

The IMPROVEDD VTE Risk Score: Incorporation of D-Dimer into the IMPROVE Score to Improve Venous Thromboembolism Risk Stratification

C. Michael Gibson¹ Alex C. Spyropoulos² Alexander T. Cohen³ Russell D. Hull⁴
 Samuel Z. Goldhaber⁵ Roger D. Yusen⁶ Adrian F. Hernandez⁷ Serge Korjian¹ Yazan Daaboul¹
 Alex Gold⁸ Robert A. Harrington⁹ Gerald Chi¹

¹Cardiovascular Division, Department of Medicine, Beth Israel Deaconess Medical Center, Harvard Medical School, Boston, Massachusetts, United States

²Department of Medicine, Anticoagulation and Clinical Thrombosis Services, Hofstra Northwell School of Medicine, Northwell Health System, Manhasset, New York, United States

³Department of Haematological Medicine, Guy's and St Thomas' Hospitals, King's College, London, United Kingdom

⁴Division of Cardiology, Faculty of Medicine, University of Calgary, Alberta, Canada

⁵Cardiovascular Division, Brigham and Women's Hospital, Harvard Medical School, Boston, Massachusetts, United States

⁶Divisions of Pulmonary and Critical Care Medicine and General Medical Sciences, Washington University School of Medicine, St. Louis, Missouri, United States

⁷Department of Medicine, Duke University and Duke Clinical Research Institute, Durham, North Carolina, United States

⁸Portola Pharmaceuticals, Inc., South San Francisco, California, United States

⁹Division of Cardiovascular Medicine, Department of Medicine, Stanford University, Stanford, California, United States

Address for correspondence Gerald Chi, MD, PERFUSE Study Group, Cardiovascular Division, Department of Medicine, Beth Israel Deaconess Medical Center, Harvard Medical School, 330 Brookline Avenue, Suite OV-540, Boston, MA 02215, United States (e-mail: geraldccchi@gmail.com).

TH Open 2017;1:e56–e65.

Abstract

Background The IMPROVE score is a validated venous thromboembolism (VTE) assessment tool to risk stratify hospitalized, medically ill patients based on clinical variables. It was hypothesized that addition of D-dimer measurement to derive a new IMPROVEDD score would improve identification of at risk of VTE.

Methods The association of the IMPROVE score and D-dimer $\geq 2 \times$ the upper limit of normal (ULN) with the risk of symptomatic deep vein thrombosis, nonfatal pulmonary embolism, or VTE-related death was evaluated in 7,441 hospitalized, medically ill patients randomized in the APEX trial. Based on the Cox regression analysis, the IMPROVEDD score was derived by adding two points to the IMPROVE score if the D-dimer was $\geq 2 \times$ ULN.

Results Baseline D-dimer was independently associated with symptomatic VTE through 77 days (adjusted HR: 2.22 [95% CI: 1.38–1.58], $p = 0.001$). Incorporation of D-dimer into the IMPROVE score improved VTE risk discrimination (Δ AUC: 0.06 [95% CI: 0.02–0.09], $p = 0.0006$) and reclassification (continuous NRI: 0.34 [95% CI: 0.17–0.51], $p = 0.001$; categorical NRI: 0.13 [95% CI: 0.03–0.23], $p = 0.0159$). Patients with an IMPROVEDD score of ≥ 2 had a greater VTE risk compared with those with an IMPROVEDD score of 0 to 1 (HR: 2.73 [95% CI: 1.52–4.90], $p = 0.0007$).

Keywords

- ▶ venous thromboembolism
- ▶ D-dimer
- ▶ risk assessment model
- ▶ thromboprophylaxis

Conclusion Incorporation of D-dimer into the IMPROVE VTE risk assessment model further improves risk stratification in hospitalized, medically ill patients who received thromboprophylaxis. An IMPROVEDD score of ≥ 2 identifies hospitalized, medically ill patients with a heightened risk for VTE through 77 days.

