STATUTORY RULES OF NORTHERN IRELAND

2001 No. 422

Biocidal Products Regulations (Northern Ireland) 2001

Part V

Packaging, Labelling and Advertisements

Labelling

31.—(1) A person shall not place on the market a biocidal product whose label—

(a) is misleading or gives an exaggerated impression of the authorised biocidal product; or

(b) contains, in relation to the authorised biocidal product, the descriptions "low-risk biocidal product", "non-toxic" or "harmless", or similar descriptions.

(2) Subject to paragraph (3), a person shall not place on the market an authorised biocidal product unless—

(a) that authorised biocidal product is labelled clearly and indelibly with the information specified in Schedule 9; and

(b) that information is in English, whether or not it is also in any other language.

(3) Subject to paragraph (4), the information referred to in sub-paragraphs 3, 5 to 12 and 14 of Schedule 9 may be given on the packaging of the authorised biocidal product or in an accompanying leaflet integral to the packaging of that authorised biocidal product.

(4) Where any information referred to in paragraph (3) is given in an accompanying leaflet, the authorised biocidal product shall be labelled clearly and indelibly with the words "Read attached instructions before use.".

(5) In this regulation, "authorised biocidal product" has the same meaning as it has in regulation 30.