

CADTH REIMBURSEMENT REVIEW

Stakeholder Feedback on Draft Recommendation

Empagliflozin (Jardiance)

(Boehringer Ingelheim (Canada) Ltd.)

Indication: Heart failure.

September 29, 2022

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CADTH Reimbursement Review

Feedback on Draft Recommendation

Stakeholder information			
CADTH project number		SR0726	
Name of the drug and		Empagliflozin (Jardiance) indicated in adults, as an adjunct to	
Indication(s)		standard of care therapy for the treatment of chronic heart failure	
Organization Providing		FWG	
Feedback			
1. Recommendation revisions Please indicate if the stakeholder requires the expert review committee to reconsider or clarify its recommendation.			
Request for Reconsideration	Major revisions: A change in recommendation category or patient population is requested		
	Minor revisions: A change in reimbursement conditions is requested		
No Request for Reconsideration	Editorial revisions: Clarifications in recommendation text are requested		х
	No req	uested revisions	

2. Change in recommendation category or conditions

Complete this section if major or minor revisions are requested

Please identify the specific text from the recommendation and provide a rationale for requesting a change in recommendation.

3. Clarity of the recommendation

Complete this section if editorial revisions are requested for the following elements

a) Recommendation rationale

- 1) Please clarify the cost condition, in that the incremental cost is related to jardiance alone, as standard of therapy is currently funded.
- 2) It would be helpful if the requirement for patients to be on standard therapies of BB+ACE/ARB/ARNI+MRA was included in the reimbursement conditions.
- 3) Further clarity around the price comparison between empagliflozin and dapagliflozin should be provided, potentially in a discussion point, as the empagliflozin is broader and includes patients with both preserved and reduced ejection fraction. Please provide details regarding the information that requires clarification.

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b) Reimbursement conditions and related reasons

Please provide details regarding the information that requires clarification.

c) Implementation guidance

Please provide high-level details regarding the information that requires clarification. You can provide specific comments in the draft recommendation found in the next section. Additional implementation questions can be raised here.