24 March 2022
EMA/184295/2022
Committee for Medicinal Products for Human Use (CHMP)

Summary of opinion[1] (initial authorisation)

Evusheld

tixagevimab / cilgavimab

On 24 March 2022, the Committee for Medicinal Products for Human Use (CHMP) adopted a positive opinion, recommending the granting of a marketing authorisation for the medicinal product Evusheld, intended for the pre-exposure prophylaxis of COVID-19 in adults and adolescents aged 12 years and older weighing at least 40 kg.

The applicant for this medicinal product is AstraZeneca AB.

Detailed recommendations for the use of this product are described in the product information (PI), which is published in English here.

The European public assessment report (EPAR) will be published after the marketing authorisation has been granted by the European Commission and will make this information available in all official European Union languages.

[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

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