



2020

CERTIFICATE OF REGISTRATION

This certifies that:

CARINE EUROPE GMBH
Corneliusstr. 44
Munich Bayern, DE 80469

is registered with the U.S. Food and Drug Administration for FY 2020 pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations:

Establishment Owner/Operator Number:
DUNS No.:
Device Classification Name:

10074960
31-321-0501
LATEX PATIENT EXAMINATION
GLOVE

Product Code:
Regulation Number:
Official Correspondent
and U.S. Agent:

LYY
880.6250
Registrar Corp
144 Research Drive, Hampton, Virginia,
23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-
224-0179

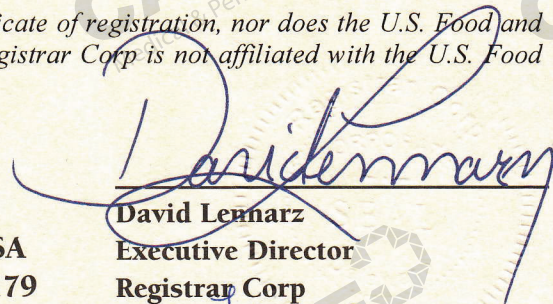
Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does 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. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Registrar Corp assumes no liability to any person or entity in connection with the foregoing.

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Registrar Corp

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David Lennarz
Executive Director
Registrar Corp

Dated: June 15, 2020