



Tel. direct: +41 22 791 5891  
Fax direct: +41 22 791 4836  
E-mail : diagnostics@who.int

In reply please  
refer to: CC/vl

Your reference: P17-370-9

SD Biosensor, Inc  
Attention: Mr Young-Gyun Kim  
C 4th and 5th, 16 Deogyong-daero, 155  
beon-gil  
Suwon-si, Geonggi-do  
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