CHAPTER 227

PHARMACY

PHARMACY (IMPORT AND EXPORT) REGULATIONS

S.I. 6/2010

(SECTION 48)

[Commencement 1st February, 2010]

Citation.

1. These Regulations may be cited as the Pharmacy (Import and Export) Regulations.

Interpretation. Ch. 227.

2. In these Regulations, "the Act" means the Pharmacy Act.

Import and export of drugs.

3. (1) Any person who desires to import or export any drugs for the purposes of the Act shall submit to the Council for approval the following supporting documentation —

(a) the name of the country and company of the original manufacturer of the drugs;

(b) certification that at the time the drugs left its country of original manufacturer, such drugs met the requisite requirements in accordance with section 27 of the Act;

(c) the name of the country and company of repackaging and transshipment;

(d) a list of all the active ingredients contained in the drugs;

(e) the categorisation and description of the drugs;

(f) the international generic nomenclature and all the applicable brand names used by the manufacturing company;

(g) the lot, batch number and expiration date of the drugs; and

(h) documentation specifying the proper storage and handling requirements.

(2) Evidence of approval granted under paragraph (1) shall be produced to the customs officer upon import or export of such drugs.

Record keeping.

4. Any person who imports or exports drugs for the purposes of the Act shall keep proper records specifying the information required under regulation 3.

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