Patients with cancer are 4–7 times more likely to acquire venous thromboembolism (VTE) than the general population,1 based on the hypercoagulable state associated with the cancer itself, patient characteristics and antineoplastic treatments.2 Despite this increased risk for VTE, clinical guidelines do not recommend the use of primary thromboprophylaxis in unselected ambulatory patients with cancer who are starting chemotherapy and are at intermediate-to-high risk of VTE. We aimed to compare the health system costs and health benefits associated with primary thromboprophylaxis using apixaban with those associated with the current standard of care (where no primary thromboprophylaxis is given), from the perspective of Canada’s publicly funded health care system in this subpopulation of patients with cancer over a lifetime horizon.

Cost–utility analysis of apixaban compared with usual care for primary thromboprophylaxis in ambulatory patients with cancer

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Abstract

Background: Apixaban (2.5 mg) taken twice daily has been shown to substantially reduce the risk of venous thromboembolism (VTE) compared with placebo for the primary thromboprophylaxis of ambulatory patients with cancer who are starting chemotherapy and are at intermediate-to-high risk of VTE. We aimed to compare the health system costs and health benefits associated with primary thromboprophylaxis using apixaban with those associated with the current standard of care (where no primary thromboprophylaxis is given), from the perspective of Canada’s publicly funded health care system in this subpopulation of patients with cancer over a lifetime horizon.

Methods: We performed a cost–utility analysis to estimate the incremental cost per quality-adjusted life-year (QALY) gained with primary thromboprophylaxis using apixaban. We obtained baseline event rates and the efficacy of apixaban from the Apixaban for the Prevention of Venous Thromboembolism in High-Risk Ambulatory Cancer Patients (AVERT) trial on apixaban prophylaxis. We estimated relative risk for bleeding, risk of complications associated with VTE treatment, mortality rates, costs and utilities from other published sources.

Results: Over a lifetime horizon, apixaban resulted in lower costs to the health system (Can$7902.98 v. Can$14 875.82) and an improvement in QALYs (9.089 v. 9.006). The key driver of cost–effectiveness results was the relative risk of VTE as a result of apixaban. Results from the probabilistic analysis showed that at a willingness to pay of Can$50 000 per QALY, the strategy with the highest probability of being most cost-effective was apixaban, with a probability of 99.87%.

Interpretation: We found that apixaban is a cost-saving option for the primary thromboprophylaxis of ambulatory patients with cancer who are starting chemotherapy and are at intermediate-to-high risk of VTE.

Patients with cancer are 4–7 times more likely to acquire venous thromboembolism (VTE) than the general population, based on the hypercoagulable state associated with the cancer itself, patient characteristics and antineoplastic treatments.2 Despite this increased risk for VTE, clinical guidelines do not recommend the use of primary thromboprophylaxis in unselected ambulatory patients with cancer,3–5 because this strategy has been associated with a small absolute reduction in symptomatic VTE and a nonstatistically significant trend in increased major bleeding events.6

The Khorana score uses the cancer type and individual patient characteristics to predict the risk of VTE in patients who are about to begin chemotherapy.7 This score has been evaluated prospectively for its capacity to identify patients with cancer who are at higher risk for VTE and, therefore, may be used to select those patients who are more likely to benefit from primary thromboprophylaxis.8,9 The 2019 Apixaban for the Prevention of Venous Thromboembolism in High-Risk Ambulatory Cancer Patients (AVERT) trial assessed the use of a low-dose direct oral factor Xa inhibitor (apixaban 2.5 mg twice daily) for primary thromboprophylaxis in ambulatory patients with cancer who were starting chemotherapy and were at intermediate-to-high risk of VTE (with a Khorana score ≥ 2).10 The study found that patients randomly assigned to apixaban had a significantly lower risk of VTE compared with placebo. The study also reported that apixaban was not associated with an increase in major bleeding during the on-treatment period. Subsequent to the published results of the AVERT trial and the Efficacy and Safety of Rivaroxaban Prophylaxis Compared with Placebo in Ambulatory Cancer Patients Initiating Systemic Cancer Therapy and at High Risk for Venous Thromboembolism (CASSINI) trial,11 clinical guideline recommendations were updated to endorse the consideration of...
primary thromboprophylaxis in high-risk ambulatory patients with cancer (Khorana score ≥ 2) before the start of chemotherapy.\textsuperscript{4,5} Given the novelty of this recommendation and supporting data, individualized discussions regarding the risk of bleeding, expected benefits and overall costs are also encouraged.

To provide a better framework to support societal discussions on primary thromboprophylaxis in this patient population, we aimed to compare the health system costs and health benefits associated with the use of apixaban primary thromboprophylaxis with those associated with the current standard of care (where no primary thromboprophylaxis is given), from the perspective of Canada’s publicly funded health care system.

