Pharmacy Act, 1991
Loi de 1991 sur les pharmaciens

ONTARIO REGULATION 202/94

GENERAL

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This Regulation is made in English only.

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PART I
INTERPRETATION

Definitions

1. In this Regulation,

"direct supervision" means supervision that is provided by a person who is physically present on the premises where the practice that is being supervised is being carried out;

"non-restricted registration" means the holding of a licence, permit, certificate or registration as a pharmacist from an authority responsible for the regulation of pharmacists in one of the states of the United States of America, or in another non-Canadian jurisdiction that has been approved by the Council, where that licence, permit, certificate or registration is not subject to any restrictions, terms, conditions or limitations, including terms, conditions or limitations that,

(a) relate to the holder's ability to practise independently,

(b) require the holder to practise under supervision or direction,

(c) require the holder to maintain a position or appointment as a condition of continued registration,

(d) require the holder to practise only in a part of the geographical area over which the authority has jurisdiction,

(e) restrict the holder to temporary or time-limited registration or practice,

(f) were imposed by that authority or any committee or panel of that authority as a result of a disciplinary, registration, fitness to practise or similar proceeding, or

(g) were placed on the holder's registration by agreement between the holder and that authority;

"pharmacy" has the same meaning as in subsection 1 (1) of the *Drug and Pharmacies Regulation Act*;

"remote dispensing location" has the same meaning as in subsection 1 (1) of the *Drug and Pharmacies Regulation Act*. O. Reg. 451/10, ss. 1, 6 (1).

PART II
GENERAL PROVISIONS RE CERTIFICATES OF REGISTRATION

Classes of Certificates of Registration

2. (1) The following are prescribed as classes of certificates of registration:

1. Pharmacist.
2. Registered pharmacy student.
3. Intern.
4. Pharmacy technician.
5. Pharmacist (emergency assignment).
6. Pharmacy technician (emergency assignment). O. Reg. 451/10, s. 1; O. Reg. 187/21, s. 1.

(2) Every certificate of registration that was in existence immediately before December 3, 2010 is continued as the equivalent certificate of registration with the same status under this Regulation until such time as it otherwise ceases to be effective. O. Reg. 451/10, s. 1.

(3) Where an application for a certificate of registration had been made but not finally dealt with before December 3, 2010, the application shall be dealt with in accordance with this Regulation as amended by Ontario Regulation 451/10. O. Reg. 451/10, s. 1.

Application for Certificate of Registration

3. A person may apply for a certificate of registration by submitting a completed application in the form provided by the Registrar together with any supporting documentation requested by the Registrar and the applicable fees. O. Reg. 451/10, s. 1.

Requirements for Issuance of Certificate of Registration, any Class

4. (1) The following are requirements for the issuance of a certificate of registration of any class:

1. The applicant must possess sufficient language proficiency, in either English or French, to be able to communicate and comprehend effectively, both orally and in writing.
2. The applicant must not have been found guilty of any offence in any jurisdiction.
3. The applicant must not be the subject of a current proceeding in respect of any offence in any jurisdiction.
4. The applicant must not have been the subject of a finding of professional misconduct, incompetence or incapacity or any like finding in Ontario or any other jurisdiction in relation to pharmacy or any other profession or occupation and must not be the subject of any current professional misconduct, incompetence, or incapacity proceeding or any like proceeding in Ontario or any other jurisdiction in relation to pharmacy or any other profession or occupation.
5. The applicant must be a Canadian citizen or permanent resident of Canada or must hold the appropriate authorization under the *Immigration and Refugee Protection Act* (Canada) to permit the applicant to engage in the practice of the profession in Ontario in the manner permitted by the certificate of registration for which he or she has applied.
6. The applicant's past and present conduct must afford reasonable grounds for the belief that the applicant,
    i. will practise pharmacy with decency, honesty and integrity, and in accordance with the law,
    ii. does not suffer from any physical or mental condition or disorder that could affect his or her ability to practise pharmacy in a safe manner,
    iii. has sufficient knowledge, skill and judgment to competently engage in the practice of pharmacy authorized by the certificate of registration, and
    iv. will display an appropriately professional attitude.
7. The applicant must provide evidence satisfactory to the Registrar that the applicant will have professional liability insurance in the amount and in the form as required by the by-laws as of the anticipated date for the issuance of his or her certificate of registration.
8. The applicant must have paid any fees required under the by-laws for the issuance of the certificate of registration for which the applicant applied. O. Reg. 451/10, s. 1; O. Reg. 187/21, s. 2 (1).

(2) The requirement under paragraph 8 of subsection (1) is non-exemptible. O. Reg. 451/10, s. 1.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, subsection 4 (2) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 1)

(2) Subject to sections 15.3 and 18.3, the requirement under paragraph 8 of subsection (1) of this section is non-exemptible. O. Reg. 295/23, s. 1.

(3) An applicant must meet all of the requirements for registration within one year following the filing his or her application, but this does not prevent the applicant from filing a new application. O. Reg. 451/10, s. 1.

(3.1) Despite subsection (3), an applicant for a certificate in the pharmacist (emergency assignment) or pharmacy technician (emergency assignment) class must meet all the requirements for registration at the time the application is filed. O. Reg. 187/21, s. 2 (2).

(4) An applicant shall be deemed not to have satisfied the requirements for the issuance of a certificate of registration if the applicant makes a false or misleading statement or representation in his or her application or supporting documentation. O. Reg. 451/10, s. 1.

Terms, etc., of Every Certificate

5. Every certificate of registration is subject to the following terms, conditions and limitations:

1. The member shall provide to the Registrar the details of any of the following that relate to the member and that occur or arise after the registration of the member:
    i. a finding of guilt arising in any jurisdiction relating to any offence,

ii. a charge arising in any jurisdiction relating to any offence,

iii. a finding of professional misconduct, incompetence or incapacity or any like finding in any jurisdiction in relation to pharmacy or any other profession or occupation,

iv. a proceeding for professional misconduct, incompetence or incapacity or any like proceeding in any jurisdiction in relation to pharmacy or any other profession or occupation.

2. The member shall not engage in the practice of pharmacy unless the member is a Canadian citizen or permanent resident of Canada or has authorization under the *Immigration and Refugee Protection Act* (Canada) permitting the member to engage in the practice of pharmacy in Ontario in the manner permitted by the certificate of registration.

3. The member shall immediately advise the Registrar in writing in the event the member ceases to be a Canadian citizen or permanent resident of Canada or to have authorization under the *Immigration and Refugee Protection Act* (Canada) permitting the member to engage in the practice of pharmacy in Ontario in the manner permitted by the certificate of registration.

4. If a member to whom paragraph 3 applies subsequently obtains Canadian citizenship or becomes a permanent resident of Canada or attains authorization under the *Immigration and Refugee Protection Act* (Canada) permitting the member to engage in the practice of pharmacy in Ontario permitted by the certificate of registration, the member shall immediately advise the Registrar in writing of that fact.

5. The member shall maintain professional liability insurance in the amount and in the form as required by the by-laws.

6. A member who fails to meet the condition in paragraph 5 shall immediately advise the Registrar in writing of that fact and immediately cease to engage in the practice of pharmacy until such time as the member obtains professional liability insurance as required in paragraph 5.

7. Where a member to whom paragraph 6 applies subsequently obtains professional liability insurance, the member shall notify the Registrar in writing of that fact and, if requested by the Registrar, shall provide details of that coverage. O. Reg. 451/10, s. 1.

PART III
REGISTRATION — PHARMACISTS

Additional Requirements

6. (1) The following are additional requirements for the issuance of a certificate of registration as a pharmacist:

1. The applicant must,

   i. have a minimum of a baccalaureate degree in pharmacy evidencing the successful completion of a program designed to educate and train persons to be practising pharmacists which was,

      A. awarded on or before December 31, 1993 by a constituent faculty of the Association of Faculties of Pharmacy of Canada, or

      B. awarded by a university as a result of successful completion of a program which was, at the time of the award, accredited by the Canadian Council for Accreditation of Pharmacy Programs or by another accrediting body approved by the Council for that purpose, or

   ii. have a university degree in pharmacy that does not meet the requirements of subparagraph i but that evidences the successful completion of a program designed to educate and train persons to be practising pharmacists, and,

      A. have successfully completed a program that, at the time the applicant commenced it, was approved by the Council as one whose graduates should possess knowledge, skill and judgment at least equal to those of current graduates of a program mentioned in sub-subparagraph i B, or

      B. have successfully completed the examination provided for in paragraph 4 on the applicant's first attempt and have successfully completed any further education or training or combination of education and training that was specified by a panel of the Registration Committee to evidence that the applicant possesses knowledge, skill and judgment at least equal to those of current graduates of a program mentioned in sub-subparagraph i B.

2. The applicant must have successfully completed an examination in pharmaceutical jurisprudence approved by the Council for applicants for a certificate of registration as a pharmacist.

3. Subject to subsection (5), the applicant must have successfully completed a structured practical training program approved by the Council, while holding a certificate of registration as an intern and while under the supervision of a preceptor approved by the Registration Committee.

4. The applicant must have successfully completed the qualifying examination of the Pharmacy Examining Board of Canada for pharmacists at a time when the examination was approved by the Council or have successfully completed another examination that has been approved by the Council for that purpose. O. Reg. 451/10, s. 1.

(2) With the approval of the Council, all or any part of a structured practical training program referred to in paragraph 3 of subsection (1) may be completed as part of a program referred to in paragraph 1 of subsection (1), even if the applicant did not hold a certificate of registration as an intern at the time. O. Reg. 451/10, s. 1.

(3) The requirement in paragraph 1 of subsection (1) must be met within two years before the date on which the applicant met all of the other requirements for the issuance of a certificate of registration as a pharmacist unless the applicant,

(a) satisfies the Registrar or a panel of the Registration Committee that the applicant has practised pharmacy within the scope of practice of the profession as set out in section 3 of the Act for at least 600 hours during the three years before the date on which the applicant met all of the other requirements for the issuance of a certificate of registration as a pharmacist in the course of providing patient care while practising as a pharmacist in Canada, the United States of America or another jurisdiction approved by the Council;

(b) undergoes a review of his or her practice conducted in a manner approved by the Registration Committee, meets any requirements regarding continuing education or remediation set by a panel of the Registration Committee within the time set by the panel, and pays the required fees; or

(c) successfully completes the examination referred to in paragraph 4 of subsection (1) within three years of the date on which he or she meets all of the other requirements for the issuance of a certificate of registration as a pharmacist. O. Reg. 451/10, s. 1.

(4) The requirement in paragraph 2 of subsection (1) shall not be considered to be met unless the applicant is issued a certificate of registration as a pharmacist within three years of meeting that requirement. O. Reg. 451/10, s. 1.

(5) An applicant is deemed to have met the requirement in paragraph 3 of subsection (1) if, at the time of application, the applicant,

(a) has successfully completed a structured practical training program which is, in the opinion of the Registration Committee at least equivalent to the program mentioned in paragraph 3 of subsection (1); or

(b) has other education, training or experience that is, in the opinion of a panel of the Registration Committee at least equivalent to the program mentioned in paragraph 3 of subsection (1). O. Reg. 451/10, s. 1.

(6) The requirement in paragraph 3 of subsection (1) shall not be considered to have been met unless the applicant is issued a certificate of registration as a pharmacist within two years of meeting the requirement or within such greater time as is specified by a panel of the Registration Committee. O. Reg. 451/10, s. 1.

(7) Subject to subsection (8), the requirement in paragraph 4 of subsection (1) is not considered to have been met unless the applicant,

(a) successfully completed the examination within three attempts; or

(b) successfully completed the examination on the applicant's fourth attempt having first successfully completed the further education or training or combination of education and training required by the examining body responsible for the administration of the examination or, if no further education or training was required by that body, the further education or training or combination of education and training, if any, specified by a panel of the Registration Committee. O. Reg. 451/10, s. 1.

(8) Where, by virtue of subsection (7), an applicant is not considered to have met the requirement in paragraph 4 of subsection (1), the applicant may not attempt the examination again until the applicant obtains a new degree mentioned in subparagraph 1 i of subsection (1). O. Reg. 451/10, s. 1.

(9) An applicant is only eligible to take the examination referred to in paragraph 2 of subsection (1) three times in any 24-month period. O. Reg. 451/10, s. 1.

(10) The requirements in paragraphs 1, 3 and 4 of subsection (1) are deemed to have been met by an applicant,

(a) who previously held a certificate of registration as a pharmacist in Ontario; and

(b) who,

(i) satisfies the Registrar or a panel of the Registration Committee that the applicant has practised pharmacy within the scope of practice of the profession as set out in section 3 of the Act for at least 600 hours during the three years before the date on which the applicant met all of the other requirements for the issuance of a certificate of registration as a pharmacist in the course of providing patient care while practising as a pharmacist in Canada, the United States of America or another jurisdiction approved by the Council, or

(ii) undergoes a review of his or her practice conducted in a manner approved by the Registration Committee, meets any requirements regarding continuing education or remediation set by a panel of the Registration Committee within the time set by the panel, and pays the required fees. O. Reg. 451/10, s. 1.

(11) An applicant who has a university degree in pharmacy mentioned in subparagraph 1 ii of subsection (1), and who successfully completes any further education or training or combination of education and training specified by a panel of the Registration Committee shall be deemed to have met the requirement in paragraph 1 of subsection (1) if the applicant,

(a) was registered as an intern on December 3, 2010; or

(b) becomes registered as an intern after December 3, 2010 but before December 3, 2011. O. Reg. 451/10, s. 1.

(12) Subject to subsections (2), (5), (10) and (11) and sections 7 and 8, the requirements in subsection (1) are non-exemptible. O. Reg. 451/10, s. 1.

(13) A reference in this section or section 7 to "all of the other requirements for the issuance of a certificate of registration" includes, without being limited to, a requirement set out in subsection 4 (1) or subsection (1) of this section. O. Reg. 451/10, s. 1.

MOBILITY FROM OUTSIDE CANADA

7. An applicant is deemed to have met the requirements in paragraph 1 of subsection 6 (1) if the applicant meets all the following non-exemptible requirements:

1. The applicant must,
   i. hold a non-restricted registration in at least one jurisdiction at the time of application and have held that registration continuously for at least two years, and
   ii. satisfy the Registrar or a panel of the Registration Committee that the applicant engaged in the full scope of practice as a pharmacist in that jurisdiction for at least 600 hours.
2. The applicant must,
   i. satisfy the Registrar or a panel of the Registration Committee that the applicant has practised pharmacy within the scope of practice of the profession as set out in section 3 of the Act for at least 600 hours during the three years before the date on which the applicant met all of the other requirements for the issuance of a certificate of registration as a pharmacist in the course of providing patient care while practising as a pharmacist in one or more of the jurisdictions where he or she held the non-restricted registration,
   ii. undergo a review of his or her practice conducted in a manner approved by the Registration Committee, meet any requirements regarding continuing education or remediation set by a panel of the Registration Committee within the time set by the panel, and pay the required fees, or
   iii. successfully complete the examination referred to in paragraph 4 of subsection 6 (1) within three years of the date on which he or she meets all of the other requirements for the issuance of a certificate of registration as a pharmacist. O. Reg. 451/10, s. 1.

