3. This Regulation comes into force on the fifteenth day following the date of its publication in the *Gazette officielle du Québec.*

102050

Draft Regulation

Veterinary Surgeons Act
(chapter M-8)

Pharmacy Act
(chapter P-10)

Terms and conditions for the sale of medications — Amendment

Notice is hereby given, in accordance with sections 10 and 11 of the Regulations Act (chapter R-18.1), that the Regulation to amend the Regulation respecting the terms and conditions for the sale of medications, made by the Office des professions du Québec, may be submitted to the Government for approval, with or without amendment, on the expiry of 45 days following this publication.

The Regulation specifies the terms and conditions for the sale of the following substances: dextromethorphan and its salts, glycosaminoglycan, and pseudoephedrine and its salts.

The Office expects that the new measures will have no impact on enterprises, including small and medium-sized businesses.

Further information may be obtained by contacting Ugo Chaillez, Direction des affaires juridiques, Office des professions du Québec, 800, place D'Youville, 10e étage, Québec (Québec) G1R 5Z3; telephone: 418 643-6912 or 1 800 643-6912; fax: 418 643-0973.

Any person wishing to comment on the matter is requested to submit written comments within the 45-day period to the Chair of the Office des professions du Québec, 800, place D'Youville, 10e étage, Québec (Québec) G1R 5Z3. The comments will be forwarded by the Office to the Minister of Justice and may also be sent to interested persons, departments and bodies.

JEAN PAUL DUTRISAC,
Chair of the Office des professions du Québec

Regulation to amend the Regulation respecting the terms and conditions for the sale of medications

Veterinary Surgeons Act
(chapter M-8, s. 9, 1st par.)

Pharmacy Act
(chapter P-10, s. 37.1)

1. The Regulation respecting the terms and conditions for the sale of medications (chapter P-10, r. 12) is amended in Schedule II

(1) by inserting the following substance and specification after "Desoxyribonuclease (Pancreatic)":

"Dextromethorphan and its salts" and "Dosage forms in packaging units containing more than 850 mg"; and

(2) by inserting the following substance and specification after "Protamine and its salts":

"Pseudoephedrine and its salts", "Dosage forms containing no other medicinal ingredient" and "Dosage forms in packaging units containing more than 1,200 mg and containing another medicinal ingredient".

2. Schedule III is amended

(1) by adding the following specification to the substance "Dextromethorphan and its salts":

"Dosage forms in packaging units containing 850 mg or less and sold in single packages containing only one packaging unit"; and

(2) by replacing the specifications of the substance "Pseudoephedrine and its salts" by the following:

"Dosage forms in packaging units containing 1,200 mg or less, sold in single packages containing only one packaging unit and containing another medicinal ingredient".

3. Schedule IV is amended by adding the following specification to the substance "Glycosaminoglycan":

"Except dosage forms for oral use".

4. Schedule V is amended by inserting the following substance and specification after "Fipronil":

"Glycosaminoglycan" and "Dosage forms for oral use".

5. This Regulation comes into force on the fifteenth day following the date of its publication in the *Gazette officielle du Québec.*

102048