HUMISS

Certificate of EU Medical Device Notification

This is to certify that, according Directive 98/79/EC of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on in vitro diagnostic medical devices,

Humiss International B.V.
Address: Joop Geesinkweg 701, 1114AB Amsterdam-Duivendrecht, the Netherlands

has fulfilled all notification responsibility and duty as the European Authorized representative of:

Manufacturer: MDHC Life Technologies (Kunshan) Co., LTD.
Address: Building A1/A5/B1, No. 188 East Shengxiang Road, Qiandeng Town, Kunshan City, Jiangsu Province, 215341, P.R. China

The manufacturer has provided with all the appropriate declaration according the Directive 98/79/EC 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 Directive 98/79/EC.

Product(s): Centrifuge Tube
CIBG number: MDHC (NL-CA002-2021-62244) , CIBG-20215733
Model(s): 15ml, 50ml, 250ml, 500ml
Classification: Other

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 INTERNATIONAL B.V.

CE

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

James St. WU

Signature of Executive Director

James St. WU

Contact HUMISS
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Cert. №: H-YQ-IVD-210906084/02
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Valid until: Sep.05, 2026