MOBILITY WITHIN CANADA

8. (1) Where section 22.18 of the Health Professions Procedural Code applies to an applicant the requirements of paragraphs 1, 3 and 4 of subsection 6 (1) are deemed to have been met by the applicant if he or she provides, for each jurisdiction where the applicant holds an out-of-province certificate, a certificate, letter or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as a pharmacist in that jurisdiction. O. Reg. 451/10, s. 1.

(2) Without in any way limiting the generality of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority that issued the applicant that out-of-province certificate as a pharmacist. O. Reg. 451/10, s. 1.

(3) An applicant referred to in subsection (1) is deemed to have met the requirements of paragraph 1 of subsection 4 (1) where the requirements for the issuance of the applicant's out-of-province certificate included language proficiency requirements equivalent to those required by that paragraph. O. Reg. 451/10, s. 1.

(4) Despite subsection (1), an applicant is not deemed to have met a requirement if that requirement is described in subsection 22.18 (3) of the Health Professions Procedural Code. O. Reg. 451/10, s. 1.

TERMS, CONDITIONS AND LIMITATIONS, PART B PHARMACIST

9. (1) Every certificate of registration of a pharmacist listed in Part B of the register is subject to the following terms, conditions and limitations:

1. The member shall not provide any care to a patient, whether direct or indirect.
2. The member shall not dispense, sell or compound drugs.
3. The member shall not supervise that part of the pharmacy where drugs are kept.
4. The member shall not be the designated manager of a pharmacy within the meaning of the *Drug and Pharmacies Regulation Act*.

5. The member shall not supervise the practice of pharmacy of an intern, registered pharmacy student or pharmacy technician.

6. The member shall, when working in a pharmacy or any other environment where patient care is being provided, clearly identify him or herself as a non-practising pharmacist. O. Reg. 451/10, s. 1.

(2) With the prior written approval of the Registrar, and despite paragraphs 1 and 2 of subsection (1), a pharmacist listed in Part B of the register may dispense, sell or compound a drug and provide care to a patient under the direct supervision of a pharmacist who is registered in Part A of the register, or of a pharmacist (emergency assignment), where the sole purpose is to assist the member in preparing to meet the requirements specified in subsection 46 (3) to transfer a member holding a certificate of registration as a pharmacist who is registered in Part B of the register to Part A of the register. O. Reg. 187/21, s. 3.

(3) Where a member wishes to seek the approval of the Registrar under subsection (2), the member shall provide to the Registrar, in writing, the name of the pharmacist or pharmacists who will be providing the required supervision, the name and address of the pharmacy or pharmacies at which the member proposes to practise under that supervision and the proposed date upon which the member wishes to commence practice. O. Reg. 451/10, s. 1.

(4) Any approval provided by the Registrar under subsection (2) must specify,

(a) the name of the pharmacist or pharmacists who will be required to supervise the member;

(b) the name and address of the pharmacy or pharmacies where the member will be practising; and

(c) the term of the approval, which must not exceed six months. O. Reg. 451/10, s. 1.

(5) Where the Registrar is satisfied that it is appropriate to do so the Registrar may extend the term of the approval provided under subsection (2) but in no case may the combined term exceed one year unless a panel of the Quality Assurance Committee approves of a further extension. O. Reg. 451/10, s. 1.

PART IV
REGISTRATION — REGISTERED PHARMACY STUDENTS

Additional Requirement

10. (1) It is an additional requirement for the issuance of a certificate of registration as a registered pharmacy student that the applicant must,

(a) have been accepted as a student in a university program referred to in subparagraph 1 i of subsection 6 (1) or in an approved program referred to in sub-subparagraph 1 ii A of that subsection;

(b) be engaged in attaining any education or training referred to in sub-subparagraph 1 ii B of subsection 6 (1); or

(c) be engaged in attaining any education or training specified by a panel of the Registration Committee as a condition for the issuance of another certificate of registration, other than a certificate of registration as a pharmacy technician. O. Reg. 451/10, s. 2.

(2) Subject to section 11, the requirement in subsection (1) is non-exemptible. O. Reg. 451/10, s. 2.

Mobility within Canada

11. (1) Where section 22.18 of the Health Professions Procedural Code applies to an applicant the requirements of subsection 10 (1) are deemed to have been met by the applicant if he or she provides, for each jurisdiction where the applicant holds an out-of-province certificate, a certificate, letter or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as a pharmacy student in that jurisdiction. O. Reg. 451/10, s. 2.

(2) Without in any way limiting the generality of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority that issued the applicant that out-of-province certificate as a registered pharmacy student. O. Reg. 451/10, s. 2.

(3) An applicant referred to in subsection (1) is deemed to have met the requirements of paragraph 1 of subsection 4 (1) where the requirements for the issuance of the applicant's out-of-province certificate included language proficiency requirements equivalent to those required by that paragraph. O. Reg. 451/10, s. 2.

(4) Despite subsection (1), an applicant is not deemed to have met a requirement if that requirement is described in subsection 22.18 (3) of the Health Professions Procedural Code. O. Reg. 451/10, s. 2.

TERMS, CONDITIONS AND LIMITATIONS

12. (1) Every certificate of registration as a registered pharmacy student is subject to the following terms, conditions and limitations:

1. The member,
   i. in the case of a member to whom subsection 11 (1) does not apply, shall only engage in the practice of pharmacy while enrolled in and actively participating in a program provided for in subsection 10 (1) or while actively engaged in any education or training specified in that subsection, and
   ii. in the case of a member to whom subsection 11 (1) applies, shall only engage in the practice of pharmacy while enrolled in and actively participating in an educational program that is a requirement for the issuance of an applicable out-of-province certificate authorizing practice as an intern or pharmacist.
2. The member may only engage in the practice of pharmacy,
   i. while under the direct supervision of a member who holds a certificate of registration as a pharmacist, or as a pharmacist (emergency assignment), or
   ii. where a program or any education or training provided for in subsection 10 (1) includes a clinical component in a premises that is not a pharmacy but at which drugs may be prescribed or dispensed, while under the direct supervision of a member of a College within the meaning of the *Regulated Health Professions Act, 1991* who has been approved for this purpose by the faculty that provides the program, education or training.
3. Despite subparagraph 2 ii, the member shall not dispense, compound or sell a drug unless under the direct supervision of a member holding a certificate of registration as a pharmacist.
4. Despite subparagraph 2 ii, the member may not supervise that part of the pharmacy where drugs are kept.
5. The member may neither delegate a controlled act nor accept the delegation of a controlled act. O. Reg. 451/10, s. 2; O. Reg. 187/21, s. 4.

(2) A certificate of registration as a registered pharmacy student automatically expires when the member is issued a certificate of registration as a pharmacist or an intern. O. Reg. 451/10, s. 2.

(3) A certificate of registration as a registered pharmacy student automatically expires,

(a) in the case of a member engaged in a program referred to in subparagraph 1 i of subsection 6 (1), when the member is refused readmission to the program, ceases to be enrolled in the program or ceases to actively participate in the program;

(b) in the case of a member engaged in an approved program referred to in sub-subparagraph 1 ii A of subsection 6 (1), two years after registration as a registered pharmacy student unless that period of time is extended by a panel of the Registration Committee;

(c) in the case of a member engaged in attaining any education or training or combination of education and training referred to in sub-subparagraph 1 ii B of subsection 6 (1) or in attaining any education or training or combination of education and training required by a panel of the Registration Committee as a condition for the issuance of another class of certificate of registration, on the date specified by the panel in its decision or, if no date was specified, one year from that decision, unless extended by a panel of the Registration Committee; and

(d) in the case of a member whose application for a certificate of registration as a registered pharmacy student was considered under subsection 11 (1), on the date on which the member ceases to hold an out-of-province certificate that is equivalent to a certificate of registration as a registered pharmacy student. O. Reg. 451/10, s. 2.

PART V
REGISTRATION — INTERNS

ADDITIONAL REQUIREMENTS

13. (1) The following are additional requirements for the issuance of a certificate of registration as an intern:

1. The applicant must,
   i. have a minimum of a baccalaureate degree in pharmacy evidencing the successful completion of a program designed to educate and train persons to be practising pharmacists which was,
      A. awarded on or before December 31, 1993 by a constituent faculty of the Association of Faculties of Pharmacy of Canada, or
      B. awarded by a university as a result of successful completion of a program which was, at the time of the award, accredited by the Canadian Council for Accreditation of Pharmacy Programs or by another accrediting body approved by the Council for that purpose, or

ii. have a university degree in pharmacy that does not meet the requirements of subparagraph i but that evidences the successful completion of a program designed to educate and train persons to be practising pharmacists, and,

A. have successfully completed a program that, at the time the applicant commenced it, was approved by the Council as one whose graduates should possess knowledge, skill and judgment at least equal to those of current graduates of a program mentioned in sub-subparagraph i B, or

B. have successfully completed the examination provided for in paragraph 4 of subsection 6 (1) on the applicant's first attempt and have successfully completed any further education or training or combination of education and training that was specified by a panel of the Registration Committee to evidence that the applicant possesses knowledge, skill and judgment at least equal to those of current graduates of a program mentioned in sub-subparagraph i B.

2. Subject to subsections (3) and (4), the applicant must have successfully completed a structured practical training program approved by the Council while holding a certificate of registration as a registered pharmacy student and while under the direct supervision of a preceptor approved by the Registration Committee. O. Reg. 451/10, s. 3.

(2) Subject to subsections (3) and (4) and section 14, the requirements in subsection (1) are non-exemptible. O. Reg. 451/10, s. 3.

(3) With the approval of the Council, all or any part of a structured practical training program referred to in paragraph 2 of subsection (1) may be completed as part of a program referred to in paragraph 1 of subsection (1), even if the applicant did not hold a certificate of registration as a registered pharmacy student at the time. O. Reg. 451/10, s. 3.

(4) An applicant shall be deemed to have met the requirement in paragraph 2 of subsection (1) if, at the time of application, the applicant holds a non-restricted registration as a pharmacist, has held that registration for at least two years and the applicant,

(a) satisfies the Registrar or a panel of the Registration Committee that the applicant engaged in the full scope of practice as a pharmacist in that jurisdiction for at least 600 hours;

(b) successfully completed a structured practical training program which is, in the opinion of a panel of the Registration Committee at least equivalent to a program mentioned in paragraph 2 of subsection (1), or has other education, training or experience that, in the opinion of a panel of the Registration Committee is at least equivalent to the program mentioned in paragraph 2 of subsection (1); or

(c) successfully completed the education and obtained the additional training or experience that a panel of the Registration Committee has specified. O. Reg. 451/10, s. 3.

(5) The requirement of paragraph 2 of subsection (1) shall not be considered to have been met unless the applicant is issued a certificate of registration as an intern within one year of meeting that requirement or within such greater time as is specified by a panel of the Registration Committee. O. Reg. 451/10, s. 3.

(6) An applicant who has a university degree in pharmacy mentioned in subparagraph 1 ii of subsection (1), and who successfully completes any further education or training or combination of education and training specified by a panel of the Registration Committee shall be deemed to have met the requirement in paragraph 1 of subsection (1) if the applicant,

(a) was registered as a registered pharmacy student on December 3, 2010; or

(b) becomes registered as a registered pharmacy student after December 3, 2010 but before December 3, 2011. O. Reg. 451/10, s. 3.

MOBILITY WITHIN CANADA

14. (1) Where section 22.18 of the Health Professions Procedural Code applies to an applicant the requirements of subsection 13 (1) are deemed to have been met by the applicant if he or she provides, for each jurisdiction where the applicant holds an out-of-province certificate, a certificate, letter or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as an intern in that jurisdiction. O. Reg. 451/10, s. 3.

(2) Without in any way limiting the generality of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority that issued the applicant that out-of-province certificate as an intern. O. Reg. 451/10, s. 3.

(3) An applicant referred to in subsection (1) is deemed to meet the requirements of paragraph 1 of subsection 4 (1) where the requirements for the issuance of the applicant's out-of-province certificate included language proficiency requirements equivalent to those required by that paragraph. O. Reg. 451/10, s. 3.

(4) Despite subsection (1), an applicant is not deemed to have met a requirement if that requirement is described in subsection 22.18 (3) of the Health Professions Procedural Code. O. Reg. 451/10, s. 3.

TERMS, CONDITIONS AND LIMITATIONS

15. (1) Every certificate of registration as an intern is subject to the following terms, conditions and limitations:

1. The member shall only engage in the practice of pharmacy,
   i. when practising in a pharmacy to which the *Drug and Pharmacies Regulation Act* applies, while under the direct supervision of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment), or
   ii. in all other cases, while under the supervision of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment).
2. The member shall not supervise that part of the pharmacy where drugs are kept.
3. The member shall not delegate a controlled act. O. Reg. 451/10, s. 3; O. Reg. 187/21, s. 5.

(2) A certificate of registration as an intern automatically expires,

(a) when the member is issued a certificate of registration as a pharmacist; or

(b) one year from the date on which it was issued unless a panel of the Registration Committee specifies otherwise. O. Reg. 451/10, s. 3.

PART V.1
REGISTRATION — PHARMACISTS (EMERGENCY ASSIGNMENT)

15.1 (1) The following are additional requirements for the issuance of a certificate of registration as a pharmacist (emergency assignment):

1. The Government of Ontario must request that the College issue certificates of registration for the pharmacist (emergency assignment) class.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, paragraph 1 of subsection 15.1 (1) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 2)

1. The Minister must have requested that the College initiate registrations under this class based on the Minister's opinion that emergency circumstances call for it or the Council must have determined, after taking into account all of the relevant circumstances that impact the ability of applicants to meet the ordinary registration requirements, that there are emergency circumstances, and that it is in the public interest that the College issue emergency certificates of registration.

