



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:

10074960

DUNS No.:

31-321-0501

Device Classification Name:

PATIENT EXAMINATION GLOVE

Product Code:

FMC

Regulation Number:

880.6250

Official Correspondent
and U.S. Agent:

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 Lemarz
Executive Director

Registrar Corp

Dated: June 15, 2020