DYNABIO S.A.

Luminy Biotech Entreprises Case 922 - 163, avenue de Luminy 13288 Marseille Cedex 9 FRANCE

Declaration of Conformity CE

In accordance with ISO standards 17050-1:2004 and 17050-2:2004

DC/CE-MPK03-EN	
Version 9	1/1

Declaration of Conformity CE

according to European Directive 98/79/CE related to in vitro diagnostic medical devices

The company

DYNABIO S.A.

Luminy Biotech Entreprises
Case 922 – 163, Avenue de Luminy
13288 Marseille cedex 9
France

ensures and declares, under its sole responsibility, that the product:

MucoPAP-F

is in conformity with applicable requirements of directive 98/79/CE, article 110 of regulation 2017/746 and that :

- there were no changes in purpose, design and content of the product,
- and that this product does not impose unacceptable or imminent safety risks for patients in their use.

This statement is based on the following:

• Technical file DT-MPK03 (version 6) demonstrating compliance with essential requirements of directive 98/79/CE (Annex I and III).

Marseille, 12/03/2025



Y. Dagorn President Dynabio S.A.

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