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.