

Certificate of EU Medical Device Notification

This is to certify that, according REGULATION (EU) 2017/745 of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on in Medical Devices,

HUMISS TRADING S. L.

Address: Calle Luis Buñuel 12, MADRID, 28018, Spain

SRN: ES-AR-000040177

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

Manufacturer: Nantong Jincheng Zencare Technology Co., Ltd.

Address: No. 8 Heli Rd, Jiuhua Town, Rugao City, Nantong, Jiangsu Province, 226541, China

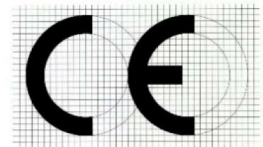
The manufacturer has provided with all the appropriate declaration according the REGULATION (EU) 2017/745 requirements including the EC Declaration of Conformity confirming that the medical devices, as stipulated here below, is fulfilling the applicable requirements of REGULATION (EU) 2017/745.

Product(s) details shown in the Annex of this certificate.

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 Spain, the Spain Competent Authority has notified the manufacturer's medical device above and has allocated registration.

You can check the authenticity of the document at the headquarters of the AEMPS (<https://localizador.aemps.es>) with CSV number listed above.



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



Signature of Executive Director

Tamara Manchón



Annex to cert No. S-YQ-MD-2510280091

Product name	Models	Activities	Registration number	CSV	Classification
<i>Adult Nursing Pad</i>	<i>M, L, XL, XXL</i>	<i>Manufacturer</i>	<i>RPS/2496/2025</i>	<i>1FLNBHFBL69MW VXBG86D</i>	<i>Class I, rule 1</i>
<i>Adult diaper</i>	<i>M, L, XL, XXL</i>	<i>Manufacturer</i>			<i>Class I, rule 1</i>

End of the annex.