Covered Equipment Attestation Letter

For Certification Service in the USA

Federal Communications Commission
Equipment Authorization Division, Application Processing Branch
7435 Oakland Mills Road
Columbia, MD 21048

To whom it may concern

Pursuant to Section 2.911(d)(5)(i)-(ii) of the Commission's Rules (47 C.F.R.) we attest as follows for the product listed below:

FCC ID	Model name
2BCAP000002	MasterX903SW, MasterX901SW, MasterX901HD, MasterX901UHD, MasterX901HGF, MasterX903HD, MasterX903UHD

We,

Applicant	
Company Name:	Medicatech USA
Address:	1245 N. Patt St 92801, Anaheom, California 92801, United States
Phone:	+1-949-679-2881

certify that the equipment for which authorization is sought is not "covered" equipment prohibited from receiving an equipment authorization pursuant to section 2.903 of the FCC rules. This includes the cybersecurity and anti-virus software produced or provided by Kaspersky Lab, Inc. or any of ist successors and assignees.
We certify that, as of the date of the filing of the application, the applicant is not identified on the Covered List as an entity producing "covered" equipment.

If you have any questions, please feel free to contact us at the address shown below.

Sincerely,

2026-03-17

Signature Date

Name / Title: George Makar / President
Company: Medicatech USA
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