

EU Declaration of Conformity



Manufacturer (SRN)	SD Biosensor, Inc. (SRN : KR-MF-000009168)
Manufacturer Address	<u>Head Office</u> C-4th&5th, 16, Deogyong-daero, 1556beon-gil, Yeongtong-gu, Suwon-si, Gyeonggi-do 16690, REPUBLIC OF KOREA <u>Manufacturing Site</u> 14, Jeungpyeongsandan-ro, Jeungpyeong-eup, Jeungpyeong-gun, Chungcheongbuk-do 27915, REPUBLIC OF KOREA
EC Representative (SRN)	MT Promedt Consulting GmbH (SRN: DE-AR-000000085)
EC Representative Address	Ernst-Heckel-Straße 7 66386 St. Ingbert Germany
Notified Body	TÜV SÜD PRODUCT SERVICE GmbH
Notified Body Address	Ridlerstr. 65, 80339 München, Germany
Notified Body No.	0123
Certificate No.	EU QMS Certificate (IVDR): V13 075369 0072 Rev. 04
Product Name	STANDARD™M10 vanA/vanB
Reference Number	M10-VAN-01
Catalogue Number	11VAN10A
Basic UDI-DI	88001117IPCA01MA20XC
Classification (Applied Rule)	Class C (Rule 3 (c)) according to Annex VIII from Regulation (EU) IVDR 2017/746
Conformity Assessment Route	Annex IX (Chapter I, III) of Regulation (EU) 2017/746 (IVDR)
Common Specification	N/A

We herewith declare that the above mentioned products meet the provisions of the following Regulations/Directives:

- Regulation (EU) 2017/746 on in vitro diagnostic medical devices (IVDR)

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: May 28, 2026



(Signed for and on behalf of SD Biosensor, Inc.)

Signature

Name : Hyo-Keun, Lee
Position : CEO