08 May 2025
EMA/PRAC/157937/2025

Timetable for the procedure

Article 20 of Regulation (EC) No 726/2004 resulting from pharmacovigilance data

IXCHIQ

Procedure number: EMA/REF/0000269473

Procedural step	**Date**
Notification:	05 May 2025
Start of the procedure (PRAC[1]):	May 2025 PRAC
List of questions:	08 May 2025
Submission of responses:	26 May 2025
Re-start of the procedure:	05 June 2025
Rapporteur/co-rapporteur assessment reports circulated to PRAC and to CHMP[2]:	17 June 2025
Comments:	24 June 2025
Updated Rapporteur/co-rapporteur assessment reports circulated to PRAC and to CHMP:	01 July 2025
PRAC list of outstanding issues or PRAC recommendation to CHMP:	July 2025 PRAC

[1] Pharmacovigilance Risk Assessment Committee
[2] Committee for Medicinal Products for Human Use

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