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)

48598-00000-45	HP TGH STR 76X26 16 DE_FR_IT
48598-00001-45	HP TGH STR 76X26 16 INT
48598-00002-45	HP TGH STR 76X26 16 EL_IT_PT
48598-00004-45	HP TGH STR 76X26 16 DE_FR_NL
48598-00004-46	HP TGH STR 76X26 16 DE_FR_NL
48598-00005-45	HP TGH STR 76X26 16 AR_EN_MS
48598-00006-45	EPL TGH STR 76X26 16 EN
48598-00006-46	EPL TGH STR 76X26 16 EN
48598-00007-45	CURI TGH STR 76X26 16 ES
48598-00010-45	HP TGH STR 76X26 16 INT
48599-00000-45	EPL TGH STR 76X26 12 PCS EN
48600-00000-45	EPL TGH STR 76X26 20 PCS EN_FR
48600-00001-45	EPL TGH STR 76X26 20 PCS FR
48600-00001-46	EPL TGH STR 76X26 20 PCS FR
48600-00002-45	HP TGH STR 76X26 20 PCS ES_PT
48601-00001-45	EPL TGH STR ASS 15 EN
48601-00001-46	EPL TGH STR ASS 15 EN
48601-00002-45	EPL TGH STR ASS 15 EN_FR
48601-00002-46	EPL TGH STR ASS 15 EN_FR
48602-00000-45	HP TGH DL 10X6 8 DE_FR_IT
48602-00001-45	HP TGH DL 10X6 8 EL_ES_IT
48602-00002-45	HP TGH DL 10X6 8 DE_FR_NL
48602-00003-45	EPL TGH DL 10X6 8 EN
48602-00003-46	EPL TGH DL 10X6 8 EN
48602-00008-45	EPL TGH DL 10X6 8 FR
48602-00008-46	EPL TGH DL 10X6 8 FR
48602-00009-46	EPL TGH DL 10X6 8 EN_FR
48619-00000-45	EPL TGH STR 50X95 10 EN
48619-00000-46	EPL TGH STR 50X95 10 EN
48619-00001-45	HP TGH STR 50X95 10 ES_PT
48619-00002-45	EPL TGH STR 50X95 10 EN_FR
48792-00000-45	EPL FAB WRPF STR 18 PCS EN
48798-00000-45	EPL TGH WRPF FMY_PCK 50PCS EN_FR
48798-00001-45	EPL TGH WRPF FMY_PCK 50PCS EN

Product Identification:	Hansaplast® / Elastoplast® / Curitas® Extra Robust / Heavy Fabric / Extra Tough
Basic-UDI-DI:	400580000000000000000001CH
Intended Use:	Covering and protection of minor, 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), 10.10.2023
(date)

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