**Methods**

**Study design and population**

We conducted a cost–utility analysis that compared health system costs and clinically relevant outcomes relating to apixaban primary thromboprophylaxis with usual care (which did not include the use of primary thromboprophylaxis) among ambulatory patients with cancer who were starting chemotherapy and were at intermediate-to-high risk of VTE. Our analysis addressed the decision problem relating to whether apixaban should be reimbursed for this subpopulation of patients within Canada’s publicly funded health care system.

Consistent with the AVERT trial, our study population included ambulatory patients aged 18 years or older, with a new diagnosis or progression of cancer, who were starting chemotherapy and had a modified Khorana score of 2 or more (modified by the inclusion of myeloma and renal cancer as high-risk, and brain cancers as very high-risk cancer types).\textsuperscript{10} In the AVERT trial, patients were randomly assigned apixaban 2.5 mg twice a day or usual care (i.e., patients received only a placebo). Our primary efficacy outcome was the first major venous thromboembolic event (defined as a proximal deep vein thrombosis [DVT] or pulmonary embolism [PE]). Our primary safety outcome was major bleeding, as defined by the International Society on Thrombosis and Haemostasis (ISTH).\textsuperscript{12} Patients took apixaban for a median of 157 days and were followed for 180 days.

**Model design and inputs**

We used a decision tree and a Markov model (Figure 1) to simulate costs and outcomes for patients who received either apixaban for...
primary thromboprophylaxis or placebo over a lifetime horizon. We conducted scenario analyses to address the model's structural uncertainty relating to the extrapolation of data from a trial with a limited follow-up period (6 mo in the AVERT trial) and the emerging use of direct oral Xa inhibitors for the treatment of cancer-associated thrombosis. Efficacy and costs associated with apixaban were applied over the trial duration (6 mo) for the base-case (lifetime model) and scenario analyses. To account for uncertainty, our model conservatively assumed that there was no difference in the risk of VTE for the 2 treatment arms beyond the trial period. We used a cycle length of 1 month based on the dosing schedules for the treatment of VTE and the expected duration of symptoms in an acute VTE event (before transitioning to more long-term symptoms in some patients). The model was run over a patient’s lifetime horizon (20.6 yr), based on the life expectancy in Canada.13

We separated the model into 2 parts to better capture the dynamic risk of VTE in patients with cancer, and to incorporate differences in costs and risk of complications during different stages of the disease. In the first part, we described the disease pathway for patients with cancer who received primary thromboprophylaxis. The second part described the disease pathway for patients with cancer who had a first VTE event and who were receiving secondary thromboprophylaxis with anticoagulants (beyond the first month of treatment for cancer-associated thrombosis).

Our model included 17 discrete health states (Appendix 1, available at www.cmaj.ca/lookup/doi/10.1503/cmaj.210523/tab-related-content) and was based on the following assumptions: patients received 1 chemotherapy regimen for 6 months before moving into complete remission; patients were assumed to be cured of cancer if they survived 5 years from the start of chemotherapy (meaning that their risk for VTE returns to that of the general population after 5 yr); primary thromboprophylaxis using apixaban was stopped if a first VTE or major bleeding event occurred; patients in the “first VTE” state received treatment-dose dalteparin (at 200 IU/kg) for cancer-associated thrombosis; patients in the “secondary thromboprophylaxis state without complications” continued to receive anticoagulation therapy with dalteparin over their remaining lifetime at a reduced dose (150 IU/kg; patients who had recurrent VTE received a dose escalation of dalteparin); and anticoagulation therapy with dalteparin was stopped if a patient had major bleeding or intracerebral hemorrhage. We chose these assumptions to best fit the current practice in the management of cancer-associated thrombosis and to coincide with the available evidence from the AVERT trial.

We identified transition probabilities, costs and utility values through a targeted literature review. The model structure and input parameters were validated by clinical experts to ensure that they coincided with current clinical practice. All costs were measured in 2020 Canadian dollars. Model input parameters are described in Table 1 and Appendix 1.

We discounted costs and quality-adjusted life-years (QALYs) annually at a discount rate of 1.5%, which is recommended by the Guidelines for the economic evaluation of health technologies: Canada.51

Sensitivity analyses
We performed a series of sensitivity analyses to assess uncertainty in study results. A one-way sensitivity analysis was performed for event rates, costs, proportion of patients with VTE who were treated as inpatients, length of stay for VTE in hospital and utility values used in the model. We also conducted a probabilistic analysis for all parameters in the model using the Monte Carlo method with 10 000 iterations. We used the results of the probabilistic analysis to create a cost-effectiveness acceptability curve, which shows the probability of apixaban being cost-effective over a range of willingness-to-pay (WTP) thresholds.