2. The applicant must,
   i. have satisfied the educational requirements of paragraph 1 of subsection 6 (1) no more than two years prior to submitting an application for the issuance of a certificate of registration as a pharmacist (emergency assignment),
   ii. currently be practising as a pharmacist in a jurisdiction approved by the Council, and provide, for each jurisdiction where the applicant holds a certificate, a letter, certificate or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as a pharmacist in that jurisdiction, or
   iii. have practised as a pharmacist in a jurisdiction approved by the Council within three years prior to the day on which the applicant met all other requirements for the issuance of a certificate of registration as a pharmacist (emergency assignment), and provide, for each jurisdiction where the applicant held a certificate, a letter, certificate or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant was in good standing as a pharmacist in that jurisdiction. O. Reg. 187/21, s. 6.

(2) Without in any way limiting the generality of subparagraphs 2 ii and 2 iii of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority that issued the applicant's certificate as a pharmacist. O. Reg. 187/21, s. 6.

TERMS, CONDITIONS AND LIMITATIONS

15.2 (1) Every certificate of registration as a pharmacist (emergency assignment) is subject to the following terms, conditions and limitations:

1. The member shall, while working in a pharmacy or any other environment where patient care is being provided, clearly identify himself or herself as a pharmacist (emergency assignment).

2. The member shall only engage in the practice of the profession while under the supervision of a member holding a certificate of registration as a pharmacist listed in Part A.

3. The member shall not be the designated manager of a pharmacy. O. Reg. 187/21, s. 6.

(2) A certificate of registration as a pharmacist (emergency assignment) expires 60 days from the date on which the certificate was issued, unless extended under subsection (3). O. Reg. 187/21, s. 6.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, subsections 15.2 (2) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 3)

(2) A certificate of registration as a pharmacist (emergency assignment) expires on the later of,

(a) 60 days from the date on which the certificate was issued, or extended under subsection (3); and

(b) three months after either the Minister or the Council declares that the emergency circumstances that gave rise to the issuance of certificates of registration in the pharmacist (emergency assignment) class have ended. O. Reg. 295/23, s. 3.

(3) The Registrar may extend a certificate of registration as a pharmacist (emergency assignment) for one or more periods, each of which is not to exceed 60 days, if, in the opinion of the Registrar, it is advisable to do so. O. Reg. 187/21, s. 6.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, subsections 15.2 (3) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 3)

(3) The Registrar may extend a certificate of registration as a pharmacist (emergency assignment) for one or more periods of 60 days as long as emergency circumstances persist. O. Reg. 295/23, s. 3.

(4) The Registrar may revoke a certificate of registration as a pharmacist (emergency assignment) prior to the expiry of the certificate if, in the opinion of the Registrar, it is advisable to do so. O. Reg. 187/21, s. 6.

(5) Where a member listed in Part B of the register also holds a certificate of registration as a pharmacist (emergency assignment), the terms, conditions and limitations listed in section 9 shall not apply to the member during the time that the member holds the emergency assignment certificate. O. Reg. 187/21, s. 6.

(6) Where a member who holds a certificate of registration as an intern also holds a certificate of registration as a pharmacist (emergency assignment), the terms, conditions and limitations listed in subsection 15 (1) shall not apply to the member during the time that the member holds the emergency assignment certificate. O. Reg. 187/21, s. 6.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, the Regulation is amended by adding the following section: (See: O. Reg. 295/23, s. 4)

TRANSFER TO OTHER CLASS OF REGISTRATION

15.3 A member who holds a certificate of registration as a pharmacist (emergency assignment) may apply for a certificate of registration in another class, and a member who does so is exempt from the requirement to pay the fee set out in paragraph 8 of subsection 4 (1). O. Reg. 295/23, s. 4.

PART VI
REGISTRATION — PHARMACY TECHNICIANS

ADDITIONAL REQUIREMENTS

16. (1) The following are additional requirements for the issuance of a certificate of registration as a pharmacy technician:

1. The applicant must,

   i. have successfully completed a pharmacy technician program designed to educate and train persons to be pharmacy technicians which was, at the time the applicant graduated, accredited by the Canadian Council for Accreditation of Pharmacy Programs or such other accrediting body approved by the Council for that purpose,

   ii. have successfully completed a pharmacy technician program designed to educate and train persons to be pharmacy technicians other than one referred to in subparagraph i or have a university degree or university diploma in pharmacy and, in either case,

      A. must have successfully completed a program that, at the time the applicant commenced was approved by the Council as one whose graduates should possess knowledge, skill and judgment at least equivalent to a current graduate of a program referred to in subparagraph i, or

      B. must have successfully completed the examination referred to in paragraph 4 on the applicant's first attempt, and have successfully completed any further education or training or combination of education and training that was specified by a panel of the Registration Committee to evidence that the applicant possesses knowledge, skill and judgment at least equivalent to a current graduate of a program referred to in subparagraph i,

iii. have successfully completed a program before January 1, 2015 that, at the time the applicant commenced was approved by the Council as one whose graduates should possess knowledge, skill and judgment at least equivalent to a current graduate of a program referred to in subparagraph i, or

iv. have met the requirements of paragraph 1 of subsection 6 (1).

2. The applicant must have successfully completed an examination in pharmaceutical jurisprudence approved by the Council for applicants for a certificate of registration as a pharmacy technician.

3. The applicant must have successfully completed a structured practical training program approved by the Council and must have done so under the direct supervision of a preceptor approved by the Registration Committee.

4. The applicant must have successfully completed the qualifying examination of the Pharmacy Examining Board of Canada for pharmacy technicians or successfully completed another examination that has been approved by the Council for that purpose. O. Reg. 451/10, s. 4.

(2) With the approval of the Council, all or any part of a structured practical training program referred to in paragraph 3 of subsection (1) may be completed as part of a program referred to in subparagraph 1i of subsection (1) or sub-subparagraph 1 ii A of subsection (1). O. Reg. 451/10, s. 4.

(3) The requirement in paragraph 1 of subsection (1) must be met within two years before the date on which the applicant met all of the other requirements for the issuance of a certificate of registration as a pharmacy technician unless the applicant,

(a) satisfies the Registrar or a panel of the Registration Committee that the applicant has practised pharmacy within the scope of practice of the profession as set out in section 3 of the Act under the supervision or direction of a pharmacist for at least 600 hours during the three years before the date on which the applicant met all the other requirements for the issuance of a certificate of registration as a pharmacy technician and did so while practising as a pharmacy technician in Canada or in another jurisdiction approved by the Council;

(b) meets any requirements regarding any further education or training or a combination of education or training, if any, set by a panel of the Registration Committee within the time set by the panel; or

(c) successfully completes the examination referred to in paragraph 4 of subsection (1) within three years of the date on which he or she meets all of the other requirements for the issuance of a certificate of registration as a pharmacy technician. O. Reg. 451/10, s. 4.

(4) The requirement of paragraph 2 of subsection (1) shall not be considered to have been met unless the applicant is issued a certificate of registration as a pharmacy technician within three years of meeting that requirement. O. Reg. 451/10, s. 4.

(5) An applicant is only eligible to take the examination referred to in paragraph 2 of subsection (1) three times in any 24-month period. O. Reg. 451/10, s. 4.

(6) The requirement in paragraph 3 of subsection (1) shall not be considered to have been met unless the applicant,

(a) is issued a certificate of registration as a pharmacy technician within two years of meeting that requirement;

(b) satisfies the Registrar or a panel of the Registration Committee that the applicant has practised pharmacy within the scope of practice of the profession as set out in section 3 of the Act under the supervision or direction of a pharmacist for at least 600 hours during the three years before the date on which the applicant met all the other requirements for the issuance of a certificate of registration as a pharmacy technician and did so while practising as a pharmacy technician in Canada or in another jurisdiction approved by the Council; or

(c) meets any requirements regarding any further education or training or a combination of education or training, if any, set by a panel of the Registration Committee within the time set by the panel. O. Reg. 451/10, s. 4.

(7) Subject to subsection (8), paragraph 4 of subsection (1) is not considered to have been met unless the applicant,

(a) successfully completed the examination within three attempts; or

(b) successfully completed the examination on the applicant's fourth attempt having first successfully completed the further education or training or combination of education and training required by the examining body responsible for the administration of the examination or, if no further education or training was required by that body, the further education or training or combination of education and training, if any, that was specified by a panel of the Registration Committee. O. Reg. 451/10, s. 4.

(8) Where, by virtue of subsection (7), an applicant is not considered to have met the requirement in paragraph 4 of subsection (1), the applicant may not attempt the examination again until the applicant successfully completes a new program mentioned in subparagraph 1 i of subsection (1). O. Reg. 451/10, s. 4.

(9) An applicant shall be deemed not to have met the requirement of subparagraph 1 iii of subsection (1) unless, before January 1, 2012 and before commencing the program referred to in that subparagraph, the applicant successfully completed,

(a) the College's Pharmacy Technician Certification Examination;

(b) the Pharmacy Technician Evaluating Examination of the Pharmacy Examining Board of Canada; or

(c) another examination approved by the Council. O. Reg. 451/10, s. 4.

(10) Subject to subsection (2) and section 17, the requirements in subsection (1) are non-exemptible. O. Reg. 451/10, s. 4.

(11) A reference in this section to "all of the other requirements for the issuance of a certificate of registration" includes, without being limited to, a requirement set out in subsection 4 (1) or subsection (1) of this section. O. Reg. 451/10, s. 4.

MOBILITY WITHIN CANADA

17. (1) Where section 22.18 of the Health Professions Procedural Code applies to an applicant the requirements of paragraphs 1, 3 and 4 of subsection 16 (1) are deemed to have been met by the applicant if he or she provides, for each jurisdiction where the applicant holds an out-of-province certificate, a certificate, letter or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as a pharmacy technician in that jurisdiction. O. Reg. 451/10, s. 4.

(2) Without in any way limiting the generality of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority in that issued the applicant that out-of-province certificate as a pharmacy technician. O. Reg. 451/10, s. 4.

(3) An applicant referred to in subsection (1) is deemed to meet the requirements of paragraph 1 of subsection 4 (1) where the requirements for the issuance of the applicant's out-of-province certificate included language proficiency requirements equivalent to those required by that paragraph. O. Reg. 451/10, s. 4.

(4) Despite subsection (1), an applicant is not deemed to have met a requirement if that requirement is described in subsection 22.18 (3) of the Health Professions Procedural Code. O. Reg. 451/10, s. 4.

TERMS, CONDITIONS AND LIMITATIONS

18. Every certificate of registration as a pharmacy technician is subject to the following terms, conditions and limitations:

1. The member shall only engage in the practice of pharmacy,
   i. when practising in a pharmacy to which the *Drug and Pharmacies Regulation Act* applies, other than a remote dispensing location, while under the direct supervision of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment), or
   ii. in all other cases, while under the supervision or direction of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment).
2. When practising in a pharmacy to which the *Drug and Pharmacies Regulation Act* applies the member shall not supervise that part of a pharmacy where drugs are kept.
3. The member shall not delegate a controlled act.
4. The member shall not provide information or education relating to drug use, either to or for a patient, where the provision of the information requires therapeutic knowledge, clinical analysis or clinical assessment. O. Reg. 451/10, ss. 4, 6 (2); O. Reg. 187/21, s. 7.

PART VI.1
REGISTRATION — PHARMACY TECHNICIANS (EMERGENCY ASSIGNMENT)

18.1 (1) The following are additional requirements for the issuance of a certificate of registration as a pharmacy technician (emergency assignment):

1. The Government of Ontario must request that the College issue certificates of registration for the pharmacy technician (emergency assignment) class.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, paragraph 1 of subsection 18.1 (1) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 5)

1. The Minister must have requested that the College initiate registrations under this class based on the Minister's opinion that emergency circumstances call for it or the Council must have determined, after taking into account all of the relevant circumstances that impact the ability of applicants to meet the ordinary registration requirements, that there are emergency circumstances, and that it is in the public interest that the College issue emergency certificates of registration.

2. The applicant must,

i. have satisfied the educational requirements of paragraph 1 of subsection 16 (1) no more than two years prior to submitting an application for the issuance of a certificate of registration as a pharmacy technician (emergency assignment),

ii. currently be practising as a pharmacy technician in a jurisdiction approved by the Council, and provide, for each jurisdiction where the applicant holds a certificate, a letter, certificate or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant is in good standing as a pharmacy technician in that jurisdiction, or

iii. have practised as a pharmacy technician in a jurisdiction approved by the Council within three years prior to the day on which the applicant met all other requirements for the issuance of a certificate of registration as a pharmacy technician (emergency assignment), and provide, for each jurisdiction where the applicant held a certificate, a letter, certificate or other evidence satisfactory to the Registrar or a panel of the Registration Committee confirming that the applicant was in good standing as a pharmacy technician in that jurisdiction. O. Reg. 187/21, s. 8.

(2) Without in any way limiting the generality of subparagraphs 2 ii or 2 iii of subsection (1), "good standing" shall include the fact that,

(a) the applicant is not the subject of any discipline or fitness to practise order or of any proceeding or ongoing investigation or of any interim order or agreement as a result of a complaint, investigation or proceeding; and

(b) the applicant has complied with the continuing competency and quality assurance requirements of the regulatory authority that issued the applicant the certificate as a pharmacy technician. O. Reg. 187/21, s. 8.

TERMS, CONDITIONS AND LIMITATIONS

18.2 (1) Every certificate of registration as a pharmacy technician (emergency assignment) is subject to the following terms, conditions and limitations:

1. The member shall, while working in a pharmacy or any other environment where patient care is being provided, clearly identify himself or herself as a pharmacy technician (emergency assignment).

2. The member shall only engage in the practice of pharmacy,

   i. when practising in a pharmacy to which the *Drug and Pharmacies Regulation Act* applies, other than a remote dispensing location, while under the direct supervision of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment), or

   ii. in all other cases, while under the supervision or direction of a member holding a certificate of registration as a pharmacist or as a pharmacist (emergency assignment).

3. When practising in a pharmacy to which the *Drug and Pharmacies Regulation Act* applies, the member shall not supervise that part of a pharmacy where drugs are kept.

4. The member shall not delegate a controlled act.

5. The member shall not provide information or education relating to drug use, either to or for a patient, where the provision of the information requires therapeutic knowledge, clinical analysis or clinical assessment. O. Reg. 187/21, s. 8.

(2) A certificate of registration as a pharmacy technician (emergency assignment) expires 60 days from the date on which the certificate was issued, unless extended under subsection (3). O. Reg. 187/21, s. 8.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, subsections 18.2 (2) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 6)

(2) A certificate of registration as a pharmacy technician (emergency assignment) expires on the later of,

(a) 60 days from the date on which the certificate was issued or extended under subsection (3); and

(b) 12 months after either the Minister or the Council declares that the emergency circumstances that gave rise to the issuance of certificates of registration in the pharmacy technician (emergency assignment) class have ended. O. Reg. 295/23, s. 6.

