The management system of

MP Biomedicals, LLC Diagnostic Division

29525 Fountain Parkway Solon, OH 44139, United States

has been assessed and certified as meeting the requirements of

ISO 13485:2016 EN ISO 13485:2016

For the following activities:

Design and development, manufacture and distribution of In-Vitro Diagnostic Test Kits for Endocrine and Neonatal Analysis.

This certificate is valid from 29 March 2019 until 15 June 2020 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 15 June 2020.

Issue 7. Certified since 22 September 2009.

Authorised by



SGS United Kingdom Ltd Rossmore Business Park Ellesmere Port Cheshire CH65 3EN UK t +44 (0)151 350-6666 f+44 (0)151 350-6600 www.sgs.com

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