

## Declaration of Conformity

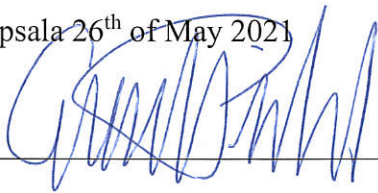
The undersigned hereby declares under the sole responsibility of ScandiDos AB (publ), that the product referenced below to which this declaration relates, is in conformity with the provisions of:

- 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 26<sup>th</sup> of May 2021

A handwritten signature in blue ink, appearing to read "Gustaf Piehl", is written over a horizontal line.

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

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