(3) The Registrar may extend a certificate of registration as a pharmacy technician (emergency assignment) for one or more periods, each of which is not to exceed 60 days, if, in the opinion of the Registrar, it is advisable to do so. O. Reg. 187/21, s. 8.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, subsections 18.2 (3) of the Regulation is revoked and the following substituted: (See: O. Reg. 295/23, s. 6)

(3) The Registrar may extend a certificate of registration as a pharmacy technician (emergency assignment) for one or more periods of 60 days as long as emergency circumstances persist. O. Reg. 295/23, s. 6.

(4) The Registrar may revoke a certificate of registration as a pharmacy technician (emergency assignment) prior to the expiry of the certificate if, in the opinion of the Registrar, it is advisable to do so. O. Reg. 187/21, s. 8.

Note: On August 31, 2023, the day subsection 3 (3) of Schedule 6 to the *Pandemic and Emergency Preparedness Act, 2022* comes into force, the Regulation is amended by adding the following section: (See: O. Reg. 295/23, s. 7)

TRANSFER TO OTHER CLASS OF REGISTRATION

18.3 A member who holds a certificate of registration as a pharmacy technician (emergency assignment) may apply for a certificate of registration in the pharmacy technician class, and a member who does so is exempt from the requirement to pay the fee set out in paragraph 8 of subsection 4 (1). O. Reg. 295/23, s. 7.

PART VII
SUSPENSIONS, RESIGNATIONS, REINSTATEMENTS, ETC.

ADMINISTRATIVE SUSPENSIONS

19. (1) If a member fails to provide information about the member in the manner and in the form as required under the by-laws, the Registrar may give the member notice of intention to suspend the member and may suspend the member's certificate of registration for failure to provide the information 60 days after notice is given. O. Reg. 451/10, s. 5.

(2) Where the Registrar suspends a member's certificate of registration under subsection (1), the Registrar shall lift the suspension upon being satisfied that the required information has been filed with the College and that any fees required for the lifting of that suspension has been paid. O. Reg. 451/10, s. 5.

20. (1) If, pursuant to the by-laws, the College requests evidence that the member holds professional liability insurance in the amount and in the form as required by the by-laws and the member fails to provide that evidence within 14 days of having been requested to do so, the Registrar shall immediately give the member notice of intention to suspend the member and may suspend the member's certificate of registration for failure to provide the evidence 30 days after notice is given. O. Reg. 451/10, s. 5.

(2) Where the Registrar suspends the member's certificate of registration under subsection (1), the Registrar shall lift that suspension upon being satisfied that the member holds professional liability insurance in the amount and in the form required by the by-laws and that any fee required for the lifting of that suspension has been paid. O. Reg. 451/10, s. 5.

21. Where the Registrar suspends a member's certificate of registration under section 24 of the Health Professions Procedural Code for failure to pay a fee, the Registrar shall lift the suspension upon being satisfied that the member,

(a) has paid all amounts owed to the College;

(b) holds professional liability insurance in the amount and in the form required by the by-laws; and

(c) pays any fees required for the lifting of that suspension. O. Reg. 451/10, s. 5.

DEEMED RESIGNATIONS

22. (1) A member shall be deemed to have resigned where,

(a) the member's certificate of registration was suspended for failure to pay a fee that the member was required to pay in accordance with the regulations or by-laws and that suspension continued for 120 days; or

(b) the member's certificate of registration was suspended pursuant to subsection 19 (1) or subsection 20 (1) and the suspension continued for 60 days. O. Reg. 451/10, s. 5.

(2) The resignation is effective,

(a) in the case of a resignation under clause (1) (a), on the 121st day following the commencement of that suspension;

(b) in the case of a suspension under clause (1) (b), on the 61st day following the commencement of the suspension. O. Reg. 451/10, s. 5.

RETURN OF CERTIFICATE, ETC.

23. A member who resigns, or whose certificate of registration is suspended or revoked shall, if so requested, immediately return to the College,

(a) his or her certificate of registration; and

(b) any card or other form of identification issued to him or her by the College for the purpose of identifying him or her as a member of the College. O. Reg. 451/10, s. 5.

REINSTATEMENT

24. (1) A former member who held a certificate of registration as a pharmacist or pharmacy technician and who resigned as a member of the College may apply for the reinstatement of his or her certificate of registration by submitting a completed application to the Registrar in the form provided by the Registrar. O. Reg. 451/10, s. 5.

(2) Subject to subsections (3), (4) and (6), the Registrar may reinstate the former member's certificate of registration if,

(a) the former member has paid,

(i) the required reinstatement fee,

(ii) the annual fee for the year in which the certificate of registration is to be reinstated, if not previously paid,

(iii) the annual fee for the year in which the former member resigned or was deemed to have resigned, if not previously paid unless the Registrar is satisfied that the former member did not engage in the practice of pharmacy in Ontario during that year, and

(iv) any other money owed by the former member to the College at the date the application for reinstatement is submitted, including, without being limited to, any penalty fees that were due at the time that he or she ceased to be a member and any costs or expenses ordered to be paid under section 53.1 of the Health Professions Procedural Code, any costs awarded to the College by a Court and any amount owing to the College under a by-law or former regulation made under the Act;

(b) the application for reinstatement was submitted to the Registrar within three years of the date on which the former member resigned or in the case of a former member who was deemed to have resigned under subsection 22 (1), three years from the date on which the former member was suspended where that suspension resulted in a deemed resignation; and

(c) the application meets the requirement set out in paragraph 7 of subsection 4 (1) with necessary modifications. O. Reg. 451/10, s. 5.

(3) A former member is ineligible for reinstatement under subsection (2) if he or she,

(a) is the subject of a proceeding for professional misconduct, incompetence or incapacity in Ontario or any like proceeding in any other jurisdiction in relation to the practice of pharmacy or another profession, or was the subject of such a proceeding, other than a proceeding that was completed on its merits;

(b) was, at the time he or she ceased to be a member or at any time since, the subject of a proceeding in respect of,

(i) any criminal offence in any jurisdiction,

(ii) any offence relating to the use, possession or sale of drugs in any jurisdiction,

(iii) any offence arising in any jurisdiction relating to the practice of pharmacy or any other profession or occupation, or

(iv) any offence under the *Controlled Drugs and Substances Act* (Canada);

(c) was, after he or she ceased to be a member, found guilty of,

(i) any criminal offence in any jurisdiction,

(ii) any offence relating to the use, possession or sale of drugs in any jurisdiction,

(iii) any offence arising in any jurisdiction relating to the practice of pharmacy or any other profession or occupation, or

(iv) any offence under the *Controlled Drugs and Substances Act* (Canada);

(d) is the subject of an inquiry or investigation by the Registrar, a committee, a panel of a committee or a board of inquiry of the College, or was the subject of such an inquiry or investigation, that was not completed on its merits or which resulted in the member's resignation;

(e) was, at the time he or she ceased to be a member, the subject of an outstanding order or requirement of a committee, a panel of a committee, or a board of inquiry of the College;

(f) was, at the time he or she ceased to be a member, in breach of an order or requirement of a committee, a panel of a committee, or a board of inquiry of the College;

(g) was, at the time he or she ceased to be a member, in violation of a decision of a panel of the Inquiries, Complaints and Reports Committee or of any predecessor committee, including a decision requiring the member to attend to be cautioned;

(h) was, at the time he or she ceased to be a member, in breach of any written agreement with or undertaking provided to the College; or

(i) had, at the time he or she ceased to be a member, terms, conditions or limitations on his or her certificate of registration, other than those applicable to all members of the class of certificate of registration he or she previously held. O. Reg. 451/10, s. 5.

(4) A former member must meet all of the requirements set out in subsection (2) within one year of submitting his or her application for reinstatement. O. Reg. 451/10, s. 5.

(5) Nothing in this section prevents a former member from making any number of applications for reinstatement or from making an application for a new certificate of registration. O. Reg. 451/10, s. 5.

(6) A former member who is seeking reinstatement of a certificate of registration as a pharmacist and who is otherwise eligible for the reinstatement shall be reinstated into Part B of the register unless the former member satisfies the Registrar that,

(a) the former member did not resign at a time when the member had been selected for but had not successfully completed a practice review under the College's Quality Assurance Program; and

(b) the member had performed at least 600 hours of patient care in Canada, the United States of America or another jurisdiction approved by the Council during the period of three years commencing immediately before the date of the member's resignation. O. Reg. 451/10, s. 5.

REINSTATEMENT, PURSUANT TO ORDER

25. If a former member's certificate of registration is ordered to be reinstated by a panel of the Discipline Committee or of the Fitness to Practise Committee, the Registrar shall reinstate the certificate of registration upon payment of,

(a) the required reinstatement fee; and

(b) the annual fee for the year in which the certificate of registration is to be reinstated, if not previously paid. O. Reg. 451/10, s. 5.

PART VII.1
NOTICES OF MEETINGS AND HEARINGS

NOTICE OF MEETINGS

26. (1) The Registrar shall ensure that notice of every Council meeting that is required to be open to the public under the Act is given in accordance with this section. O. Reg. 451/10, s. 5.

(2) The notice must be published at least 14 days before the date of the meeting in a daily newspaper of general circulation throughout Ontario. O. Reg. 451/10, s. 5.

(3) The notice must be in English and French. O. Reg. 451/10, s. 5.

(4) The notice must contain the following information:

1. The date, time and place of the meeting.

2. A statement of the purpose of the meeting. O. Reg. 451/10, s. 5.

(5) The Registrar shall provide the information contained in the notice to every person who requests it by telephone. O. Reg. 451/10, s. 5.

NOTICE OF HEARINGS

27. (1) The Registrar shall ensure that the information concerning an impending hearing by a panel of the Discipline Committee to deal with allegations of professional misconduct or incompetence made against a member is given, in accordance with this section, to a person who requests the information. O. Reg. 451/10, s. 5.

(2) The information shall be given,

(a) at least 14 days before the date of the hearing, if the request is received 14 days before the date of the hearing; or

(b) as soon as possible after the request is made, if the request is received after that time but before the date of the hearing. O. Reg. 451/10, s. 5.

(3) The information given shall be as follows:

1. The name of the member against whom the allegations have been made.

2. The member's principal place of practice.

3. The date, time and place of the hearing.

4. A statement of the purpose of the hearing. O. Reg. 451/10, s. 5.

(4) The Registrar shall provide the information in French to a person who requests that the information be provided in French, wherever reasonably possible. O. Reg. 451/10, s. 5.

PART VII.2
ADVERTISING

ADVERTISING

28. (1) In this section,

"advertisement" includes an announcement, directory listing or other form of communication similar to an advertisement;

"drug services" means one or more of the compounding, dispensing or sale by retail of drugs or the provision of information or advice with respect to drugs. O. Reg. 451/10, s. 5; O. Reg. 59/11, s. 1 (1, 2).

(2) A member shall not, through any medium, publish, display, distribute or use, or permit, directly or indirectly, the publication, display, distribution or use through any medium of, an advertisement relating to drug services that,

(a) is false, misleading or deceptive, whether as a result of the inclusion of information or the omission of information;

(b) is not readily comprehensible to the persons to whom it is directed;

(c) is not dignified and in good taste;

(d) contains anything that cannot be verified;

(e) contains testimonials, comparative statements or endorsements;

(f) contains a reference to a member's area of practice or to a procedure or treatment available from a member practising in the pharmacy, unless the advertisement discloses whether or not the member has an area of expertise and, if the member does have such an area of expertise, the particular expertise;

(g) contains references to a particular brand of equipment used to assist in providing drug services;

(h) contains information that is not relevant to the choice of a pharmacist; or

(i) contains any representations as to the safety or effectiveness or an indication for use of any drug referred to in Schedule I established by the regulations under the *Drug and Pharmacies Regulation Act*.

(j) REVOKED: O. Reg. 59/11, s. 1 (4).

O. Reg. 451/10, s. 5; O. Reg. 59/11, s. 1 (3, 4).

(3) An advertisement by a member that includes price information relating to drugs referred to in Schedule I established by the regulations under the Drug and Pharmacies Regulation Act shall include the price information for at least 15 different drugs, 10 of which each belong to a different one of the following drug classifications:

1. Anti-infective agents.
2. Antineoplastic agents.
3. Autonomic agents.
4. Blood formation and coagulation drugs.
5. Cardiovascular drugs.
6. Central nervous system drugs.
7. Diagnostic agents.
8. Electrolytic, caloric and water balance drugs.
9. Cough preparations.
10. Eye, ear, nose and throat preparations.
11. Gastrointestinal drugs.
12. Gold compounds.
13. Heavy metal antagonists.
14. Hormones and substitutes.
15. Oxytocics.
16. Skin and mucous membrane preparations.
17. Spasmolytics.
18. Unclassified therapeutic agents.
19. Vitamins. O. Reg. 451/10, s. 5; O. Reg. 59/11, s. 1 (5).

(4) Where an advertisement by a member includes price information relating to drugs referred to in Schedule I established by the regulations under the *Drug and Pharmacies Regulation Act*, the advertisement shall include at a minimum the following information with respect to each drug:

1. The quantity of the drug being advertised at the advertised price.

2. The total cost for the drug to the purchaser including any dispensing fee.

3. The time period during which the advertised price will be available. O. Reg. 59/11, s. 1 (6).

(5) An advertisement by a member that includes price information relating to drugs referred to in Schedule I established by the regulations under the *Drug and Pharmacies Regulation Act* shall include, in addition to the price information referred to in subsection (4), the following information with respect to each advertised drug:

1. The strength of the drug.

2. The brand name of the drug.

3. The dosage form of the drug. O. Reg. 59/11, s. 1 (6).

(6) Where an advertisement by a member includes price information relating to drugs referred to in Schedule I established by the regulations under the *Drug and Pharmacies Regulation Act,* equal prominence shall be given to each drug and, for each of those drugs, equal prominence shall be given to all the information required under subsections (4) and (5). O. Reg. 59/11, s. 1 (6).

(7), (8) REVOKED: O. Reg. 59/11, s. 1 (6).

PROFESSIONAL MISCONDUCT RE ADVERTISING

29. It is professional misconduct for the purposes of clause 51 (1) (c) of the Health Professions Procedural Code for a member who advertises price information with respect to a drug referred to in Schedule I established by the regulations under the *Drug and Pharmacies Regulation Act* to charge any purchaser, including the executive officer under the *Ontario Drug Benefit Act* more for the drug than the member has advertised, pursuant to paragraph 2 of subsection 28 (4), as the total cost for the drug to the purchaser including any dispensing fee. O. Reg. 59/11, s. 2.

