

We,

**Beiersdorf AG**

**Unnastr. 48  
20253 Hamburg**

are exclusively responsible for this declaration of conformity and hereby declare, that the following products

01861-00000-45	HP SPB 32,5ML DE_FR_IT
01861-00001-45	HP SPB 32,5ML DE_FR_NL
01861-00002-45	HP SPB 32,5ML EL_EN_IT
01861-00003-45	HP SPB 32,5ML ES_PT
01861-00004-45	HP SPB 32,5ML DA_FI_SV
01861-00005-45	EPL SPB 32,5ML FR
48605-00000-45	EPL SPP 40ML EN
48605-00001-45	EPL SPP 40ML EN_ZH

within the

**class IIa**

comply with

**Medical Device Directive 93/42/EEC, annex I.**

Conformity assessment procedure follows

**Medical Device Directive 93/42/EEC, annex II.**

<b>Beiersdorf</b>	<b>Beiersdorf international documentation system</b> <b>BEC.10245887.000.01</b> <b>DoC Spray Bandage Forward</b>	<b>Released:</b> 28.10.20-... Page 2 of 2
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Notified Body for Conformity Assessment Procedure:

DEKRA Certification GmbH  
Handwerkstraße 15  
70565 STUTTGART  
Country: Germany

Notified Body number : 0124

20253 Hamburg (Germany)  
(place of issuance)

28.10.20  
(date)

  
Hedda Siebrecht  
Senior Regulatory Affairs Manager

**Expiry date of this document:**  
valid until 26.05.2024