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Tel. direct: Fax direct: E-mail :	+41 22 791 5891 +41 22 791 4836 diagnostics@who.int	SD Biosensor, Inc Attention: Mr Young-Gyun Kim C 4th and 5th, 16 Deogyeong-daero, 155
In reply please refer to:	CC/vl	beon-gil Suwon-si, Geonggi-do
Your reference:	P17-370-9	16690 République de Corée

11 June 2020

Dear Mr Kim,

Subject: WHO Prequalification of In Vitro Diagnostics – Final Public Report

Product name: STANDARD Q HIV 1/2 Ab 3-Line Test Product codes: 09HIV30D and 09HIV30DM Regulatory version: Rest-of-World regulatory version Manufacturer: SD Biosensor Inc PQDx Reference Number: PQDx 0383-117-00

We are pleased to inform you that the above-referenced product was prequalified on 10 June 2020 and listed on the World Health Organization (WHO) list of prequalified in vitro diagnostic products.

The following post-prequalification activities are required to maintain the prequalification status:

- 1. Notification to WHO of any planned changes to a prequalified product, in accordance with "*WHO procedure for changes to a WHO prequalified in vitro diagnostic*" (document number PQDx 121); and
- 2. Post-market surveillance activities, in accordance with "*WHO guidance on post*market surveillance of in vitro diagnostics" (ISBN 978 92 4 150921 3).

You are also required to submit an annual report that details sales data and all categories of complaints in a summarized form. There are certain categories of complaints and changes to the product that must be notified immediately to WHO, as per the above-mentioned documents. The sales data will serve as denominator data to guide the frequency of re-inspection.

Failure to comply with any of the above-mentioned post-prequalification requirements may lead to remedial action by WHO, including but not limited to, de-listing from the WHO list of prequalified in vitro diagnostic products.

If you have any questions, please do not hesitate to contact us by email (diagnostics@who.int) or by telephone (+4122 791 5891).

Yours sincerely,

Mr Deus Mubangizi Unit Head Prequalification Unit Regulation and Prequalification Department Access to Medicines and other Health Products Division