Common Drug Review 1					
Canadian Agency for Drugs and Technologies In Health Product:	Submission Status				
Generic Name:					
	Pfizer Canada I				
Indication: Heart failure, NYHA class II					
Submission Type:	New Indication				
Date Submission Received:	2013-Jul-12	Da	te NOC Issued:	2012-Aug-30	
Orginal Targeted CDEC Meeting:	2013-Nov-20 Priority Review Granted:			Not Requested	
Phase	Target Time (Business Days)	Target Date ²	Actual CDR Date	Comment	S
Submission deemed complete	5	2013-Jul-19	2013-Jul-19	 Submission placed in queue in accordance with CDR procedures. Review to be initiated pending availability of resources and target dates will be updated. Review has been initiated 2013-Aug-20 	
Patient group input submission received ³		2013-Aug-02	2013-Aug-02	 Call for patient input posted on 2013-Jul-12 Patient input deadline: 2013-Aug-02 Patient input submission received 	
Patient group input summary comments received	5	2013-Aug-29	2013-Aug-29	 Patient input summary sent for review on 2013-Aug-22 Patient input summary feedback deadline: 2013-Aug-29 Patient input summary feedback received 	
CADTH Reviewers' Reports sent to Manufacturer ⁴	45	2013-Oct-04	2013-Nov-04	- New target date: 2013-Nov-04	
Comments from Manufacturer on Reviewers' Reports Received by CADTH	7	2013-Oct-16	2013-Nov-14	- New target date: 2013-Nov-13 - New target date: 2013-Nov-14	
Redaction response from Manufacturer on Reviewers' Reports Received by CADTH	3	2013-Oct-21	2013-Nov-19	- New target date: 2013-Nov-18 - New target date: 2013-Nov-19	
CDEC Meeting		2013-Nov-20	2014-Jan-15	- New target date: 2014-Jan-15	
CDEC recommendation & redacted CDR review reports sent to drug plans and manufacturer	5 to 7	2013-Nov-27	2014-Jan-22	- New target date: 2014-Jan-22	
Embargo period and validation of redacted CDR review reports ⁵ Manufacturers may make a request for reconsideration and drug plans may make a request for clarification of the recommendation	10	2013-Dec-11	2014-Feb-05	- New target date: 2014-Feb-05	
Final recommendation sent to drug plans and manufacturer (No requests for clarification are made AND no request for reconsideration is made or request for reconsideration is resolved)	5				
CDEC final recommendation posted ⁶	variable				
Final CDR review reports posted ⁷	variable				
OR					
Clarification and final recommendation sent to drug plans and manufacturer (Clarification requested, no request for reconsideration made)	5				
CDEC final recommendation posted ⁶	variable				
Final CDR review reports posted 7	variable				
OR					
Placed on CDEC agenda for reconsideration (At manufacturer's request)	25 Depends on Meeting Dates	2014-Apr-16	2014-Apr-16	- Revised CDR review reports sent to manufacturer 2014-Mar- 14 - Comments from manufacturer on revised CDR review reports received 2014-Mar-19	
Final recommendation sent to drug plans and manufacturer	5	2014-Apr-24	2014-Apr-24	- Notice of final recommendation issue	d
CDEC final recommendation posted ⁶	variable		2014-Apr-28		
Final CDR review reports posted ⁷	variable				

¹ Refer to the Procedure for Common Drug Review on the Common Drug Review section of <u>www.cadth.ca</u> for more details.

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

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 in the redaction 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 in the redaction 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 in the redaction 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 in the redaction of the posting of the posting of CDR review reports dependent on the redaction of the posting of the

This Submission Status Report reflects status as of Thursday noon.

³ The deadline for Patient Group Input is 15 business days after CADTH receives the Submission or up to 25 business days if advance notice 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 binders (5 business days) and time allocated for distribution of binders to reviewers (3 business days).