

EU DECLARATION OF CONFORMITY

Product:

QS II & Spiroguide II

Manufacturer:

Interspiro AB, Kemistvägen 21, S-183 79 TÄBY, Sweden

Type:

Self-contained open-circuit compressed air breathing apparatus with full face mask

Interspiro hereafter declares that the new product described above,

complies with the applicable essential health and safety requirements set out in Annex II of Personal Protective Equipment Regulation (EU) 2016/425:

- Conformity based on the compliance to standard EN 137:2006 (type 2)
- Module B - EU type-examination certificate 11255 A/24/29 PSA issued by Notified Body DEKRA Testing and Certification GmbH (No 0158) Handwerkstraße 15, 70565 Stuttgart, Germany
- Module D - Quality assurance of the production process under the surveillance of Notified Body SGS FIMKO OY (No 0598) Takomotie 8, 00380 HELSINKI, Finland
- The component BAC-III.v5 (as used for Spiroguide II) and the Wireless Heads-Up Display are compliant with Ex category Ia IIC T4. Conformity based on the requirements of standards EN 60079-0, EN 60079-11, and EN 60079-26
- If fitted with the Inposition belt, compliance to standard EN358

complies with the essential safety requirements set out in Annex I of the Pressure Equipment Directive 2014/68/EU for an assembly:

- Module B Production type of pressure cylinders and valves assembly - EU type-examination certificate 16-1007542-101 issued by Notified Body KIWA Sweden AB (No 0409)
- Module D - Quality assurance of the production process of pressure cylinders and valves assembly under the surveillance of Notified Body KIWA Inspecta AB (No 0409)

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.



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