



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

36, Jalan Universiti, 46200 Petaling Jaya, Selangor, Malaysia

Tel : +603-78835400
Fax: +603-79562924 / 79581312
<http://npa.moh.gov.my>

GMP Certificate No. 1214/19

Our Ref. : KKM/NPRA.PKP/600-2/5 (160)

Date : 24 May 2019

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part I

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

The Manufacturer : **ExcelVite Sdn. Bhd.**
Site Address : **Lot 56442, 7 1/2 Mile,
Jalan Ipoh / Chemor,
31200 Chemor,
Perak,
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 **8 - 9 April 2019**, 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.

(NOR HAFIZAH MOHD POTRI) RPh. 3916
Senior Principal Assistant Director
Centre for Compliance and Licensing

Page 1 of 3



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

3. MANUFACTURING OPERATIONS – ACTIVE SUBSTANCES

3.2	Extraction of active substance from natural sources
3.2.1	Extraction of substance from palm oil source (esterification)
3.2.6	Purification of extracted substance (molecular distillation)
3.5	General Finishing Steps
3.5.1	Physical processing steps (blending and spray drying)
3.5.2	Primary packaging
3.5.3	Secondary packaging
3.6	Quality Control Testing
3.6.1	Physical/chemical testing
3.6.2	Microbiological testing (non-sterility testing)

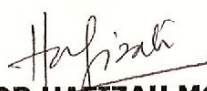
Active Substance(s):

Powders

- 1) EVTeneSol™ Range of Products
- 2) EVNoIMax™ Range of Products
- 3) EVNoIMax™ (T) Range of Products
- 4) EVRoI™ Range of Products
- 5) EVNoIBev™ 15%

Liquid (oil based):

- 1) EVNoI™ Range of Products
- 2) EVNoI™ 50%C (N) / EVNoI™ 50%C (L)
- 3) EVNoIEgg™ 20% / EVNoIEgg™ 20% (V)
- 4) EVNoI SupraBio™ Range of Products
- 5) EVNoI SupraBio™ 20% (N)
- 6) EVNoI SupraBio™ 20% (P)
- 7) EVTene™ Range of Products
- 8) EVOlein™
- 9) EVPene™ Range of Products
- 10) EVSpectra™ Range of Products
- 11) EVTene™ 3% (T) / EVTene Feed™ 3%
- 12) Palm Oil Carotene Interim Product 5%


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Page 2 of 3



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Co-Products:

- 1) EVOleo™ & EVFuel™
- 2) GlyceRaw™ 85%

Any restrictions or clarifying remarks related to the scope of this certificate: -None-

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Hafizah

(NOR HAFIZAH MOHD POTRI) RPh. 3916

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Page 3 of 3



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