

EU Declaration of Conformity

We as the manufacturer, MICROS Produktions- und Handelsges.m.b.H. with registered office in Hunnenbrunn - Gewerbezone 5, 9300 St. Veit an der Glan, Austria, hereby declare under our sole responsibility that the devices covered by this declaration comply with Regulation (EU) 2017/746, General Safety and Performance Requirements, and the relevant Union legislation.

SRN (Single Registration Number): AT-MF-000039061

Product category: Pathological Instruments
Product groups: Microtome Blades, Trimming Knives
Classification: Class A, rule 5 (acc. to Annex VIII of Regulation (EU) 2017/746)
EMDN code: W0202059085

Ref. No.	Basis UDI-DI	Product name	Product specifications
110888	9120131868889	Routine LP 35	Low Profile Disposable Microtome Blade, angle 35°
110999	9120131869992	Routine HP 35	High Profile Disposable Microtome Blade, angle 35°

Intend of use: These disposable, non-sterile in vitro diagnostic devices for clinical use are intended for thin sectioning of tissue prepared for pathological examination.

EC regulation: Regulation (EU) 2017/746 (In Vitro Diagnostics Medical Devices)

Conf. Assessment Route: Annex II and Annex III of this Regulation



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This declaration is effective for products placed on the market as of the date of issue. Any modification of the devices not authorized by MICROS Produktions- und Handelsges.m.b.H will invalidate this declaration.

St. Veit an der Glan, 26.11.2025



Herbert Luttenberger
Quality Manager & PRRC

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