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STATUTORY INSTRUMENTS

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**2001 No. 880**

**HEALTH AND SAFETY**

**The Biocidal Products Regulations 2001**

*Made* - - - - - *7th March 2001*  
*Laid before Parliament* *16th March 2001*  
*Coming into force* *6th April 2001*

ARRANGEMENT OF REGULATIONS

PART I

General

1. Citation and commencement
2. Interpretation
3. Application

PART II

Active Substances

4. Placing on the market of active substances
5. Applications concerning new active substances
6. Assessment of applications concerning new active substances
7. Applications for variation or renewal of the inclusion of active substances in Annex I, IA or IB

PART III

Biocidal Products

8. Prohibitions
9. Authorisation of a biocidal product
10. Registration of a low-risk biocidal product
11. Mutual recognition of authorisations
12. Mutual recognition of registrations
13. Provisional authorisation
14. Provisional registration
15. Emergency authorisation
16. Research and development

17. Experimental authorisation
18. Frame-formulations
19. Revocation of authorisations and registrations
20. Modification and review of authorisations and registrations
21. Notification of new information
22. Emergency prohibition or restriction

#### PART IV

##### Use of Information

23. Data protection for active substances
24. Data protection for biocidal products
25. Co-operation in the use of information
26. Confidentiality
27. Treatment of confidential information
28. Exchange of information
29. Notification of information to the National Poisons Information Service

#### PART V

##### Packaging, Labelling and Advertisements

30. Packaging
31. Labelling
32. Samples, models and drafts
33. Advertisements

#### PART VI

##### Miscellaneous and general

34. General provisions on applications for authorisations and registrations
35. Files on applications
36. Appeals
37. Tests
38. Enforcement, offences and civil liability
39. Fees
40. Transitional provisions
41. Extension outside Great Britain
42. Amendments
- Signature

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#### SCHEDULES

- 1 — Biocidal product types and their descriptions
- 2 — Regulations relating to biocidal products
- 3 — Determinations of the Ministers
- 4 — Information to be contained in a dossier submitted in support of an application for the registration of a biocidal product

- 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
- 6 — Non-confidential information
- 7 — Information relating to biocidal products to be given to the Commission and to the competent authorities
- 8 — Information to be notified to the National Poisons Information Service
- 9 — Information to be included on labels
- 10 — Appeals
- 11 — Enforcement, offences and civil liability
- 12 — Fees
- 13 — Transitional provisions

Explanatory Note