

Declaration of Conformity

The undersigned hereby declares, on behalf of ScandiDos AB (publ), that the product referenced below to which this declaration relates, is in conformity with the provisions of:

- LVFS 2003:11, the Swedish implementation of the COUNCIL DIRECTIVE 93/42/EEC on Medical Devices, by means of Annex II.
- RoHS 2 Directive (2011/65/EU)

The Technical Construction File required by the MDD Directive is maintained at the headquarter of ScandiDos AB (publ) in Uppsala (Sweden).

Manufacturer: ScandiDos AB (publ), Uppsala (Sweden)

Product: Pre-treatment Verification System

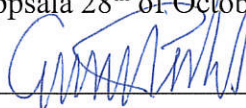
Model: Delta⁴ Phantom+
Delta⁴ Phantom+ IMRT
Delta⁴ Phantom+ Wired
Delta⁴ Phantom+ MR

Classification: Class I with measuring function

Serial Number: D0171 0000 – D0171 1000
D0172 0000 – D0172 2000
D0173 0000 – D0173 1000
D0174 0000 – D0174 2000
D0175 0000 – D0175 1000
D0176 0000 – D0176 2000
D0177 0000 – D0177 1000

Notified Body: Intertek Semko AB, CE 0413

Uppsala 28th of October 2022



Gustaf Piehl
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