GÉPTESZT Kft.
Notified Body No. 2233
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MODULE B - EU TYPE-EXAMINATION CERTIFICATE EXTENSION

The PPE type complies with the applicable essential health and safety requirements of Regulation (EU) 2016/425.

*Present EU type-examination certificate is valid only with**

Document No: **TD11/GT553-X1/534/2212/EN/2233**

Category: III - Personal protective equipment providing respiratory system protection

Designation: **Particle filtering half mask**

Model: **AllProMask AIR, AllProMask AIR H, AllProMask AIR H+**

Classification: **FFP2 NR**

Description: *AllProMask AIR: FFP2 NR particle filtering half mask without valves in white colour with earloop.*
AllProMask AIR H: FFP2 NR particle filtering half mask without valves in white colour with head loop.
AllProMask AIR H+: FFP2 NR particle filtering half mask without valves in white colour, with head loop and with nose foam.

Applicant: **PAUL ALBRECHTS VERLAG GMBH**
Hamburger Strasse 6, 22952 Lütjensee, Germany

Reference of applied standard(s) / other technical specification(s):

EN 149:2001+A1:2009 Respiratory protective devices. Filtering half masks to protect against particles.

Test report / document reference(s):

*VD35/GT553-01/2212/EN/2233 & TD11/GT553-01/532/2212/EN/2233

VD36/GT553-X1/2022/EN

The certificate shall only be used in conjunction with one of the conformity assessment procedures referred to in point (c) of Article 19. of Regulation (EU) 2016/425.

Date of issue: Budapest, HUN - 15/12/2022
Date of expiry: 02/12/2027

GÉPTESZT Kft.
EVE Tanusító Szervezet
NB 2233
1037 Budapest, Jablonka u.79.

Andrea Nagy
Certification manager

Marking and instructions have been assessed in the English language only. It is the Manufacturer's/Authorised Representative's responsibility to obtain and supply language versions acceptable to the country where the product is to be sold.

The EU type-examination certificate remains the property of GÉPTESZT Kft. and will be withdrawn in case of existence of conditions stated in Article 32 point 5 and in Annex V. point 7.7 of Regulation (EU) 2016/425 of the European Parliament of the Council.

The manufacturer shall inform the notified body of all modifications to the approved type and of all modifications of the technical documentation that may affect the conformity of the PPE with the applicable essential health and safety requirements or the conditions for validity of that certificate. (Annex V. point 7.2 of Regulation (EU) 2016/425)

Legal remedy can be applied against the condition stated in the EU type-examination certificate. The application for appeal should be submitted to the Director Manager of GÉPTESZT Kft. and the application will be judged by the board of GÉPTESZT Kft. Certificate Body.

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