# 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.

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|---|---|
| **Product:** | Rap//Tac/CsA Control |
| **Catalog Number (REF):** | 290 - Tri-level package<br>290-1 – Level 1 package<br>290-2 – Level 2 package<br>290-3 – Level 3 package |
| **Manufacturer Address:** | More Diagnostics, Inc.<br>2020 11th Street<br>Los Osos, CA 93402 USA |
| **EU Authorized Representative:** | Siemens Healthcare Diagnostics Manufacturing Ltd<br>Chapel Lane, Swords<br>Co. Dublin, Ireland |
| **Date:** | 2019-1-31 |

**Authorization:**

Signature: Harry R Holden

Harry R. Holden
RA/QA Management Representative
More Diagnostics Inc.

More Diagnostics Inc.
P.O. Box 6714
Los Osos, CA
93402 USA
2020 11th Street