Delta4
by ScandiDos

Doc No: D009 04 009 05

Declaration of Conformity

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

- Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices .

Manufacturer: **ScandiDos AB**
Dag Hammarskjölds väg 52A, third floor SE-752 37, Uppsala, Sweden

Single Registration Number: SE-MF-000035494

Product Name	Intended Use
HexaMotion	The intended use of the device is to realize a multi-dimensional movement pattern, e.g. of the Delta4PT or the Delta4 Phantom+.

Basic UDI-DI: 73500051800001EE

Classification: Class I

Conformity Assessment: Annex II & III

Uppsala, 2025-09-05

Signature on behalf of the manufacturer:

Gustaf Piehl, *CEO and President*

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