CLARIFICATION RE APPLICATION OF PART

30. Nothing in this Part prohibits a member from publishing, displaying, distributing or using, or permitting directly or indirectly the publication, display, distribution or use of, an advertisement that relates solely to the co-payment or dispensing fee charged by the member for supplying a drug that is a listed drug product under the *Ontario Drug Benefit Act* to an eligible person under that Act. O. Reg. 451/10, s. 5.

PART VII.3
CONTROLLED ACTS

INTERPRETATION

31. (1) In this Part,

"adapt" means, subject to subsection (2), to change a patient's prescription respecting,

(a) the dose of the prescribed drug,

(b) the dosage form of the prescribed drug,

(c) the directions for use of the prescribed drug, or

(d) the route of administration for taking the prescribed drug,

but does not include therapeutic substitution;

Note: On September 30, 2026, the definition of "adapt" in section 31 of the Regulation is amended by striking out "subject to subsection (2)" in the portion before clause (a). (See: O. Reg. 126/20, s. 1 (2) and O. Reg. 766/21, s. 2)

"coronavirus exemption" means the exemption issued by the Minister of Health for Canada on March 19, 2020 under subsection 56 (1) of the *Controlled Drugs and Substances Act* (Canada) entitled "Subsection 56(1) Class Exemption for Patients, Practitioners and Pharmacists Prescribing and Providing Controlled Substances in Canada During the Coronavirus Pandemic", available on a website of the Government of Ontario, including any renewal or replacement of the exemption;

Note: On September 30, 2026, the definition of "coronavirus exemption" in section 31 of the Regulation is revoked. (See: O. Reg. 126/20, s. 1 (4) and O. Reg. 766/21, s. 2)

"Part A pharmacist" means a member who holds a certificate of registration as a pharmacist and who is listed in Part A of the register;

"point-of-care test" means a test that employs a medical device authorized by the Minister of Health for Canada for point-of-care use;

"prescriber" means a person who is authorized under the laws of a province or territory of Canada to give a prescription within the scope of his or her practice of a health profession;

"prescription" means a direction from a prescriber directing the dispensing of a drug or mixture of drugs for a specific patient;

"renew" means to provide a patient with a prescription that repeats a prescription previously provided to that patient;

"therapeutic substitution" means the substitution of a drug that contains chemically different active ingredients that are considered to be therapeutically equivalent. O. Reg. 302/12, s. 1; O. Reg. 126/20, s. 1 (1, 3); O. Reg. 46/22, s. 1.

(2) While the coronavirus exemption is in effect, in this Part,

"adapt", in relation to the adaptation of a prescription for a controlled substance under the *Controlled Drugs and Substances Act* (Canada), means to change the prescription respecting,

(a) the dose and regime of the prescribed drug,

(b) the dosage form of the prescribed drug,

(c) the de-prescribing of the prescribed drug, or

(d) the part-filling of the prescription,

but does not include therapeutic substitution. O. Reg. 126/20, s. 1 (5)

Note: On September 30, 2026, subsection 31 (2) of the Regulation is revoked. (See: O. Reg. 126/20, s. 1 (6) and O. Reg. 766/21, s. 2)

32. (1) Where the provisions of this Part are inconsistent with a law of Canada respecting prescriptions, including those related to a targeted substance, the law of Canada shall prevail and the provisions of this Part to the extent they are inconsistent with that law shall not apply. O. Reg. 302/12, s. 1.

(2) Where the provisions of this Part are inconsistent with the provisions of the *Narcotics Safety and Awareness Act, 2010*, the provisions of that Act shall prevail and the provisions of this Part, to the extent they are inconsistent with that Act, shall not apply. O. Reg. 302/12, s. 1.

(3) In this Part,

(a) a reference to a part A pharmacist includes a member who holds a certificate of registration as a pharmacist (emergency assignment); and

(b) a reference to a pharmacy technician includes a member who holds a certificate of registration as a pharmacy technician (emergency assignment). O. Reg. 187/21, s. 9.

CONTROLLED ACTS

33. A member shall not perform a controlled act under paragraph 2, 3, 4 or 5 of subsection 4 (1) of the Act except in accordance with this Part. O. Reg. 302/12, s. 1.

34. (1) For the purposes of paragraph 2 of subsection 4 (1) of the Act, a member referred to in subsection (2) who meets all the requirements in subsection (3) is authorized to perform the following acts:

1. Administering a substance specified in Schedule 1 by injection ~~to a patient~~.
2. Administering a substance specified in Schedule 2 by inhalation ~~to a patient. O. Reg. 452/16, s. 1 (1).~~.
3. Administering an influenza vaccine by injection.
4. Administering one of the vaccines specified in Schedule 3 by injection.

(2) A Part A pharmacist, an intern or a registered pharmacy student is authorized to perform an act provided for in ~~subsections~~subsection (1~~), (4) and (5~~), subject to the terms, conditions and limitations imposed on his or her certificate of registration. O. Reg. 302/12, s. 1; O. Reg. 452/16, s. 1 (2).

(2.1) A pharmacy technician who meets all the requirements in subsection (4) is authorized to perform an act provided for in paragraphs 3 and 4 of subsection (~~4.1~~1), subject to the terms, conditions and limitations imposed on ~~their~~his or her certificate of registration. O. Reg. 766/21, s. 1.

(3) A member referred to in subsection (2) may only perform an act provided for in subsection (1) if he or she complies with the following:

1. Before performing the act, the member must receive an informed consent from the patient or the patient's authorized agent.
   i. ~~must explain that purpose to the patient or his or her authorized agent, and~~
   ii. ~~must receive an informed consent from the patient or his or her authorized agent.~~
2. The member shall ensure that he or she only performs the act in an environment that is clean, safe, private and comfortable for the patient.
3. The member shall ensure that appropriate infection control procedures are in place.
4. The member must possess sufficient knowledge, skill and judgment respecting the substance to be administered, and sufficient understanding of the condition of the patient, to be able to administer the substance safely.

5. The member must consider whether administering a substance by injection or inhalation to the patient is appropriate, given the known risks and benefits to the patient and the safeguards and resources available to safely manage the outcome after administration and any other relevant circumstances.
6. The member must maintain a patient record that includes,
   i. the name and address of the patient,
   ii. the name and address of the member,
   iii. the date the act was performed,
   iv. the name, strength (where applicable) and quantity of the substance that the member administered to the patient,
   v. the circumstances relating to the administration of the substance to the patient and any adverse reaction experienced by the patient, and
   vi. confirmation that an informed consent was given by the patient or his or her authorized agent.
7. Where administering a substance specified in Schedule 1 by injection to a patient through an established central or peripheral venous access device, the member must only do so in collaboration with a member of the College of Nurses of Ontario who is a registered nurse in the extended class or a member of the College of Physicians and Surgeons of Ontario.
8. Where the act is performed for a purpose other than that of patient education or demonstration the member must, within a reasonable time after performing the act, notify the following persons that the member performed the act, and provide details respecting the act:
   i. The prescriber, if any, of the substance that was administered.
   ii. The patient's primary care provider, where the member knows that the patient has such a care provider other than the prescriber. O. Reg. 302/12, s. 1; O. Reg. 95/23, s. 1 (1, 2).
9. Where administering an influenza vaccine by injection, the member must administer the vaccine in accordance with Ontario's Universal Influenza Immunization Program as described on the Ministry's website.

(3.1) Where a limitation or a route of administration is indicated with respect to a substance listed in Schedule 1, a member shall only administer the substance in compliance with the limitation and in accordance with the route of administration specified. O. Reg. 95/23, s. 1 (3).

(4) ~~For the purposes of paragraph 2 of subsection 4 (1) of the Act, a~~A member referred to in subsection (~~2) is authorized to administer influenza vaccine by injection to a patient who is two years of age or older, if the member~~2.1) may only perform an act provided for in paragraphs 3 or 4 of subsection (1) if he or she,

~~(a) administers the vaccine in accordance with Ontario's Universal Influenza Immunization Program as described on the Ministry's website;~~

~~(b) receives an informed consent from the patient or his or her authorized agent; and~~

~~(c) meets all the requirements in paragraphs 2 to 6 of subsection (3). O. Reg. 302/12, s. 1; O. Reg. 452/16, s. 1 (3); O. Reg. 742/20, s. 1.~~

~~(4.1) For the purpose of paragraph 2 of subsection 4 (1) of the Act, a member referred to in subsection (2.1) is authorized to administer influenza vaccine by injection to a patient who is two years of age or older, if the member,~~

~~(a) administers the vaccine in accordance with Ontario's Universal Influenza Immunization Program as described on the Ministry's website;~~

(~~b~~a) possesses sufficient knowledge, skill and judgment to be able to administer the ~~influenza~~ vaccine safely;

(~~c~~b) meets all the requirements in paragraphs 2, 3, and 6 of subsection (3); ~~and~~

(c) meets the requirement in paragraph 9 of subsection (3), where the member is administering an influenza vaccine by injection; and

(d) has confirmed that a member referred to in subsection (2), or another regulated health professional authorized to administer the ~~influenza~~ vaccine by injection, has,

(i) received an informed consent from the patient or the patient's authorized agent,

(ii) a sufficient understanding of the ~~influenza~~ vaccine and condition of the patient for the ~~influenza~~ vaccine to be administered safely, and

(iii) considered whether administering the ~~influenza~~ vaccine by injection to the patient is appropriate, given the known risks and benefits to the patient and the safeguards and resources available to safely manage the outcome and any other relevant circumstances. O. Reg. 766/21, s. 1.

~~(5) For the purposes of paragraph 2 of subsection 4 (1) of the Act, a member referred to in subsection (2) is authorized to administer a vaccine from one of the vaccines specified in Schedule 3 by injection to a patient who is five years of age or older, if the member,~~

~~(a) receives an informed consent from the patient or his or her authorized agent;~~

~~(b) meets all the requirements in paragraphs 2 to 6 of subsection (3); and~~

~~(c) notifies the patient's primary care provider (if any) within a reasonable time that the member administered a vaccine to the patient and provides details respecting the administration. O. Reg. 452/16, s. 1 (4).~~

35. (1) For the purposes of paragraph 3 of subsection 4 (1) of the Act, a member referred to in subsection (3) who ~~complies with the other~~meets all the requirements ~~of this section~~in subsection (4) is authorized to prescribe the following drugs:

1. For the sole purpose of smoking cessation, the following specified drugs:
   i. Varenicline Tartrate.
   ii. Bupropion Hydrochloride.
2. For the sole purpose of treating a minor ailment listed in Column 1 of the Table to Schedule 4~~,~~: a drug in a class of drugs listed opposite the minor ailment in Column 2 of that Table. O. Reg. 460/22, s. 1 (1).
3. For the sole purpose of treating COVID-19: Nirmatrelvir/ritonavir.
4. For the sole purpose of treating influenza: Oseltamivir.

Note: On October 1, 2023, paragraph 2 of subsection 35 (1) of the Regulation is amended by striking out "in a class of drugs opposite the minor ailment in Column 2" and substituting "opposite the minor ailment in Column 3". (See: O. Reg. 179/23, s. 1)

(2) REVOKED: O. Reg. 460/22, s. 1 (2).

(3) A Part A pharmacist is authorized to perform an act provided for in subsection (1), subject to the terms, conditions and limitations imposed on his or her certificate of registration.

~~(3) A Part A pharmacist, an~~3.1) An intern or a registered pharmacy student who meets all the requirements in subsection (4) is authorized to perform ~~the~~an act provided for in paragraphs 1, 2 and 4 of subsection (1), subject to the terms, conditions and limitations imposed on his or her certificate of registration. O. Reg. 302/12, s. 1.

(4) A member may only prescribe a drug under this section if he or she,

(a) possesses sufficient knowledge, skill and judgment respecting the drug and the patient's condition to prescribe the drug for the patient;

(b) has considered whether prescribing the drug for the patient is appropriate, given the known risks and benefits of prescribing the drug for the patient and other relevant factors respecting the patient;

(c) gives the prescription to the patient or his or her authorized agent;

(d) advises the patient or his or her authorized agent, at the time of giving the prescription, that he or she may elect to take it to a pharmacy of his or her choosing for dispensing;

(e) notifies the patient's primary care provider (if any) within a reasonable time, that the member prescribed a drug for the patient and provides details respecting the prescription;

(f) complies with the additional requirements under sections 37 and 38; and

(g) ~~in the case of a drug referred to in paragraph 2 of subsection (1),~~ has determined, through a therapeutic assessment, that the drug is the most appropriate treatment for the patient's ~~minor ailment~~condition. O. Reg. 302/12, s. 1; O. Reg. 460/22, s. 1 (3).

36. (1) For the purposes of paragraph 4 of subsection 4 (1) of the Act, a member referred to in subsection (3) who complies with the other provisions of this section is authorized to perform the following acts:

1. Adapting a patient's prescription.
2. Renewing a patient's prescription for the purpose of continuity of care. O. Reg. 302/12, s. 1.

(2) Subject to subsection (2.1), subsection (1) does not authorize a member referred to in subsection (3) to adapt or renew a prescription for a controlled substance as defined in the *Controlled Drugs and Substances Act* (Canada) or a drug designated as a monitored drug by the regulations under the *Narcotics Safety and Awareness Act, 2010*. O. Reg. 302/12, s. 1; O. Reg. 126/20, s. 2 (1).

Note: On September 30, 2026, subsection 36 (2) of the Regulation is amended by striking out "Subject to subsection (2.1)" at the beginning. (See: O. Reg. 126/20, s. 2 (2) and O. Reg. 766/21, s. 2)

(2.1) During the period of time in which the coronavirus exemption is in effect, subsection (2) does not apply to the extent that the coronavirus exemption or the *Controlled Drugs and Substances Act* (Canada) authorizes the member to adapt or renew a prescription for a controlled substance under that Act. O. Reg. 126/20, s. 2 (3).

Note: On September 30, 2026, subsection 36 (2.1) of the Regulation is revoked. (See: O. Reg. 126/20, s. 2 (4) and O. Reg. 766/21, s. 2)

(3) A Part A pharmacist, an intern or a registered pharmacy student is authorized to perform an act provided for in subsection (1), subject to the terms, conditions and limitations imposed on his or her certificate of registration. O. Reg. 302/12, s. 1.

(4) A member may only perform an act provided for in subsection (1) if he or she complies with the following:

1. The member must either possess the patient's prescription to be adapted or renewed or,
    i. receive a copy of the prescription directly from the pharmacy where the prescription was dispensed to the patient,
    ii. be satisfied based on verbal confirmation from a pharmacist at the pharmacy where the prescription was dispensed to the patient as to the existence and details of the prescription,
    iii. have access to the medical record that contains information about the prescription, or
    iv. during the period of time in which the coronavirus exemption is in effect, if the criteria set out in subparagraphs i, ii and iii cannot be met, be satisfied as to the existence and details of the prescription from an alternative source, including, but not limited to, the prescription label, the prescription receipt with medication history, a photograph of the prescription or a facsimile of the prescription.

