SCHEDULE 2 Regulation 11

CONSENT TO MARKET

PART I

GENERAL INFORMATION

1. The name of the product and the name of the genetically modified organisms in the product.

2. The name and address in the Community of the manufacturer or distributor of the product.

3. The specificity of the product and the exact conditions of use including, where appropriate, the type of environment and/or the geographical areas within the Community for which the product is suited.

4. The type of expected use of the product and the description of the persons who are expected to use the product.

PART II

ADDITIONAL RELEVANT INFORMATION

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

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

7. The estimated level and amount of production of the product within the Community and the estimated level and amount of imports of the product into the Community.

8. Information regarding the proposed packaging for the product and its appropriateness so as to avoid the escape of genetically modified organisms during storage or at a later stage.

9. Information regarding proposed labelling including the proposals for stating, in full or summarised form, the information prescribed in paragraphs 1 to 3, 5 and 6 above.