**Common Drug Review**

**Submission Status**

<table>
<thead>
<tr>
<th>Phase</th>
<th>Target Time (Business Days)</th>
<th>Target Date</th>
<th>Actual CDR Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Submission deemed complete</strong></td>
<td>5</td>
<td>2014-Mar-06</td>
<td>2014-Mar-14</td>
<td>- Submission deemed incomplete: 2014-Mar-06&lt;br&gt;- Submission placed in queue in accordance with CDR procedures. Review to be initiated pending availability of resources and target dates will be updated.&lt;br&gt;- Priority review request under assessment.&lt;br&gt;- Priority review not granted.&lt;br&gt;- Review has been initiated 2014-Sep-12</td>
</tr>
<tr>
<td><strong>Patient group input submission received</strong>&lt;sup&gt;3&lt;/sup&gt;</td>
<td>2014-Aug-28</td>
<td>2014-Aug-28</td>
<td>- Patient input invitations will be posted at a later date (please refer to CDR Update 95)&lt;br&gt;- Call for patient input posted on 2014-Jul-09&lt;br&gt;- Patient group input deadline: 2014-Aug-28&lt;br&gt;- Patient input submission received</td>
<td></td>
</tr>
<tr>
<td><strong>Patient group input summary comments received</strong></td>
<td>5</td>
<td>2014-Oct-07</td>
<td>2014-Oct-07</td>
<td>- Patient input summary sent for review on 2014-Sep-30&lt;br&gt;- Patient input summary feedback deadline: 2014-Oct-07&lt;br&gt;- Patient input summary feedback received</td>
</tr>
<tr>
<td><strong>CDR review reports sent to manufacturer</strong>&lt;sup&gt;4&lt;/sup&gt;</td>
<td>45</td>
<td>2014-May-30</td>
<td>2014-Nov-26</td>
<td>- New target date: 2014-Nov-26</td>
</tr>
<tr>
<td><strong>Comments from manufacturer on CDR review reports received by CADTH</strong></td>
<td>7</td>
<td>2014-Jun-10</td>
<td>2014-Dec-05</td>
<td>- New target date: 2014-Dec-05</td>
</tr>
<tr>
<td><strong>Redaction response from manufacturer on CDR review reports received by CADTH</strong></td>
<td>5</td>
<td>2014-Jun-17</td>
<td>2014-Dec-12</td>
<td>- New target date: 2014-Dec-12</td>
</tr>
<tr>
<td><strong>CDEC meeting</strong></td>
<td>2014-Sep-17</td>
<td>2015-Jan-21</td>
<td>- New target date: 2015-Jan-21</td>
<td></td>
</tr>
<tr>
<td><strong>CDEC recommendation &amp; redacted CDR review reports sent to drug plans and manufacturer</strong>&lt;sup&gt;5&lt;/sup&gt;</td>
<td>5 to 7</td>
<td>2014-Sep-24</td>
<td>2015-Jan-30</td>
<td>- New target date: 2015-Jan-28&lt;br&gt;- New target date: 2015-Jan-30</td>
</tr>
<tr>
<td><strong>Final recommendation sent to drug plans and manufacturer</strong>&lt;sup&gt;7&lt;/sup&gt;</td>
<td>5</td>
<td>2014-Oct-16</td>
<td>2015-Feb-24</td>
<td>- New target date: 2015-Feb-23&lt;br&gt;- New target date: 2015-Feb-24&lt;br&gt;- Notice of final recommendation issued</td>
</tr>
<tr>
<td><strong>CDEC final recommendation posted</strong>&lt;sup&gt;6&lt;/sup&gt;</td>
<td>variable</td>
<td>2015-Feb-26</td>
<td>2015-Feb-26</td>
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<tr>
<td><strong>Final CDR review reports and patient input posted</strong>&lt;sup&gt;7&lt;/sup&gt;</td>
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<tr>
<td><strong>Clarification and final recommendation sent to drug plans and manufacturer</strong>&lt;sup&gt;5&lt;/sup&gt;</td>
<td>- Clarification requested, no request for reconsideration made&lt;br&gt;- Clarification request sent to drug plans and manufacturer</td>
<td>5</td>
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<tr>
<td><strong>Placed on CDEC agenda for reconsideration</strong>&lt;sup&gt;8&lt;/sup&gt;</td>
<td>- (At manufacturer’s request)&lt;br&gt;- 25 depends on Meeting Dates</td>
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1 Refer to the Procedure for Common Drug Review on the Common Drug Review section of www.cadth.ca for more details.

2 The target dates for this report are based on the CDEC meeting schedule, which is posted on www.cadth.ca.

3 The deadline for patient group input is 15 business days after CADTH receives the submission or up to 35 business days if advance notice (20 business days maximum) of a submission is received from the manufacturer.

4 Target time is calculated, based on the date the reviewers receive copies of the manufacturer’s submission. Target time does not include the time allocated for receipt of manufacturer’s additional electronic copies (5 business days) and time allocated for distribution of electronic copies to reviewers (3 business days).

5 The Recommendation is held in confidence by all stakeholders and not acted upon until after CADTH has issued the notice of final recommendation.

6 The target date for posting the CDEC final recommendation depends on several factors including the need for consultation with the manufacturer regarding redaction issues.

7 The timing of the posting of CDR review reports depends on several factors, including the need for consultation with the manufacturer in case of disagreement with regard to redactions made.

8 This Submission Status Report reflects status as of Thursday noon.