Introduction

Venous thromboembolism (VTE) is a major contributor to the global disease burden with an estimated incidence of 3.0 to 3.3 cases per 100 hospitalizations per year.¹ Hospitalized, medically ill patients represent a population with heterogeneous predisposition to VTE for which risk assessment is recommended prior to thromboprophylaxis.^{2–4} However, existing risk assessment models (RAMs) may not adequately identify at-risk subsets, and risk stratification remains an ongoing challenge and imprecise science.^{5,6} D-dimer, a biomarker for fibrinolysis, has been associated with heightened VTE risk among patients hospitalized for an acute medical illness.^{7–9} The International Medical Prevention Registry on Venous Thromboembolism (IMPROVE) VTE RAM has undergone extensive external validation in the medically ill population.^{6,10,11} In this article, D-dimer was incorporated into the IMPROVE VTE RAM to derive the IMPROVEDD VTE risk score.¹² The model-based performance was tested in a cohort of hospitalized medical patients receiving primary pharmacologic prophylaxis. It was hypothesized that incorporation of the biomarker D-dimer would provide incremental prognostic value to the IMPROVE RAM in identifying patients at risk of developing symptomatic deep vein thrombosis, non-fatal pulmonary embolism, and VTE-related death.

Methods

Study Design

The Acute Medically Ill VTE (Venous Thromboembolism) Prevention with Extended Duration Betrixaban Trial (APEX; ClinicalTrials.gov identifier: NCT01583218) was a randomized, double-blind, multinational clinical trial that compared extended-duration betrixaban (80 mg once daily for 35–42 days) to standard-duration enoxaparin (40 mg once daily for 10 ± 4 days) among hospitalized medical patients.^{9,13} Four principal enrollment criteria were as follows: (1) hospitalization for acute medical illness, including heart failure, respiratory failure, infection, ischemic stroke, or rheumatic disorder; (2) age ≥ 75 years, age 60 to 74 years with D-dimer $\geq 2 \times$ the upper limit of normal (ULN), or age 40 to 59 years with D-dimer $\geq 2 \times$ ULN and history of VTE or cancer; (3) anticipated severe immobilization for ≥ 24 hours followed by moderate or severe immobilization for 3 or more days; and (4) anticipated hospitalization for 3 or more days. Serum samples for D-dimer were obtained at the time of screening and sent to the central laboratory for analysis using the quantitative STA Liatest D-Di immunoturbidimetric assay (Diagnostica Stago, Asnières-sur-Seine, France). Endpoints were assessed at 42 and 77 days after randomiza-

tion to approximate the 90-day endpoint from the original IMPROVE VTE RAM. All VTE events were adjudicated by an independent clinical events committee blinded to thromboprophylaxis allocation based on the documentation from the case report form, the narratives prepared by the study sites, and any other available supporting source documentation. The process of events adjudication occurred in two phases (Phase I and Phase II). Phase I review was conducted by two independent physicians. If the Phase I reviewers agreed in the adjudication of the event, the process was complete. If the Phase I reviewers did not agree, the event was submitted for adjudication by Phase II physician committee. In the Phase II meeting, each case was decided by majority rule of the Phase II reviewers. The enrollment period was from March 2012 to October 2015 and the follow-up of the last patient was completed in January 2016.

Statistical Analysis

All randomized patients fulfilling the enrollment criteria were included in the analyses. Baseline characteristics among patients with and without events were compared using the χ^2 test for categorical variables and the one-way ANOVA or Kruskal-Wallis test for continuous variables, as appropriate. Cumulative incidence of symptomatic VTE from randomization to 42 and 77 days was estimated by the Kaplan-Meier method. To test the association between D-dimer and VTE-related events, univariate and multivariate regression analyses evaluated variables included in the IMPROVE associative model (i.e., previous VTE, known thrombophilia, current lower-limb paralysis, current cancer, immobilized ≥ 7 days, intensive care unit (ICU) or coronary care unit (CCU) stay, and age >60 years). An additional sensitivity analysis that considered the confounding effect of thromboprophylaxis on this association was performed. To determine the appropriate weight for D-dimer, the risk estimate of D-dimer relative to the per point increase in the IMPROVE score was calculated from the adjusted Cox proportional hazards model. Consequently, the IMPROVEDD score was calculated by adding two points to the IMPROVE score if the D-dimer level was $\geq 2 \times$ ULN.

