

Inspecta Sertifiointi Oy has granted this certificate as proof  
that the quality system of

## **Screentec Oy** **Oulu**

complies with the requirements of the standard

### **ISO 13485:2016**

Certification covers

**Contract manufacturing and manufacturing process development of  
professional electronics and medical devices subassemblies.**

**Main Technical Area: Active Medical Devices (Non-implantable)**  
**Technical Area: General active medical devices**

The certificate is issued on 2017-10-13.

The certificate is valid until 2020-10-13.



Mikko Törmänen, Managing Director

The certificate is valid on condition that the quality system of the organization  
remains in compliance with the aforementioned standard and the General Regulations ABC 200.  
The validity of the certificate can be verified on the Internet at [www.inspecta.fi](http://www.inspecta.fi)



ISO 13485

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