



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Oxbryta

Procedural steps taken and scientific information after the authorisation

Application number	Scope	Opinion/ Notification ¹ issued on	Commission Decision Issued ² / amended on	Product Information affected ³	Summary
A20/0014		16/10/2025	09/12/2025		
PSUSA/10983 /202408	Periodic Safety Update EU Single assessment - voxelotor	13/03/2025	n/a		PRAC Recommendation - maintenance
PSUSA/10983 /202402	Periodic Safety Update EU Single assessment - voxelotor	05/09/2024	n/a		PRAC Recommendation - maintenance

¹ Notifications are issued for type I variations and Article 61(3) notifications (unless part of a group including a type II variation or extension application or a worksharing application). Opinions are issued for all other procedures.

² A Commission decision (CD) is issued for procedures that affect the terms of the marketing authorisation (e.g. summary of product characteristics, annex II, labelling, package leaflet). The CD is issued within two months of the opinion for variations falling under the scope of Article 23.1a(a) of Regulation (EU) No. 712/2012, or within one year for other procedures.

³ SmPC (Summary of Product Characteristics), Annex II, Labelling, PL (Package Leaflet).



II/0011	C.I.11.b - Introduction of, or change(s) to, the obligations and conditions of a marketing authorisation, including the RMP - Implementation of change(s) which require to be further substantiated by new additional data to be submitted by the MAH where significant assessment is required	05/09/2024	n/a		
PSUSA/10983/202308	Periodic Safety Update EU Single assessment - voxelotor	21/03/2024	17/05/2024	SmPC and PL	Refer to Scientific conclusions and grounds recommending the variation to terms of the Marketing Authorisation(s)' for PSUSA/10983/202308.
IA/0012	B.II.d.2.a - Change in test procedure for the finished product - Minor changes to an approved test procedure	20/02/2024	n/a		
IB/0009	B.I.a.1.z - Change in the manufacturer of AS or of a starting material/reagent/intermediate for AS - Other variation	09/11/2023	n/a		
PSUSA/10983/202302	Periodic Safety Update EU Single assessment - voxelotor	28/09/2023	n/a		PRAC Recommendation - maintenance
T/0007	Transfer of Marketing Authorisation	24/04/2023	08/06/2023	SmPC, Labelling and PL	
PSUSA/10983/202208	Periodic Safety Update EU Single assessment - voxelotor	30/03/2023	26/05/2023	SmPC and PL	Refer to Scientific conclusions and grounds recommending the variation to terms of the Marketing Authorisation(s)' for PSUSA/10983/202208.
IAIN/0006	B.II.b.2.c.1 - Change to importer, batch release arrangements and quality control testing of the FP - Replacement or addition of a manufacturer	17/02/2023	26/05/2023	Annex II and PL	

	responsible for importation and/or batch release - Not including batch control/testing				
IA/0004/G	This was an application for a group of variations. B.I.a.2.a - Changes in the manufacturing process of the AS - Minor change in the manufacturing process of the AS B.I.a.4.a - Change to in-process tests or limits applied during the manufacture of the AS - Tightening of in-process limits	15/06/2022	n/a		
IAIN/0003	B.II.b.1.a - Replacement or addition of a manufacturing site for the FP - Secondary packaging site	16/05/2022	n/a		
IA/0002	B.I.a.1.f - Change in the manufacturer of AS or of a starting material/reagent/intermediate for AS - Changes to quality control testing arrangements for the AS -replacement or addition of a site where batch control/testing takes place	19/04/2022	n/a		
IAIN/0001	B.II.b.2.c.1 - Change to importer, batch release arrangements and quality control testing of the FP - Replacement or addition of a manufacturer responsible for importation and/or batch release - Not including batch control/testing	12/04/2022	26/05/2023	SmPC, Annex II, Labelling and PL	