[reg. 29(6)]
[第 29(6) 條]

PHARMACY AND POISONS ORDINANCE
藥劑業及毒藥條例
(Chapter 138)
(第 138 章)
CERTIFICATE OF PHARMACEUTICAL PRODUCT
藥劑製品證明書

Name and dosage form of product (specify strength) :
製品的名稱及劑型 (指明劑量) :

.....

.....

.....

.....

Name and amount of each active ingredient (as provided by manufacturer) :
每種有效成分的名稱及分量 (按製造商所提供的資料) :

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.....

Manufacturer, and/or when applicable, the person responsible for placing the product on the market :
製造商及／或 (如適用) 負責將該製品推出市場出售的人 :

.....

.....

.....

Address(es):
地址 :

.....

.....

.....

It is certified that —
現證明 —

(*a*) this product has been registered with the Pharmacy and Poisons Board;
該製品已向藥劑業及毒藥管理局註冊 ;

(*b*) this product has been authorised to be placed on the market for use in Hong Kong —
該製品已獲准推出市場出售以供在香港使用 —

Number of permit: .....
許可證編號 :

Date of issue: .....
發出日期 :

(*c*) the manufacturing plant in which the product is produced is subject to inspection at suitable intervals.
製造該製品的製造廠每隔適當的期間即受到檢查。

This certificate is valid for one year from the date of issue.
本證明書自簽發日期起計有效期為一年。

HONG KONG
香 港

..... *(Date)*
日期 :

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for *Pharmacy and Poisons Board*
藥劑業及毒藥管理局 代行