

May 20, 2026

To: *Department of Service Nova Scotia, Business Licensing and Compliance (SNS-BLC)*

**RE: Review of Registration Practices for “Hearing Aid Salespersons”
Under the [Fair Registration Practices Act](#)**

Thank you for submitting the FRPA Review Report (appended below), received by this office on May 14, 2026, as required under Section 16 of the [Fair Registration Practices Act \(FRPA\)](#). Any statistical information provided through the annual FRPA survey and as published in the FRPA Annual Report, which can be found on the [FRPA website](#), has been accepted as forming part of this report.

This letter pertains only to SNS-BLC’s compliance status with the FRPA. It does not speak to any other enactment.

Upon review of this report, including information found through links contained within, no FRPA compliance issues were identified. Therefore, no further follow-up is required in relation to this FRPA review report.

Pursuant to Section 16, the next FRPA review report is due on or before **May 14, 2031**, which is five years from the date the most recent report was received.

Thank you for your cooperation. Compliance with the FRPA helps to ensure fair access to Nova Scotia’s labour force of regulated professions.

Sincerely,



Frank Reinhardt, Review Officer, Fair Registration Practices Act
Department of Labour, Skills and Immigration

Enclosure: FRPA review Report as prepared and submitted by SNS-BLC.

Service Nova Scotia as the body responsible for the registration of Hearing Aid Salespersons

Report on Registration Practices under Nova Scotia's

Fair Registration Practices Act (FRPA)

April 15, 2026

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1. Requirement and Purpose of this Report under the FRPA:

This report has been submitted to the “*Review Officer*”, appointed under Section 13 of the [Fair Registration Practices Act](#) (FRPA or “the Act”), in fulfillment of the requirements of Section 16 of the Act, which states, in part that:

16(2) Every regulating body shall review its registration practices in accordance with this Section and shall file a report on the results of the review with the Review Officer...(8) every five years ...unless the Review Officer, based on an assessment of the information provided in a report, specifies a more frequent reporting...

Section 6 of the Act summarizes the “**Duty**” of each “regulating body” as:

A regulating body has a duty to carry out registration practices that are transparent, objective, impartial and procedurally fair.

Sections 5 to 12 of the Act, formally referred to as the “*Fair Registration Practices Code*”, details the specific legislative requirements that must be met in fulfillment of this “**duty**”. This FRPA review report, which is subject to review by the FRPA “*Review Officer*”, details the registration practices of this regulating body in relation to those specific requirements and others detailed under Section 16.

2. Overview of the Regulating Body:

Name of Regulating Body:	Service Nova Scotia In this report, “regulating body” refers to this organization.
Homepage of Regulating Body:	https://www.novascotia.ca/hearing-aid-salesperson-permit
List of Occupations Regulated by the Regulating Body and that are being reported on in this report:	Hearing Aid Salespersons
List of types of Licences / Certificates / Registrations Issued (eg. Full, Conditional, Temporary, Student) by the Regulating Body:	Licence
Name of the authorizing legislation (include link(s)):	https://novascotia.ca/just/regulations/regs/dsregs.htm https://nslegislature.ca/sites/default/files/legc/statutes/direct%20sellers'%20regulation.pdf
The following information is accessible through the Regulating Body’s home page (Yes/No):	YES
<ul style="list-style-type: none"> • The role of the regulating body. • Descriptions of the occupations and licence types listed above. 	
If no , please provide that information here:	

Click or tap here to enter text.

3. Reporting on Registration Practices:

This report has been submitted by the regulating body using a template provided by the FRPA “Review Officer”. It details the registration practices of this regulating body in relation to the requirements under the Act for all three applicant types:

“New” applicants: *Those who only include Canadian credentials in their application to apply for registration (licensure) and are not currently registered (licensed) to practice anywhere in Canada.*

“Interprovincial” applicants: *Those who are currently registered (licensed) to practice elsewhere in Canada (outside of Nova Scotia); and*

“International” applicants: *Those who are including international credentials in their application to apply for registration (licensure) and are not currently registered (licensed) to practice anywhere in Canada.*

