

**L.N. 51 of 2016**

**Pharmacy and Poisons (Amendment) (No. 2) Regulation  
2016**

(Made by the Pharmacy and Poisons Board under section 29(1B) of the Pharmacy and Poisons Ordinance (Cap. 138) subject to the approval of the Secretary for Food and Health)

**1. Pharmacy and Poisons Regulations amended**

The Pharmacy and Poisons Regulations (Cap. 138 sub. leg. A) are amended as set out in sections 2, 3 and 4.

**2. Schedule 1 amended (substances to which certain restrictions with respect to the sale, supply, labelling and storage apply under regulations 3, 5, 6, 22 and 24)**

- (1) Schedule 1, Division A, after item “Acitretin; its salts; its esters”—

**Add**

“Aclidinium; its salts”.

- (2) Schedule 1, Division A, after item “Asenapine; its salts; its isomers”—

**Add**

“Asunaprevir; its salts”.

- (3) Schedule 1, Division A, after item “Dacarbazine”—

**Add**

“Daclatasvir; its salts”.

- (4) Schedule 1, Division A, after item “Everolimus; its salts; its esters; their salts”—

**Add**

“Evolocumab”.

- (5) Schedule 1, Division A, after item “Lenalidomide; its salts”—

**Add**

“Lenvatinib; its salts”.

- (6) Schedule 1, Division A, after item “Sildenafil; its salts; any compound containing the chemical structure of 5-(2-ethoxyphenyl)-1-methyl-3-propyl-1*H*-pyrazolo[4,3-*d*]pyrimidin-7(6*H*)-one substituted to any degree or without substitution; its salts”—

**Add**

“Siltuximab”.

**3. Schedule 3 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) Schedule 3, Division A, after item “Acitretin; its salts; its esters”—

**Add**

“Acridinium; its salts”.

- (2) Schedule 3, Division A, after item “Asenapine; its salts; its isomers”—

**Add**

“Asunaprevir; its salts”.

- (3) Schedule 3, Division A, after item “Dacarbazine”—

**Add**

“Daclatasvir; its salts”.

- (4) Schedule 3, Division A, after item “Everolimus; its salts; its esters; their salts”—

**Add**

“Evolocumab”.

- (5) Schedule 3, Division A, after item “Lenalidomide; its salts”—

**Add**

“Lenvatinib; its salts”.

- (6) Schedule 3, Division A, after item “Sildenafil; its salts; any compound containing the chemical structure of 5-(2-ethoxyphenyl)-1-methyl-3-propyl-1*H*-pyrazolo[4,3-*d*]pyrimidin-7(6*H*)-one substituted to any degree or without substitution; its salts”—

**Add**

“Siltuximab”.

**4. Schedule 10 amended (Poisons List)**

- (1) Schedule 10, section 2, Table, Part 1, Division A, after item “Acitretin; its salts; its esters”—

**Add**

“Acridinium; its salts”.

- (2) Schedule 10, section 2, Table, Part 1, Division A, after item “Asenapine; its salts; its isomers”—

**Add**

“Asunaprevir; its salts”.

- (3) Schedule 10, section 2, Table, Part 1, Division A, after item “Dacarbazine”—

**Add**

“Daclatasvir; its salts”.

- (4) Schedule 10, section 2, Table, Part 1, Division A, after item “Everolimus; its salts; its esters; their salts”—

**Add**

“Evolocumab”.

- (5) Schedule 10, section 2, Table, Part 1, Division A, after item “Lenalidomide; its salts”—

**Add**

“Lenvatinib; its salts”.

- (6) Schedule 10, section 2, Table, Part 1, Division A, after item “Sildenafil; its salts; any compound containing the chemical structure of 5-(2-ethoxyphenyl)-1-methyl-3-propyl-1*H*-pyrazolo[4,3-*d*]pyrimidin-7(6*H*)-one substituted to any degree or without substitution; its salts”—

**Add**

“Siltuximab”.

Dr. Constance CHAN  
Chairman,  
Pharmacy and Poisons Board

25 April 2016

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### Explanatory Note

This Regulation—

- (a) adds 6 substances to Division A of Schedule 1 and Division A of Schedule 3 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) adds 6 substances to Division A of Part 1 of the Poisons List set out in Schedule 10 to the principal Regulations so that, among other applicable requirements, those substances can only be sold on registered premises of an authorized seller of poisons by a registered pharmacist or in the presence and under the supervision of a registered pharmacist.