Common Drug Review *

Submission Status

Product: Ralivia

Generic Name: tramadol hydrochloride

Manufacturer: Biovail Pharmaceuticals Canada

Submission Type: New

Date Submission Received: 2007-Oct-29 **Date NOC Issued:** 2007-Aug-31

Targeted CEDAC Meeting: 2008-Mar-19 **Priority Review Granted:** Not Requested

	Phase	Target Time (Business Days)	Target Date**	Actual CDR Date	Comments
1	Submission Assessment	5	2007-Nov-05	2007-Nov-05	
	Submission deemed complete			2007-Nov-05	Submission deemed complete.
2	CDR Reviewers' Reports Completed • Reviewers selected and contracted • Literature search and selection completed • Systematic review of clinical data completed • Critical appraisal of pharmacoeconomic (PE) data completed • Clinical and PE reports written • Reports edited and finalized • Reviewers' reports sent to manufacturer	45	2008-Jan-25	2008-Jan-25	Additional information requested November 19, 2007. Additional information received November 23, 2007.
3	Comments from Manufacturer on Reviewers' Reports Received by CDR	7	2008-Feb-05	2008-Feb-07	Request for extension received February 4, 2008. Extension granted, new due date for manufacturer's comments is February 7, 2008.
4	Reviewers' Reply to Manufacturer's Comments Completed	7	2008-Feb-14	2008-Feb-19	Due date for reviewers' reply February 19, 2008.
5	CEDAC Brief Completed and Sent to CEDAC Members	5	2008-Mar-05	2008-Mar-05	
6	CEDAC Meeting		2008-Mar-19	2008-Mar-19	
7	CEDAC Recommendation and Reasons for Recommendation Sent to Drug Plans, ACP and Manufacturer	5	2008-Mar-27	2008-Mar-27	
8	Embargo Period*** Manufacturers may make a Request for Reconsideration and Drug Plans may make a Reques for Clarification of the Recommendation and Reasons for Recommendation	10	2008-Apr-10	2008-May-08	Request for extension of Embargo received April 9, 2008. Extension to May 8, 2008 granted. Request for Reconsideration received May 8, 2008. Request for Clarification received from ACP April 10, 2008.
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			
	OR				
9 (b)	Clarification and Final Recommendation sent to Drug Plans, ACP, and Manufacturer (Clarification Requested, no Request for Reconsideration made)	5			
	OR				
9 (c)	Placed on CEDAC Agenda For Reconsideration (At Manufacturer's request)	25 Depends on Meeting Dates	2008-Jun-18	2008-Jun-18	
10	Final Recommendation sent to Drug Plans, ACP, and Manufacturer	5	2008-Jun-25	2008-Jun-25	Notice of Final Recommendation issued.

* Refer to the Procedure for Common Drug Review on the Common Drug Review section of 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.

*** 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.