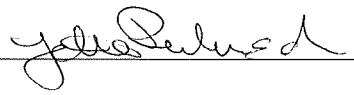
Declaration of conformity
Medical Devices, class I

Legal Manufacturer:	Abena A/S including Abena International Egelund 35 DK 6200 Aabenraa
Conformity assessment procedure	Annex VII of the Medical Devices Directive 93/42/EEC as amended by the Council Directive 2007/47/EEC.
Classification and harmonized standards	Class I
Product	Curi-Med Film roll, non-sterile item 220947, 220948, 220949
We, the legal manufacturer hereby declare that the above-mentioned product complies with the European Medical Device Directive 93/42/EEC as amended by the Council Directive 2007/47/EEC and its relevant transposition into all national laws of the member states into which we place the devices.	
Signed in Aabenraa	30.10.2020
Name and authority	Jette Rubusch Global Category Manager
Signature	

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