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 devices

48736-00001-45	HP SO DL 5MX4CM TA 1 PCS INT
48737-00001-45	HP SO DL 5MX6CM TA 1 PCS INT
48738-00001-45	HP SO DL 5MX8CM TA 1 PCS INT
48739-00001-45	HP SO STR 19X72 TA 100 PCS INT
48740-00001-45	HP SO STR 30X72 TA 100 PCS INT
48741-00001-45	HP SO IJP 40X19 TA 100 PCS INT
48741-00002-45	EPL SO IJP 40X19 TA 100 PCS FR
48777-00001-45	HP SO DL 80CMX6CM INT

Product Identification:	Hansaplast/ Elastoplast/Curitas Soft New Backing
Basic-UDI-DI:	40058000000000000000035D2
Intended Use:	For covering and protection of minor, everyday wounds

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).

Hamburg (Germany) 10.10.2023
(date)

M. Fraenkel
Martina Fraenkel

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