



Declaration of Conformity



Manufacturer Name SD Biosensor, Inc.

Manufacturer Address **Head Office**

C-4th&5th, 16, Deogyeong-daero, 1556beon-gil, Yeongtong-gu,

Suwon-si, Gyeonggi-do 16690, KOREA

Manufacturing Site

74, Osongsaengmyeong 4-ro, Osong-eup, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do 28161, KOREA

EC Representative Name MT Promedt Consulting GmbH

Altenhofstrasse 80 66386 St. Ingbert Germany **EC Representative Address**

Common Name Immunoassay Test Kit

Product Name STANDARDTM F Chikungunya IgM/IgG FIA

*Please refer to "Annex I. Product List" on page 2 in more detail.

FCHK01B Reference Number

Classification Others not covered by Annex II and self-testing according to

Directive 98/79/EC

Annex III of Directive 98/79/EC (EC Declaration of Conformity) **Conformity Assessment Route**

Applied Standards EN ISO 13485:2016 EN ISO 18113-1:2011

EN ISO 14971:2012 EN ISO 18113-2:2011 EN ISO 23640:2015 EN ISO 15223-1:2016 EN ISO 17511:2003 EN 62366:2008

EN 13612:2002

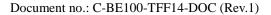
We herewith declare that the above mentioned products meet the provisions of the Council Directive 98/79/EC for in vitro diagnostic medical devices. All supporting documentations are retained under the premises of the manufacturer. SD Biosensor, Inc. is exclusively responsible for the declaration of conformity.

Place: Suwon-si, Republic of Korea

Valid from: October 7, 2019

Signature

Hvo-Keun, Lee CEO / President





Annex I. Product List

<u>FCHK01B</u> STANDARDTM F Chikungunya IgM/IgG FIA

- Test Device
- Assay diluent
- STANDARDTM Ezi tube+ (10ul)

EDMA	Code
15 70 9	0 90 0

Description of EDMA code Other Other Virology Rapid Tests