submitted in Maltese or English for the purpose of marketing. For the purpose of the Commission Decision, full Product Information should be submitted in Maltese.

**** German Labelling and Package Leaflet must be provided for those medicinal products marketed in the Bolzano region (art. 80.1 D. Lvo 219/2006).

Iceland	Icelandic
Norway	Norwegian

