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Press Office

Start of union reviews

CHMP meeting of 21-24 July 2025

Table 1. Start of reviews for centrally authorised medicines

Name	INN	Type of procedure	Scope
Tecovirimat SIGA	tecovirimat	Article 20 of Regulation (EC) No 726/2004	Procedure triggered by the European Commission asking for a review of Tecovirimat SIGA. This was prompted by emerging data from clinical trials raising concerns over potential lack of efficacy, and the need to determine whether this impacts the benefit-risk balance of Tecovirimat SIGA in its authorised indications.