23 July 2020
EMA/381090/2020
Committee for Medicinal Products for Human Use (CHMP)

Summary of opinion[1] (post authorisation)

Crysvita

burosumab

On 23 July 2020, the Committee for Medicinal Products for Human Use (CHMP) adopted a positive opinion recommending a change to the terms of the marketing authorisation for the medicinal product Crysvita. The marketing authorisation holder for this medicinal product is Kyowa Kirin Holdings B.V.

The CHMP adopted a change and an extension to the existing indication as follows:[2]

> "Crysvita is indicated for the treatment of X-linked hypophosphataemia**, in children and adolescents aged 1 to 17 years** with radiographic evidence of bone disease, **and in adults.** ~~in children 1 year of age and older and adolescents with growing skeletons.~~".

Detailed recommendations for the use of this product will be described in the updated summary of product characteristics (SmPC), which will be published in the revised European public assessment report (EPAR), and will be available in all official European Union languages after a decision on this change to the marketing authorisation has been granted by the European Commission.

[1] Summaries of positive opinion are published without prejudice to the Commission decision, which will normally be issued 67 days from adoption of the opinion
[2] New text in bold, removed text as strikethrough

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