



CERTIFICATION

To Whom This May Concern:

This is to certify that the **Panbio COVID-19 Ag Rapid Test Device (NASAL)** manufactured by Abbott Rapid Diagnostics Jena GmbH – Orlaweg 1 07743 Jena, Germany has complied with all the requirements for the special certification of COVID-19 Diagnostic Kits. The product is listed on WHO Emergency Use Listing (EUL) - Product eligible for listing approved by the World Health Organization (WHO). With this approval, the company is required to indicate in the product label or in the accompanying product insert the following statement:

“This product is strictly for medical professional use only and not intended for personal use. The administration of the test and the interpretation of the results should be done by a trained health professional. The result of this test should not be the sole basis for the diagnosis; confirmatory testing is required”

This certification is issued upon the request of **SUNFU SOLUTIONS INC.** with business address at Unit 615 City and Land Megaplaza, ADB Ave., Ortigas Center, Pasig City for whatever legal purpose this may serve.

This certificate cannot be used for advertising purposes in whatever medium and neither can this certificate be construed as an endorsement by the Center for Device Regulation, Radiation Health, and Research.

Done this 4th February 2021 at Alabang, Muntinlupa City.

BY AUTHORITY OF THE DIRECTOR GENERAL


MARIA CECILIA C. MATIENZO
Director IV

Center for Device Regulation, Radiation Health, and Research

Not valid without FDA Seal

Landbank Seq No. : 12921233495
Amount : PhP 510.00
Date : 29 January 2021
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DTN: 20210129092338
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