The model-based probability for the IMPROVE and IMPROVEDD score was estimated using logistic regression analysis. Metrics of model discrimination and reclassification were computed to assess the improvement in VTE predictability by the IMPROVEDD score, including area under the receiver-operating-characteristic curve (AUC), integrated discrimination improvement (IDI), and net reclassification improvement (NRI).¹⁴ Model calibration was assessed by comparing the observed and predicted risk for

each IMPROVEDD score category. In addition, decision curve analysis was performed to compare the net benefit of the IMPROVE and IMPROVEDD score in VTE prediction.¹⁵ A cutoff for the IMPROVEDD score corresponding to an event rate of $\geq 1\%$, as recommended for warranting pharmacologic prophylaxis by the American College of Chest Physicians (ACCP) guidelines, was used to dichotomize patients as at-risk versus low-risk category. Accordingly, the risk for symptomatic VTE was compared between the at-risk warranting prophylaxis (≥ 2 points) and low-risk (0–1 points) categories. Analyses were performed independently by an academic research organization, Percutaneous-Pharmacologic Endoluminal Revascularization for Unstable Syndromes Evaluation (PERFUSE) Study Group, using SAS software version 9.4 (SAS Institute, Inc., Cary, North Carolina, United States).

Results

Baseline Characteristics

Patients who developed symptomatic VTE at 77 days were hospitalized longer and were more likely to have had a previous VTE, ICU or CCU stay, higher IMPROVE score, and D-dimer $\geq 2 \times$ ULN (**Table 1**). Age, sex, race, weight, height, body mass index, creatinine clearance, and acute medical condition were balanced between patients with events and without events.

Risk Stratification by D-Dimer

Baseline D-dimer levels were measured in 7,235 patients. The risk for symptomatic VTE was significantly higher among patients with D-dimer $\geq 2 \times$ ULN compared with $< 2 \times$ ULN, with respective rates of 1.11 versus 0.57% at 42 days and 2.37 versus 1.00% at 77 days (HR: 2.26 [95% CI: 1.41–3.64], $p = 0.0008$; **Fig. 1**). Of the seven risk factors identified in the IMPROVE RAM study, previous VTE and ICU or CCU stay were statistically associated with symptomatic VTE in the study population (**Table 2**). Multivariate analysis confirmed that D-dimer was independently associated with symptomatic VTE at 42 days (adjusted HR: 2.28 [95% CI: 1.35–3.85], $p = 0.0020$) and at 77 days (adjusted HR: 2.22 [95% CI: 1.38–3.58], $p = 0.0010$). When thromboprophylaxis allocation was considered in the model, estimates for the VTE risk factors do not alter substantially (**Table S1**, supplementary table available in the online version only).

Derivation of the IMPROVEDD VTE Risk Score

In the Cox proportional hazards model, symptomatic VTE risk was approximately 2.26 to 2.33 times greater among patients with D-dimer $\geq 2 \times$ ULN compared with $< 2 \times$ ULN (**Table 3**). The risk was approximately 1.22 to 1.26 times greater for each point increase in the IMPROVE score. There was no significant interaction between D-dimer and the IMPROVE score in the model ($p = 0.41$ at 42 days and $p = 0.61$ at 77 days). Based on the relative size of the risk estimates, the IMPROVEDD VTE risk score was derived by adding two points to the IMPROVE score for patients with D-dimer $\geq 2 \times$ ULN (**Table 4**).

Risk Discrimination and Reclassification by the IMPROVEDD Score

Metrics of model discrimination and reclassification are summarized in **Tables 4** and **5**. The AUC of the D-dimer, IMPROVE score, and IMPROVEDD score were 0.588, 0.560, and 0.621 at 42 days and 0.584, 0.568, and 0.625 at 77 days, respectively (**Fig. 2**). Addition of D-dimer to the IMPROVE score significantly improved the risk discrimination and reclassification at 42 and 77 days.

