EU – Declaration of Conformity

We, the company,

Beiersdorf AG
SRN: DE-MF-000014151

20245 Hamburg
Germany

hereby declare in our sole responsibility as legal manufacturer that the device(s)

02728-00000-45	HP SCPL 21 PCS DE_FR_IT
02728-00002-45	HP SCPL 21 PCS ES_PT
02728-00003-45	HP SCPL 21 PCS DE_FR_NL
02728-00006-45	HP SCPL 21 PCS IT
02728-00006-46	HP SCPL 21 PCS EL_IT
02728-00009-46	CURI SCPL 21 PCS ES
02728-00011-45	EPL SCPL 21 PCS EN_FR
02728-00011-46	EPL SCPL 21 PCS EN_FR
02728-00012-45	EPL SCPL 21 PCS EN_ZH
02728-00013-45	HP SCPL 21 PCS AR_EN
48731-00000-45	HP SCPL L 21 PCS DE_FR_IT
48731-00000-46	HP SCPL L 21 PCS DE_FR_IT
48731-00001-45	HP SCPL L 21 PCS ES_HU_PT
48731-00002-45	HP SCPL L 21 PCS DE_FR_NL
48731-00003-45	EPL SCPL L 21 PCS EN_FR
48731-00003-46	EPL SCPL L 21 PCS EN_FR

Product Identification:	Hansaplast® / Elastoplast® Scar Reducer
Basic-UDI-DI:	4005800000000000000037D6
Intended Use:	For treatment of raised and coloured hypertrophic scars or keloid scars

is/are in compliance with Regulation (EU) 2017/745 concerning medical devices.

Classification
according to Annex VIII of the Regulation (EU) named above.

class I

Conformity Assessment Procedure
according to Article 52(7) and Annex IV of the Regulation (EU) named above.

Notified Body
not applicable for medical device class I.

This declaration is also valid for products which may carry additional labels to comply with local market and regulatory requirements (relabeling).

20245 Hamburg (Germany), 10.10.2023
(date)

Name
Senior Regulatory Affairs Manager
(Signed for and on behalf of Beiersdorf AG)