EC CERTIFICATION

EU Quality Assurance System Certificate

Regulation (EU) 2017/745 for Medical Devices, Annex XI Part A

We hereby declare that a conformity assessment based on a production quality assurance system, restricted to the aspects relating to the conformity of the devices with reusability requirements, has been carried out following the requirements of Regulation (EU) 2017/745 for Medical Devices.

We certify that the documentation conforms to the relevant provisions of the aforementioned regulation, and the result entitles the organization to use the CE 2862 marking on the products listed below.

Norfolk International

Hussaini Road, Muzafar Pur, Sialkot (51310), Pakistan

Manufacturer SRN: PK-MF-000033919

Authorised Representative:
CS Active
Usmas iela, 19 dz. 24, 1083, Riga, Latvia

Scope:

Class I reusable surgical instruments

Certificate Number:
28620248199

Revision:
00

Initial Certification Date:
15 June 2026

Certificate Decision Date:
15 June 2026

Certificate Issue Date:
15 June 2026

Certificate Expiry Date:
14 June 2031

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.

PRODUCT LIST FOR CERTIFICATE

See attached product list

EXAMINATION AND TESTS PERFORMED

Last Audit report reference	**Stage 1 ACTY-2025-370567**
	Stage 2 ACTY-2025-379963

CONDITIONS FOR OR LIMITATIONS TO VALIDITY OF CERTIFICATE

None

CERTIFICATE HISTORY

CERTIFICATE NUMBER	DATE OF ISSUE	IDENTIFICATION OF CHANGES
28620248199	15 June 2026	Initial certificate

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28620248199

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