SCHEDULE 7 Regulation 28(1)

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 whom, the authorisation or registration was granted.

2. The trade name of the biocidal product.

3. The name and amount of each active substance which the biocidal product contains.

4. The name and amount of each substance which the biocidal product contains which is a substance dangerous for supply within the meaning of regulation 2(1) of the 1995 Regulations and its classification.

5. The product-type for the biocidal product and the use for which it is authorised or registered, as the case may be.

6. The type of formulation of the biocidal product, namely whether it is in the form of a powder, granules, a solid, a liquid concentrate or some other form.

7. Any proposed limits on residues which have been determined by the Executive in accordance with paragraph 3(b) of Schedule 3.

8. Any conditions subject to which the authorisation or registration was granted.

9. The reasons for the modification or cancellation of an authorisation or registration.

10. Whether the biocidal product is a low-risk biocidal product or within a frame-formulation.