

## SCHEDULE 3

### **Information to be included in applications for consent to market genetically modified organisms**

#### **PART II**

##### **Additional relevant information**

**9.** The measures to be taken in the event of the escape of the organisms in the product or misuse of the product.

**10.** Specific instructions or recommendations for storage and handling of the product.

**11.** Specific instructions for carrying out monitoring and reporting to the applicant and, if required, the Secretary of State, which are consistent with Part C of Annex VII of the Deliberate Release Directive.

**12.** The proposed restrictions in the approved use of the genetically modified organism, such as where the product may be used and for what purposes.

**13.** The proposed packaging.

**14.** The estimated production in and/or imports to the Community.

**15.** Any proposed additional labelling, which may include, at least in summarised form, the information referred to in paragraphs 4 and 5 of Part I of this Schedule, or paragraphs 9 to 12 of this Part.