Results

Over a lifetime horizon, we found that apixaban resulted in lower health system costs (Can$7902.98 v. Can$14 875.82) and an improvement in QALYs (9.089 v. 9.006) in patients with cancer who received chemotherapy, from the perspective of Canada’s health system (Table 2). The key driver of the cost-effectiveness results was the relative risk of VTE when receiving apixaban (Figure 2). Apixaban remained cost-effective compared with usual care across all 1-way sensitivity analyses.

In a cohort of 1000 patients, we found that apixaban resulted in fewer VTE events (57 v. 131), but an increase in major bleeding (129 v. 111) and clinically relevant non-major bleeding (1242 v. 1133) over a lifetime horizon.

Results from the probabilistic analysis showed most iterations lead to cost savings and improved QALYs (Figure 3). At a commonly used WTP of Can$50 000 per QALY, the strategy with the highest probability of being most cost-effective was apixaban (probability of 99.87%; Figure 4).

Our results are robust to changes in time horizon and model assumptions. We performed a scenario analysis over the trial follow-up period of 6 months (scenario analysis 1). Over 6 months, apixaban was associated with a reduction in costs to the health system of Can$257.37 and an improvement in QALYs by 0.001 units. However, we observed wide confidence intervals around the mean incremental QALYs over a 6-month time horizon.

We also performed scenario analyses by varying the proportion of patients with cancer and a first VTE who received direct oral anticoagulants for treatment of cancer-associated thrombosis: 50% of patients received dalteparin and 50% of patients received edoxaban (scenario analysis 2), and 100% of patients received edoxaban (scenario analysis 3). Results from these scenario analyses were consistent with our base case results, showing that apixaban lowered health system costs and improved QALYs over a lifetime horizon in all scenarios (Appendix 2, available at www.cmaj.ca/lookup/doi/10.1503/cmaj.210523/tab-related-content).

Interpretation

We found that the use of apixaban resulted in lower health system costs by Can$6973 and improved QALYs by 0.083 units (30.40 quality-adjusted life-days) compared with usual care. The
The primary driver of the results was the relative efficacy of apixaban. The relative safety of apixaban, costs and utility values had minimal effect on the cost-effectiveness findings. We also performed scenario analyses by varying the time horizon from lifetime to the trial follow-up period (6 mo) and the regimen used for the treatment of cancer-associated thrombosis. Apixaban remained dominant across all scenario analyses. However, owing to the limited data available about apixaban, we observed a high level of uncertainty around the estimates of cost-effectiveness for apixaban over the 6-month time horizon.

