Declaration of Conformity

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

- EU Regulation 2017/745, Medical Device Regulation (MDR)
- RoHS Directive 2011/65/EU and amendment (EU) 2015/863

The Technical Documentation required by the Medical Device Regulation is maintained at the headquarter of ScandiDos AB (publ) in Uppsala (Sweden).

Manufacturer:	ScandiDos AB (publ), Uppsala (Sweden)
Basic UDI-DI:	73500051800001EE
Model:	HexaMotion
Classification:	Class I
Serial Number:	D009 0054 – D009 0254

Uppsala 26th of May 2021

Gustaf Piehl
CEO and President
on behalf of ScandiDos AB (publ)

Production Site	*Main Office*		
ScandiDos	*ScandiDos*	Tel: +46-18-60 22 05	
Dag Hammarskjölds väg 52A	Dag Hammarskjölds väg 52A	Fax: +46-18-10 74 02	
SE-752 37 Uppsala	SE-752 37 Uppsala	Info@scandidos.com	Org. Nr. 556613-0927
Sweden	Sweden	www.scandiDos.com	VAT Nr. SE556613092701