Council Directive of 26 March 1990 laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Community (90/167/EEC)

ANNEX A

..... (Copy for the manufacturer or authorized distributor) (1)

..... (to be kept for .....) (2)

Surname, forename and address of the prescribing veterinarian :

PRESCRIPTION FOR A MEDICATED FEEDINGSTUFF

This prescription may not be re-used

Name or business name and address of the manufacturer or supplier of the medicated feedingstuff :

.....

.....

Name and address of the stockfarmer or holder of the animals : .....

.....

Identification and number of animals : .....

Disease to be treated (3) : .....

Designation of the authorized medicated pre-mixes :

.....

.....

.....

.....

Quantity of medicated feedingstuff : ..... kg

Special instructions for the stockfarmer :

Percentage of medicated feedingstuff in the daily ration, frequency and duration of treatment : .....

.....

Withdrawal time before slaughtering, or waiting time before placing on the market products from treated animals : .....

.....

.....

Personal signature of veterinarian

To be completed by the manufacturer or authorized distributor :

Date of delivery : .....

To be used before : .....

.....

Signature of manufacturer or supplier

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(1) To be filled in in accordance with Article 8 (1) (b).
(2) To be specified by the competent national authorities.
(3) To be entered only on the copy for the veterinarian.