ScandiDos

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.

The Technical Construction File required by the 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+
Classification:	Class I with measuring function
Serial Number:	D0171 0000 – D0171 0500 D0172 0000 – D0172 1000 D0173 0000 – D0173 0500

Uppsala 30th of June 2015

Görgen Nilsson
CEO and president

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