CADTH Non-Sponsored Reimbursement Review

**Industry Input Template**

Instructions

This form is to be used by eligible industry stakeholders (i.e., any current or future Drug Identification Number [DIN] holders for the drug under review) for the purpose of providing initial input to CADTH regarding a non-sponsored reimbursement review.

Before Completing the Template

Please review the following documents to ensure an understanding of CADTH’s procedures:

* [Procedures for Non-sponsored Reimbursement Reviews](https://www.cadth.ca/sites/default/files/Drug_Review_Process/CADTH_Non-Sponsored_Reviews.pdf)
* [Procedures for CADTH Reimbursement Reviews](https://cadth.ca/sites/default/files/Drug_Review_Process/CADTH_Drug_Reimbursement_Review_Procedures.pdf)
* [CADTH Pharmaceutical Review Updates](https://www.cadth.ca/search?f%5B0%5D=project_sub_line%3A108404) for any applicable information.

Completing the Template

Please complete all applicable sections of the industry input template. The maximum size is 5 pages (this total does not include the reference list). References must be provided in the following format: In-text citations must be numbered in order of appearance and a numbered reference list must be provided in the JAMA Oncology format.

As stated in the *Procedures for Non-sponsored Reimbursement Reviews,* industry participants will not have the opportunity to review and request redactions of CADTH reports or recommendations before they are posted on the CADTH website. Input from industry **must not contain any confidential information** (all information included in the template will be considered disclosable by CADTH).

Filing the Completed Template

**Delete** first page of this template and all red font instructions once document is complete.

**Send**the completed template by using the Submit link next to the drug listed on the [Open Calls](https://www.cadth.ca/open-calls-input-and-feedback-0) page. The input must be filed as a Microsoft Word document by the posted deadline date for the information to be used by CADTH.

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Industry Input

CADTH Project Number: <Enter Response here>

Generic Drug Name: <Enter Response here>

Indication: <Enter Response here>

Name of Organization: <Enter Response here>

Author of Submission: <Enter Response here>

1. Does the proposed project scope accurately reflect the treatment landscape?

<Enter Response Here>

1. Are you aware of relevant published studies that you would like considered in the clinical review?

<Enter Response Here>

1. Do you have additional comments that you feel are pertinent to this review?

<Enter Response Here>