## **GRIFOLS**

### **Declaration of Conformity**

**Technical File Reference:** 

SDTF-002, Rev. 002

Issuer's Name:

Grifols Diagnostic Solutions Inc.

**Issuer's Contact Information:** 

4560 Horton Street

Emeryville, CA 94608, USA

**Authorized Representative:** 

Diagnostic Grifols, S.A.

Passeig Fluvial, 24

08150 Parets del Vallès, Spain

### Object of the Declaration:

Catalog No.	Description
301120	Procleix® Auto Detect Reagents Kit (Procleix® and Procleix® Tigris
	Systems)
303345	Procleix® Auto Detect Reagents Kit (Procleix® Panther System)

Grifols Diagnostic Solutions Inc. declares that the above mentioned object of the declaration meets the provision of the Council Directive 98/79/EC for the In Vitro Diagnostic Medical Devices and the IVDD Directive 98/79/EC as transposed in the national laws of the Member States.

The object of the declaration described above is in conformity with the requirements of the following standards:

Standard	Revision	Title
EN ISO 13485	2016	Medical devices-Quality management systems-Requirements for regulatory purposes
EN 13975	2003	Sampling procedures used for acceptance testing of <i>in vitro</i> diagnostic medical devices – Statistical aspects
EN ISO 14971	2012	Medical devices-Application of risk management to medical devices – Rationale for requirements
EN ISO 15223-1	2016	Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements
EN ISO 18113-1 and 18113-2	2011	In Vitro Diagnostic Medical Devices-Information Supplied by the Manufacturer. Part 1: Terms, definitions and general requirements. (Part 2: IVD Reagents for Professional Use)
ISO 23640	2015	In vitro diagnostic medical devices- Evaluation of stability of in vitro diagnostic reagents

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#### **Additional Information:**

Classification/

Conformity Assessment: Self-Certified, Annex III

Date of Initial CE mark: March 29, 2004
Date of Current CE Mark: May 15, 2019

Signed for and on behalf of: Grifols Diagnostic Solutions Inc.

Amanda Doe, Manager Regulatory Affairs