3.1 Registration Process – Requirements, Criteria Communications:

Sections of FRPA: 7, 9(a) and 16 Indicate which of the following aspects of the registration process is described <i>“in a clear and understandable form”</i> on, or accessible through, the regulating body’s website, including any differences, if any, for each applicant type (“New”, “Interprovincial” and “International”):	YES / NO
a) The step-by-step process that applicants must follow to apply for registration. [Sections 16(3)(a), 7(a) and (c)]	<input type="checkbox"/> YES
b) “Requirements for registration” (including qualifications and required documentation) [Sections 16(3)(a) and (c), 7(c), 9(a)]	<input type="checkbox"/> YES
c) “the criteria used to assess whether the requirements for registration have been met” [Sections 16(3)(b), 7(d)];	<input type="checkbox"/> YES
d) “the fees charged for registration” (if any) [(Sections 16(3)(d), 7(f)];	<input type="checkbox"/> YES
e) “information about the length of time that the registration process for that regulating body usually takes” [Sections 16(3)(l), 7(b)];	<input type="checkbox"/> YES

For each item above, for which the answer is **“YES”**, the FRPA Review Officer will review the regulating body’s website to confirm compliance with the above cited FRPA sections.

For each item above for which the answer is **“NO”** (if any), in order to determine compliance with the cited FRPA sections, please provide a description of that item in the space provided below as well as a description of how this information is (or is not) provided to the unidentified individuals who only *“...intend to apply for registration”* (e.g. How it is made available to the public):

Click or tap here to enter text.

3.2 Communicating Registration Decisions:

Sections of FRPA: 8, 10(1)

For each of the following statements, indicate whether, or not, it accurately describes the registration practices of the regulating body ('Yes' or 'No').

Accurate?
(YES/NO)

a) Where registration **is granted**, written confirmation is provided to applicants within a reasonable time.

YES

b) Where registration **is not granted**, the regulating body:

- provides written decisions that include reasons to applicants within a reasonable time respecting registration decisions;

YES

- provides, if/where practical, information respecting measures or programs that may be available to assist unsuccessful applicants in obtaining registration at a later date; and

YES

- informs the applicant of the internal review process and of the procedures and time frames for the internal review.

YES

c) (Optional) Please use the space below to provide any further details as/if necessary:

Click or tap here to enter text.

3.3 Allowance for Alternative Information

<p>Sections of FRPA: 7(c), 8(a), 9 and 16(3)(a), (c) and (g)</p> <p>Note: FRPA compliance does not require that a regulating body accept “alternative information”. It requires that, if there are such policies (there exists some known circumstances where alternative information may be considered), that such policies be communicated to applicants and unidentified “potential applicants” (e.g. by making the information publicly available).</p>	(YES/NO)
<p>a) “Where documentation cannot be obtained by an applicant for reasons beyond the applicant’s control...”, does there exist any “...<i>alternative information</i> [which] <i>may be supplied by the applicant that may be acceptable to the regulating body?</i>”</p>	<div style="border: 2px solid black; padding: 5px; width: 40px; margin: 0 auto;">NO</div>
<p>b) If ‘Yes’ to part ‘(a)’, and these “alternative information” policies, are detailed on the regulating body’s website, including what alternative information may be acceptable, and under what circumstances, please provide the direct weblink in the space provided below:</p> <div style="border: 1px solid black; padding: 2px; margin-top: 5px;"> <p style="background-color: yellow; display: inline-block;">Click or tap here to enter text.</p> </div>	
<p>c) If ‘Yes’ to part ‘(a)’, and these “alternative information” policies are not detailed on the regulating body’s website, in the space provided below, please detail these “alternative information” policies, including what alternative information may be acceptable, under what circumstances, and how this information is (or is not) provided to the unidentified individuals who only “...<i>intend to apply for registration</i>” (e.g. How it is made available to the public) (Section 7c):</p> <div style="border: 1px solid black; padding: 2px; margin-top: 5px;"> <p style="background-color: yellow; display: inline-block;">Click or tap here to enter text.</p> </div>	
<p>d) If an applicant informs the regulating body that they are unable to provide the standard required information or documentation, and the Regulating Body determines that there is no acceptable “alternative information”, does the regulating body inform the applicant of this “<i>within a reasonable time</i>”?</p>	<div style="border: 2px solid black; padding: 5px; width: 40px; margin: 0 auto;">YES</div>
<p>a) (<u>Optional</u>) Please use the space below to provide any further details as/if necessary:</p> <div style="border: 1px solid black; padding: 5px; margin-top: 5px;"> <p>If they can’t provide the necessary educational documentation, they can take the practical tests or exams requirements.</p> </div>	

3.4 Accommodation Policies for Applicants with A Physical or Mental Disability

Sections of FRPA: 7(e), 8(a) and 16(3)(h)

**Accurate?
(YES/NO)**

Note: FRPA compliance does not require that a regulating body provide accommodations for applicants with disabilities. It requires that, if there are such policies (there exists some known circumstances where accommodations may be provided), that such policies be communicated to applicants and unidentified individuals who are only *“intending to apply”* (e.g. by making the information publicly available). It also requires that any requests for accommodations be responded to in a timely manner (whether granted or not).

