Certificate of EU Medical Device Notification

This is to certify that, according REGULATION (EU) 2017/746 of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on in vitro diagnostic medical devices,

Humiss Beratung GmbH
Address: Gneisenaustraße 8. 40477, Düesseldorf, Deutschland
TEL: +49-211-90760042, FAX: +49-211-90760043, E-mail: eurep@humiss.com
SRN: DE-AR-000023447

has fulfilled all notification responsibility and duty as the European Authorized representative of:
Manufacturer: Shanghai Tongbai Healthcare Technology Company
Address: Room 307, Building C, No. 898, Jiujing Road, Songjiang District, Shanghai City, 201615, China

The manufacturer has provided with all the appropriate declaration according the REGULATION (EU) 2017/746 requirements including the EC Declaration of Conformity confirming that the in vitro diagnostic medical devices, as stipulated here below, is fulfilling the applicable requirements of REGULATION (EU) 2017/746.

Product(s):	Microtome Blades
Notification number:	DE/CA20/00124881
Model(s):	CR75, CW75
Classification:	Class A, Rule 5 (Annex VIII)

Where then manufacturer affixes the CE marking to the product listed, they must ensure that all the requirements of the appropriate EU regulation(s) have and continue to be met.

The notification of aforementioned device(s) has been completed by European Authorized representative in Netherlands, the Netherlands Competent Authority has notified the manufacturer's in vitro diagnostic medical device above and has allocated registration.

Humiss Beratung GmbH

This is only a CE mark sample which is only use for reference

Signature of Executive Director
James St. WU

Cert. №: D-YQ-IVD-260126228
Issue date:Jan.26, 2026
Valid until: Jan.25, 2031

Contact HUMISS
E-mail: ce-tech@humiss.com
http://www.humiss.com