Insuman

Procedural steps taken and scientific information after the authorisation*

*Due to the Agency`s update of its procedure management systems, an additional document, reflecting the historical lifecycle may be available in the 'Assessment history' section. For the complete product lifecycle procedures, you may need to also refer to **EPAR - Procedural steps taken and scientific information after authorisation (archive)**.

Application number	Scope	Opinion/ Notification [1] issued on	Commission Decision Issued [2] / amended on	Product Information affected [3]	Summary
Variation type IB /	C.I HUMAN AND VETERINARY MEDICINAL	17/10/2025		SmPC,	

[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).

EMA/VR/0000295999	PRODUCTS - C.I.z Other variation - Accepted C.I.z - to update section 6.3 of the SmPC, section 5 of the Package Leaflet and outer packaging section in the labelling (8. Expiry date) for additional information for shelf life in the product information of Insuman Implantable 400 IU/ml solution for infusion (EU/1/97/030/202 and EU/1/97/030/203). Additionally, the MAH updated the local representative information in Iceland (name change) and Italy (telephone number change), and deletion of local representative in UK-NI.			Labelling and PL	