Indicate whether, or not, the following statements accurately describes the registration practices of the regulating body (‘Yes’ or ‘No’).

a) “A description of existing accommodation policies for applicants with a physical disability or mental disability” is accessible “in a clear and understandable form” on, or through, the regulating body’s website.

NO

If ‘Yes’, please provide a weblink to this description in the space provided below (then skip to the next question, 3.5):

[Click or tap here to enter text.](#)

If ‘No’, indicate which of the following (b, c, or d) accurately describes this regulating body’s registration practices:

b) Although no formal “accommodation policies” exist, the regulating body considers, and provides timely responses to, any request from an applicant, or potential applicant, for accommodations for a disability (physical or mental). Such responses would detail any accommodations that may be provided (if any) and reasons for any specific requests not granted.

YES

c) No accommodations for physical or mental disabilities are currently offered under any circumstances by the regulating body. Any applicant who requests such accommodations are informed of this in a timely manner.

select

d) The regulating body has policies related to accommodations for applicants with a physical or mental disability that are different than those described in parts ‘b’ and ‘c’. However, descriptions of these policies are not posted on the regulating body’s website.

select

If yes, in the space provided below, please provide *“a description of existing accommodation policies for applicants with a physical disability or mental disability”* and describe how the regulating body provides this information to the unidentified individuals who only *“...intend to apply for registration”* (e.g. How it is made available to the public):

[Click or tap here to enter text.](#)

3.5 Other Support Provided to Applicants During the Registration Process	
Sections of FRPA: 7(e) and 16(3)(k)	Accurate? (YES/NO)
For the following statement, indicate whether, or not, it accurately describes the registration practices of the regulating body ('Yes' or 'No'):	YES
<p>a) The regulating body's website informs applicants that general support (e.g. to answer questions they may have about the registration process or requirements) is available upon request throughout the registration process by using the contact information provided on the website.</p> <p>b) If there are any other supports available to applicants (which have not already been described within this template), in the space provided below, please list those supports and, for each, either: 1) provide a direct weblink to its description; or 2) provide that description including how this information is (or is not) provided to the unidentified individuals who only <i>"...intend to apply for registration"</i> (e.g. How it is made available to the public):</p> <div style="border: 1px solid black; padding: 5px; margin-top: 10px;"> <p style="background-color: yellow;">Click or tap here to enter text.</p> </div>	

3.6 Role of Third-Party Assessors	
Sections of FRPA: 16(3)(i)	YES/NO
A "third-party assessor" is defined in the Act as: <i>"a body external to a regulating body relied on by the regulating body to assess the equivalence of the qualifications of an applicant for registration."</i>	
Are "third-party assessors" involved in the registration process?	YES
<p>If yes, compliance with the Section 16(3)(i) of the FRPA requires that <i>"an outline of the role of third-party assessors"</i> be provided. Therefore, in the space provided below, please provide that outline/description or, if this is described on the regulating body's website, provide a weblink to this information:</p> <div style="border: 1px solid black; padding: 10px; margin-top: 10px;"> <p>https://www.novascotia.ca/hearing-aid-salesperson-permit</p> <p>You also need to meet 1 of the following educational standards:</p> <p>master's degree in audiology and be registered with the Nova Scotia College of Audiologists and Speech-Language Pathologists</p> <p>6-month apprenticeship with a licensed hearing aid dealer and successful completion of both the International Licensing Exam (ILE) and the practical exam Applying with a 6-month apprenticeship with a licensed hearing aid dealer</p> <p>Complete the application form.</p> <p>Check the application for details on all required supporting documents.</p> <p>Include a letter of completion for your apprenticeship with a licensed hearing aid dealer. A representative of the licensed hearing aid company needs to write the letter. The letter needs to state your name, the name of the hearing aid company and confirmation that you competed at</p> </div>	

least 6 months in an apprenticeship role.

Include payment with your application.

Send your completed application, supporting documents and payment by mail.

Department of Service Nova Scotia and Internal Services arranges for you to take the written International Licensing Exam through the International Hearing Society and the practical exam through a speech clinic at Nova Scotia Hearing and Speech Centres.