Note: On September 30, 2026, paragraph 1 of subsection 36 (4) of the Regulation is amended by adding "or" at the end of subparagraph ii, by striking out "or" at the end of subparagraph iii and by revoking subparagraph iv. (See: O. Reg. 126/20, s. 2 (6) and O. Reg. 766/21, s. 2)

2. If the member is renewing a prescription, he or she must not prescribe a quantity of the drug that exceeds the lesser of,
    i. the quantity that was originally prescribed, including any refills that were authorized by the prescriber, and
    ii. a 12 months' supply.
3. The member must, within a reasonable time, notify the prescriber identified on the prescription adapted or renewed by the member, as well as the patient's primary care provider if the member knows that the patient has such a care provider other than the prescriber, providing details about the patient's prescription, when the member,
    i. renews a patient's prescription, or
    ii. adapts a patient's prescription, if, in the member's opinion,
        A. adapting the prescription is clinically significant in relation to the patient, or
        B. the notification is necessary to support the patient's care.
4. At the time that the member adapts or renews the patient's prescription, the member must advise the patient or his or her authorized agent,
    i. that he or she is entitled to the prescription, and
    ii. that he or she may take the prescription to a pharmacy of his or her choosing for dispensing.
5. The member must comply with the additional requirements under sections 37 and 38. O. Reg. 302/12, s. 1; O. Reg. 126/20, s. 2 (5); O. Reg. 742/20, s. 2.

37. A member who performs an act provided for in section 35 or 36 must ensure that the following information is recorded on the prescription:

1. The name and address of the patient for whom the drug is prescribed.
2. The name, strength (where applicable) and quantity of the prescribed drug.
3. Directions for the use of the drug, including its dose, frequency, route of administration and any special instructions.
4. The name, address, telephone number and College registration number of the member issuing the prescription.
5. The date the prescription was issued by the member.
6. If applicable, reference to the prescription that the member adapted or renewed, including the name and contact details of the original prescriber.
7. The number of refills that the member authorized, if applicable.
8. Any other information required by law. O. Reg. 302/12, s. 1.

38. A member who performs an act under section 35 or 36 must maintain a patient record that includes details of the member's rationale for his or her decision to act under section 35 or 36 and the following information, if applicable:

1. Reference to, or a copy of, the patient's prescription that the member renewed or adapted, including the name and contact information of the prescriber.

2. A copy of the prescription that the member gave to the patient or his or her authorized agent ~~under clause 35 (4) (c) or that the member gave to the patient or his or her authorized agent to take to a pharmacy of their choosing under clause 35 (4) (d) or paragraph 4 of subsection 36 (4)~~.

3. A record of the results of laboratory or other tests that the member considered in making the decision to act under section 35 or 36.

4. The date on which the member notified the following persons, as applicable, and the method by which the notification occurred:

   i. The patient's primary care provider ~~notified under clause 35 (4) (e) or paragraph 3 of subsection 36 (4)~~.

   ii. The patient's prescriber ~~notified under paragraph 3 of subsection 36 (4)~~. O. Reg. 302/12, s. 1; O. Reg. 460/22, s. 2.

39. (1) For the purposes of paragraph 5 of subsection 4 (1) of the Act, a member referred to in subsection (2) who meets all the requirements in subsection (4) is authorized to perform the act of piercing a patient's dermis with a lancet-type device to obtain blood. O. Reg. 302/12, s. 1.

(2) A member who is a Part A pharmacist, an intern, a registered pharmacy student or a pharmacy technician is authorized to perform the act provided for in subsection (1), subject to the terms, conditions and limitations imposed on his or her certificate of registration. O. Reg. 302/12, s. 1.

(3) A pharmacy technician shall not perform the act provided for in subsection (1) unless,

(a) a Part A pharmacist is physically present on the premises at the time when the pharmacy technician performs the act;

(b) the pharmacy technician is under the direction of a Part A pharmacist at the time when the pharmacy technician performs the act; and

(c) if the act is performed to administer a point-of-care test, a Part A pharmacist interprets the results of the test and makes any professional decision arising from those results. O. Reg. 302/12, s. 1; O. Reg. 46/22, s. 2 (1).

(4) A member may only perform the act provided for in subsection (1) if he or she complies with the following:

1. The member may only perform the act for the purpose of demonstrating the appropriate use of lancet-type devices for the patient's self-care and education or for the patient's self-monitoring of his or her chronic disease, unless the act is performed to administer a point-of-care test.

1.1 The member may only perform the act to administer a point-of-care test if the test is listed in subsection 28 (2) of Ontario Regulation 45/22 and if it is administered for the purpose of assisting patients with the management of their medication to treat chronic disease.

1.2 Before performing an act described in paragraphs 1 or 1.1, the member must,

   i. explain the purpose to the patient or his or her authorized agent, and

   ii. receive an informed consent from the patient or his or her authorized agent.

2. The member shall ensure that he or she only performs the act in an environment that is clean, safe, private and comfortable for the patient.

3. The member shall ensure that appropriate infection control procedures are in place.

4. The member must possess the knowledge, skill and judgment respecting the performance of the act and understand the condition of the patient, to perform it safely and effectively.

5. The member must consider whether performing the act on the patient is appropriate, given the known risks and benefits to the patient and the safeguards and resources available to safely manage the outcome and any other relevant circumstances.

6. The member must maintain a patient record that includes,

   i. the name and address of the patient,

   ii. the name and work address of the member,

   iii. the date the act was performed,

   iv. the circumstances relating to the performance of the act and any adverse reaction experienced by the patient,

   v. confirmation that an informed consent was given by the patient or his or her agent, and

   vi. if the act was performed to administer a point-of-care test,

A. the results of the test, and

B. the professional decision arising from the results of the test and the rationale for the decision.

7. If the act is performed to administer a point-of-care test, the member must notify the patient's primary care provider (if any) within a reasonable time that the member performed the act and provide details respecting the act. O. Reg. 302/12, s. 1; O. Reg. 46/22, s. 2 (2, 3).

40. REVOKED: O. Reg. 451/10, s. 5.

PART VIII
QUALITY ASSURANCE

GENERAL

41. In this Part,

"assessor" means an assessor appointed under section 81 of the Health Professions Procedural Code;

"Committee" means the Quality Assurance Committee. O. Reg. 98/98, s. 2.

42. The Committee shall administer the quality assurance program, which shall include the following components:

1. Maintenance of a portfolio of continuous learning.
2. Maintenance of a two-part register for pharmacist members.
3. Practice review and remediation.
4. Remediation of behaviour and remarks of a sexual nature. O. Reg. 98/98, s. 2.

CONTINUOUS LEARNING PORTFOLIO

43. (1) A pharmacist shall maintain a portfolio of continuous learning activities in accordance with guidelines on such activities published by the College and distributed to the members.

(2) A pharmacist shall submit the portfolio to the College on request. O. Reg. 98/98, s. 2.

TWO-PART REGISTER FOR PHARMACISTS

44. (1) The part of the College's register that lists pharmacists shall have a Part A (patient care) and a Part B (no patient care). O. Reg. 451/10, s. 7.

(2) Every pharmacist shall be listed in either Part A or Part B. O. Reg. 451/10, s. 7.

45. (1) Upon being issued a certificate of registration as a pharmacist for the first time, the member shall ask to be listed in Part A or Part B of the register by completing and submitting the form provided by the Registrar. O. Reg. 451/10, s. 7.

(2) Every year at the time of paying the annual membership fee, a pharmacist shall ask for a renewal of his or her listing in Part A or Part B or for a transfer to the other Part. O. Reg. 451/10, s. 7.

(3) A member who asks for a renewal of a listing in Part A after the third anniversary of being issued a certificate of registration as a pharmacist for the first time shall not be listed in that Part unless he or she has dispensed, sold or compounded drugs, provided non-prescription drugs, health care aids and devices or information related to drug use for at least 600 hours during the preceding three years in the course of providing patient care while practising the profession in Canada. O. Reg. 451/10, s. 7.

46. (1) A pharmacist may ask for a transfer from Part A of the register to Part B or from Part B to Part A at any time. O. Reg. 451/10, s. 7.

(2) If a member listed in Part A asks for a transfer to Part B, the member shall be transferred to Part B. O. Reg. 451/10, s. 7.

(3) If a member listed in Part B asks for a transfer to Part A, the member shall be transferred to Part A if he or she,

(a) undergoes a practice review in accordance with section 47; and

(b) satisfies the educational and practice requirements that may be specified by the Quality Assurance Committee. O. Reg. 451/10, s. 7.

(4) If the Registrar proposes to reject a request for a transfer to Part A, the request shall be referred to a panel of the Quality Assurance Committee. O. Reg. 451/10, s. 7.

(5) The member shall be given a reasonable opportunity to make written submissions to the panel before it makes a decision. O. Reg. 451/10, s. 7.

(6) A member whose request to be listed in Part A is rejected by the panel may appeal to another panel of the Quality Assurance Committee. O. Reg. 451/10, s. 7.

(7) No member of a panel that rejects a request to be listed in Part A shall sit on a panel hearing an appeal of that decision. O. Reg. 451/10, s. 7.

(8) On an appeal, the member shall be given a reasonable opportunity to make written submissions to the panel before it makes a decision. O. Reg. 451/10, s. 7.

PRACTICE REVIEW AND REMEDIATION

47. (1) Each year the College shall select at random the names of pharmacists required to undergo a practice review.

(2) A pharmacist listed in Part A is required to undergo a practice review if his or her name is selected at random or the member is referred to the Committee by the Complaints Committee or Executive Committee.

(3) If a pharmacist listed in Part A fails to undergo a required practice review, the Committee may transfer the pharmacist to Part B after giving him or her a reasonable opportunity to make written submissions.

(4) A pharmacist listed in Part B is required to undergo a practice review if he or she is referred to the Committee by the Complaints Committee or Executive Committee or if the pharmacist has asked to be listed in Part A under subsection 46 (3).

(5) The Committee shall appoint an assessor to conduct a practice review.

(6) The assessor shall prepare a written report on the review and submit it to the Committee.

(7) After considering the report, the Committee may decide,

(a) that no further action is required;

(b) that the pharmacist is required to undertake the remediation specified by the Committee to correct any deficiency in his or knowledge, skills or judgment identified by the review; or

(c) that the pharmacist is to be listed in Part A where the review took place pursuant to a request to be listed in Part A.

(8) If the Committee proposes to require a pharmacist to undertake remediation under clause (7) (b), it shall not do so unless,

(a) the pharmacist has been given a report of the results of the review;

(b) the pharmacist has been given written notice of the Committee's intention to require him or her to undertake remediation;

(c) the pharmacist has been given at least 14 days from receipt of the notice to make written submissions to the Committee; and

(d) the Committee has considered any such submissions.

(9) After the pharmacist undertakes the specified remediation, the Committee may require him or her to undergo another practice review by an assessor, and subsections (6), (7) and (8) apply to that review. O. Reg. 98/98, s. 2.

48. (1) If the Committee requires a pharmacist to undertake remediation under section 47 and the pharmacist either fails to do so or fails to successfully complete the remediation, the Committee may direct the Registrar to impose terms, conditions or limitations on the pharmacist's certificate of registration for a specified period not exceeding six months.

(2) If the Committee proposes to make a direction under subsection (1), it shall not do so unless,

(a) the pharmacist has been given written notice of its intention;

(b) the pharmacist has been given at least 14 days from receipt of the notice to make written submissions to the Committee or to request an appearance before the Committee in order to make oral submissions; and

(c) the Committee has considered any such submissions.

(3) A pharmacist who requests an appearance under clause (2) (b) shall be given a reasonable opportunity to appear but the Committee may dispose of the matter if he or she has been given a reasonable opportunity to appear and does not.

(4) If the period specified under subsection (1) expires and the pharmacist still has not undertaken or successfully completed the remediation, the Committee may report him or her to the Executive Committee and provide it with such information as it considers appropriate, except information that may not be disclosed under section 83 of the Health Professions Procedural Code.

(5) If the Registrar imposes terms, conditions or limitations on a pharmacist's certificate of registration for a specified period pursuant to a direction given by the Committee under subsection (1), the Committee may direct the Registrar to remove the terms, conditions or limitations before the end of the specified period if the Committee is satisfied that they are no longer needed.

(6) After directing the imposition of terms, conditions or limitations on a pharmacist's certificate of registration for a specified period not exceeding six months under subsection (1), the Committee may direct the imposition of terms, conditions or limitation on the pharmacist's certificate of registration for a second specified period not exceeding six months under

subsection (1) but, after having done so, the Committee shall not direct the imposition of terms, conditions or limitations on the pharmacist's certificate of registration for any further specified period.

(7) If the Committee directs a second imposition of terms, conditions or limitations on the pharmacist's certificate, subsections (2), (3), (4) and (5) apply with respect to the second imposition. O. Reg. 98/98, s. 2.

REMEDIATION OF BEHAVIOUR AND REMARKS OF A SEXUAL NATURE

49. (1) This section applies to matters referred to the Committee by,

(a) a panel of the Complaints Committee under subsection 26 (3) of the Health Professions Procedural Code; and

(b) the Executive Committee under section 79.1 of the Code.

(2) The chair of the Committee shall establish a panel from among the members of the Committee for the purpose of considering a matter referred to in subsection (1).

(3) The chair of the Committee shall appoint a mediator to attempt to resolve the matter.

(4) If the mediator is unable to resolve the matter within 90 days after being appointed, the mediator shall report the failure to the chair without delay and provide the chair with a written report on the mediation.

(5) The chair shall give the member complained against a copy of the mediator's report and a notice advising him or her of the right to make written submissions to the panel.

(6) The member shall be given at least 14 days after receipt of the mediator's report and recommendations to make written submissions to the panel or to request an appearance before the panel to make oral submissions, or to do both.

(7) A member who requests an appearance shall be given a reasonable opportunity to make an appearance, but the panel may dispose of the matter without such appearance if the member has been given a reasonable opportunity to appear.

(8) If the mediation concerns a matter referred by the Complaints Committee, the chair shall give the complainant a copy of the mediator's report.

(9) A mediator's proposed resolution of a matter referred to the Committee by the Complaints Committee must be acceptable to the complainant, the member complained against and the panel.

(10) A mediator's proposed resolution of a matter referred to the Committee by the Executive Committee must be acceptable to the member complained against and the panel.

