

PHARMACY AND POISONS ORDINANCE

RESOLUTION

(Under section 29 of the Pharmacy and Poisons Ordinance (Cap. 138))

RESOLVED that the following Regulations, made by the Pharmacy and Poisons Board on 21 June 2010, be approved –

- (a) the Pharmacy and Poisons (Amendment) (No. 3) Regulation 2010; and
- (b) the Poisons List (Amendment) (No. 3) Regulation 2010.

PHARMACY AND POISONS (AMENDMENT) (NO. 3) REGULATION 2010

(Made by the Pharmacy and Poisons Board under section 29 of the
Pharmacy and Poisons Ordinance (Cap. 138) subject to the approval of the
Legislative Council)

1. First Schedule amended

The First Schedule to the Pharmacy and Poisons Regulations (Cap. 138 sub.
leg. A) is amended, in Division A –

- (a) by adding “Agomelatine; its salts”;
- (b) by adding “Galsulfase”;
- (c) by adding “Golimumab”;
- (d) by adding “Nebivolol; its salts”;
- (e) by adding “Saxagliptin; its salts”;
- (f) by adding “Tolvaptan”.

2. Third Schedule amended

The Third Schedule is amended, in Division A –

- (a) by adding “Agomelatine; its salts”;
- (b) by adding “Galsulfase”;
- (c) by adding “Golimumab”;
- (d) by adding “Nebivolol; its salts”;
- (e) by adding “Saxagliptin; its salts”;
- (f) by adding “Tolvaptan”.

Chairman,
Pharmacy and Poisons Board

21 June 2010

Explanatory Note

This Regulation adds 6 substances to Divisions A of the First and Third Schedules to the Pharmacy and Poisons Regulations (Cap. 138 sub. leg. A) (“principal Regulations”) respectively so that the sale, supply, labelling and storage of those substances are subject to the restrictions imposed under the Pharmacy and Poisons Ordinance (Cap. 138) and the principal Regulations.

POISONS LIST (AMENDMENT) (NO. 3) REGULATION 2010

(Made by the Pharmacy and Poisons Board under section 29 of the
Pharmacy and Poisons Ordinance (Cap. 138) subject to the approval of the
Legislative Council)

1. The Poisons List

The Schedule to the Poisons List Regulations (Cap. 138 sub. leg. B) is amended, in Part I, in Division A –

- (a) by adding “Agomelatine; its salts”;
- (b) by adding “Galsulfase”;
- (c) by adding “Golimumab”;
- (d) by adding “Nebivolol; its salts”;
- (e) by adding “Saxagliptin; its salts”;
- (f) by adding “Tolvaptan”.

Chairman,
Pharmacy and Poisons Board

21 June 2010

Explanatory Note

This Regulation adds 6 substances to Division A of Part I of the Poisons List set out in the Schedule to the Poisons List Regulations (Cap. 138 sub. leg. B) so that, among other applicable requirements, poisons containing those substances can only be sold on registered premises of an authorized seller of

poisons by a registered pharmacist or in the pharmacist's presence and under the pharmacist's supervision.