MORE DIAGNOSTICS, INC.

Declaration of Conformity

We hereby declare that the in vitro diagnostic devices and the accessories described below conform to all of the applicable essential requirements contained in the Annex I Essential Requirements of In Vitro Diagnostic Device Directive 98/79/EC as published in the European Official Journal on 1999-12-07.

This conformance is supported by the technical documentation which is maintained by MORE DIAGNOSTICS, Inc.

Product:	MPA Control
Catalog Number (REF):	240 – 4- level package 240-1 – Level 1 package 240-2 – Level 2 package 240-3 – Level 3 package 240-4 – Level 4 package
Manufacturer Address:	More Diagnostics, Inc. 2020 11th Street Los Osos, CA 93402
EC Authorized Representative:	Siemens Healthcare Diagnostics Limited Sir William Siemens Square Frimley Camberley GU16 8QD United Kingdom
Date:	2018-3-26

Authorization:

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