Caution:

This consultation draft is intended to facilitate dialogue concerning its contents. Should the decision be made to proceed with the proposal, the comments received during consultation will be considered during the final preparation of the regulation. The content, structure, form and wording of the consultation draft are subject to change as a result of the consultation process and as a result of review, editing and correction by the Office of Legislative Counsel.

CONSULTATION DRAFT

ONTARIO REGULATION

to be made under the

LIVESTOCK MEDICINES ACT

Amending Reg. 730 of R.R.O. 1990

(GENERAL)

1. (1) The definition of “biological” in section 1 of the Regulation 730 of the Revised Regulations of Ontario, 1990 is revoked.

(2) Section 1 of the Regulation is amended by adding the following definition:

“veterinary biologic” has the same meaning as in the Health of Animals Act (Canada).

2. Section 2 of the Regulation is revoked and the following substituted:

2. (1) Subject to subsections (4) and (5), the following drugs are designated as livestock medicines for the purposes of the Act and this Regulation:

1. Modified-live virus and live virus vaccines for use in poultry.

2. Preparations for the control of external and internal parasites and insect pests of livestock.
3. Oral preparations labelled by the manufacturer for the prevention or treatment of disease of the digestive system in livestock, including bloat, colic, indigestion, diarrhea, constipation and impaction.

4. Preparations labelled by the manufacturer for the treatment of surface wounds and lacerations, wire cuts and burns in livestock.

5. Preparations labelled by the manufacturer for the treatment of skin diseases in livestock, including topical hoof care products.

6. Vitamins for injection or oral administration to livestock, including injectable vitamin A, not to exceed 500,000 I.U.’s per millilitre and injectable vitamin D, not to exceed 75,000 I.U.’s per millilitre.

7. Preparations for the prevention or treatment of deficiencies in livestock, including hematinics for horses, containing,

   i. minerals for oral administration,

   ii. selenium for injection, or

   iii. iron for injection.

8. Injectable epinephrine for treatment of anaphylactic reactions in livestock.

9. Propylene glycol and preparations, including dextrose, calcium, phosphorus and magnesium preparations, labelled by the manufacturer for the treatment and prevention of acetonemia and hypocalcemia in livestock or otherwise intended as an aid in the supportive treatment of nutritional deficiencies in livestock.

10. Implants and feed additives labelled by the manufacturer as growth promotants for use in livestock.

11. Anti-cannibalism compounds for poultry.
12. Topical preparations labelled by the manufacturer as liniments, counterirritants and poultices for the treatment of joint pain and swollen ligaments, tendons or muscles.

13. Oral or topical preparations labelled by the manufacturer as antitussives, decongestants, bronchodilators and expectorants.


15. Disinfectants, udder washes, teat dips and sanitizers.

(2) Subject to subsections (4) and (5), a veterinary biologic is designated as a livestock medicine for the purposes of the Act and this Regulation, unless it is exempted under subsection (3), if the veterinary biologic was,

(a) imported into Canada pursuant to a permit issued under section 121 of the Health of Animals Regulation (Canada); or

(b) manufactured pursuant to a product licence issued under section 124 of the Health of Animals Regulation (Canada) and it is permitted to be sold under that Regulation.

(3) The following veterinary biologics are not designated as livestock medicines:

1. Anthrax vaccine.

2. Brucella vaccine.

3. Rabies vaccine.


(4) A drug shall not be designated as a livestock medicine unless it has been assigned a drug identification number pursuant to subsection C.01.014.2 (1) of the Food and Drug Regulations (Canada).
(5) A drug that is prescribed by subsection 3 (2) of Regulation 264/16 (General) made under the *Drug and Pharmacies Regulation Act* as being in Schedule I to the *Drug and Pharmacies Regulation Act* is not designated as a livestock medicine.

3. Subsection 3 (2) of the Regulation is amended by striking out “the Schedule” and substituting “section 2”.

4. Clause 9 (1) (d) of the Regulation is amended by striking out “biologics” at the end and substituting “veterinary biologics”.

5. Clause 11 (1) (a) of the Regulation is revoked and the following substituted:

   (a) attach a tag to the livestock medicine bearing a serial number and the words “Ont. Detained/(French)”;

6. The Schedule to the Regulation is revoked.

7. The Regulation is amended by adding the following French version:

   (French)

   Commencement

   8. [commencement]