(11) After considering the mediator's report and any written or oral submissions, the panel may require the member to undergo an assessment for the purpose of establishing if he or she requires education with respect to sexual abuse.

(12) The assessment shall be carried out by an assessor appointed by the Committee.

(13) The assessor shall provide a written report to the panel and shall make such recommendations as the assessor considers appropriate about the member's need for education with respect to sexual abuse.

(14) A copy of the report and recommendations, and a notice informing him or her of the right to make submissions in accordance with subsections (6) and (7), shall be provided to the member.

(15) After considering the assessor's report and recommendations and the member's submissions, if any, the panel may require the member to attend or participate in a sexual abuse education program.

(16) If the panel proposes to take action under subsection (15), the member has the right to make submissions in accordance with subsections (6) and (7). O. Reg. 98/98, s. 2.

50. (1) If a member refuses to undergo an assessment under subsection 49 (11) or to attend or participate in a program under subsection 49 (15), the panel may direct the Registrar to impose terms, conditions or limitations on the member's certificate of registration for a specified period not exceeding six months.

(2) If the panel proposes to take action under subsection (1), the member has the right to make submissions in accordance with subsections 49 (6) and (7).

(3) If the panel is satisfied that the terms, conditions and limitations imposed on a member's certificate or registration are no longer needed, it shall direct the Registrar to remove them before the end of the specified period.

(4) If, at the end of the specified period, the member continues to refuse to undergo the required assessment or to attend or participate in the program, the panel shall refer the matter to the Executive Committee. O. Reg. 98/98, s. 2.

PANEL REQUIREMENTS

51. (1) The Committee may sit as a panel to consider a report on a practice review or any matter arising out of a practice review, a matter relating to the imposition of terms, conditions or limitations on a member's registration under section 48 or a matter under section 49.

(2) A panel shall have at least three members appointed by the chair of the Committee from among the Committee members; at least one member of the panel shall be a member appointed to the Committee by the Lieutenant Governor in Council.

(3) Three members of a panel constitute a quorum. O. Reg. 98/98, s. 2.

PART IX
INSPECTION OF DRUG PREPARATION PREMISES

Temporal Application

52. This Part applies to the College and members as of the day that it comes into force, except that,

(a) sections 54, 55, 56, 59 and 60 apply as of 90 days from the day that this Part comes into force; and

(b) the requirements in subsection 57 (1) and section 58 apply as of 30 days from the day that this Part comes into force. O. Reg. 154/13, s. 1.

Interpretation

53. (1) In this Part,

"designated member" means,

(a) the member designated for a drug preparation premises in accordance with section 58, or

(b) where only one member engages in or supervises drug preparation activities at or in connection with a drug preparation premises, that member;

"drug" means a substance or a preparation containing a substance referred to in clauses (a) to (d) of the definition of "drug" in subsection 1 (1) of the *Drug and Pharmacies Regulation Act*, but does not include,

(a) a substance or preparation referred to in those clauses that is manufactured, sold or represented for use in animals or fowl, or

(b) a substance or preparation referred to in clause (e), (f), (g), (h) or (i) of that definition;

"drug preparation activities" means reconstituting, diluting or otherwise preparing a drug or combining, admixing or mixing together two or more substances, at least one of which is a drug, to create a final product for the purposes of the sale or provision to another person, other than pursuant to or in anticipation of a prescription;

"drug preparation premises" means any place where a member engages in drug preparation activities, or where drug preparation activities take place that a member supervises, but does not include,

(a) a pharmacy in respect of which a valid certificate of accreditation has been issued under the *Drug and Pharmacies Regulation Act*,

(b) a premises in respect of which a valid establishment licence has been issued under the *Food and Drugs Act* (Canada), or

(c) a hospital or a health or custodial institution approved or licensed under any general or special Act;

"inspector" means a person appointed by the College to carry out an inspection on behalf of the College;

"supervise" means to supervise either directly or indirectly. O. Reg. 154/13, s. 1.

(2) Anything that may be done by the College under this Part may be done by the Council or by a committee established under clause 94 (1) (i) of the Health Professions Procedural Code. O. Reg. 154/13, s. 1.

Inspection

54. (1) All drug preparation premises are subject to inspection by the College in accordance with this Part. O. Reg. 154/13, s. 1.

(2) In carrying out an inspection of a drug preparation premises under subsection (1), the College may also require any or all of the following:

1. Inspection, examination or testing regarding any equipment, instrument, materials or any other thing that may be used in the drug preparation premises.
2. Examination and copying of books, accounts, reports, records or similar documents that are, in the opinion of the College, relevant to the member's practice with respect to the drug preparation activities at or in connection with the drug preparation premises.
3. Inquiries or questions to be answered by the member that are relevant to the member's practice with respect to the drug preparation activities at or in connection with the drug preparation premises.

4. Direct observation of a member in his or her practice with respect to drug preparation activities at or in connection with the drug preparation premises. O. Reg. 154/13, s. 1.

55. An inspector may, on the production of information identifying him or her as an inspector, enter and have access to any drug preparation premises at reasonable times and may inspect the drug preparation premises and do any of the things mentioned in subsection 54 (2) on behalf of the College. O. Reg. 154/13, s. 1.

56. (1) It is the duty of every member engaging in or supervising drug preparation activities at or in connection with drug preparation premises that are subject to an inspection to,

(a) submit to an inspection of the drug preparation premises in accordance with this Part;

(b) promptly answer a question or comply with a requirement of the inspector that is relevant to an inspection under this Part; and

(c) co-operate fully with the College and the inspector who is conducting an inspection of a drug preparation premises in accordance with this Part. O. Reg. 154/13, s. 1.

(2) A member shall not engage in or supervise drug preparation activities at or in connection with a drug preparation premises where an inspector has been denied entry or access. O. Reg. 154/13, s. 1.

57. (1) No member shall commence engaging in or supervising drug preparation activities at or in connection with drug preparation premises unless the member has previously given notice in writing to the College in accordance with subsection (5) of the member's intention to do so. O. Reg. 154/13, s. 1.

(2) Where a member has provided notice in writing to the College in accordance with subsection (1) and the drug preparation premises have not passed an inspection or passed an inspection with conditions within the previous five years, the College shall ensure that an inspection of the drug preparation premises is performed within 60 days from the day that the College receives the member's notice or 150 days from the day this Part comes into force, whichever is later. O. Reg. 154/13, s. 1.

(3) A member who engages in or supervises drug preparation activities at or in connection with a drug preparation premises as of the day that is 30 days from the day this Part comes into force shall give notice in writing to the College in accordance with subsection (5) within 90 days from the day this Part comes into force. O. Reg. 154/13, s. 1.

(4) The College shall ensure that an inspection of the drug preparation premises with respect to which a member gives notice under subsection (3) is performed within 150 days from the day this Part comes into force. O. Reg. 154/13, s. 1.

(5) The notice required in subsections (1) and (3) shall include the following information, submitted in the form and manner required by the College:

1. The full name of the member giving the notice and the full name of the individual or corporation who is the owner or occupier of the drug preparation premises, if he or she is not the member who is required to give notice under this section.
2. The full address of the drug preparation premises.
3. The date when the member first began engaging in or supervising drug preparation activities at or in connection with the drug preparation premises or the proposed date when the member intends to begin engaging in or supervising drug preparation activities at or in connection with the drug preparation premises.
4. Any other information the College requires that is relevant to an inspection of the drug preparation premises conducted under this Part. O. Reg. 154/13, s. 1.

58. Where two or more members engage in or supervise drug preparation activities at or in connection with a drug preparation premises, the members shall designate a member as the designated member for the drug preparation premises, and shall immediately notify the College of the designated member's identity. O. Reg. 154/13, s. 1.

59. All drug preparation premises are subject to an inspection by the College once every five years after the initial inspection of the premises or more often if, in the opinion of the College, it is necessary or advisable to do so. O. Reg. 154/13, s. 1.

60. (1) After an inspection of a drug preparation premises, the College shall determine, in accordance with the accepted standards of practice, whether the drug preparation premises pass, pass with conditions or fail. O. Reg. 154/13, s. 1.

(2) In determining whether drug preparation premises pass, pass with conditions or fail an inspection, the College may consider,

(a) the inspection results provided to the College by the inspector;

(b) information provided by one or more members engaging in or supervising drug preparation activities at or in connection with the drug preparation premises respecting the inspection, including the answers given by them in response to inquiries or questions asked by the inspector;

(c) the information contained in a notice given by a member under subsection 57 (1) or (3);

(d) any submissions made by the member or members engaging in or supervising drug preparation activities at or in connection with the drug preparation premises that are relevant to the inspection; and

(e) any other information that is directly relevant to the inspection of the drug preparation premises conducted under this Part. O. Reg. 154/13, s. 1.

(3) The College shall deliver a report, in writing and in accordance with section 39 of the *Regulated Health Professions Act, 1991*, to the individual or corporation that is the owner or occupier of the drug preparation premises and to the designated member for the drug preparation premises, within a reasonable time after the inspection is completed. O. Reg. 154/13, s. 1.

(4) Any report made by the College respecting an inspection of drug preparation premises where a member is engaging in or in respect of which the member is supervising drug preparation activities shall make a finding that the drug preparation premises passed, passed with conditions or failed the inspection and shall provide reasons where the drug preparation premises passed with conditions or failed the inspection. O. Reg. 154/13, s. 1.

(5) Any report made by the College that finds that drug preparation premises failed an inspection or passed with conditions is effective on the day that it is received, in accordance with section 39 of the *Regulated Health Professions Act, 1991*, by the designated member for the drug preparation premises. O. Reg. 154/13, s. 1.

(6) The designated member who receives a report made by the College that finds that a drug preparation premises failed an inspection or passed with conditions shall promptly provide copies of the report to all members engaging in or supervising drug preparation activities at or in connection with the drug preparation premises. O. Reg. 154/13, s. 1.

(7) A member shall not engage in or supervise drug preparation activities at or in connection with a drug preparation premises that fail an inspection until,

(a) the College delivers a report indicating that the drug preparation premises passed a subsequent inspection, or passed with conditions; or

(b) after considering submissions under subsection (9), the College substitutes a finding that the drug preparation premises pass or pass with conditions. O. Reg. 154/13, s. 1.

(8) A member shall not engage in or supervise drug preparation activities at or in connection with drug preparation premises that pass an inspection with conditions except in accordance with the conditions set out in the report until,

(a) the College delivers a report indicating that the drug preparation premises passed a subsequent inspection; or

(b) after considering submissions under subsection (9), the College substitutes a finding that the drug preparation premises pass. O. Reg. 154/13, s. 1.

(9) A member may make submissions in writing to the College within 14 days from the date on which a report made by the College that finds that the drug preparation premises passed with conditions or failed the inspection becomes effective in accordance with subsection (5). O. Reg. 154/13, s. 1.

(10) The College may or may not elect to reinspect the drug preparation premises after receiving a member's submissions, but no more than 60 days after a member provides his or her submissions, the College shall do one or more of the following:

1. Confirm its finding that the drug preparation premises failed the inspection or passed with conditions.

2. Make a report and find that the drug preparation premises passed with conditions.

3. Make a report and find that the drug preparation premises passed the inspection. O. Reg. 154/13, s. 1.

(11) Drug preparation premises that fail an inspection or pass with conditions may be subject to one or more further inspections within a reasonable time after the College delivers its report, at the request of a member, any other person to whom the College gave the report, or at any time at the discretion of the College. O. Reg. 154/13, s. 1.

(12) Where, as a result of an inspection carried out under this Part, a report made by the College finds that a member's knowledge, skill or judgment is unsatisfactory, the College may direct the Registrar to refer the report to the Quality Assurance Committee. O. Reg. 154/13, s. 1.

(13) Where, as a result of an inspection carried out under this Part, a report made by the College finds that a member may have committed an act of professional misconduct or may be incompetent or incapacitated, the College may direct the Registrar to refer the report to the Inquiries, Complaints and Reports Committee. O. Reg. 154/13, s. 1.

PART X
FUNDING FOR THERAPY AND COUNSELLING

61. In this Part,

"member" includes a former member. O. Reg. 225/13, s. 1.

62. (1) The alternative requirements that must be satisfied in order for a person to be eligible for funding under clause 85.7 (4) (b) of the Health Professions Procedural Code are prescribed in this section. O. Reg. 225/13, s. 1.

(2) A person is eligible for funding for therapy or counselling if,

(a) there is an admission made by a member in a statement to the College or in an agreement with the College that he or she sexually abused the person while the person was a patient of the member;

(b) a member has been found guilty under the *Criminal Code* (Canada) of sexually assaulting the person while the person was a patient of the member;

(c) there is a statement, contained in the written reasons of a committee of the College given after a hearing, that the person, while a patient, was sexually abused by a member; or

(d) there is sufficient evidence presented to the Patient Relations Committee to support a reasonable belief that the person, while a patient, was sexually abused by a member. O. Reg. 225/13, s. 1.

(3) For the purposes of clause (2) (d), and without limiting the generality of that clause, the following kinds of evidence may support a reasonable belief that a person, while a patient, was sexually abused by a member:

1. Evidence of reports made with respect to the member under subsection 85.1 (1) or 85.2 (1) of the Health Professions Procedural Code.

2. Evidence that corroborates the person's allegations of sexual abuse by the member. O. Reg. 225/13, s. 1.

(4) A person is not eligible under subsection (2) unless, at the time the sexual abuse occurred, the person was a patient of the member and the member was practising in Ontario. O. Reg. 225/13, s. 1.

(5) Despite subsections (2), (3) and (4), a person is eligible for funding for therapy or counselling under this Part only if,

(a) the person submits an application for funding to the Patient Relations Committee in the form provided by the College and, in the application, the person names the member who is alleged to have sexually abused the person;

(b) the person adheres to the procedures followed by the Patient Relations Committee when determining whether the person has satisfied the requirements for eligibility for funding; and

(c) the person provides such other information as is required by the Patient Relations Committee. O. Reg. 225/13, s. 1.

(6) A decision by the Patient Relations Committee that a person is eligible for funding for therapy or counselling does not constitute a finding against the member and shall not be considered by any other committee of the College dealing with the member. O. Reg. 225/13, s. 1.

TABLES 1, 2 REVOKED: O. Reg. 452/16, s. 2.