The observed and predicted risk for each IMPROVEDD score category is provided in **Table 6**. The rates were generally comparable in the two categories that comprise the majority of patients: IMPROVEDD scores of 1 (28.4%) and 3 (48.7%). Calibration of the IMPROVEDD score was suboptimal in the other categories: overestimation was noted in the score of 0 and ≥ 5 , whereas underestimation was noted in the scores of 2 and 4. Decision curves of the IMPROVE and IMPROVEDD scores in VTE prediction at 42 and 77 days were shown in **Figs. S1** and **S2** (supplementary figures available in the online version only). With the VTE threshold of 1% that corresponds to the cutoff warranting pharmacologic prophylaxis, IMPROVE score is associated with minimal net benefit (0.002 at both 42 and 77 days) against the “treat all” strategy, whereas the use of IMPROVEDD score would avoid undue thromboprophylaxis in 15 and 11 per 100 patients, respectively.

Identifying At-Risk Patients by the IMPROVEDD Score

The American College of Chest Physicians² selected a cutoff corresponding to an event rate of $\geq 1\%$ to dichotomize individuals as at-risk and in need of prophylaxis versus low-risk. Consequently, patients with an IMPROVEDD score of ≥ 2 were deemed as at-risk, whereas those with a score of 0 to 1 were deemed to be at low-risk (**Table 7**). Compared with low-risk patients, at-risk patients had a higher rate of symptomatic VTE at 42 and 77 days. The Kaplan–Meier rates of symptomatic VTE were 1.11 versus 0.39% at 42 days and 2.22 versus 0.91% at 77 days (HR: 2.73 [95% CI: 1.52–4.90], $p = 0.0007$; **Fig. 3**).

Discussion

Incorporation of the quantitative D-dimer level, which increases as overall hypercoagulability increases, provides incremental prognostic value to the IMPROVE VTE RAM at 42 and 77 days. Incorporating D-dimer into the IMPROVE VTE risk score significantly improves VTE risk discrimination and reclassification. An IMPROVEDD score of ≥ 2 identified a subset of hospitalized, medically ill patients at a sustained, heightened symptomatic VTE risk through 77 days.

VTE risk assessment for hospitalized patients has been associated with reduced morbidity, mortality, and incidence of hospital-acquired thrombosis.^{16–18} To identify the at-risk population and to guide appropriate thromboprophylaxis among hospitalized patients, several RAMs have been developed using clinical parameters.^{12,19–21} Research efforts have focused on exploring biomarkers associated with VTE in various populations. D-dimer has been considered as one

Table 1 Patient characteristics in the study population with and without symptomatic VTE

Characteristic	With symptomatic VTE (N = 104)	Without symptomatic VTE (N = 7,337)
Age, mean (SD)—y	77.0 (9.0)	76.4 (8.4)
Male sex, n (%)	43 (41.3)	3,349 (45.6)
Race, n (%)		
White	94 (90.4)	6,868 (93.6)
Black/African American	3 (2.9)	137 (1.9)
Asian	0 (0.0)	16 (0.2)
Others	7 (6.7)	316 (4.3)
Weight, mean (SD)—kg	81.8 (19.8)	80.3 (19.3)
Height, mean (SD)—cm	166.4 (8.4)	165.3 (9.1)
Body mass index, mean (SD)—kg/m ²	29.5 (6.9)	29.4 (6.6)
Creatinine clearance, n (%)		
< 30 mL/min	7 (6.7)	316 (4.3)
≥ 30 to < 60 mL/min	46 (44.2)	3,055 (41.7)
≥ 60 to < 90 mL/min	32 (30.8)	2,595 (35.5)
≥ 90 mL/min	19 (18.3)	1,352 (18.5)
Duration of hospitalization, median (Q1, Q3) ^a	12.0 (7.0, 17.0)	10.0 (8.0, 14.0)
Acute medical condition, n (%)		
Heart failure	34 (32.7)	3,304 (45.0)
Respiratory failure	20 (19.2)	885 (12.1)
Infection	33 (31.7)	2,103 (28.7)
Rheumatic disorder	3 (2.9)	219 (3.0)
Ischemic stroke	14 (13.5)	824 (11.2)
IMPROVE VTE risk factor, n (%)		
Previous VTE ^a	17 (16.3)	581 (7.9)
Known thrombophilia ^b	0 (0.0)	8 (0.1)
Current lower-limb paralysis	7 (6.7)	559 (7.6)
Current cancer	3 (2.9)	284 (3.9)
Immobilized ≥7 d	0 (0.0)	0 (0.0)
ICU or CCU stay ^a	21 (20.2)	682 (9.3)
Age >60 y	99 (95.2)	7,037 (95.9)
IMPROVE score, n (%) ^a		
0	1 (1.0)	119 (1.6)
1	63 (60.6)	5,322 (72.5)
2	14 (13.5)	561 (7.6)
3	8 (7.7)	777 (10.6)
4	15 (14.4)	463 (6.3)
≥ 5	3 (2.9)	95 (1.3)
IMPROVE score, median (Q1, Q3) ^a	1.0 (1.0, 2.5)	1.0 (1.0, 2.0)
D-dimer ≥ 2 × ULN, n (%) ^a	75 (77.3)	4,315 (60.5)

