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Canadian Coordinating Office for Health Technology Assessment (CCOHTA)

Common Drug Review *

Submission Status

Product: Strattera

Generic Name: atomoxetine hydrochloride

Manufacturer: Eli Lilly Canada Inc.

Submission Type: NEW

Date Submission Received: 2005-Jan-25 **Date NOC Issued:** 2004-Dec-24

Targeted CEDAC Meeting: 2005-May-18 **Priority Review Granted:** Not Requested

	Phase	Target Time (Business Days)	Target Date**	Actual CDR Date	Comments
1	Submission Deemed Complete	5	2005-Feb-01	2005-Feb-01	
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	2005-Apr-06	2005-Apr-18	Additional information requested February 18, 2005. Additional information received February 21, 2005. Additional information requested February 28, 2005. Additional information received March 1, 2005. Additional information requested March 7, 2005. Additional information received March 9, 2005. Additional information requested March 30, 2005. Additional information received April 4 & 5 & 6, 2005. Additional information requested April 11, 2005. Additional information received April 11, 2005.
3	Comments from Manufacturer on Reviewers' Reports Received by CDR	7	2005-Apr-15	2005-Apr-25	Due date for manufacturer's comments April 25, 2005.
4	Reviewers' Reply to Manufacturer's Comments Completed	7	2005-Apr-26	2005-Apr-28	Due date for reviewer's reply April 28, 2005. Additional information requested April 25, 2005. Additional information received April 26, 2005.
5	CEDAC Brief Completed and Sent to CEDAC Members	5	2005-May-04	2005-May-05	
6	CEDAC Meeting		2005-May-18	2005-May-18	
7	CEDAC Recommendation and Reasons for Recommendation Sent to Drug Plans, ACP and Manufacturer	5	2005-May-26	2005-May-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	2005-Jun-09	2005-Jun-09	Request for reconsideration received June 9, 2005. Additional information requested July 7, 2005. Additional information received July 8, 2005.
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	2005-Sep-21	2005-Sep-21	Discussed at July 27, 2005 CEDAC. Notice of Final Recommendation delayed. Deferred to September 21, 2005 CEDAC.
10	Final Recommendation sent to Drug Plans, ACP, and Manufacturer	5	2005-Sep-28	2005-Sep-28	Notice of Final Recommendation Issued.

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