SCHEDULE 8 Regulation 29(2) and (7)

Information to be notified to the Poison Information Service

1. The name of the biocidal product.

2. If the biocidal product is authorised or registered under these Regulations—

   (a) the use for which it is so authorised or registered; and

   (b) the name, address and telephone number and any e-mail address and any fax number of the person to whom the authorisation or registration was granted.

3. The date on which the biocidal product was first placed on the market in Northern Ireland.

4. The name, address and telephone number and any e-mail address and any fax number of—

   (a) the manufacturer of the biocidal product;

   (b) any importer of the biocidal product; and

   (c) the individual to be contacted in an emergency in the event of an individual being affected by the biocidal product.

5. A description of the packaging of the biocidal product, including its size and type.

6. The pH, physical state and colour of the biocidal product.

7. The identity of the ingredients of the biocidal product, and their concentration in metric units.

8. The effects on human health of contact with the biocidal product.

9. Particulars of the likely direct or indirect adverse side effects of the biocidal product and any directions for first aid.

10. Any other information relating to the health and safety of humans which is given on the label of the biocidal product.