Abbreviations: CCU, coronary care unit; ICU, intensive care unit; ULN, upper limit of normal; VTE, venous thromboembolism.

^a*p* < 0.05.

^bDefined as inherited or acquired disorder of hemostasis including antithrombin III deficiency, protein C deficiency, and protein S deficiency.

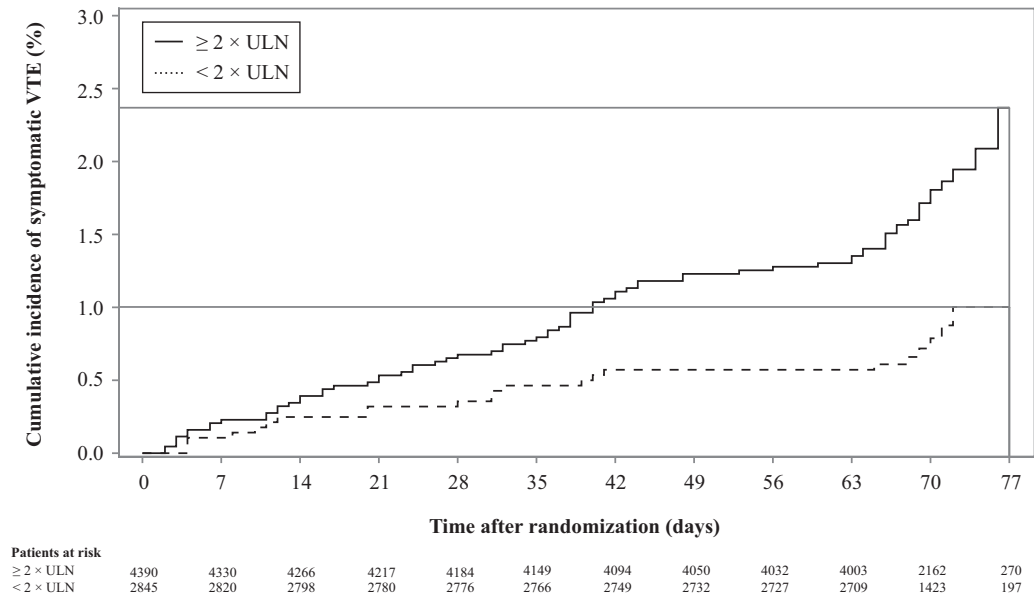


Fig. 1 Kaplan–Meier curves for symptomatic VTE stratified by D-dimer concentration.

