



## Common Drug Review \*

### Submission Status

**Product:**   
**Generic Name:**   
**Manufacturer:**   
**Submission Type:**   
**Date Submission Received:**       **Date NOC Issued:**   
**Targeted CEDAC Meeting:**       **Priority Review Granted:**

Phase	Target Time (Business Days)	Target Date**	Actual CDR Date	Comments	
1	Submission Assessment	5	2009-Sep-11	2009-Sep-11	Category 1 requirements deemed incomplete.
	Submission deemed complete			2009-Sep-15	Submission deemed complete.
2	CDR Reviewers' Reports Completed <ul style="list-style-type: none"> <li>• Reviewers selected and contracted</li> <li>• Literature search and selection completed</li> <li>• Systematic review of clinical data completed</li> <li>• Critical appraisal of pharmaco-economic (PE) data completed</li> <li>• Clinical and PE reports written</li> <li>• Reports edited and finalized</li> <li>• Reviewers' reports sent to manufacturer</li> </ul>	45	2009-Nov-30	2009-Dec-07	Additional information requested September 16, 2009. Additional information received September 18, 2009. Additional information received September 23, 2009. Additional information received October 1, 2009. Additional information requested October 2, 2009. Additional information received October 8, 2009. Additional information requested October 9, 2009. Additional information received October 13, 2009. Additional information requested October 15, 2009. Additional information received October 16, 2009. Additional information received October 19, 2009. Additional information requested October 20, 2009. Additional information requested November 13, 2009. Additional information received November 25, 2009. Additional information requested December 7, 2009.
3	Comments from Manufacturer on Reviewers' Reports Received by CDR	7	2009-Dec-09	2009-Dec-16	Due date for manufacturer comments December 16, 2009.
4	Reviewers' Reply to Manufacturer's Comments Completed	7	2009-Dec-18	2010-Jan-04	Due date for reviewers' reply January 4, 2010.
5	CEDAC Brief Completed and Sent to CEDAC Members	5	2010-Feb-02	2010-Feb-02	
6	CEDAC Meeting		2010-Feb-17	2010-Feb-17	
7	CEDAC Recommendation and Reasons for Recommendation Sent to Drug Plans, ACP and Manufacturer	5	2010-Feb-26	2010-Feb-26	
8	Embargo Period*** Manufacturers may make a Request for Reconsideration and Drug Plans may make a Request for Clarification of the Recommendation and Reasons for Recommendation	10	2010-Mar-12	2010-Apr-12	Request for extension of Embargo period received March 3, 2010. Extension granted, new end date for Embargo Period is April 12, 2010. Request for reconsideration received April 12, 2010. Request for reconsideration granted April 20, 2010.
9 (a)	Final Recommendation sent to Drug Plans, ACP, and Manufacturer (No Requests for Clarification are made AND no Request for Reconsideration is made or Request for Reconsideration is Resolved)	5			
<b>OR</b>					
9 (b)	Clarification and Final Recommendation sent to Drug Plans, ACP, and Manufacturer (Clarification Requested, no Request for Reconsideration made)	5			
<b>OR</b>					
9 (c)	Placed on CEDAC Agenda For Reconsideration (At Manufacturer's request)	25 Depends on Meeting Dates	2010-May-19	2010-May-19	
10	Final Recommendation sent to Drug Plans, ACP, and Manufacturer	5	2010-May-27	2010-May-27	Notice of Final Recommendation issued.

\* Refer to the Procedure for Common Drug Review on the Common Drug Review section of [www.cadth.ca](http://www.cadth.ca) for more details.

\*\* The CDR review process is initiated AFTER a submission is deemed complete. The target dates for this report are based on the CEDAC meeting schedule, which is posted on [www.cadth.ca](http://www.cadth.ca).

\*\*\* The Recommendation and Reasons for Recommendation are to be held in confidence and not acted upon until after the CDR Directorate has issued the notice of Final Recommendation.

Reflects updates as of Thursday noon.