

OP134 Pan-Canadian Oncology Drug Review Decisions And Access To Anticancer Treatments In Canada

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INTRODUCTION:

The Canadian Agency for Drugs and Technologies in Health (CADTH) pan-Canadian Oncology Drug Review (pCODR) plays an important role in public reimbursement decision-making for oncology drugs in Canada. This research studies the relation of positive pCODR decisions to new cancer treatment and their subsequent inclusion in Canada's public drug plans.

METHODS:

We studied all oncology drugs that received an approval from Health Canada and were reviewed by the pCODR from inception till 26th Sep, 2017. The data was obtained from CADTH and Health Canada. Data such as indication, submission type and date, recommendation date, final recommendation, and subsequent provincial funding status was extracted and analyzed. Impact was evaluated by analyzing the percentage of drug submissions with assessment outcome (positive recommendation rate and conditional recommendation rate) and time taken for the final decision (recommendation gap). The percentage of drugs included in public formulary after positive recommendation by pCODR (coverage rate) and the gap in days from positive recommendation to subsequent coverage in provinces (coverage gap) was also assessed.

RESULTS:

Among 119 drugs reviewed by pCODR, the positive recommendation rate was eight percent. Nine applications comprising seven drugs for six indications received positive recommendations, and genitourinary treatments received maximum positive recommendations. The conditional recommendation rate was 52 percent; 62 applications of 45 drugs for 46 indications received conditional recommendation. Lymphoma and myeloma treatments received maximum conditional recommendations. The average recommendation gap for positive and conditional recommendations was 180 and 172 days, respectively. The coverage rate for drugs with positive recommendation was 100 percent for all provinces except 89 percent for

Newfoundland and Labrador, and 67 percent for Prince Edward Island. Among the provinces, British Columbia had a maximum of 433 days and Saskatchewan has the minimum of 165 days coverage gap.

CONCLUSIONS:

Despite Health Canada's approval, only a fraction of oncology drugs receive positive pCODR recommendation; furthermore, provincial drug plans take time to include these in the reimbursement formularies. While health technology assessment is crucial for appropriate allocation of limited resources, efforts should also be made to reduce access barriers, particularly to positively recommended oncology drugs inclusion in provincial formularies.

OP136 Full Texts Versus Conference Abstract Data: How Does The Message Change?

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INTRODUCTION:

High quality evidence for test accuracy can be scarce. We assessed the test accuracy of two tests (Actim Partus and PartoSure) for the prediction of preterm birth. Twenty published full-text papers were included whilst conference abstracts were excluded. Since systematic reviews of diagnostic tests on other topics may need to rely on data from conference abstracts, we test whether the findings of our review would change with conference abstracts included.

METHODS:

Conference citations previously excluded (n=108) were re-screened for inclusion using the following criteria: i) the diagnostic test was Actim Partus or PartoSure ii) test accuracy data of preterm delivery within seven days was reported iii) the population was women with signs/symptoms of preterm labor with intact membranes. Relevant test accuracy data were extracted and used to calculate sensitivity and specificity. Pooled sensitivity and specificity for each test were run using data from full-text papers and conference abstracts combined.