Table 2 Univariable and multivariable regression analysis of VTE risk factors^a

Variable	Comparison	Univariable model		Multivariable model	
		HR (95% CI)	p-Value	Adjusted HR (95% CI)	p-Value
At 42 d					
D-dimer	≥ 2 × ULN vs. < 2 × ULN	2.33 (1.38–3.94)	0.0015	2.28 (1.35–3.85)	0.0020
IMPROVE VTE risk factor					
Previous VTE	Yes vs. no	2.31 (1.32–4.02)	0.0032	2.20 (1.16–4.17)	0.0155
Known thrombophilia	Yes vs. no	–	–	–	–
Current lower-limb paralysis	Yes vs. no	0.58 (0.21–1.59)	0.29	0.66 (0.24–1.79)	0.41
Current cancer	Yes vs. no	0.58 (0.14–2.35)	0.44	0.63 (0.16–2.57)	0.52
Immobilization	≥ 7 d vs. < 7 d	–	–	–	–
ICU or CCU stay	Yes vs. no	2.60 (1.55–4.37)	0.0003	2.95 (1.75–4.99)	< 0.0001
Age	> 60 y vs. ≤ 60 y	0.72 (0.29–1.78)	0.48	0.99 (0.37–2.62)	0.98
At 77 d					
D-dimer	≥ 2 × ULN vs. < 2 × ULN	2.26 (1.41–3.64)	0.0008	2.22 (1.38–5.38)	0.0010
IMPROVE VTE risk factor					
Previous VTE	Yes vs. no	2.21 (1.32–3.72)	0.0028	2.20 (1.22–3.97)	0.0084
Known thrombophilia	Yes vs. no	–	–	–	–
Current lower-limb paralysis	Yes vs. no	0.91 (0.42–1.96)	0.81	1.01 (0.47–2.18)	0.98
Current cancer	Yes vs. no	0.75 (0.24–2.36)	0.62	0.82 (0.26–2.59)	0.73
Immobilization	≥ 7 vs. < 7 d	–	–	–	–
ICU or CCU stay	Yes vs. no	2.69 (1.67–4.34)	<0.0001	2.98 (1.84–4.84)	< 0.0001
Age	> 60 vs. ≤ 60 y	0.86 (0.35–2.11)	0.74	1.20 (0.46–3.14)	0.71

Abbreviations: CCU, coronary care unit; HR, hazard ratio; ICU, intensive care unit; ULN, upper limit of normal; VTE, venous thromboembolism. ^aA total of 206 patients with incomplete covariate information were dropped from the model.

Table 3 Cox proportional hazards model for symptomatic VTE

Variable	Comparison	HR (95% CI)	p-Value
At 42 d			
D-dimer	$\geq 2 \times \text{ULN}$ vs. $< 2 \times \text{ULN}$	2.33 (1.38–3.94)	0.0015
IMPROVE score	Per point increase	1.22 (1.03–1.45)	0.0223
At 77 d			
D-dimer	$\geq 2 \times \text{ULN}$ vs. $< 2 \times \text{ULN}$	2.26 (1.41–3.64)	0.0008
IMPROVE score	Per point increase	1.26 (1.09–1.47)	0.0026

Abbreviations: HR, hazard ratio; ULN, upper limit of normal.

Table 4 Improvement in model performance by the IMPROVEDD VTE risk score

Metric	Value	p-Value
At 42 d		
ΔAUC	0.061 (0.026–0.097)	0.0008
IDI		
Absolute	0.0012 (0.0005–0.0019)	0.0004
Relative	1.71	
NRI, continuous		
Overall	0.346 (0.162–0.530)	0.0018
Events correctly reclassified	54%	< 0.0001
Nonevents correctly reclassified	–19%	< 0.0001
NRI, categorical		
Overall	0.215 (0.111–0.319)	< 0.0001
Events correctly reclassified	–6%	0.25
Nonevents correctly reclassified	28%	< 0.0001
At 77 d		
ΔAUC	0.057 (0.024–0.090)	0.0006
IDI		
Absolute	0.0015 (0.0007–0.0023)	0.0002
Relative	1.25	
NRI, continuous		
Overall	0.337 (0.169–0.506)	0.0010
Events correctly reclassified	55%	< 0.0001
Nonevents correctly reclassified	–21%	< 0.0001
NRI, categorical		
Overall	0.125 (0.026–0.225)	0.0159
Events correctly reclassified	–11%	0.0278
Nonevents correctly reclassified	24%	< 0.0001

