THE GLOBAL FUND

Version 1
30-June-2025

List of eligible Quality Control Laboratories compliant with the Global Fund QA requirements for testing Medical Devices and PPE

Disclaimer

1. This document does not constitute an endorsement, or warranty of any laboratory's fitness for a particular purpose.
2. The Global Fund does not guarantee that the list(s) below are complete or error-free, nor that the laboratories meeting the standards will continue to do so. Laboratories that no longer meet the required standards will be removed from the list.
3. The list is not exhaustive. Principal Recipients may select laboratories not listed below, provided they comply with the respective Global Fund Quality Policies and Global Fund's PSM Guide.
4. This list is updated on a regular basis. Users should refer to the most recent version available on the website when conducting testing.

Section A: WHO-Prequalified Quality Control Laboratories

There are currently no WHO prequalified quality control laboratories for Medical Devices.

Section B: WHO-Prequalified and/or ISO 17025 certified Quality Control Laboratories that were found by the Global Fund to be capable of performing quality control testing as part of the monitoring activities outlined in the Global Fund's QA Policy

The Quality Control Laboratories listed below responded to a Request For Proposal (RFP) initiated by the Global Fund and were evaluated to meet its requirements. These laboratories are capable of performing quality control testing as part of the monitoring activities outlined in the Global Fund's QA Policy.

Country	Name of laboratory	Address	Contact person	E-Mail	Testing capabilities
Zimbabwe	Medicines Control Authority of Zimbabwe	106 Baines Avenue P. O. Box 10559 Harare, Zimbabwe	Tinashe Gono	tgono@mcaz.co.zw	- Male Condoms

The Quality Control Laboratories listed below responded to a Request For Proposal (RFP) initiated by the Global Fund and were evaluated to meet its requirements. These laboratories are capable of performing quality control testing as part of the monitoring activities outlined in the Global Fund's QA Policy.

Country	Name of laboratory	Address	Contact person	E-Mail	Testing capabilities
Uganda	National Drug Authority	Plot 93, Buganda Road, P.O. Box 23096 Kampala, Uganda	Naluyima Amoreen	namoreen@nda.or.ug	- Male Condoms - PPE (gloves only)
Singapore	TUV SUD PSB Pte Ltd	15 International Business Park TÜV SÜD @ IBP Singapore 609937	Sale team: - Loo Jia Jian - Gao Xiao fei Technical team: - Wong Bee Hui - James Huang - Shirley Tjoa	Jia-Jian.LOO@tuvsud.com XIAOFEI.GAO@tuvsud.com bee-hui.wong@tuvsud.com James.HUANG@tuvsud.com shirley.tjoa@tuvsud.com	- Medical Devices - Male and female Condoms - PPE (gloves and masks)

Section C: Other ISO 17025 Quality Control Laboratories

Global Fund Grant Recipients shall utilize any ISO 17025 accredited laboratory, provided that the necessary test methods are included within the laboratory's accreditation certificate. The Principal Recipient must verify the suitability and competence of the laboratory before utilizing it for Quality Control activities.

The following steps should be considered for verification:

1. Request the ISO 17025 or Good Laboratory Practice certificate of the laboratory in question.
2. Verify the authenticity of the ISO 17025 certificate on the website of the accreditation body that issued the certificate.
3. Check the following parameters:
    a. The Health Product(s) to be tested is listed in the scope.
    b. The test(s) to be performed is listed in the scope of accreditation/certification.
    c. Test(s) is conducted according to the methods and specifications approved by the relevant pharmacopeia, WHO Prequalification Programme, applicable standards and/or by the ERP.
    d. The ISO 17025 certificate is valid.