



Health Sciences Authority
Republic of Singapore

ORIGINAL
CERTIFICATE NO.
GMPCT2400014Z

HEALTH PRODUCTS ACT 2007

GOOD MANUFACTURING PRACTICE CERTIFICATE

A certificate is hereby granted under the Health Products Act 2007 to
Novabay Pte Ltd

With manufacturing premises at
20 TEMBUSU AVENUE SINGAPORE 627536

With store at: See attached.

From the knowledge gained during the audit/conformity assessment performed by the Health Sciences Authority, Singapore from 10 June 2024 to 13 June 2024, it is considered that the manufacturer has maintained an overall acceptable level of compliance with the Pharmaceutical Inspection Convention/Co-operation Scheme (PIC/S) Guide to Good Manufacturing Practice (GMP) for Medicinal Products (Part II) relating to active pharmaceutical ingredients, which encompasses all the recommendations of the World Health Organization (WHO) in relation to GMP.

Active Ingredient(s) in respect of which the manufacturer is certified: See attached.

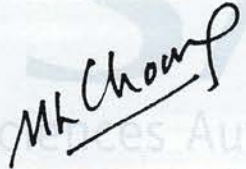
Any restrictions or clarifying remarks related to the scope of this certificate: See attached.

This certificate shall not be reproduced except in full. It remains the property of the Health Sciences Authority and must be returned upon written demand. The authenticity and validity of this certificate may be verified by contacting the certifying Authority stated on this certificate.

This certificate takes effect from 10 June 2024, and unless revoked shall expire on 09 June 2027.

Date of Issue : 14 November 2024
Application : 2441521A

AUDIT AND LICENSING DIVISION
Health Products Regulation Group
Health Sciences Authority
11 Biopolis Way
#11-01 Helios, Singapore 138667
Email: HSA_Certification@hsa.gov.sg


Dr CHOONG May Ling, Mimi
for Licensing Authority

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ACTIVE INGREDIENTS IN RESPECT OF WHICH THE MANUFACTURER IS CERTIFIED:

Manufacture

Active Ingredients

1. Non-Sterile Chemical Active Ingredients

Manufacturing Processes

1. Non-Sterile Active Ingredients
 - 1.1. Chemical Synthesis
 - 1.1.1. Dissolution
 - 1.1.2. Crystallisation
 - 1.2. Physical Processing Steps
 - 1.2.1. Drying
 - 1.2.2. Sieving, Filtration, Centrifugation

Quality Control Testing

1. Chemical/Physical
2. Microbiological

To conduct primary packaging of the following:

1. Chemical Active Ingredients

To conduct secondary packaging of the following:

1. Chemical Active Ingredients

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List of Active Ingredients

1. Sodium Bicarbonate

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

1. This certification applies to the manufacture, including the receipt of materials, production, packaging, labelling, quality control, release, storage and distribution of the above-mentioned active pharmaceutical ingredient(s) for use in human drug (medicinal) products.
2. This certification is a statement of the manufacturing quality system standard of the factory and is not an endorsement of the quality of the active ingredients manufactured.

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