Abbreviations: ΔAUC , improvement in the area under ROC curve; IDI, integrated discrimination improvement; NRI, net reclassification improvement.

of the most promising, well-validated, readily available markers for VTE prediction, particularly among cancer patients.²² D-dimer has also been used in combination with gender and the location of VTE to predict recurrence after discontinuing anticoagulation.²³ This is the first study to demonstrate the complementary nature of combining a

biomarker such as D-dimer with clinical variables for the risk stratification of hospitalized, medically ill patients. Results from this study demonstrate that there is a spectrum of thromboembolic risk across the IMPROVEDD score categories, with a higher score indicating a greater VTE risk that persists beyond the course of hospitalization. At-risk

Table 5 Reclassification by the IMPROVEDD VTE risk score^a

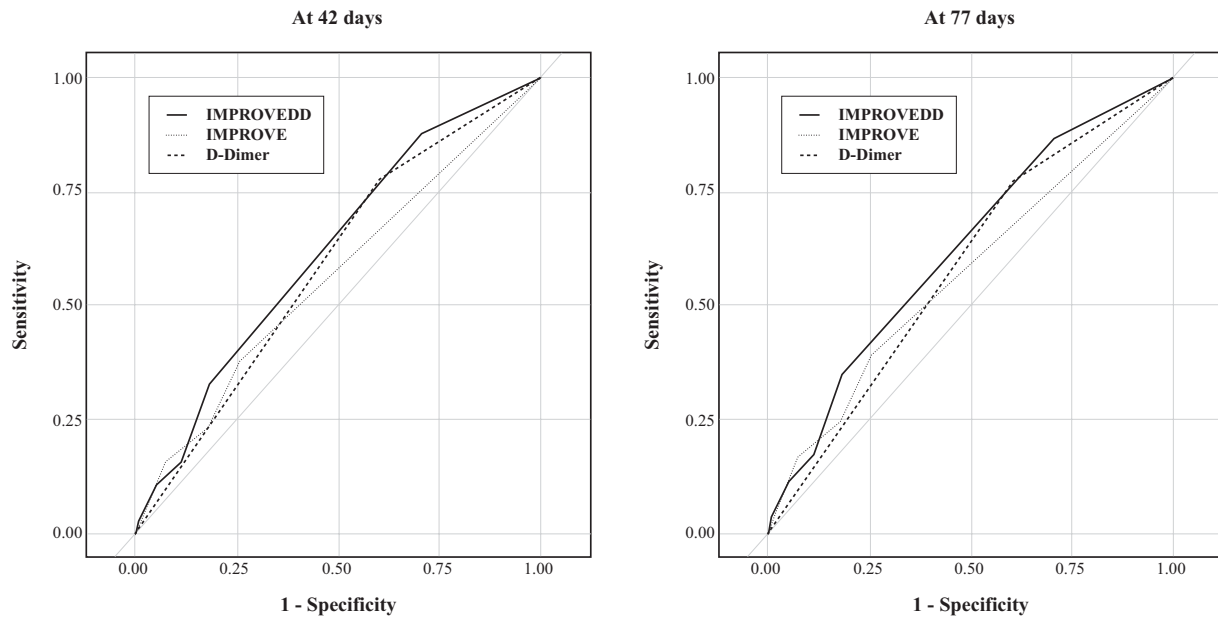
Estimated risk by the IMPROVE score	Estimated risk by the IMPROVEDD score			Total
	< 1%	≥ 1% to < 2%	≥ 2%	
At 42 d				
Overall				
< 1%	117 (100.0)	0 (0.0)	0 (0.0)	117
≥ 1% to < 2%	2,263 (32.2)	4,461 (63.5)	301 (4.3)	7,025
≥ 2%	0 (0.0)	15 (16.1)	78 (83.9)	93
Total	2,380	4,476	379	7,235
Events				
< 1%	1 (0.0)	0 (0.0)	0 (0.0)	1
≥ 1% to < 2%	12 (15.2)	60 (75.9)	7 (8.9)	79
≥ 2%	0 (0.0)	0 (0.0)	2 (100.0)	2
Total	13	60	9	82
Nonevents				
< 1%	116 (100.0)	0 (0.0)	0 (0.0)	116
≥ 1% to < 2%	2,251 (32.4)	4,401 (63.4)	294 (4.2)	6,946
≥ 2%	0 (0.0)	15 (16.5)	76 (83.5)	91
Total	2,367	4,416	370	7,153
At 77 d				
Overall				
< 1%	48 (41.0)	69 (59.0)	0 (0.0)	117
≥ 1% to < 2%	2,053 (31.3)	4,077 (62.1)	431 (6.6)	6,561
≥ 2%	0 (0.0)	163 (29.3)	394 (70.7)	557
Total	2,101	4,309	825	7,235
Events				
< 1%	0 (0.0)	1 (100.0)	0 (0.0)	1
≥ 1% to < 2%	13 (16.3)	61 (76.3)	6 (7.5)	80
≥ 2%	0 (0.0)	5 (31.3)	11 (68.8)	16
Total	13	67	17	97
Nonevents				
< 1%	48 (41.4)	68 (58.6)	0 (0.0)	116
≥ 1% to < 2%	2,040 (31.5)	4,016 (62.0)	425 (6.6)	6,481
≥ 2%	0 (0.0)	158 (29.2)	383 (70.8)	541
Total	2,088	4,242	808	7,138

