



Ver: CERT-202110.V1

**CERTIFICATE**  
ECREP20220526.43



## CMC MEDICAL DEVICES & DRUGS S.L.

CONFIRMED THAT CMC MEDICAL DEVICES & DRUGS S.L. Is the  
European Authorized Representative of

**Hunan Runmei Gene Technology Co., Ltd.**  
**Room 401-1, Building 3, Shanhe Medical and Health**  
**Industrial Park, No. 1048,Zhongqing Road, Shaping**  
**Street, Kaifu District**

The certificate remains valid until the expiration agreement of EC REP, manufacturing conditions, the quality system or relevant legislation are changed. The validity is conditioned by positive results of periodic surveillance audits.

The product liability rests with the manufacturer in accordance with applicable directive/regulation and standard mention in Annex I of this certificate, after fulfilling of the relevant EU legislation requirements, the manufacturer shall affix relevant CE marking to all below mentioned models of the medical device.



  
Authorized Signature

Issue date: 26/05/2022

Expiration date: 26/04/2027

Verification Code

CMC Medical Devices & Drugs S.L.  
C/ Horacio Lengo Nº18, CP29006, Málaga-Spain  
[www.cmcmedicaldevices.com](http://www.cmcmedicaldevices.com)





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## ANNEX I

Product Name	CLASSIFICATION	REGULATION	RPS (AEMPS)	Incluir en Certificado
Monkeypox Virus Detection Kit(PCR-Fluorescent Probe)	Others	IVDD	RPS/3935/2022	Si



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