MedPath

EC-Registration Certificate

Directive 93/42/EEC on Medical Devices (MDD), Article 14

No. R A002 87/A Rev. 01

Manufacturer: FUJIAN SHANDESET ENVIROMENTAL PROTECTION TECHNOLOGY CO.,LTD

SONG TANG ROAD 838, BAO GAI TECHNOLOGY PARK,

SHISHI, QUAN ZHOU CITY, FUJIAN PROVINCE, CHINA

Product Category(ies): See Appendix A

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This is to certify that, in accordance of the Medical Device Directive 93/42/EEC (amended by 2007/47/EC), MedPath GmbH agrees to perform all duties and responsibilities as the Authorized Representative for the aforementioned manufacturer as stipulated and demanded by the aforementioned Directive. The German Competent Authority is notified of the manufacturer's medical device(s) shown in Appendix A. The manufacturer has provided MedPath GmbH with the appropriate Declaration(s) of Conformity confirming that the medical device(s) fulfills/fulfill the applicable requirements of the aforementioned Directive.

MedPath GmbH
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Date, 2020-05-05 MedPath GmbH

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Appendix A: Product Category(ies)

Name	Classification	UMDNS Code	Form No.
Single-use medical face mask	I	12-447	00304332

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