Once you pass the International Licensing Exam, you're cleared to take the practical exam at Nova Scotia Hearing and Speech Centres. The Department of Service Nova Scotia and Internal Services notifies the speech clinic to make arrangements for your practical exam.

When the speech clinic confirms you passed the practical exam, the Department of Service Nova Scotia and Internal Services issues the permit.

3.7 Access to Registration Records

Sections of FRPA: Sections 12 and 16(3)(j).

**Accurate?
(YES/NO)**

For the following statement, indicate whether, or not, it accurately describes the registration practices of the regulating body ('Yes' or 'No'):

“Upon the written request from an applicant, ...the regulating body ...provides the applicant with access to [any and all] records held by it that are related to the application, other than any records or portions of records (if any) that Section 12 of the [FRPA](#) specifically permits regulating bodies to refuse to provide (eg. those protected by legal privilege or other existing legislation or to protect the identities of other individuals, or to avoid negative impacts on public safety or the integrity of the registration process).

YES

In the space provided below, please provide **either**: a description; **or** a link to a “description of the process under which requests for access to records are considered”; **or** both if/as deemed necessary.

Licensing would advise if credentials aren't acceptable and provided with other options. The application and website also note that any applicant requiring accommodation may request them such as providing any records they submitted with the application. Test results from the practical exam at Nova Scotia Hearing and Speech Centres or written International Licensing exam through the International Hearing Society would have to be provided by these organizations.

3.8 Internal Review Process

Sections of FRPA: 7(a), 8(a), 10, 16(3)(m) and (n)

For each of the following statements, indicate whether, or not, it accurately describes the registration practices of the regulating body ('Yes' or 'No').

**Accurate?
(YES/NO)**

a) An ***“internal review process”***, is available to unsuccessful applicants to appeal their registration decision.

YES

b) The internal review process includes the following features:

- Applicants appealing a registration decision are provided an opportunity to provide new information and to make submissions with respect to an internal review in such a manner as determined by the internal review decisionmaker.
- An internal review decision-maker provides applicants with a written decision that includes reasons within a reasonable time.
- No one who acted as a decision-maker in respect of a registration decision may act as a decision-maker in an internal review in respect of that registration decision.

YES

YES

YES

c) ***“A description of the internal review process”*** is accessible ***“in a clear and understandable form”*** on the regulating body’s website.

YES

If yes, in the space provided below, please provide a link to this description and the FRPA Review Officer will review to confirm compliance with the cited FRPA sections:

<https://www.novascotia.ca/hearing-aid-salesperson-permit>

If no, in the space provided below, please provide ***“a description of the internal review process”*** and describe how the regulating body provides this information to the unidentified individuals who only ***“...intend to apply for registration”*** (e.g. How it is made available to the public):

<https://novascotia.ca/just/regulations/regs/dsregs.htm>

Section 8 of the Direct Sellers Regulations states:

8 (1) In determining whether to grant, refuse, cancel or reinstate a Category H1 or H2 direct seller’s permit or a salesperson’s permit, the Registrar may, if the Registrar considers it necessary, establish an Advisory Board consisting of 4 members, as follows:

- (a) the Registrar, or another person appointed by the Registrar as the representative of the Registrar;
- (b) 1 member selected and appointed by the Registrar to represent the hearing-aid industry;
- (c) 1 member selected and appointed by the Registrar to represent the medical profession;
- (d) 1 member selected and appointed by the Registrar to represent hearing-aid users.

(2) After the Advisory Board referred to in subsection (1) considers any matter submitted to it by the Registrar, the Board shall make a recommendation to the Registrar in writing, and the Registrar shall take such recommendation into consideration when making his decision.

With this authority in place, SNS-IS will develop and document a plain language process which details

the steps applicants may take to request a review and submit additional documentation. Instructions and timelines for the process will be publicly available and communicated to denied applicants. Decisions and rationale will be provided to applicants in writing.

Under Subsection 3(4) of the Direct Sellers' Regulation Act, a deputy registrar may perform any of the duties and exercise any of the powers of the Registrar as directed by the Registrar. A policy and process will be created and documented to ensure a Registrar or Deputy Registrar does not participate in an internal review of their own decision.

All interested parties would be provided the link to the legislation.

