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)

48575-00000-48	HP BLPL S 6 DE_FR_NL
48575-00001-48	HP BLPL S 6 DE_FR_IT
48575-00002-48	HP BLPL S 6 EL_EN_ES
48575-00005-48	EPL BLPL S 6 EN_FR
48584-00000-48	HP BLPL L 5 DE_FR_NL
48584-00001-48	HP BLPL L 5 EL_ES_PT
48584-00002-48	HP BLPL L 5 EN_IT
48584-00003-48	HP BLPL L 5 INT
48584-00005-48	EPL BLPL L 5 EN_FR
48584-00007-48	HP BLPL L 5 DE_FR_IT
48834-00000-48	HP BLPL MIX 5PCS DE_ES
48834-00001-48	HP BLPL MIX 5PCS DE_FR_NL
48834-00002-48	EPL BLPL MIX 5PCS EN_FR

Product Identification:	Hansaplast®/Elastoplast® Blister Plaster / Blister Plaster Mix
Basic-UDI-DI:	4005800000000000000050CW
Intended Use:	For the use on blisters, pressure and chafe marks on hands and feet. Suitable for open and closed blisters. / Prevention of blisters, pressure and chafe marks / Plasters for prevention, protection and treatment of blisters. For heels and bunions / For toes and fingers

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

Classification

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

class IIa

Conformity Assessment Procedure

according to Annex IX chapter I and III of the Regulation (EU) named above.

Notified Body

DEKRA Certification GmbH
Handwerkstraße 15
70565 STUTTGART
Germany

Notified Body number: 0124

EU Certificate no. 50533-60

This declaration remains valid also in case the above-mentioned EU Certificate for the Quality Management System of Beiersdorf AG is renewed or revised.

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

Hamburg (Germany), 03.12.2024
(date)

M. Fraenkel
Martina Fraenkel
Regulatory Affairs Manager
(Signed for and on behalf of Beiersdorf AG)