Table 1 (part 1 of 2): Model input parameters

| Parameter                                                                 | Mean ± SE*               | Source                                      |
|---------------------------------------------------------------------------|--------------------------|---------------------------------------------|
| **Transition probability**                                                |                          |                                             |
| Baseline risk of primary VTE (0–6 mo)                                     | Time variant risk        | Carrier et al.10                            |
| Baseline risk of primary VTE (6 mo–5 yr)                                  | Time variant risk        | Blix et al.14                               |
| Annual risk of primary VTE (> 5 yr), mean                                 | 0.0001                   | Alostaibi et al.16                          |
| Six-month risk of major bleeding (0–5 yr)                                 | 0.0109 ± 0.0063          | Carrier et al.14                            |
| Annual risk of major bleeding (> 5 yr)                                    | 0.0045 ± 0.0009          | Castellucci et al.15                        |
| Six-month risk of CRNMB                                                   | 0.0509 ± 0.0133          | Carrier et al.14                            |
| Monthly drug stoppage rate (unrelated to death, VTE or bleed)             | 0.0590                   | Carrier et al.14                            |
| **For cancer patients with a history of VTE, who are receiving treatment for secondary thromboprophylaxis** |                          |                                             |
| Annual risk of CRNMB                                                     | 0.1709 ± 0.0167          | Raksob et al.,17 Ihaddadene et al.18        |
| Annual risk of major bleeding (non-ICH)                                   | 0.0495 ± 0.0096          | Raksob et al.,17 Ihaddadene et al.18        |
| Six-month risk of ICH                                                     | 0.0036 ± 0.0026          | Roja-Hernandez et al.,19 Ihaddadene et al.18|
| Annual risk of recurrent VTE                                             | 0.1345 ± 0.0151          | Raksob et al.17                             |
| Two-year risk of CTEPH                                                    | 0.0320 ± 0.0061          | Ende-Verhaar et al.20                       |
| Two-year risk of PTS                                                      | 0.1270 ± 0.0168          | Kahn et al.21                               |
| **For cancer patients with a history of VTE, who are not receiving treatment for secondary thromboprophylaxis** |                          |                                             |
| Six-month risk of CRNMB                                                  | 0.0509 ± 0.0133          | Carrier et al.14                            |
| Six-month risk of major bleeding (non-ICH)                               | 0.0109 ± 0.0063          | Carrier et al.14                            |
| Annual risk of ICH                                                        | 0.0003 ± 0.0001          | Sacco et al.22                              |
| Annual risk of recurrent VTE                                             | 0.0838 ± 0.0086          | Cohen et al.23                              |
| **Mortality**                                                             |                          |                                             |
| Baseline age-adjusted mortality for general population                    | 10.97                    | Statistics Canada24                          |
| Excess mortality due to cancer, hazard ratio                              |                          | Canadian Cancer Society25                    |
| Excess mortality due to VTE, mean (95% CI)                                | 2.20 (2.05 to 2.40)      | Sorensen et al.16                           |
| Excess mortality due to major bleeding, mean (95% CI)                     | 2.10 (1.60 to 2.90)      | Nagata et al.27                             |
| Excess mortality due to ICH, mean (95% CI)                                | 2.60 (2.09 to 3.24)      | Gonzalez-Perez et al.28                     |
| Excess mortality due to CTEPH, mean (95% CI)                              | 12.25 (10.27 to 14.31)   | Delcroix et al.29                           |
| **Relative risk due to apixaban**                                         |                          |                                             |
| CRNMB, mean (95% CI)                                                      | 1.296 (0.663 to 2.533)   | Carrier et al.15                            |
| Major bleeding, mean (95% CI)                                             | 1.960 (0.800 to 4.820)   | Pooled from AVERT10 and CASSINI11 trials; Li et al.34|
| VTE, mean (95% CI)                                                       | 0.143 (0.043 to 0.477)   | Carrier et al.29                            |
| Proportion of patients with ICH who have a major ICH                     | 0.50                     | Murthy et al.31                             |
| Proportion of patients who experience major bleeding and resume anticoagulation treatment | 0.00                     | Li et al.32                                 |
| **DVT and PE management**                                                |                          |                                             |
| DVT length of stay, d; mean (95%CI)                                       | 6.70 (5.00 to 8.00)      | CADTH13                                     |
| DVT proportion managed as inpatient, mean (95%CI)                         | 0.19 (0.00 to 0.40)      | CADTH13                                     |
| PE length of stay, d; mean (95%CI)                                       | 7.80 (6.00 to 9.00)      | CADTH13                                     |
| PE proportion managed as inpatient, mean (95%CI)                         | 0.67 (0.30 to 0.75)      | CADTH13                                     |
## Table 1 (part 2 of 2): Model input parameters

