Health Sciences Authority

Licence Summary for Therapeutic Products - Importer's Licence

1. Licence/Permit/Certificate/Listing/Notification Summary	
1.1 Licence/Permit/Certificate/Listing Notification Type :	Therapeutic Products - Importer's Licence
1.2 Licence/Permit/Certificate/Listing Notification No :	IMTPT1600059
1.3 Company Name :	HEALTHCARE ASIA PTE. LTD.
1.4 Approved Date :	01/11/2016
1.5 Expiry Date :	31/10/2018
1.6 Licence Type :	Limited Scope - Term (Annual)

2. Licence Information	
2.1 Approved Business premise :	3025, Ubi Road 3, #04-117, SINGAPORE408653
2.2 Approved warehouse(s) (Storage Condition):	1) 3025 UBI ROAD 3 #04-117 KAMPONG UBI INDUSTRIAL ESTATE, SINGAPORE 408653 (Cold Chain, Non-Cold Chain)
2.3 Aspect of Importation Activity :	1) Therapeutic products solely for export only
2.4 Product(s) approved to be imported	
NIL	
2.5 Responsible Person(s) in this Licence(Defined Scope as in application) :	1) Goh Chun Ping

All persons issued with a Importer's Licence under the Health Products Act (HPA) must comply with the HPA 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 health products which have not been registered or licensed, HSA has not assessed their safety, quality and efficacy.

3. The Licence/Permit/Certificate/Listing/Notification is issued subject to the following condition	
3.1	The licence is restricted to the importation of therapeutic products for the scope of activities approved in the licence only.
3.2	The importer shall obtain prior approval from the Authority for each consignment of therapeutic product that contains a psychotropic substance to be imported.
3.3	For unregistered therapeutic products that are imported solely for the purpose of re-exportation out of Singapore, they are subject to the applicable provisions under the Therapeutic Products Regulations and the following conditions: (a) Unless otherwise approved by the Authority, the unregistered therapeutic product(s) shall not be supplied for use in Singapore. (b) Unless otherwise approved by the Authority, the importer shall ensure that the unregistered therapeutic product(s) be completely re-exported whether by the importer directly or through another licensed wholesaler, within a period of 2 years from the date on which the therapeutic product(s) is imported. (c) Unregistered therapeutic product(s) imported for re-export shall be supplied to the recipient for the purpose of his/her trade, business or profession only and not be supplied to individual end users for personal consumption.

3.4	The granting of licence fee exemption is contingent on your company's declaration to be dealing in any one of, or any combination of the following activities (i) unregistered therapeutic products importation solely for the purpose of re-exportation out of Singapore; and/or ii) unregistered therapeutic products importation for direct supply to vessels or aircraft, and/or (iii) importation for non-clinical purposes; after the implementation date of Nov 2016 for a period of 3 years. By the end of this 3-year period from the date on which the Therapeutic Products Regulations 2016 comes into force, the licensee shall ensure that the quality system meets the Good Distribution Practice standard as stipulated by the Authority. The continued validity of the licence shall be subject to the conditions determined as such by the Authority.

The above information is current as of the date of print : 10/09/2017

The validity of the information on this document may be verified with the licensing authority stated on this document

This is a computer generated document, no signature is required.

Audit & Licensing Division

Health Products Regulation Group

Health Sciences Authority

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