

# EU type-examination certificate

The undersigned Notified Body declares, on the basis of the assessment of the tests (the results of which are referenced in the report) and the technical documentation provided mentioned below by the applicant

**PEKA Medical GmbH**  
**Konrad-Zuse-Straße 2**  
**42551 Velbert**  
**Germany**

that the tested product complies with the essential safety requirements of Annex II of Regulation 2016/425 (personal protective equipment) and Annex V (Module B)

The certificate shall only be used in conjunction with one of the conformity assessment procedures based on Annex VII (Module C2) or Annex VIII (Module D)

Manufacturing plant	<b>PEKA Medical GmbH</b> <b>Konrad-Zuse-Straße 2</b> <b>42551 Velbert</b> <b>Germany</b>
Type	<b>Filtering Half Mask FFP2 NR 318</b>
Description of product:	<b>See annex 1</b>
Test in accordance with	<b>EN 149:2001+A1:2009</b>

Registration No. 44 206 21151001  
Test Report No. 3530 1201  
File reference 8003034954

Valid from 2021-12-15  
Valid until 2026-12-14



TÜV NORD CERT GmbH  
Zertifizierungsstelle Konsumgüter  
Certification Body Consumer Products  
Benannte Stelle 0044 / Notified Body 0044

Essen, 2021-12-15

Bei PSA der Kategorie III folgt auf die CE-Kennzeichnung die Kennnummer der notifizierten Stelle, die in dem Verfahren nach Anhang VII oder VIII tätig war.



For Category III PPE, the CE marking shall be followed by the identification number of the notified body involved in the procedure set out in Annex VII or VIII.



# ANNEX

Annex 1, page 1 of 1

to Certificate Registration No. 44 206 21151001

Description	Filtering Half Mask
Type:	FFP2 NR
Model:	318

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