

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 14 February 2014, be approved—

- (a) the Pharmacy and Poisons (Amendment) Regulation 2014; and
- (b) the Poisons List (Amendment) Regulation 2014.

Pharmacy and Poisons (Amendment) Regulation 2014

(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. Pharmacy and Poisons Regulations amended

The Pharmacy and Poisons Regulations (Cap. 138 sub. leg. A) are amended as set out in sections 2 and 3.

2. First Schedule amended (substances to which certain restrictions with respect to the sale, supply, labelling and storage apply under regulations 3, 5, 6, 15, 19, 22, 23 and 24)

- (1) First Schedule, Chinese text, Division A, item “洋地黃的甙類；洋地黃的其他有效成份”—

Repeal

“成份”

Substitute

“成分”.

- (2) First Schedule, Division A—

- (a) After item “Aminoglutethimide”—

Add

“5-Aminolevulinic acid; its salts; its derivatives; their salts”;

- (b) After item “Clozapine; its salts”—

Add

“Cobicistat; its salts”;

- (c) After item “Eltrombopag; its salts; its esters; their salts”—

Add

“Elvitegravir; its salts”;

- (d) After item “Lithium Sulphate”—

Add

“Lixisenatide”;

- (e) After item “Midodrine; its salts”—

Add

“Mifepristone; its salts; its esters; their salts”;

- (f) After item “Pentolinium; its salts”—

Add

“Perampanel”;

- (g) After item “Perindoprilat; its salts; its esters; their salts”—

Add

“Pertuzumab”.

3. Third Schedule amended (substances required by regulation 9 to be sold by retail only upon a prescription given by a registered medical practitioner, registered dentist or registered veterinary surgeon)

- (1) Third Schedule, Division A—

Repeal

“DIVISION A”.

- (2) Third Schedule, after “VETERINARY SURGEON”—

Add

“Division A”.

- (3) Third Schedule, Division A—

- (a) After item “Aminoglutethimide”—
Add
“5-Aminolevulinic acid; its salts; its derivatives; their salts”;
- (b) After item “Clozapine; its salts”—
Add
“Cobicistat; its salts”;
- (c) After item “Eltrombopag; its salts; its esters; their salts”—
Add
“Elvitegravir; its salts”;
- (d) After item “Lithium Sulphate”—
Add
“Lixisenatide”;
- (e) After item “Midodrine; its salts”—
Add
“Mifepristone; its salts; its esters; their salts”;
- (f) After item “Pentolinium; its salts”—
Add
“Perampanel”;
- (g) After item “Perindoprilat; its salts; its esters; their salts”—
Add
“Pertuzumab”.



Chairman,
Pharmacy and Poisons Board

14 February 2014

Explanatory Note

This Regulation—

- (a) adds 7 substances to Division A of the First Schedule and Division A of the Third Schedule to the Pharmacy and Poisons Regulations (Cap. 138 sub. leg. A) (*principal Regulations*) 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; and
- (b) makes technical amendments to the 2 Schedules.

Poisons List (Amendment) Regulation 2014

(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. Poisons List Regulations amended

The Poisons List Regulations (Cap. 138 sub. leg. B) are amended as set out in section 2.

2. Schedule amended (the Poisons List)

The Schedule, Part I, Division A—

- (a) After item “Aminogluthimide”—

Add

“5-Aminolevulinic acid; its salts; its derivatives; their salts”;

- (b) After item “Clozapine; its salts”—

Add

“Cobicistat; its salts”;

- (c) After item “Eltrombopag; its salts; its esters; their salts”—

Add

“Elvitegravir; its salts”;

- (d) After item “Lithium Sulphate”—

Add

“Lixisenatide”;

- (e) After item “Midodrine; its salts”—

Add

“Mifepristone; its salts; its esters; their salts”;

- (f) After item “Pentolinium; its salts”—

Add

“Perampanel”;

- (g) After item “Perindoprilat; its salts; its esters; their salts”—

Add

“Pertuzumab”.



Chairman,
Pharmacy and Poisons Board

14 February 2014

Explanatory Note

This Regulation adds 7 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.