

---

First published in the Government *Gazette*, Electronic Edition, on 31st October 2016 at 5:00 pm.

---

**No. S 555**

**POISONS ACT  
(CHAPTER 234)**

**POISONS (AMENDMENT NO. 2) RULES 2016**

In exercise of the powers conferred by section 20(1) of the Poisons Act, the Minister for Health makes the following Rules:

**Citation and commencement**

1. These Rules are the Poisons (Amendment No. 2) Rules 2016 and come into operation on 1 November 2016.

**Amendment of rule 2**

2. Rule 2(1) of the Poisons Rules (R 1) is amended —

- (a) by deleting the definition of “codeine cough preparation”;
- (b) by inserting, immediately after the definition of “licensed person”, the following definition:

““licensed retail pharmacy” means premises specified in a pharmacy licence issued under the Health Products (Licensing of Retail Pharmacies) Regulations 2016 (G.N. No. S 330/2016);”;

- (c) by deleting the definition of “registered pharmacy”; and
- (d) by deleting the full-stop at the end of the definition of “sell by retail” and substituting a semi-colon, and by inserting immediately thereafter the following definition:

““therapeutic product” means a health product under the Health Products Act which is categorised as a therapeutic product in the First Schedule to that Act.”.

---

---

**Deletion and substitution of rule 14**

3. Rule 14 of the Poisons Rules is deleted and the following rule substituted therefor:

**“Complete exemption of persons dealing with certain poisons**

**14.** Nothing in the Act or these Rules applies to —

- (a) a person who deals with or possesses —
  - (i) any article specified in Group I of the Second Schedule; or
  - (ii) any poison specified in the first column of Group II of the Second Schedule when contained in or consisting of the article or substance specified opposite that poison in the second column;
- (b) a public officer or an officer of the Authority who imports any poison on account of the Government in the course of that officer’s duty; or
- (c) a holder of a manufacturer’s licence for a therapeutic product under the Health Products Act (Cap. 122D) who imports a substance specified in the First Schedule if that substance is required for the manufacture of the therapeutic product.”.

**Amendment of rule 14A**

4. Rule 14A of the Poisons Rules is amended by deleting the words “registered pharmacy” in paragraphs (1) and (2) and substituting in each case the words “licensed retail pharmacy”.

**Deletion of rule 17**

5. Rule 17 of the Poisons Rules is deleted.