Pharmacy and Poisons (Amendment) (No. 2) 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)

First Schedule, Division A—

(a) After item "Dalteparin; its salts"—

Add

"Dapagliflozin; its salts";

(b) After item "Recombinant human erythropoietin"—

Add

"Regorafenib; its salts";

(c) After item "Todralazine; its salts"—

Add

"Tofacitinib; its salts";

(d) After item "Vigabatrin"—

Add

"Vilanterol; its salts".

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)

Third Schedule, Division A—

(a) After item "Dalteparin; its salts"—

Add

"Dapagliflozin; its salts";

(b) After item "Recombinant human erythropoietin"—

Add

"Regorafenib; its salts";

(c) After item "Todralazine; its salts"—

Add

"Tofacitinib; its salts";

(d) After item "Vigabatrin"—

Add

"Vilanterol; its salts".

Chairman,
Pharmacy and Poisons Board

16 June 2014

Appendix II

Explanatory Note

This Regulation adds 4 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.