

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)

Status: EU Directives are being published on this site to aid cross referencing from UK legislation. After IP completion day (31 December 2020 11pm) no further amendments will be applied to this version.

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

(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.