



CERTIFICATE

EC Certificate No. 1434-IVDD-154/2022
Full Quality Assurance System
Directive 98/79/EC concerning
***in vitro* diagnostic medical devices**

Polish Centre for Testing and Certification certifies
that the quality assurance system in the organization:

Benetech Inc.

555 Richmond Street West Suite 500
Toronto, Ontario M5V 3B1 CANADA

for the design, manufacture and final inspection of *in vitro* diagnostic medical device
List B

Benetech PRA Software, 3.x.x.x

complies with requirements
of Annex IV (excluding Section 4, 6) to Directive 98/79/EC (as amended)
implemented into Polish law,
as evidenced by the audit conducted by the PCBC

Validity of the Certificate: from **25.05.2022** to **27.05.2025**

The date of issue of the Certificate: **25.05.2022**

The date of the first issue of the Certificate: **29.06.2020**



Issued under the Contract No. MD-016/2022
Application No: 527/2022
Certificate bears the qualified signature.
Warsaw, 25/05/2022
Module H7

Director
Medical Device Certification
Department