# PRODUCT LIST FOR CERTIFICATE

**Issued to:** ScandiDos AB

**Certificate number:** 28620160804

**Certificate valid from:** 2023-11-13

**Product List Issue Date:**
13 November 2023

| Product | Classification and EMDN | Intended use[1] | Date Added |
|---|---|---|---|
| **Class I devices with a measuring function** | | | |
| ***Basic UDI-DI: 73500051800002EG*** | | | |
| Delta4 Phantom+ | Class I(m)<br>Z11019099 | | 2023-11-13 |
| Delta4 Phantom+ IMRT | Class I(m)<br>Z11019099 | | 2023-11-13 |
| Delta4 Phantom+ MR | Class I(m)<br>Z11019099 | | 2023-11-13 |
| Delta4 Phantom+ Wired | Class I(m)<br>Z11019099 | | 2023-11-13 |

**Mikael Hagelin**
Certification Authority, MDR
Intertek Medical Notified Body AB, Torshamnsgatan 43,
Box 1103, SE-164 22 Kista, Sweden

Intertek Medical Notified Body AB is a Notified Body in accordance with the requirements set out in EU Regulation 2017/745 on medical devices, with the identification number 2862.

[1]The intended use is only included for class IIb devices and devices covered by an EU technical documentation certificate.