| Parameter                              | Mean ± SE* | Source                                                                 |
|----------------------------------------|------------|------------------------------------------------------------------------|
| **Cost, $**                            |            |                                                                        |
| CRNMB treatment                        | 383 ± 122  | CADTH33                                                               |
| Major bleeding treatment (non-ICH)     | 9191 ± 2424| MOHLTC24 (OSoB), MOHLTC25 (OCCI)                                      |
| ICH, mean ± SD                         | 16 962 ± 16 705 | Specogna et al.36                                                   |
| Post-ICH                               | 756 ± 25% of base case | CADTH,23 Goeree et al.37                                               |
| CTEPH treatment                        | 91 412 ± 25% of base case | CADTH,22 Delcroix et al.28                                            |
| Post-CTEPH management                  | 140 ± 25% of base case | CADTH23                                                               |
| PTS treatment                          | 8181 ± 25% of base case | CADTH,22 Caprini et al.35                                              |
| Post-PTS management                    | 299 ± 25% of base case | CADTH,23 Caprini et al.35                                              |
| **Primary VTE treatment**              |            |                                                                        |
| DVT outpatient                         | 759        | CADTH,22 MOHLTC (OSoB),24 MOHLTC25 (OCCI)                             |
| DVT per inpatient day, mean (95%CI)    | 15 58 (1000 to 1947) | MOHLTC26 (OCCI)                                                      |
| PE outpatient                          | 1551       | CADTH,22 MOHLTC (OSoB),24 MOHLTC25 (OCCI)                             |
| PE per inpatient day, mean (95%CI)     | 1655 (1000 to 2563) | MOHLTC25 (OCCI)                                                      |
| Medication — LMWH                      | 1221.58    | MOHLTC28 (ODBF)                                                       |
| Medication — DOAC                      | 274.60     | MOHLTC28 (ODBF)                                                       |
| Recurrent VTE treatment                | 8083       | †                                                                     |
| Post VTE management — LMWH             | 937 ± 25% of base case | MOHLTC24 (OSoB), Dranitsaris et al.,40 MOHLTC28 (ODBF)               |
| Post VTE management — DOAC             | 144 ± 25% of base case | MOHLTC24 (OSoB), MOHLTC28 (ODBF)                                     |
| Apixaban per month                     | 98.02      | MOHLTC28 (ODBF)                                                       |
| **Utility value**                      |            |                                                                        |
| Baseline health utility value for patients with cancer | 0.824 ± 0.045 | Sullivan et al.,41 McCarter et al.,42 Allareddy et al.,43 Best et al.,44 Curran et al.,45 Doyle et al.,46 Fossà et al.,47 Uyl-de Groot et al.,48 Jewell et al.,49 Klahn et al.,50 Kulkarni et al.,51 Papaioannou et al.,52 Pelligrina et al.,53 Rogers et al.,54 Romanus et al.,55 Shiroiwa et al.,56 Stewart et al.57 |
| Disutility: primary or recurrent VTE   | 0.142 ± 0.022 | Hogg et al.18                                                          |
| Disutility: CRNMB                      | 0.013 ± 0.003 | Sullivan et al.41                                                      |
| Disutility: MB (non-ICH)               | 0.270 ± 0.024 | Hogg et al.18                                                          |
| Disutility: major ICH                  | 0.770 ± 0.166 | Hogg et al.18                                                          |
| Disutility: minor ICH                  | 0.170 ± 0.094 | Hogg et al.18                                                          |
| Disutility: ICH (weighted average of major and minor ICH) | 0.470 ± 0.130 | Hogg et al.18                                                          |
| Utility: Post ICH                      | 0.150 ± 0.166 | Hogg et al.18                                                          |
| Disutility: CTEPH                      | 0.360 ± 0.016 | CADTH,33 Meads et al.59                                                |
| Utility: Post CTEPH                    | 0.560 ± 0.016 | CADTH,33 Meads et al.59                                                |
| Disutility: PTS                        | 0.050 ± 0.022 | Li et al.,32 Lenert et al.60                                            |
| Utility: Post PTS state                | 0.774 ± 0.045 | Lenert et al.60                                                        |

Note: Detailed description for each input parameter presented in Appendix 1, Table A1 (available at www.cmaj.ca/lookup/doi/10.1503/cmaj.210523/tab-related-content). CADTH = Canadian Agency for Drugs and Technologies in Health, CI = confidence interval, CRNMB = clinically relevant non-major bleeding, CTEPH = chronic thromboembolic pulmonary hypertension, DOAC = direct oral anticoagulants, DVT = deep vein thrombosis, ICH = intracerebral hemorrhage, LMWH = low-molecular-weight heparin, MOHLTC = Ontario Ministry of Health and Long-Term Care, OCCI = Ontario Case Costing Initiative, ODBF = Ontario Drug Benefit Formulary, OSO = Ontario Schedule of Benefits, PE = pulmonary embolism, PTS = post-thrombotic syndrome, SE = standard error, VTE = venous thromboembolism.

*Unless stated otherwise.

†Sources for resource use and unit costs for recurrent VTE are the same as those for primary VTE. Dosage for patients with recurrent VTE was assumed to be 120% of that for primary VTE.
Table 2: Results from base-case analysis

| Treatment          | Mean cost (95% CI), (Can$) | Mean QALYs (95% CI) | Mean LYs (95% CI) |
|--------------------|-----------------------------|---------------------|-------------------|
| Usual care         | 14,875.82 (10,511.47 to 21,952.46) | 9.006 (8.150 to 9.613) | 12.658 (12.606 to 12.697) |
| Apixaban           | 7,902.98 (5,500.67 to 13,216.39) | 9.089 (8.177 to 9.732) | 12.738 (12.685 to 12.760) |
| Incremental difference (apixaban v. usual care) | –6,972.84 (–11,324.68 to –3,697.13) | 0.083 (0.013 to 0.157) | 0.080 (0.044 to 0.114) |

Note: CI = confidence interval, LY = life-year, QALY = quality-adjusted life-year.

