

EU DECLARATION OF CONFORMITY

Product: Spiromatic 90U

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 11041 A/23/35 PSA issued by Notified Body DEKRA Testing and Certification GmbH (No 0158)
- Module D - Quality assurance of the production process under the surveillance of Notified Body SGS FIMKO OY (No 0598)

complies with the Marine Equipment Directive 2014/90/EU:

- Conformity based on the compliance to standard and/or technical specification(s) EN 136:1998, ISO 23269-2:2011, ISO 23269-3:2011, EN 137:2006, MED/3.7, SOLAS 74, Regulation II-2/10 & X/3, 2000 HSC Code 7, FSS Code 3, IBC Code 11, IGC Code 11 and IMO MSC/Circ.1499 & Circ.1555
- Module B - EC type-examination certificate MEDB000034D issued by Notified Body DNV AS (No 0575)
- Module D - Quality assurance of the production process under the surveillance of Notified Body DNV AS (No 0575)

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.



Roland Tschöp
CEO
Interspiro Group
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Interspiro AB