3.9 Training for Internal Reviewers

Sections of FRPA: 11, 16(3)(p)

Accurate?
(YES/NO)

YES

- a) The regulating body ensures that any individuals acting as decisionmakers in internal reviews receive training on conducting an internal review and that training includes the following features:
- Structured/formalized (expectations are clearly defined);
 - Specific to the process of conducting an internal review; and
 - Includes a means of verifying that the training was “received” (eg. attendance tracking, signed declarations by trainees and/or some form of course assessment such as a test or assignment).

- b) In the space provided below, please describe the training provided to individuals who make internal review decisions, including the three required features noted in part ‘a’ above. Alternatively, if this description happens to be available on or through the regulating body’s website (not required), you may provide a link to this description:

Under Subsection 3(4) of the Direct Sellers’ Regulation Act, a deputy registrar may perform any of the duties and exercise any of the powers of the Registrar as directed by the Registrar. A policy and process will be created and documented to ensure a Registrar or Deputy Registrar does not participate in an internal review of their own decision.

POLICY ON INTERNAL REVIEWS OF HEARING AID REGISTRATION DECISIONS

Business Licensing for SNS provides the following process for appeal of its registration decisions. Instructions and timelines for the process will be publicly available and communicated to denied applicants. Decisions and rationale will be provided to applicants in writing. This process will include the requirement that no one who acted as a decision-maker will take part in the internal review process and will contain a clear statement to that effect.

Definitions

*Committee means the Internal Review Committee
Parties refers to the applicant and the Registrar
Policy is the Policy on Internal Reviews of Registration Decisions*

Composition of the Internal Review Committee

1. *An Internal Review Committee will be established from the following groups:*
 - *[1-2] [Service Nova Scotia staff]*
 - *[1-2] [Third Party assessor]*
2. *No one who acted as a decision-maker in reaching the original registration decision may serve on the Internal Review Committee.*
3. *Internal Review Committee members will participate in training prior to participating in an internal review hearing.*

Notification of Right to Internal Review

4. *When an applicant is not granted registration with SNS, they will be informed of the decision by letter.*
5. *The letter shall include:*
 - *reasons for the registration decision;*
 - *a statement that indicates the applicant’s right to internal review; and*

- *a statement indicating the applicant may exercise their right to an internal review by informing the Registrar [in writing] within [30 days] of this letter's date.*

6. *This Policy shall be attached to the decision letter.*

Requesting the Internal Review

7. *Within [30 days] of the letter's date, the applicant may submit a written request for internal review of a registration decision via [email or mail].*

8. *The applicant's request for internal review should outline the basis for the review.*

Scheduling the Internal Review

9. *Upon receipt of a request for internal review, the Registrar will notify the Internal Review Committee and provide all records relating to the applicant's application to the Committee within [7 days].*

10. *The Internal Review Committee will contact the applicant within [14 days] of receipt of the application records to schedule the date for the Internal Review Hearing.*

11. *The Internal Review Hearing will take place within [60 days] of the receipt of application records by the Committee.*

Opportunity to Make Submissions

12. *Either party may make submissions [in writing] to the Committee within 30 days after the review has been scheduled.*

13. *The Committee will share submissions with both parties to review. The parties will be given at least [14 days] to review submissions in advance of the hearing date.*

Internal Review Hearing

14. *At the Internal Review Hearing, the Committee will review all submissions made by the parties.*

15. *Both Parties will have the opportunity to appear and speak before the committee.*

16. *Parties may appear with or without legal counsel.*

The Internal Review Decision

17. *The Committee will render its decision based on careful consideration of all written submissions and oral presentations at the Hearing.*

18. *The Committee may uphold the rejection of the applicant or order the Registrar to reverse the decision and register the applicant. The Committee may require terms, conditions, or restrictions be imposed on the applicant's license as a condition of licensure.*

19. *A decision will be reached within [14 days] of the hearing, at which point both Parties will be informed in writing.*

20. *The decision will be issued to the applicant via [mail or email] with reasons.*

21. *If the Committee issues a decision to reverse the Registrar's decision and license the applicant, the Registrar must register the applicant within [7 days].*

22. *[The decision of the Internal Review Committee is final.]*

3.10 “Interprovincial” Applicants under the Canadian Free Trade Agreement (CFTA):

Sections of FRPA: 3, 7, and 16(3)

Context:

Section 3 of the FRPA “...recognizes the commitments ...made under the Canadian Free Trade Agreement (CFTA)...” and Section 16 requires that the registration practices of this regulating body, for all applicants, including “Interprovincial” applicants, be detailed in this FRPA review report.

