Veterinary Prescription for Medicated Feed – POULTRY

FEEDMILL INFORMATION

Feed Mill Name & Address:	Telephone:
	Fax:
	Email:

CLIENT AND VETERINARY CONTACT INFORMATION

Client Name: Telephone:	Veterinarian Name: Clinic Name: Address:
Manager Name: Telephone:	
Farm Address: Telephone:	Telephone:

ANIMALS TO BE TREATED

Species	Production Type	Number of Animals	Placement Date (MM/DD/YYY)	Processing Date (MM/DD/YYYY)	Location of Animals

TREATMENT DURATION

	Type of Feed to be Medicated *(Complete/Supplement/Macro OR Name of Feed)*	Total Amount of Feed *(Tonnes)*	Duration of Feeding	Withdrawal	CgFARAD#
Feed #1					
Feed #2					
Feed #3					

MEDICATED FEED INFORMATION

Brand Substitution Acceptable? Yes ◯ No ◯

Feed to be Medicated *(As stated above)*	CMIB code	CMIB Claim #	DIN product BRAND NAME *(Active ingredient)*	g of AI/kg of DIN product	g of AI/tonne complete feed	g of DIN product/ tonne of complete feed

Manufacturing Instructions:	On-Farm Mixing & Feeding Directions:
Warning(s):	Caution(s):
Veterinarian Name (printed):	Signature:
Date:	License No.:
Prescription Expiry Date:	

OPTIONAL TEMPLATE

Instruction for Completing the Veterinary Prescription Template for Medicated Feed - POULTRY

(*Note: The use of this template is optional. It has been prepared as an additional prescribing tool for veterinarians.)*

Feed Mill Name and Address where the feed is manufactured.

Name and address of animal owner: provide the name and address of the person for whom the feed is to be manufactured or sold. The Client is the person who owns the animals.

Name of animal manager (if different from above): provide the name of the person responsible for the management of the animals on the premises (*Feeds Regs 5.2(g)(iii)(B); Food and Drug Regs C.08.012.2(d)(i)*).

Veterinarian's name and contact information: name and contact information for the prescribing veterinarian.

Information on the animals to be medicated: include the species, production type and placement date of the animals to be treated with the medicated feed (*Feeds Regs 5.2(g)(iii)(E); Food and Drug Regs C.08.012.2(d)(ii)*). Include the expected processing date, if known.

Location of animals to be medicated: provide the location where the animals to be medicated are housed, including both the address of the premises and the specific location of the animals on the site (e.g. Barn 2, Heifer barn). As the name, ID number or tag number for each animal to be medicated is not indicated on a prescription for a medicated feed, the location of the animals help to distinguish which animals on the farm are intended to be treated.

Indicate the Treatment Duration in the number of days or the age of the birds for each medicated feed.

Type of feed to be medicated (i.e. complete feed, supplement, micro/macro premix) **and** the total amount of feed to be manufactured under that prescription for the number of animals and the duration of treatment indicated. (*Feeds Regs 5.2(g)(iii)(D); Food and Drug Regs C.08.012.2(d)(iii & iv)*). There is space on the template to prescribe up to 3 rations for the same group of animals to account for changing needs as an animal moves through the feeding cycle. There is no regulatory requirement that limits the number of rations to 3. The age of the birds is used to identify the group to be fed each ration and the treatment duration.

Name of the medicating ingredient(s) to be added to each feed, as indicated in the form above, by indicating the proper name, or the common name if there is no proper name, of the drug or each of the drugs, as the case may be, to be used as medicating ingredients in the preparation of the medicated feed. (*Feeds Regs 5.2(g)(iii)(C); Food and Drug Regs C.08.012.2(d)(iv)*) . If prescribing a feed in accordance with the Compendium of Medicated Ingredients Brochure (CMIB), indicate the CMIB medicated ingredient code and the appropriate claim number. If prescribing feeds off-label, leave the CMIB fields blank.

Amount of the medicating ingredient(s) to be added (*Feeds Regs 5.2(g)(iii)(C)*), including at minimum:

a) Amount of active in the DIN product in g of active per kg of premix (often included in the Brand Name)

b) Amount of **active ingredient** in mg per tonne of medicated feed

c) Amount of **DIN product** in mg per tonne of medicated feed

* Including all three pieces of information will ensure that the calculations have been conducted properly and should reduce potential mixing errors

Any special mixing instructions, including any special manufacturing or mixing instructions (*Feeds Regs 5.2(g)(iii)(F); Food and Drug Regs C.08.012.2(d)(v)*).

On-Farm Mixing and Feeding Directions provides the directions for use on-farm, including frequency of feeding, any other special feeding instructions and if further mixing is required on-farm (*Feeds Regs 5.2(g)(iii)(G); Food and Drug Regs C.08.012.2(d)(vi)(A)*).

Warnings and Cautions: If feed is being prescribed as per the CMIB, the prescription can indicate the CMIB code and particular claim for the warnings and cautions to be populated as per the CMIB. If prescribing a medicated feed in a manner that is not consistent with the CMIB (off-label), the prescription must contain all pertinent warnings pertaining to human health, and cautions related to animal health, which are present on the approved drug label, and any additional warnings or cautions that the veterinarian deem necessary (*Feeds Regs 5.2(g)(iii)(H); Food and Drug Regs C.08.012.2(d)(vi)(C)*).

Withdrawal Period: If prescribing a medicated feed as per CMIB, the labelled withdrawal period will be copied into the Warnings section as it is listed in the CMIB. If prescribing a medicated feed in a manner that is not consistent with the CMIB (off-label), it is the veterinarian's responsibility to indicate the appropriate withdrawal period on the prescription (*Feeds Regs 5.2(g)(iii)(H); Food and Drug Regs C.08.012.2(d)(vi)(B)*).

Date: provide the date on which the prescription is written.

Veterinary Signature: The prescription must be physically signed by the prescribing veterinarian or provided electronically following best practices for electronic signatures (*Feeds Regs 5.2(g)(iii); Food and Drug Regs C.08.012.2(b)*).