

**NOTICE TO MANUFACTURERS OF DRUGS, MEDICAL DEVICES RELATED TO DRUG ADMINISTRATION, AND
BLOOD SYSTEM PRODUCTS**

DATE: October 13, 2026

SUBJECT: Important Information Regarding Registration Applications

Update to the “Brief Description of the File” Forms

INESSS would like to inform you that the “Summary Description of the Dossier” forms for [drugs](#), [medical devices](#), and [blood system products](#) have been updated. This revision is intended, in particular, to obtain information on clinical studies conducted in Canada and Québec regarding the product under review.

Both French and English versions of these documents are available on the [website](#). We encourage you to **start** using them **immediately**.

New Interactive Tool Now Available to Assist with Registration Applications

INESSS has developed an interactive tool designed to guide manufacturers in selecting the appropriate registration form for their application and outlining the required supporting documents.

This tool was tested with a select group of manufacturers as part of the validation process and subsequently enhanced. It is available in French and English and is aligned with current reference documents, notably the [Guide for Submitting an Application to INESSS \(June 2026\)](#).

Starting October 14, the tool will be available at the following link: [Submission Guide - Selection Tool.html](#). It will also be available on the following webpage: [INESSS: Registration Application - Medication](#).

To support the ongoing improvement of this tool, we invite you to use it and submit any observations, comments, or suggestions to the following address: inscription@inesss.qc.ca.

Thank you for your cooperation.

Direction de l'évaluation des médicaments et des technologies à des fins de remboursement,
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