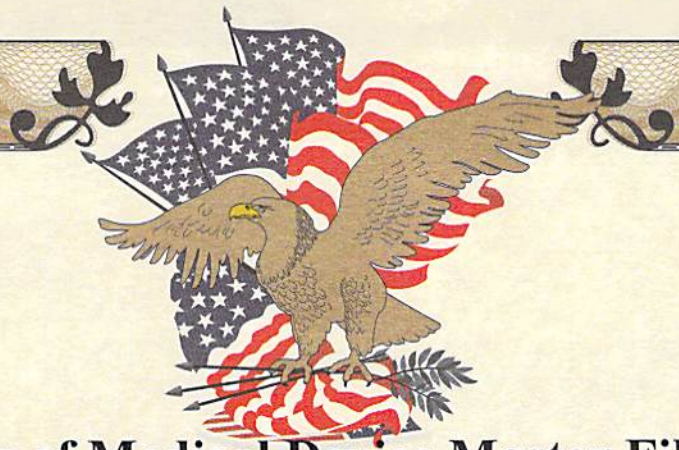

Certificate of Medical Device Master File (MAF)

Service Period
June 1, 2021 – May 31, 2022

This certifies that:

Kimmerling Holdings Group
3698 Largent Way
Suite 202
Marietta, GA 30064
United States

is a Medical Device Master File holder with the U.S. Food and Drug Administration pursuant to part 814 of Title 21, US Code of Federal Regulations, such filing having been verified as currently effective on the date hereof by Registrar Corp.

Medical Device Master File Number:	**2174**
Status:	**Active**
Subject:	**K-18 QAMS and K-21 QAS**

Registrar Corp will confirm that such filing remains effective upon request and presentation of this certificate until May 31, 2022, unless terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. Filing of a Medical Device Master File does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of filing of Medical Device Master File is misleading. The U.S. Food and Drug Administration does not issue a certificate of Medical Device Master Files. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lemarz
Executive Director
Registrar Corp
Dated: June 10, 2021