

## 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                                     |
|----------|---------------|-----------------------------|------------------------------------------------------------|
| 109214   | 9120131862146 | <b>MS22</b>                 | Low Profile Disposable Microtome Blade, angle 22°          |
| 109206   | 9120131862061 | <b>MS33</b>                 | Low Profile Disposable Microtome Blade, angle 30°          |
| 109207   | 9120131862078 | <b>MS24</b>                 | Low Profile Disposable Microtome Blade, angle 35°          |
| 109211   | 9120131862115 | <b>MS200</b>                | Low Profile Disposable Microtome Blade, angle 35°          |
| 109212   | 9120131862122 | <b>MS300</b>                | Low Profile Disposable Microtome Blade, angle 35°          |
| 109213   | 9120131862139 | <b>HP</b>                   | High Profile Disposable Microtome Blade, angle 35°         |
| 109220   | 9120131862207 | <b>130mm Trimming Knife</b> | Sectioning of hard tissues with fibers, blade length 130mm |
| 109221   | 9120131862214 | <b>260mm Trimming Knife</b> | Sectioning of hard tissues with fibers, blade length 260mm |

**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

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, 30.11.2023



**Herbert Luttenberger**  
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