STATUTORY RULES OF NORTHERN IRELAND

2001 No. 422

HEALTH AND SAFETY

Biocidal Products Regulations (Northern Ireland) 2001

Made - - - - 5th December 2001

Coming into operation 16th January 2002

BIOCIDAL PRODUCTS REGULATIONS (NORTHERN IRELAND) 2001

Part I

General

Part II

Active Substances

Part III

Biocidal Products

4. The Executive has determined— (a) the physical and chemical properties...

SCHEDULE 4 **Information to be contained in a dossier submitted in support of an application for the registration of a biocidal product**

1. The name and address of the applicant.
2. The name and address of the manufacturer of the biocidal...
3. The name and address of the manufacturer of the active...
4. The trade name of the biocidal product.
5. The name of each substance in the biocidal product, including...
6. The physical and chemical properties of the biocidal product relating...
7. The product-type and field of use of the biocidal product....
8. The intended category of users.
9. The intended method of use.
10. Efficacy data.
11. Analytical methods.
12. The classification, packaging and labelling of the biocidal product, including...
13. Where the biocidal product is a substance or preparation dangerous...

SCHEDULE 5 **Matters in respect of which additional conditions may be imposed on the mutual recognition of an authorisation or a registration of a biocidal product**

1. Directions for use of the biocidal product in question, including...
2. Particulars of any likely direct or indirect adverse side effects...
3. Directions for safe disposal of the biocidal product in question...
4. The period of time needed for the biocidal effect.
5. The interval to be observed between— (a) applications of the...
6. Particulars for adequate cleaning of equipment.
7. Particulars concerning precautionary measures during use, storage and transport, such...
8. Information on any specific dangers to the environment, including protection...

SCHEDULE 6 **Non-confidential information**

1. The name and address of the applicant for the authorisation...
2. The name of the biocidal product.
3. The name and address of the manufacturer of the biocidal...
4. The name and address of the manufacturer of the active...
5. The name and content of the active substance in the...
6. The name of any other substance in the biocidal product...
7. Physical and chemical data concerning the biocidal product and the...
8. Any ways of rendering harmless the biocidal product and the...
9. A summary of the results of the tests, referred to...
10. Recommended methods and precautions to reduce dangers from handling, storage,...
11. Safety data sheets.
12. Methods of analysis necessary to enable the Executive to make...
13. Methods of disposal of the biocidal product and its packaging....
14. Procedures to be followed and measures to be taken in...
15. First aid and medical advice to be given in the...

SCHEDULE 7 **Information relating to Biocidal products to be given to the Commission and to the competent authorities**

1. The name of the applicant for, or the person to...