

Certificate

We hereby certify the company

NG Biotech SAS
Atelier Relais Le Tremplin
Parc d'Act.de Courbouton Sect 1
35480 Guipry
France

the introduction and application of a

Quality management system according to EN ISO 13485

in the scope

Design, Development, Manufacturing and Sales of in vitro diagnostic devices based on lateral flow technology

An audit by mdc has proven that this quality management system meets the requirements of the following standard:

EN ISO 13485:2016 + AC:2018 + A11:2021 - ISO 13485:2016
Medical devices – Quality management systems – Requirements for regulatory purposes

Valid from 2024-11-14
Valid until 2027-11-13

Registration No. D1395900020
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Stuttgart, 2024-11-14



Certification Body