ORIGINAL
LICENCE NO
IMCD2000158

Health Sciences Authority
Republic of Singapore

THE MISUSE OF DRUGS ACT
THE MISUSE OF DRUGS REGULATIONS

LICENCE TO IMPORT CONTROLLED DRUGS

Distribution:
Original - To Importer
Duplicate (Copy 1) - To Exporter
Duplicate (Copy 2) - To Competent Authority of Exporting Country
Duplicate (Copy 3) - To Immigration & Checkpoints Authority (ICA), Republic of Singapore
Duplicate (Copy 4) - To Health Products Regulation Group, Health Sciences Authority, Republic of Singapore

Name and Address of Importer	Name and Address of Exporter
KELLY PNG JIA XING GOVIN HOLDINGS PTE LTD 21 BUKIT BATOK CRESCENT #05-74 WCEGA TOWER SINGAPORE 658065	M/S VERVE HUMAN CARE LABORATORIES A-43 OFFICE NO.102-103 G.T.KARNAL ROAD INDUSTRIAL AREA DELHI, INDIA, 110033

Name and Quantity of Substance or Preparation, and if applicable, Pharmaceutical Dosage Form and Strengths, to be imported:

VERMOR-10 MORPHINE SULPHATE INJECTION BP 10mg/ml. Each 1 ml ampoule contains 10 mg of Morphine Sulphate.
Quantity: 1050 ampoules

Total Base Drug Content:

Total Morphine Base = 7.875 gram(g)

Purpose of Import: For Local Consumption	Method of Import: AIR

Remarks: The consignment shall be delivered to the warehouse address: 10 PIONEER CRESCENT 01-00, SINGAPORE LOGISTICS HUB, SINGAPORE 628566 c/o KUEHNE + NAGEL PTE LTD (UEN:199400013D)

The above-mentioned Importer is hereby licensed to import the substance(s) specified in this Licence from the Exporter named. This Licence is issued subject to the Conditions stipulated in this licence.

This Licence takes effect from 07 July 2020 and, unless revoked, shall expire on 06 January 2021.

Date of Issue : 07 July 2020
Fee : $ 103
Application Number : 2075781M

All persons issued with a Licence to Import Controlled Drugs under the Misuse of Drugs Act (MDA) must comply with the MDA and their regulations. This is to ensure that all health products in Singapore meet the required standards of safety, quality and efficacy. Licensees must also comply with all other applicable laws and their regulations.

For medicinal or health products which have not been registered or licensed, HSA has not assessed their safety, quality and efficacy.

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

Group Director
Health Products Regulation Group
Health Sciences Authority

ORIGINAL
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CONDITIONS

1 The import shall be made in ONE consignment before the expiry of this licence.

2 The consignment shall not be imported through the post.

3 This Licence is not a Licence to be in possession of or to supply the drugs imported.

4 This Licence does not relieve the importer from compliance with any Customs regulations in force for the time being relating

5 This Licence is only valid for the importer and may be revoked at any time by the licensing authority, to whom it shall in that

6 This Licence unless sooner revoked shall be produced to the Customs Officer at the time of importation for verification and

7 If the importation of the drugs specified in this Licence is not effected before the expiry date of this Licence, the Licence

8 This Licence is not to leave the possession of the importer until it is surrendered to the licensing authority, or the Customs

9 The copy of the Export Licence, if any which accompanies the drugs shall be forwarded to the licensing authority,

10 The importer shall notify the licensing authority within seven (7) days of import of the consignment of controlled drugs stated

11 The importer shall advise the licensing authority in writing at the end of each quarter of every import made detailing in

12 The importer shall submit yearly returns of stocks of controlled drugs held as at 31 December of each year to the licensing

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ENDORSEMENT BY IMMIGRATION & CHECKPOINTS AUTHORITY (ICA) OFFICER AT THE TIME OF IMPORTATION

Date	Description of Drugs Imported	Number and Date of Export Licence	Quantity	Mode of Importation	Cargo Clearance Permit No	Signature, mark and station of ICA Officer

This licence, when all drugs to which it relates have been imported, must be returned by the ICA Officer to the Deputy Group Director, Health Products Regulation Group, Health Sciences Authority.

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Republic of Singapore

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