STATUTORY INSTRUMENTS

1997 No. 1830

The Prescription Only Medicines (Human Use) Order 1997

Medicinal products on prescription only

3. Subject to article 6, the following descriptions and classes of medicinal products are specified for the purposes of section 58, namely–

(a) medicinal products consisting of or containing a substance listed in column 1 of Schedule 1;

(b) medicinal products that are controlled drugs;

(c) medicinal products that are for parenteral administration, other than preparations of insulin for parenteral administration;

(d) cyanogenetic substances, other than preparations for external use;

(e) medicinal products that on administration emit radiation, or contain or generate any substance which emits radiation, in order that radiation may be used;

(f) medicinal products for human use which are classified as subject to medical prescription in marketing authorizations granted under Council Regulation 2309/93(**1**);

(g) medicinal products–

(i) which are not of a description and do not fall within a class specified in subparagraphs (a) to (f),

(ii) which are of a description in respect of which the conditions specified in section 59(1) are satisfied, and

(iii) in respect of which a product licence or marketing authorization has been granted which contains a provision to the effect that the method of sale or supply of the medicinal product is to be only in accordance with a prescription given by an appropriate practitioner.

(**1**) OJ No. L214, 24.8.93, p. 1.