Delta^4
by ScandiDos

D017 04 020 05

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:	Delta^4 Phantom+ Delta^4 Phantom+ IMRT Delta^4 Phantom+ Wired Delta^4 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
CEO and President

ScandiDos AB
Dag Hammarskjölds väg 52A,
SE-752 37, Uppsala, Sweden

e: info@scandidos.com
t: +46 18 472 30 30
w: delta4family.com