MedPath GmbH, Mies-van-der-Rohe-Straße 8, 80807 Munich, Germany

To whom it may concern

MedPath GmbH

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Confirmation of EU Medical Device(s) Notification

This is to confirm that in accordance of Article 14 of Directive 93/42/EEC concerning Medical Devices (MDD), MedPath GmbH agrees to perform all duties and responsibilities as the Authorized Representative for the manufacturer

LHM Medical Technology (Hong Kong) Limited

Unit No. 2, 3/Floor, Block A, Ko Fai Industrial Building, 7 Ko Fai Road, Yau Tong, Kowloon, Hong Kong 999077

as stipulated and demanded by the aforementioned Directive. The German Competent Authority is notified of the manufacturer's medical device(s)

- **Fluid Resistant Procedure Mask**

The manufacturer has provided MedPath GmbH with the appropriate Declaration(s) of Conformity claiming that the medical device(s) fulfills/fulfill the applicable requirements of the aforementioned Directive.

Safety Officer
MedPath GmbH

MedPath GmbH
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