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)

01145-00000-45	HP CLS DL 1X6 10 DE
01145-00002-45	HP CLS DL 1X6 10 DE_FR_NL
01145-00002-46	HP CLS DL 1X6 10 DE_FR_NL
01145-00005-45	HP CLS DL 1X6 10 INT
01145-00006-45	HP CLS DL 1X6 10 CS_HU_SK
01265-00000-45	HP CLS DL 2X6 TA 20 DE
01265-00002-45	HP CLS DL 2X6 TA 20 DE_FR_NL
01273-00000-45	HP CLS DL 1X8 TA 10 DE
01273-00001-45	HP CLS DL 1X8 TA 10 DE_FR_NL
48688-00001-45	HP CLS 5MX4CM TA 1 INT
48689-00001-45	HP CLS 5MX6CM TA 1 INT
48690-00001-45	HP CLS 5MX8CM TA 1 INT

Product Identification:	Hansaplast® Classic
Basic-UDI-DI:	400580000000000000000006CT
Intended Use:	For covering and protection of everyday wounds such as scratches, cuts and grazes.

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), 25.10.2022
(date)

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