BODY PRODUCTS

Declaration of Conformity

We,

BODY PRODUCTS RELAX Pharma und Kosmetik GmbH
Alfred-Nobel-Straße 1-3
D-50226 Frechen, Germany

declare on our sole responsibility that the Personal Protective Equipment (PPE) described below

Item code	N19034-XS/-S/-M/-L/-XL
Description	MEDI-INN GREEN Nitril
Type	Nitrile protective glove, green, powder-free, biodegradable

complies with the provisions of Regulation (EU) 2016/425 as amended and supplemented.

Furthermore, the PPE complies with the following applied specifications:

- EN ISO 374-1:2016+A1:2018
- EN ISO 374-4:2019
- EN ISO 374-5:2016
- EN ISO 21420:2020
- EN 421:2010 (excluding clause 4.3)

In accordance with Articles 18 and 19 and Annex I, the product is assigned to Category III and is identical to the PPE for EU Type-Examination (Module-B), which is the subject of EU Type-Examination certificate No. 2777/14052-03/E02-01 issued by the body indicated below:

Notified Body	**SATRA Technology Europe Limited** Bracetown Business Park, Clonee, D15YN2P Republic of Ireland

The PPE is subject to the procedure set out in Annex VIII (Module D), of Regulation (EU) 2016/425 under the supervision and control of the notified body SATRA Technology Europe Limited - European Notified Body 2777.

For the use and application of the aforementioned product, we explicitly draw your attention to the fact that the instructions for use and the relevant application and safety regulations must be observed. The user alone bears responsibility for any improper use and/or use contrary to the intended purpose. For the purpose of proof of conformity, all relevant material and manufacturing certificates within the scope of the applicable regulations are stored in our premises and can be presented to the authorities.

This declaration is valid until 17.11.2028

Frechen, 24.01.2024
Place, Date

Yorck von Kries, Managing Director