

EC - DECLARATION OF CONFORMITY EG – KONFORMITÄTSERKLÄRUNG

We
Wir

ATMOS MedizinTechnik GmbH & Co. KG
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declare under our sole responsibility that the medical device(s), classified as
erklären in alleiniger Verantwortung, dass das/die Medizinprodukt(e), klassifiziert als

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Ear/Nose/Throat Treatment Units HNO-Behandlungseinheit

ATMOS S 61 Servant system 2 modules ATMOS S 61 Servant system 2 Module	REF 534.1005.0
ATMOS S 61 Servant system 3 modules ATMOS S 61 Servant system 3 Module	REF 534.1010.0
ATMOS S 61 Servant system 4 modules XXL ATMOS S 61 Servant system 4 Module XXL	REF 534.0161.0
ATMOS S 61 Servant system 3 modules XXL ATMOS S 61 Servant system 3 Module XXL	REF 534.0160.0
ATMOS S 61 Servant system 4 modules ATMOS S 61 Servant system 4 Module	REF 534.1020.0
ATMOS S 61 Servant vision ATMOS S 61 Servant vision	REF 531.0000.0
ATMOS S 61 Servant instruments ATMOS S 61 Servant instruments	REF 532.2100.0
ATMOS S 61 Servant instruments XL ATMOS S 61 Servant instruments XL	REF 532.0500.0
ATMOS S 61 Servant instruments XXL ATMOS S 61 Servant instruments XXL	REF 532.0400.0
ATMOS S 61 CORIAN® instruments ATMOS S 61 CORIAN® instruments	REF 532.1000.0
ATMOS S 61 CORIAN® instruments XXL ATMOS S 61 CORIAN® instruments XXL	REF 532.0900.0
ATMOS S 61 CORIAN® integral ATMOS S 61 CORIAN® integral	REF 532.0800.0

meet(s) all applicable requirements of the Directive 93/42/EEC.
allen anwendbaren Anforderungen der Richtlinie 93/42/EWG entspricht/ entsprechen.

Conformity assessment procedure: Directive 93/42/EEC Annex VII on medical products, passed by the
commission on 14th June 1993, last amended on 5th September 2007
Konformitätsbewertungsverfahren: Richtlinie 93/42/EWG Anhang VII des Rates über Medizinprodukte
vom 14. Juni 1993, zuletzt geändert am 5. September 2007

Lenzkirch, 01.04.2020
Place and date of issue



i.V. Andreas Heer
Head of Quality Management



i.V. Steffi Focke
Quality Management