2. Paragraphs *b* and *c* of section 1.07, replaced by section 1 of this Regulation, remain applicable to persons who, on 22 October 2009, hold the diplomas referred to in the replaced paragraphs or are registered in a program enabling them to obtain such diplomas.

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

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Gouvernement du Québec

O.C. 1024-2009, 23 September 2009

Pharmacy Act
(R.S.Q., c. P-10)

Terms and conditions for the sale of medications — Amendments

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

WHEREAS, under section 37.1 of the Pharmacy Act (R.S.Q., c. P-10), the Office des professions du Québec, after consultation with the Conseil du médicament, the Ordre professionnel des médecins du Québec, the Ordre professionnel des médecins vétérinaires du Québec and the Ordre des pharmaciens du Québec, may, by regulation, establish categories of medications and determine, for each category, if need be, by whom and subject to what terms and conditions the medications may be sold;

WHEREAS the Office, after carrying out the consultations required by that section, adopted the Regulation to amend the Regulation respecting the terms and conditions for the sale of medications at its sitting of 26 February 2009;

WHEREAS, in accordance with sections 10 and 11 of the Regulations Act (R.S.Q., c. R-18.1), the draft Regulation to amend the Regulation respecting the terms and conditions for the sale of medications was published in Part 2 of the *Gazette officielle du Québec* of 15 April 2009 with a notice that it could be submitted to the Government for approval on the expiry of 45 days following that publication;

WHEREAS the Office has received no comments following the publication of the Regulation;

WHEREAS, under section 13 of the Professional Code (R.S.Q., c. C-26), every regulation adopted by the Office under the Code or under an Act constituting a professional order must be submitted to the Government, which may approve it with or without amendment;

WHEREAS it is expedient to approve the Regulation without amendment;

IT IS ORDERED, therefore, on the recommendation of the Minister of Justice:

THAT the Regulation to amend the Regulation respecting the terms and conditions for the sale of medications, attached to this Order in Council, be approved.

GÉRARD BIBEAU,
Clerk of the Conseil exécutif

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

Pharmacy Act
(R.S.Q., c. P-10, s. 37.1)

1. The Regulation respecting the terms and conditions for the sale of medications is amended in Schedule III by adding the following paragraph at the end of the specification of the substance "Famotidine and its salts":

"Dosage forms for oral use containing more than 10 mg and not more than 20 mg per dosage unit, in package units containing less than 51 dosage units".

2. Schedule III is amended by inserting ", lozenges" after "inhalers" in the specification of the substance "Nicotine and its salts".

3. Schedule III is amended by adding the following paragraph at the end of the specification of the substance "Ranitidine and its salts":

"Dosage forms for oral use containing more than 75 mg and not more than 150 mg per dosage unit, in package units containing less than 51 dosage units".

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

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* The Regulation respecting the terms and conditions for the sale of medications, approved by Order in Council 712-98 dated 27 May 1998 (1998, *G.O.* 2, 2149), was last amended by the regulation approved by Order in Council 539-2008 dated 28 May 2008 (2008, *G.O.* 2, 2113). For previous amendments, refer to the *Tableau des modifications et Index sommaire*, Québec Official Publisher, 2009, updated to 1 March 2009.