EU Declaration of Conformity

according to EU regulation 2017/746 relating to in vitro diagnostic medical devices

The company:

Company name	Single registration number	Address
DYNABIO S.A.	Awaiting issuance by ANSM	Luminy Biotech Entreprises Case 922 – 163, Avenue de Luminy 13288 Marseille cedex 9 France

certifies under its sole responsibility that the device:

Product name	Product code	Destination	Risk class according to appendix VIII	Basic UDI-DI
MucoPAP-F	MPK03	Cystic fibrosis newborn screening	C (3m ruler)	3770026964DYNABIOMPK03E4

complies with European regulation 2017/746 relating to *in vitro diagnostic medical devices* and complies with the French Public Health Code.

Done in Marseille, on 08/18/2022

Yann Dagorn
President, Dynabio S.A.