Lumicare Medtech Co., Ltd

Covered Equipment Certification Attestation Letter

3/20/2025

Nemko North America, Inc.
303 River Road
Ottawa
K1V 1H2
Canada

ATTN.: Reviewing Engineer

FCC ID: **2BM9Y0001**

Lumicare Medtech Co., Ltd ("the applicant") certifies 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.

> Note: If the equipment for which the applicant seeks authorization is produced by any of the entities identified on the current Covered List, the applicant should include an explanation on why the equipment is not "covered" equipment.

Lumicare Medtech Co., Ltd ("the applicant") certifies 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.

Lumicare Medtech Co., Ltd ("the applicant") certifies that the equipment for which authorization is sought does not include cybersecurity or anti-virus software produced or provided by Kaspersky Lab, Inc. or any of its successors and assignees, including equipment with integrated Kaspersky Lab, Inc. (or any of its successors and assignees) cybersecurity or anti-virus software pursuant to DA-24-886 and KDB 986446 D01 Covered Equipment Guidance section B(2a).

Signed: 刘彤 LiuTong.

Printed name: **Tong Liu**
Title: **R&D Manager**
Company Name: **Lumicare Medtech Co., Ltd**
Date: **3/20/2025**