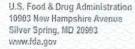
APOSTILLE

(Convention de La Haye du 5 octobre 1961)

1. Country:	United States of America
This public document 2. has been signed by	CDR Cesar A. Perez, PhD, Director
3. acting in the capacity of	DRP2: Division of Establishment Support
4. bears the seal/stamp of	U. S. Department of Health and Human Services
Certified	
5. at Washington, D.C.	
6. the twenty-eighth of January, 2020	
7. by Assistant Authentication Officer, United States Department of State	
8. No. 20018405 2	
9. Seel/Stamp:	10. Signature:
Sonya N. Johnson	





Certificate No. 3854-1-2020

CERTIFICATE TO FOREIGN GOVERNMENT

In order to allow the importation of United States products into foreign countries, the U.S. Food and Drug Administration (FDA) certifies the following information concerning the product(s) to be exported listed below:

Name of Product(s)

Name of Manufacturer/Distributor, Address

See Attached List

See Attached List

(Two Pages)

(One Page)

The product(s) described above (and the manufacturing/distribution site(s) which produces/distributes it) is subject to the jurisdiction of the FDA under the Federal Food, Drug, and Cosmetic Act.

It is certified that the above product(s) may be marketed in, and legally exported from, the United States of America at this time. The manufacturing plant(s) in which the product(s) is produced is subject to periodic inspections. The last such inspection showed that the plant(s), at that time, appeared to be in substantial compliance with current good manufacturing practice requirements for the product(s) listed above.

Sincerely,

CDR Cesar A. Perez, PhD, Director DRP2: Division of Establishment Support Office of Regulatory Programs Office of Product Evaluation and Quality Center for Devices and Radiological Health

U.S. Food and Drug Administration, DHHS

This certificate is valid from January 09, 2020 to January 08, 2022.





U.S. Food & Drug Administration 10903 New Hampshire Avenue Silver Spring, MD 20993 www.fda.gov

Certificate No. 3854-1-2020
Certificate to Foreign Government - Name of Manufacturer/Distributor Attachment Page 1 of 1

Name of Manufacturer

Legal Manufacturer

3M COMPANY 3M Center, 2510 Conway Ave. Bldg. 275-5W-06 Saint Paul, MN USA 55144

Manufacturing Facility

3M CANADA CO. 400 RT. 100 MORDEN, Manitoba CANADA R6M 1Z9

Name of Distributor

Distributor 3M COMPANY 806 W. CRYSTAL LAKE RD. FOREST CITY, IA USA 50436

----END OF MANUFACTURER/DISTRIBUTOR LIST----





U.S. Food & Drug Administration 10903 New Hampshire Avenue Silver Spring, MD 20993 www.fda.gov

Certificate No. 3854-1-2020 Certificate to Foreign Government - Name of Product(s) Attachment Page 1 of 2 Name of Manufacturer

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Manufacturing Facility

3M CANADA CO. 400 RT. 100 MORDEN, Manitoba CANADA R6M 1Z9

Name of Distributor

Distributor

3M COMPANY 806 W. CRYSTAL LAKE RD. FOREST CITY, IA USA 50436

Name of Product(s)

3M™ Double-Stick Discs 2181

3M[™] Red Dot[™] Monitoring Electrode with Foam Tape 2228, 2228-3, 2228-5, 2228BA

3M[™] Red Dot[™] Monitoring Electrode 2230, 2235, 2237, 2237-3, 2237-5, 2238, 2238-3, 2238-5, 2239 2245-50, 2248-50, 2249, 2249-50, 2255-50, 2259-50 2270-5, 2270-50, 9642

3M[™] Red Dot [™] Diaphoretic Monitoring Electrode 2231, 2271-50

3M™ Red Dot™ Trace Prep

2236

3M™ Red Dot™ Radiolucent Monitoring Electrode with Foam Tape

3M™ Red Dot™ Wet Gel Monitoring Electrode 2256-50

3M™ Red Dot™ ECG Monitoring Electrode 2268-3, 2268-5

3M[™] Red Dot[™] Neonatal Monitoring Electrode with Pre-Attached Lead Wire 2269T

3M™ Red Dot™ Prewired Monitoring Electrode 2289PA, 2289PAL, 2289T

 $3M^{\intercal\!M}$ Red Dot $^{\intercal\!M}$ Monitoring Electrode with Foam Tape and Sticky Gel 2560, 2560-3, 2560-5

3M™ Red Dot™ Radiolucent Monitoring Electrode with Foam Tape, Sticky Gel and Abrader Pad 2570, 2570-3, 2570-5





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Certificate No. 3854-1-2020 Certificate to Foreign Government - Name of Product(s) Attachment Page 2 of 2

3M™ Red Dot™ Repositionable Monitoring Electrode 2660-3, 2660-5, 2670-3, 2670-5 -----END OF PRODUCT LIST-------

