



BAHAGIAN REGULATORI FARMASI NEGARA

National Pharmaceutical Regulatory Agency

KEMENTERIAN KESIHATAN MALAYSIA

Ministry of Health Malaysia

Lot 36, Jalan Prof. Diraja Ungku Aziz, 46200 Petaling Jaya, Selangor, Malaysia

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GMP Certificate No. 3251/23

Our Ref. : KKM/NPRA.PKP/600-2/5 (22) s/d 34.
Date : 31 October 2023

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

The National Pharmaceutical Regulatory Agency, Ministry of Health Malaysia hereby certifies that:

The Manufacturer : **Orient Laboratories Sdn. Bhd.**

Site Address : **No. 37, Jalan PS 3,
Taman Industri Prima Selayang,
68100 Batu Caves,
Selangor,
Malaysia.**

Has been subjected to a Good Manufacturing Practices (GMP) audit for the manufacture of traditional medicines and health supplements by officer from the National Pharmaceutical Regulatory Agency.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **10 May 2022**, it is certified that the manufacturing plant conforms to the requirement of GMP in accordance with the Guidelines on Good Manufacturing Practice for Traditional Medicines and Health Supplements, 1st Edition, 2008.

The manufacturer is authorised to manufacture the following registered traditional medicines and health supplements:

Traditional Medicines

1. Capsules (hard shell)
2. Capsules (soft shell)
3. Powders
4. Tablets
5. Liquids for internal use

Health Supplements

1. Capsules (hard shell)
2. Capsules (soft shell)
3. Tablets

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection, after which time the issuing authority should be consulted.

The authenticity of this certificate may be verified with the issuing authority.

(DR. NORAIIDA MOHAMAD ZAINOOR) RPh. 2289

Head of Centre for Compliance & Quality Control
National Pharmaceutical Regulatory Agency



Certified to ISO 9001:2015
Cert. No. QMS 00894



Member of
Pharmaceutical Inspection
Cooperation Scheme



Non Member Adherence to
Mutual Acceptance of
Data for GLP