SCHEDULE 1
INJECTED SUBSTANCES

Analgesics and Antipyretics

Codeine — For patient education and demonstration only

Hydromorphone — For patient education and demonstration only

Morphine — For patient education and demonstration only

Nalbuphine — For patient education and demonstration only

Antibacterials

Amikacin

Ampicillin

Cefazolin

Cefepime

Cefotaxime

Cefoxitin

Ceftazidime

Ceftriaxone

Clindamycin

Cloxacillin

Ertapenem

Gentamicin

- Penicillin G
- Anticholinergic Agents
  - Scopolamine — Must not be administered intravenously
  - Hyoscine — Must not be administered intravenously
  - Glycopyrrolate — Must not be administered intravenously
- Anticoagulants
  - Dalteparin — Must not be administered intravenously
  - Danaparoid — Must not be administered intravenously
  - Enoxaparin — Must not be administered intravenously
  - Fondaparinux — Must not be administered intravenously
  - Heparin — For patient education and demonstration only
  - Nadroparin — Must not be administered intravenously
  - Tinazaparin
- Antidiabetic Agents
  - Exenatide
  - Insulins
  - Liraglutide
  - Dulaglutide
  - Lixisenatide
  - Semaglutide
- Antihemorrhagic Agents
  - Emicizumab
- Antihistamines
  - Diphenhydramine — Only for monitoring and management of allergic reactions
  - Dimenhydrinate — Must not be administered intravenously
- Antimigraine Agents
  - Sumatriptan
  - Erenumab
- Antiparkinsonian Agents
  - Apomorphine
  - Benztropine
- Antivirals
  - Enfuvirtide
  - Interferons
  - Peginterferon alfa-2a
- Central Nervous System Agents, Miscellaneous
  - Inotersen
- Complement Inhibitors
  - Icatibant
  - Lanadelumab
- Disease-modifying Antirheumatic Drugs
  - Abatacept

- Adalimumab
- Anakinra
- Etanercept
- Golimumab — Must not be administered intravenously
- Ustekinumab — Must not be administered intravenously
- Methotrexate — Must not be administered intravenously
- Sarilumab
- Tocilizumab — Must not be administered intravenously

Enzymes
- Asfotase Alfa

GI Drugs, Miscellaneous
- Certolizumab Pegol
- Methylnaltrexone

Gonadotropins and Antigonadotropins
- Follitropin-alpha
- Follitropin-beta
- Follitropin-delta
- Gonadotropin-chorionic
- Gonadotropin-chorionic-alfa
- Lutropin-alfa
- Menotropins
- Goserelin — For patient education and demonstration only
- Triptorelin acetate

Gonadotropin-releasing Hormone Antagonists
- Cetrorelix
- Ganirelix

Heavy Metal Antagonists
- Deferoxamine — For patient education and demonstration only

Hematopoietic Agents
- Darbepoetin alfa — Must not be administered intravenously
- Epoetin alfa — Must not be administered intravenously
- Filgrastim — Must not be administered intravenously
- Pegfilgrastim
- Romiplostim — For patient education and demonstration only

Immunomodulatory Agents
- Denosumab
- Glatiramer
- Interferon-Beta-1A
- Interferon-Beta-1B
- Natalizumab

Immunosuppressive Agents
- Belimumab — Must not be administered intravenously

- Mepolizumab

Miscellaneous Agents
- Sterile Water for Injection (Diluent)
- Sodium Chloride

Parathyroid
- Calcitonin Salmon — For patient education and demonstration only
- Teriparatide

Pituitary
- Desmopressin — For patient education and demonstration only
- Vasopressin — For patient education and demonstration only

Progestins
- Medroxyprogesterone
- Progesterone

Prokinetic Agents
- Metoclopramide

Proprotein Convertase Subtilisin Kexin Type 9 (Pcsk9) Inhibitors
- Alirocumab
- Evolocuma

Psychotherapeutic Agents
- Haloperidol — For patient education and demonstration only
- Methotrimeprazine — For patient education and demonstration only

Respiratory Tract Agents
- Omalizumab

Skin And Mucous Membrane Agents
- Brodalumab
- Dupilumab
- Guselkumab
- Ixekizumab
- Risankizumab — Must not be administered intravenously
- Secukinumab

Somatostatin Agonists and Antagonists
- Pasireotide
- Octreotide — Must not be administered intravenously
- Lanreotide

Somatotropin Agonists and Antagonists
- Somatropin
- Pegvisomant
- Tesamorelin

Sympatholytic (Adrenergic Blocking) Agents
- Dihydroergotamine — Must not be administered intravenously

Vitamins
- Cyanocobalamin

Folic Acid — Must not be administered intravenously

Pyridoxine — Must not be administered intravenously

Thiamine — Must not be administered intravenously

Ascorbic Acid — Must not be administered intravenously

Vitamin K

O. Reg. 95/23, s. 2.

SCHEDULE 2
INHALED SUBSTANCES

Anticholinergic Agents

- Ipratropium
- Tiotropium
- Aclidinium
- Glycopyrronium
- Umeclidinium
- Formoterol
- Indacaterol
- Olodaterol
- Salbutamol
- Salmeterol
- Terbutaline
- Vilanterol

Anti-infective Agents

- Zanamivir
- Levofloxacin
- Tobramycin
- Aztreonam

Autonomic Drugs, Miscellaneous

- Nicotine

Eye, Ear, Nose and Throat (EENT) Preparations

- Sodium Cromoglycate
- Beclomethasone
- Budesonide
- Ciclesonide
- Fluticasone
- Mometasone

Miscellaneous Agents

- Sodium chloride
- Sterile water for inhalation

Respiratory Tract Agents

- Acetylcysteine
- Dornase alfa

O. Reg. 95/23, s. 2.

SCHEDULE 3
VACCINES

1. Bacille Calmette-Guerin (BCG) Vaccines
2. Haemophilus Influenzae type b (Hib) Vaccines
3. Meningococcal Vaccines
4. Pneumococcal Vaccines
5. Typhoid Vaccines
6. Combined Typhoid and Hepatitis A Vaccines
7. Hepatitis A Vaccines
8. Hepatitis B Vaccines
9. Hepatitis A and B combined Vaccines
10. Herpes Zoster Vaccines
11. Human Papillomavirus (HPV) Vaccines
12. Japanese Encephalitis Vaccines
13. Rabies Vaccines
14. Varicella Vaccines
15. Yellow Fever Vaccines
16. Coronavirus (COVID-19) Vaccines
17. Respiratory Syncytial Virus (RSV) Vaccines.

O. Reg. 452/16, s. 3.

SCHEDULE 4
DRUGS — MINOR AILMENTS

(Arranged by American Hospital Formulary Service (AHFS) Pharmacologic-Therapeutic Classification)

Item	Column 1 Minor Ailment	Column 2 AHFS Classification
1.	Allergic rhinitis	4:08 Second Generation Antihistamines 52:02 Eye, Ear, Nose and Throat (EENT) Preparations — Antiallergic Agents 52:08.08 Eye, Ear, Nose and Throat (EENT) Preparations — Anti-inflammatory Agents — Corticosteroids
2.	Candidal stomatitis	8:14.28 Anti-infectives — Antifungals — Polyenes
3.	Conjunctivitis (bacterial, allergic or viral)	04:04.20 Propylamine Derivatives 52:32 Eye, Ear, Nose and Throat (EENT) Preparations — Vasoconstrictors 52:04.04 Eye, Ear, Nose and Throat (EENT) Preparations — Anti-infectives — Antibacterials 52:02 Eye, Ear, Nose and Throat (EENT) Preparations — Antiallergic Agents
4.	Dermatitis (atopic/eczema, allergic or contact)	84:06 Skin and Mucous Membrane Agents — Anti-inflammatory Agents
5.	Dysmenorrhea	28:08.04 Central Nervous System Agents — Analgesics and Antipyretics — Nonsteroidal Anti-inflammatory Agents
6.	Gastroesophageal reflux disease (GERD)	56:04 Gastrointestinal Drugs — Antacids and Adsorbents 56:28.12 Gastrointestinal Drugs — Antiulcer Agents and Acid Suppressants — Histamine H_2-Antagonists 56:28.36 Gastrointestinal Drugs — Antinuclear Agents and Acid Suppressants — Proton-Pump Inhibitors

7.	Hemorrhoids	12:12.04 Autonomic Drugs — Sympathomimetic (Adrenergic) Agents — Alpha-Adrenergic Agonists 52:04.92 Eye, Ear, Nose and Throat (EENT) Anti-infectives — Miscellaneous 84:06 Skin and Mucous Membrane Agents — Anti-inflammatory Agents 84:08 Skin and Mucous Membrane Agents — Antipruritics and Local Anesthetics 84:04.04 Skin and Mucous Membrane Agents — Anti-infectives — Antibacterials
8.	Herpes labialis	8:18.32 Anti-infective Agents — Antivirals — Nucleosides and Nucleotides 84:06 Skin and Mucous Member Agents — Anti-inflammatory Agents 84:04.06 Skin and Mucous Membrane Agents — Anti-infectives — Antivirals
9.	Impetigo	84:04.04 Skin and Mucous Membrane Agents — Anti-infectives — Antibacterials 84:06 Skin and Mucous Member Agents - Anti-inflammatory Agents
10.	Insect bites and urticaria	4:04 Antihistamine Drugs — First Generation Antihistamines 4:08 Antihistamine Drugs — Second Generation Antihistamines 84:06 Skin and Mucous Member Agents — Anti-inflammatory Agents 84:08 Skin and Mucous Membrane Agents — Antipruritics and Local Anesthetics
11.	Tick bites, post-exposure prophylaxis to prevent Lyme disease	8.12.24 Anti-infective Agents — Antibacterials — Tetracyclines
12.	Musculoskeletal sprains and strains	28:08.04 Central Nervous System Agents — Analgesics and Antipyretics — Nonsteroidal Anti-inflammatory Agents 28.08.92 Central Nervous System Agents — Analgesics and Antipyretics — Miscellaneous
13.	Urinary Tract Infection (uncomplicated)	8:12.20 Anti-infective Agents — Antibacterials — Sulfonamides 8:36 Anti-infective Agents — Urinary Anti-infectives

O. Reg. 460/22, s. 3.

Note: On October 1, 2023, Schedule 4 to the Regulation is revoked and the following substituted: (See: O. Reg. 179/23, s. 2)

SCHEDULE 4
DRUGS — MINOR AILMENTS

Item	Column 1 Minor Ailment	Column 2 Drug Classes	Column 3 Specified Drugs
1.	Acne (mild)	Skin and Mucous Membrane Agents	Adapalene Azelaic acid Benzoyl peroxide Clindamycin Dapsone Erythromycin Glycolic acid Salicylic acid Tazarotene Tretinoin Trifarotene
2a.	Allergic rhinitis	Antihistamines	Azelastine Bilastine Cetirizine Cyproheptadine Desloratadine Fexofenadine Loratadine

			Olopatadine Rupatadine
2b.	Allergic rhinitis	Corticosteroids	Beclomethasone Budesonide Ciclesonide Fluticasone Mometasone Triamcinolone
3.	Candidal stomatitis	Antifungals	Nystatin
4a.	Conjunctivitis (bacterial, allergic or viral)	Antiallergic Agents	Antazoline Bepotastine Cromolyn sodium (Sodium cromoglycate) Ketotifen Lodoxamide Olopatadine Pheniramine
4b.	Conjunctivitis (bacterial, allergic or viral)	Antibacterials	Erythromycin Fusidic acid Gramicidin Polymyxin B Tobramycin Trimethoprim
4c.	Conjunctivitis (bacterial, allergic or viral)	Vasoconstrictors	Naphazoline Oxymetazoline Phenylephrine Tetrahydrozoline
5.	Dermatitis (atopic/eczema, allergic or contact)	Anti-inflammatory Agents	Beclomethasone Betamethasone valerate Clobetasone Crisaborole Desonide Fluocinolone Hydrocortisone Prednicarbate Triamcinolone
6a.	Dermatitis (diaper)	Antifungals	Ciclopirox Clotrimazole Ketoconazole Miconazole Nystatin
6b.	Dermatitis (diaper)	Anti-inflammatory Agents	Desonide Hydrocortisone
7.	Dysmenorrhea	Nonsteroidal Anti-inflammatory Agents	Acetylsalicylic acid (ASA) Celecoxib Diclofenac Flurbiprofen Ibuprofen Ketoprofen Mefenamic acid Naproxen
8a.	Gastroesophageal reflux disease (GERD)	Antacids and Adsorbents	Alginic acid Aluminum hydroxide Calcium carbonate Magnesium salts
8b.	Gastroesophageal reflux disease (GERD)	Histamine H2-Antagonists	Cimetidine Famotidine Nizatidine Ranitidine
8c.	Gastroesophageal reflux disease (GERD)	Proton-Pump Inhibitors	Dexlansoprazole Esomeprazole Lansoprazole Omeprazole Pantoprazole Rabeprazole
9.	Hemorrhoids	Skin and Mucous Membrane Agents	Dibucaine (Cinchocaine)

			Esculin (Aesculin) Framycetin (Neomycin B) Hydrocortisone Phenylephrine Pramoxine Zinc sulfate
10a.	Herpes labialis	Anti-inflammatory Agents	Hydrocortisone
10b.	Herpes labialis	Antivirals	Acyclovir Docosanol Famciclovir Valacyclovir
11.	Impetigo	Antibacterials	Bacitracin Fusidic acid (Sodium fusidate) Gramicidin Mupirocin Ozenoxacin Polymyxin B
12a.	Insect bites and urticaria	Antihistamines	Bilastine Cetirizine Chlorpheniramine Cyproheptadine Desloratadine Diphenhydramine Fexofenadine Hydroxyzine Loratadine Rupatadine
12b.	Insect bites and urticaria	Antipruritics and Anti-inflammatory Agents	Benzocaine Calamine Camphor Desonide Hydrocortisone Lidocaine Menthol Pramoxine Zinc oxide
13a.	Musculoskeletal sprains and strains	Analgesics	Acetaminophen
13b.	Musculoskeletal sprains and strains	Nonsteroidal Anti-inflammatory Agents	Acetylsalicylic acid (ASA) Celecoxib Diclofenac Flurbiprofen Ibuprofen Ketoprofen Mefenamic acid Naproxen
14.	Nausea and vomiting of pregnancy	Antiemetics and Antinauseants	Dimenhydrinate Diphenhydramine Doxylamine Promethazine Pyridoxine
15.	Oral aphthae	Anti-inflammatory Agents	Triamcinolone
16.	Pinworms/Threadworms	Anthelmintics	Mebendazole Pyrantel pamoate
17.	Tick bites, post-exposure prophylaxis to prevent Lyme disease	Antibacterials	Doxycycline
18.	Urinary tract infection (uncomplicated)	Urinary Anti-infectives	Fosfomycin Nitrofurantoin Sulfamethoxazole Trimethoprim
19.	Vulvovaginal candidiasis	Antifungals	Clotrimazole Fluconazole Miconazole Terconazole

O. Reg. 179/23, s. 2.

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