Figure 2: One-way sensitivity analyses. Note: CRNMB = clinically relevant non-major bleeding, CTEPH = chronic thromboembolic pulmonary hypertension, DVT = deep vein thrombosis, ICH = intracranial hemorrhage, PE = pulmonary embolism, PTS = postthrombotic syndrome, RR = relative risk, VTE = venous thromboembolism.
In 2020, a cost-effectiveness analysis involving a hypothetical cohort in the United States suggested that low-dose direct oral Xa inhibitors were cost-effective for the prevention of VTE in ambulatory patients with cancer who started chemotherapy and were at increased risk of VTE (Khorana score ≥ 2), with an incremental cost-effectiveness ratio of US$11,947 per QALY gained. This analysis was based on the pooled efficacy and safety results of both the AVERT and CASSINI trials. Our results are consistent with those from the US study. However, our cost-effectiveness analysis differs from the US study in several noteworthy ways. First, our analysis involved Canada’s publicly funded health care system. Therefore, costs and health system usage are reflective of this environment. Second, we made use of patient-level data from the AVERT trial, instead of summary results, which increases the specificity of our results for this patient population in Canada. Third, although combining the efficacy and safety results from the AVERT and CASSINI trials has the advantage of decreasing the uncertainty around these estimates, there are important differences between the 2 trials, which should lead to a more nuanced interpretation of each trial’s results and caution when combining their data. The patient populations in the 2 trials differed, most likely because of the use of the modified Khorana score in the AVERT trial and the Khorana score in the CASSINI trial to identify potential participants and the exclusion of patients with intracranial disease in the CASSINI trial. In addition, the CASSINI trial mandated the use of serial screening compression ultrasonography of the lower extremities throughout the follow-up period, which is not standard of care in ambulatory patients with cancer. The cost, impact on patients’ quality of life and most appropriate use of screening compression ultrasonography are unknown. Finally, the definition of an outcome was inconsistent between the 2 studies. Although both the AVERT and CASSINI trials used a composite outcome (by combining end points with varying degrees of clinical severity), the CASSINI trial included VTE events that were of lesser or unknown clinical significance, such as symptomatic distal or proximal DVT found on
screening compression ultrasonography. These differences in patient populations, use of screening compression ultrasonography to identify asymptomatic DVT, and the divergence in composite outcomes to include less clinically significant and screening-detected events may account for the lack of significant efficacy of rivaroxaban in the CASSINI trial, which is contrary to the findings in the AVERT trial.

Although additional data on the optimal duration and effectiveness of primary thromboprophylaxis using apixaban are needed, our results support the coverage of low-dose apixaban by Canada’s publicly funded health care system for primary thromboprophylaxis in ambulatory patients with cancer who are starting chemotherapy and have a modified Khorana score of 2 or more. The prevention of VTE during the first 6 months after the start of chemotherapy translates to cost savings, which persist throughout the patient’s lifetime, by avoiding future complications including VTE, chronic complications of VTE and increased bleeding risk when receiving anticoagulation for the management of VTE. The use of apixaban for primary thromboprophylaxis appears to be a well-tolerated intervention, which further increases quality of life for patients with cancer. Given the importance and increasing costs associated with novel cancer treatment, strategies that alleviate the financial impact of providing high-quality patient-centred care to patients with cancer must be sought and adopted to promote the sustainability of the health care system. Primary thromboprophylaxis in appropriately chosen ambulatory patients with cancer appears to be such an intervention.

Limitations
There is uncertainty regarding the efficacy of apixaban beyond the trial follow-up period because data are lacking on long-term outcomes of thromboprophylaxis using apixaban. To account for this uncertainty, our model conservatively assumed that there was no difference in the risk of VTE for the 2 treatment arms beyond the trial period. In addition, there was a high level of uncertainty around the estimate of the relative safety of apixaban derived from the AVERT trial because of the small number of outcomes. Therefore, our study used synthesis-based estimates to supplement the patient-level data from the AVERT trial for the relative safety of apixaban by combining the major bleeding events from the AVERT and CASSINI trials that evaluated the use of oral anticoagulants for thromboprophylaxis among this patient population. This practice was justified because both trials used the same definitions for bleeding outcomes. Furthermore, we were unable to perform subgroup analysis by Khorana score or tumour type because the AVERT trial was not sufficiently powered to detect statistically significant differences by patient subgroups. Our study used data on resource use for the treatment of VTE, clinically relevant non-major bleeding and major bleeding derived from populations of patients without cancer. Previous research has shown that length of stay for treatment of VTE among patient populations with cancer is longer than that for those without cancer. This suggests that our finding of cost savings due to reduction in incidence of VTE attributable to apixaban may be an underestimation. In addition, we did not consider low-molecular-weight heparin as a comparator because of the lack of direct or indirect evidence comparing the efficacy of apixaban to low-molecular-weight heparin among ambulatory patients with cancer. Finally, our results are not generalizable to patients with cancer who have undergone surgery but have not received chemotherapy because only patients who started a new course of chemotherapy were eligible for the AVERT trial.

