EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

1 January 2021
EMA/315185/2010 *rev. 3*#
Human Medicines Division

List of official languages per country

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

Iceland	Icelandic
Norway	Norwegian

* 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).

#Rev. 3 Changes since last revision: For the UK, as from 1.1.2021, EU Law applies only to the territory of Northern Ireland (NI) to the extent foreseen in the Protocol on Ireland/NI.

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