Nova Scotia’s Canadian Free Trade Agreement Implementation Act requires that regulating bodies comply with Chapter Seven, “Labour Mobility”, of the CFTA.

This Section reports on the regulating body’s registration practices for “Interprovincial Applicants” as they relate to Chapter 7 (Labour Mobility) of the CFTA.

Instructions:

Indicate which of the following documents/items are required from Interprovincial Applicants before registration (licensure) may be granted. (For each item, answer either ‘Yes’ or ‘No’. Do not leave blank.):

Note: Items ‘a’ to ‘c’ are specifically permitted under Article 705 the CFTA. Items 3d to 3j are also permitted but only if they “...are the same as, or substantially similar to, but no more onerous than, those imposed by the regulatory authority on its own workers as part of the normal certification process; and ...the requirement does not create a disguised restriction on labour mobility”.

Sections of FRPA: 3 and 16(3)

Indicate whether, or not, each of the following items are required of interprovincial applicants before registration may be granted (Yes or No):

**Required?
(YES/NO)**

a) An application form.	YES
b) Proof of current licensure in one or more Canadian jurisdictions (outside of Nova Scotia), for the same occupation, for which the applicant is applying.	NO
c) A certificate, letter, or other evidence from the regulatory authority (or authorities) that issued the applicant’s current license, confirming that their current license is in good standing.	YES
d) Any items that are referred to as “ <i>Exceptions to Labour Mobility</i> ” to satisfy one or more “ <i>Legitimate Objectives</i> ”, or (LOEs - “ <i>Legitimate Objective Exceptions</i> ”) which are approved by the Province of Nova Scotia and listed here: https://workersmobility.ca/exceptions-by-jurisdiction/ (Click on Nova Scotia – if none are relevant, chose “no”)	NO
e) an application or processing fee.	YES
f) insurance, malpractice coverage, or similar protection	NO
g) requirement to post a bond.	NO
h) a criminal background check.	NO
i) evidence of good character (besides any item already listed above). If yes , please provide a brief description:	NO

Click or tap here to enter text.

<p>j) demonstrated knowledge of measures maintained by Nova Scotia to practice the occupation in Nova Scotia (eg. jurisprudence exam)</p> <p>If yes, please provide a brief description:</p>	<div style="border: 2px solid black; padding: 5px; width: fit-content; margin: auto;">YES</div>
<p>master's degree in audiology and be registered with the Nova Scotia College of Audiologists and Speech-Language Pathologists or 6-month apprenticeship with a licensed hearing aid dealer and successful completion of both the International Licensing Exam (ILE) and the practical exam</p>	
<p>k) demonstrated proficiency in either English or French for at least some interprovincial applicants:</p> <p>If yes, is this only required if no equivalent language proficiency requirement was imposed on, and satisfied by, the worker as a condition of the worker's certification in his or her current certifying jurisdiction?</p>	<div style="border: 2px solid black; padding: 5px; width: fit-content; margin: auto;">NO</div>
<p style="text-align: center; background-color: yellow;">select</p>	
<p>l) Any other document(s)/item(s), not covered by the above categories (items 'a' to 'k') (eg. proof of education or other credentials or additional training requirements):</p> <p>If yes, list these items in the space provided below, the criteria used to assess them, and describe why current licensure from another Canadian jurisdiction is not accepted as confirmation of meeting these criteria:</p>	<div style="border: 2px solid black; padding: 5px; width: fit-content; margin: auto;">NO</div>
<p style="background-color: yellow;">Click or tap here to enter text.</p>	
<p>m) If you answered 'Yes' to any of the items from 'e' to 'k', are all of those items "...<i>the same as, or substantially similar to, but no more onerous than, those imposed...</i>" on "New" applicants (unlicensed with only Canadian credentials)?</p> <p>If no (or "Not Sure"), please list each item, from 'e' to 'k', that are, or may, not be "...<i>the same as, or substantially similar to...</i>" those imposed on non-CFTA applicants and describe those differences:</p>	<div style="border: 2px solid black; padding: 5px; width: fit-content; margin: auto;">YES</div>
<p style="background-color: yellow;">Click or tap here to enter text.</p>	

4. Declaration by Regulating Body:

The *Regulatory Body* hereby declares that the information contained in this report, including any information provided through weblinks contained in this report, is a true and accurate representation of its current registration practices.

SIGNATURE OF THE AUTHORIZED MEMBER OF THE REGULATING BODY:



Name (print): Jim Kochanoff

DATE: 2026-04-15