WHO Emergency Use Assessment Coronavirus disease (COVID-19) IVDs
PUBLIC REPORT

Product: Novel Coronavirus (2019-nCoV) IgG/IgM Test Kit (Colloidal gold)
EUL Number: EUL 0540-216-00
Outcome: Not Recommended.

The EUL process is intended to expedite the availability of in vitro diagnostics needed in public health emergency situations and to assist interested UN procurement agencies and Member States in determining the acceptability of using specific products in the context of a Public Health Emergency of International Concern (PHEIC), based on an essential set of available quality, safety and performance data. The EUL procedure includes the following:


Novel Coronavirus (2019-nCoV) IgG/IgM Test Kit (Colloidal gold) with product code 52026069, manufactured by Genrui Biotech Inc, 4-10F, Building 3, Geya Technology Park, Guangming District Shenzhen 518106, China is not eligible for WHO procurement.

Quality Management Systems Review

To establish the eligibility for WHO procurement, Genrui Biotech Inc was asked to provide up-to-date information about the status of their quality management system.

Based on the review of the submitted quality management system documentation by WHO staff and external technical experts (assessors), it was established that insufficient information was provided by Genrui Biotech Inc to fulfil the requirements described in the “Instructions for Submission Requirements: In vitro diagnostics (IVDs) Detecting SARS-CoV-2 Nucleic Acid, PQDx_ 347”. 