EU Declaration of Conformity

Annex IX PPE Regulation (EU) 2016/425

Date of issue: 2022-05-02 Revision: 1

(1) Product name	Type	Batch number or Serial number or Identifier
Hengst P3 Respirator	Re-usable filtered respiratory half mask	See individual packaging

(2) Manufacturer name and address is as follows:-
Hengst SE
Nienkamp 55-85
48147 Münster
Germany
Tel: +49 (0) 251 20 20 2

(3) This declaration of conformity is issued under the sole responsibility of the manufacturer

(4) Detailed description of the PPE to allow traceability/identification of the PPE

RE-USABLE FILTERED RESPIRATORY HALF MASKS
PRODUCT REF: F01.1.030 & F01.1.031

(5) The article identified in (4) above is in conformance with the relevant Union Harmonisation Legislation Regulation (EU) 2016/425

(6) References to the relevant harmonised standards used, including the date of the standard, or references to the other technical specifications, including the date of the specification, in relation to which conformity is declared: EN 140:1998

(7) BSI Netherlands, company registration number 33264284, John M. Keynesplein 9, 1066 EP, Amsterdam, The Netherlands Notifed Body CE 2797 performed the EU Type Examination (Module B) and issued the Type Examination Certificate Number CE 766364

(8) This product is category III and is subject to Module D Conformity to type based on quality assurance of the production process and is under the surveillance of BSI Netherlands, company registration number 33264284, John M. Keynesplein 9, 1066 EP, Amsterdam, The Netherlands Notifed Body CE 2797

(9) Additional information
No further additional information is available

Signed for and on behalf of

Place: Münster, Germany	Date: 2022-05-05
Name: Dr. Sebastian Stühle	Job title: Head of Business Unit Healthcare Filtration
Signature: S. Stühle	
Place: Münster, Germany	Date: 2022-05-05
Name:	Job title: Group Vice President
Signature: ppa.	