

Beiersdorf	Beiersdorf international documentation system BEC.10329232.000.01 DoC Spray plaster	Released: 25.11.25-... Page 1 of 2
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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)

48844-01010-46	HP SPY_PL 40 ML DE_FR_IT
48844-01020-46	HP SPY_PL 40 ML EL_IT
48844-01030-46	HP SPY_PL 40 ML ES_PT
48844-01040-46	HP SPY_PL 40 ML AR_EN_MY
48844-01050-46	HP SPY_PL 40 ML ID
48844-01060-46	HP SPY_PL 40 ML INT
48844-01070-46	HP SPY_PL 40 ML INT
48844-01080-46	HP SPY_PL 40 ML DE_FR_NL
48844-02010-46	EPL SPY_PL 40 ML EN
48844-02011-46	EPL SPY_PL 40 ML EN
48844-02020-46	EPL SPY_PL 40 ML EN_FR
48844-02021-46	EPL SPY_PL 40 ML FR
48844-03010-46	CURI SPY_PL 40 ML ES

Product Identification:	Hansaplast® / Elastoplast® / Curitas® Spray Plaster
Basic-UDI-DI:	40058000000000000000069DK
Intended Use:	Spray plaster to cover and protect minor, superficial, dry wounds / For covering and protection of minor, superficial, dry 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.

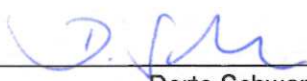
Notified Body

not applicable for medical device class I.

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

Hamburg (Germany), 25.11.2025
(date)


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