



Bahagian Regulatori Farmasi Negara
National Pharmaceutical Regulatory Agency
KEMENTERIAN KESIHATAN MALAYSIA
MINISTRY OF HEALTH MALAYSIA

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GMP Certificate No. 1839/20

Our Ref. : KKM/NPRA.PKP/600-2/5 (14) JLD 15
Date : 9 October 2020

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part I

The National Pharmaceutical Regulatory Agency, Ministry of Health Malaysia confirms the following:

The Manufacturer : **SteriPack Asia Sdn. Bhd.**

Site Address : **Lot 119992,
Jalan Canang Emas 8, Telok Gong,
42000 Klang,
Selangor,
Malaysia.**

Has been inspected in accordance with Malaysian Control of Drugs and Cosmetics Regulations 1984 and Malaysian Drug Registration Guidance Document (DRGD).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **7 June 2018**, it is considered that it complies with the principles and guidelines of the current Pharmaceutical Inspection Co-Operation Scheme (PIC/S) GMP Guides.

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.

This certificate is valid only when presented with all pages and both Parts I and II.

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

(DR. ROSHAYATI MOHAMAD SANI) RPh. 1449
Head of Centre for Compliance & Quality Control
National Pharmaceutical Regulatory Agency

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Certified to ISO 9001: 2008
Cert. No. AR 2293



Member of
Pharmaceutical Inspection
Cooperation Scheme



Non Member Adherence to
Mutual Acceptance of
Data for GLP



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Part II

✓ Human Medicinal Products

1. MANUFACTURING OPERATIONS – MEDICINAL PRODUCTS*	
1.2	Non-sterile Products
	1.2.1 Non-sterile products (processing operations for the following dosage forms)
	1.2.1.1 Capsules, hard shell
	1.2.1.2 Capsules, soft shell
	1.2.1.13 Tablets
	1.2.2 Batch certification
1.4	Other products or processing activity
	1.4.1 Manufacture of:
	1.4.1.3 Others – Health Supplements (capsules, hard shell; capsules, soft shell; Tablets),
1.5	Packaging
	1.5.1 Primary packing
	1.5.1.1 Capsules, hard shell
	1.5.1.2 Capsules, soft shell
	1.5.1.13 Tablets
	1.5.2 Secondary packing
1.6	Quality Control Testing
	1.6.2 Microbiological: non-sterility
	1.6.3 Chemical/Physical

Any restrictions or clarifying remarks related to the scope of this certificate:

-None-

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(DR. ROSHAYATI MOHAMAD SANI) RPh. 1449
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