



# Medicines Act 1968

## 1968 CHAPTER 67

### PART III

#### FURTHER PROVISIONS RELATING TO DEALINGS WITH MEDICINAL PRODUCTS

##### *Additional provisions*

#### **66 Further powers to regulate dealings with medicinal products**

- (1) The appropriate Ministers may by regulations prescribe such requirements as they may consider necessary or expedient with respect to any of the following matters, that is to say—
- (a) the manner in which, or persons under whose supervision, medicinal products may be prepared or may be dispensed;
  - (b) the amount of space to be provided in any premises for persons preparing or dispensing medicinal products, the separation of any such space from the remainder of the premises, and the facilities to be provided in any premises for such persons ;
  - (c) the amount of space to be provided in any premises for the sale or supply of medicinal products ;
  - (d) the accommodation (including the amount of space) to be provided in any premises for members of the public to whom medicinal products are sold or supplied or for whom medicinal products are being prepared or assembled;
  - (e) the amount of space to be provided in any premises for the storage of medicinal products ;
  - (f) the safekeeping of medicinal products;
  - (g) the disposal of medicinal products which have become unusable or otherwise unwanted ;
  - (h) precautions to be observed before medicinal products are sold or supplied;
  - (i) the keeping of records relating to the sale or supply of medicinal products;
  - (j) the supply of medicinal products distributed as samples;

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*Status: This is the original version (as it was originally enacted).*

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- (k) sanitation, cleanliness, temperature, humidity or other factors relating to the risks of deterioration or contamination in connection with the manufacture, storage, transportation, sale or supply of medicinal products;
  - (l) the construction, location and use of automatic machines for the sale of medicinal products.
- (2) Without prejudice to the generality of the preceding subsection, regulations made under subsection (1) of this section may prescribe requirements in respect of—
- (a) the construction, lay-out, drainage, equipment, maintenance, ventilation, lighting and water supply of premises at or from which medicinal products are manufactured, stored, transported, sold or supplied;
  - (b) the disposal of refuse at or from any such premises; and
  - (c) any apparatus, equipment, furnishings or utensils used at any such premises.