Conclusion
We found that within Canada’s publicly funded health system, primary thromboprophylaxis with apixaban in selected ambulatory patients with cancer is cost saving.

References
1. Timp JF, Braekkan SK, Versteeg HH, et al. Epidemiology of cancer-associated venous thrombosis. Blood 2013;122:1712–23.
2. Ay C, Fablinger I, Cohen AT. Cancer-associated venous thromboembolism: burden, mechanisms, and management. Thromb Haemost 2017;117:219–30.
3. Mandalà M, Falanga A, Rolla F; ESMO Guidelines Working Group. Management of venous thromboembolism (VTE) in cancer patients: ESMO Clinical Practice Guidelines, Ann Oncol 2011;22(Suppl 6):vi85–92.
4. Key NS, Khorana AA, Kuderer NM, et al. Venous thromboembolism prophylaxis and treatment in patients with cancer: ASCO clinical practice guideline update. J Clin Oncol 2020;38:496–520.
5. Lyman GH, Carrier M, Ay C, et al. American Society of Hematology 2021 guidelines for management of venous thromboembolism: prevention and treatment in patients with cancer. Blood Adv 2021;5:927–74.
6. Di Nisio M, Porreca E, Candeloro M, et al. Primary prophylaxis for venous thromboembolism in ambulatory cancer patients receiving chemotherapy. Cochrane Database Syst Rev 2016;12:CD008500.
7. Khorana AA, Kuderer NM, Culakova E, et al. Development and validation of a predictive model for chemotherapy-associated thrombosis. Blood 2008;111:4902–7.
8. Ay C, Dunkler D, Marosi C, et al. Prediction of venous thromboembolism in cancer patients. Blood 2010;116:5377–82.
9. Khorana AA, Francis CW. Risk prediction of cancer-associated thrombosis: appraising the first decade and developing the future. Thromb Res 2018;164(Suppl 1):S70–6.
10. Carrier M, Abou-Nassar K, Mallick R, et al.; AVERT Investigators. Apixaban to prevent venous thromboembolism in patients with cancer. N Engl J Med 2019;380:711–9.
11. Khorana AA, Soff GA, Kakkar AK, et al.; CASSINI Investigators. Rivaroxaban for thromboprophylaxis in high-risk ambulatory patients with cancer. N Engl J Med 2019;380:720–8.
12. Schulman S, Kearon C; Subcommittee on Control of Anticoagulation of the Scientific and Standardization Committee of the International Society on Thrombosis and Haemostasis. Definition of major bleeding in clinical investigations of antithrombotic medicinal products in non-surgical patients. J Thromb Haemost 2005;3:692–4.
13. Table 13-10-0389-01: Life expectancy, at birth and at age 65, by sex, three-year average, Canada, provinces, territories, health regions and peer groups. Ottawa: Statistics Canada. Available: https://www150.statcan.gc.ca/t1/tbl1/en/tv.action?pid=1310038901 (accessed 2019 Oct. 19).
14. Blix K, Gran OV, Severinsen MT, et al. Impact of time since diagnosis and mortality rate on cancer-associated venous thromboembolism: the Scandinavian Thrombosis and Cancer (STAC) cohort. J Thromb Haemost 2018;16:1327–35.
15. Alotaibi GS, Wu C, Senthilvelan A, et al. Secular trends in incidence and mortality of acute venous thromboembolism: the AB-VTE population-based study. Am J Med 2016;129:879.e19–25.
16. Castellucci LA, Le Gal G, Rodger MA, et al. Major bleeding during secondary prevention of venous thromboembolism in patients who have completed anticoagulation: a systematic review and meta-analysis. J Thromb Haemost 2014;12:344–8.
17. Raskob GE, van Es N, Verhamme P, et al. Edoxaban for the treatment of cancer-associated venous thromboembolism. N Engl J Med 2018;378:615-24.

18. Ihaddadene R, Le Gal G, Delluc A, et al. Dose escalation of low molecular weight heparin in patients with recurrent cancer-associated thrombosis. Thromb Res 2014;134:53-9.

19. Rojas-Hernandez CM, Oo TH, Garcia-Perdomo HA. Risk of intracranial hemorrhage associated with therapeutic anticoagulation for venous thromboembolism in cancer patients: a systematic review and meta-analysis. J Thromb Thrombolysis 2017;43:233-40.

20. Ende-Verhaar YM, Cannegie SC, Vonk Noordegraaf A, et al. Incidence of chronic thromboembolic pulmonary hypertension after acute pulmonary embolism: a contemporary view of the published literature. Eur Respir J 2017;49:1610792.

21. Kahn SR, Shapiro S, Wells PS, et al.; SOX trial investigators. Compression stockings to prevent post-thrombotic syndrome: a randomised placebo-controlled trial. Lancet 2014;383:880-8.

22. Sacco S, Marini C, Toni D, et al. Incidence and 10-year survival of intracerebral hemorrhage in a population-based registry. Stroke 2009;40:394-9.

23. Cohen AT, Katholing A, Rietbrock S, et al. Epidemiology of first and recurrent venous thromboembolism in patients with active cancer. A population-based cohort study. Thromb Haemost 2017;117:57-65.

24. Tables: Life tables, Canada, provinces, and territories. Tables 84-537-X. Ottawa: Statistics Canada. Available: https://www150.statcan.gc.ca/n1/en/catalogue-84-537-X (accessed 2019 Oct. 18).

25. Canadian Cancer Statistics Advisory Committee, Canadian Cancer Statistics: a 2018 special report on cancer incidence by stage. Toronto: Canadian Cancer Society; 2018. Available: https://www.canada.ca/en/cancer/cancer%20information/cancer%20incidence/cancer%20incidence-by-stage.html (accessed 2019 Nov. 22).

26. Sørensen HT, Mellemkjaer L, Olsen JH, et al. Prognosis of cancers associated with venous thromboembolism. N Engl J Med 2000;343:1846-50.

27. Nagata N, Sakurai T, Shimbo T, et al. Acute severe gastrointestinal tract bleed with venous thromboembolism in patients with advanced adenocarcinoma of the stomach or esophagogastric junction treated with irinotecan combined with 5-fluorouracil and folinic acid: results of a randomised phase III trial. Qual Life Res 2009;18:853-61.

28. Sullivan PW, Slejkof JF, Sculpher MJ, et al. Catalogue of EQ-5D scores for the United Kingdom. Med Decis Making 2011;31:800-4.

29. Dranitsaris G, Shane LG, Crowther M, et al. Value Health 2006;9:243-54.

30. Caprini JA, Botteman MF, Stephens JM, et al. Economic burden of lethal complications of deep vein thrombosis after total hip replacement surgery in the United States. Value Health 2003;6:59-74.

31. Murthy SB, Shastri A, Merkler AE, et al. Intracerebral hemorrhage outcomes in patients with multiple myeloma undergoing a double transplantation. Eur J Haematol 2005;74:136-43.

32. Jewell EL, Smythe M, Brodbarow G, et al. Utility scores and treatment preferences for clinical early-stage cervical cancer. Value Health 2011;14:582-6.

33. poker G, Bremner KE, Alihah SMH, et al. A reference set of health utilities for long-term survivors of prostate cancer: population-based data from Ontario, Canada. Qual Life Res 2013;22:2951-62.

34. Kulkarni GS, Alihah SMH, Finelli A, et al. Cost-effectiveness analysis of immediate radical cystectomy versus intravesical Bacillus Calmette-Guerin therapy for high-risk, high-grade (T1G3) bladder cancer. Cancer 2009;115:5450-9.

35. Sullivan PW, Slejkof JF, Sculpher MJ, et al. Catalogue of EQ-5D scores for the United Kingdom. Med Decis Making 2011;31:800-4.

36. Sacco S, Marini C, Toni D, et al. Incidence and 10-year survival of intracerebral hemorrhage in a population-based registry. Stroke 2009;40:394-9.

37. Raskob GE, van Es N, Verhamme P, et al. Edoxaban for the treatment of cancer-associated venous thromboembolism. N Engl J Med 2018;378:615-24.

38. Ihaddadene R, Le Gal G, Delluc A, et al. Dose escalation of low molecular weight heparin in patients with recurrent cancer-associated thrombosis. Thromb Res 2014;134:53-9.

39. Rojas-Hernandez CM, Oo TH, Garcia-Perdomo HA. Risk of intracranial hemorrhage associated with therapeutic anticoagulation for venous thromboembolism in cancer patients: a systematic review and meta-analysis. J Thromb Thrombolysis 2017;43:233-40.

40. Ende-Verhaar YM, Cannegie SC, Vonk Noordegraaf A, et al. Incidence of chronic thromboembolic pulmonary hypertension after acute pulmonary embolism: a contemporary view of the published literature. Eur Respir J 2017;49:1610792.

41. Kahn SR, Shapiro S, Wells PS, et al.; SOX trial investigators. Compression stockings to prevent post-thrombotic syndrome: a randomised placebo-controlled trial. Lancet 2014;383:880-8.

42. Sacco S, Marini C, Toni D, et al. Incidence and 10-year survival of intracerebral hemorrhage in a population-based registry. Stroke 2009;40:394-9.
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