^aValues expressed as number of patients (row percentage).

patients, as identified by the IMPROVEDD score, may be appropriate candidates for extended thromboprophylaxis to minimize the risk of symptomatic VTE.

With the intent of maintaining generalizability from the IMPROVE derivation cohort, the weights for each VTE risk factor were unchanged, and two points were added to the original score for patients with D-dimer $\geq 2 \times$ ULN. It is notable that adding D-dimer measurement to the IMPROVE score has been implemented in the enrollment criteria of an ongoing trial for preventing hospital-associated VTE (Medically Ill Patient Assessment of Rivaroxaban Versus Placebo in

Reducing Post-Discharge Venous Thrombo-Embolism Risk [MARINER]).²⁴ In the MARINER study, eligible patients (i.e., “high-risk” patients) must have a total modified IMPROVE VTE risk score of ≥ 4 , or a risk score of 2 or 3 with a plasma D-dimer level of more than twice the ULN. The relative weight for D-dimer of $\geq 2 \times$ ULN is equivalent to two points when taken together with the IMPROVE VTE risk score and matches the rationale in this analysis. Furthermore, it supports the use of D-dimer measurement in conjunction with a standardized RAM for optimizing VTE risk assessment among acutely ill, hospitalized medical patients. However, although



Likelihood Ratio Test Statistics

Model	Comparison	Chi-Square	P-value
At 42 days			
D-dimer	$\geq 2 \times \text{ULN}$ vs. $< 2 \times \text{ULN}$	11.36	0.0008
IMPROVE score	Per point increase	4.36	0.0366
IMPROVEDD score	Per point increase	13.66	0.0002
At 77 days			
D-dimer	$\geq 2 \times \text{ULN}$ vs. $< 2 \times \text{ULN}$	12.30	0.0005
IMPROVE score	Per point increase	7.43	0.0064
IMPROVEDD score	Per point increase	18.33	< 0.0001

Fig. 2 Receiver–operating–characteristic (ROC) curves for D-dimer, IMPROVE, and IMPROVEDD models in predicting symptomatic VTE.

Table 6 Observed and predicted risk by the IMPROVEDD VTE risk score

IMPROVEDD score	Patients, n (%)	At 42 d			At 77 d		
		Event, n	Observed risk, %	Predicted risk, %	Event, n	Observed risk, %	Predicted risk, %
0	48 (0.7)	0	0.0	0.4	0	0.0	0.5
1	2,053 (28.4)	10	0.5	0.6	13	0.6	0.7
2	279 (3.9)	3	1.1	0.8	3	1.1	1.0
3	3,520 (48.7)	42	1.2	1.2	47	1.3	1.4
4	510 (7.0)	14	2.7	1.6	17	3.3	1.9
≥ 5	825 (11.4)	13	1.6	2.2	17	2.1	2.7

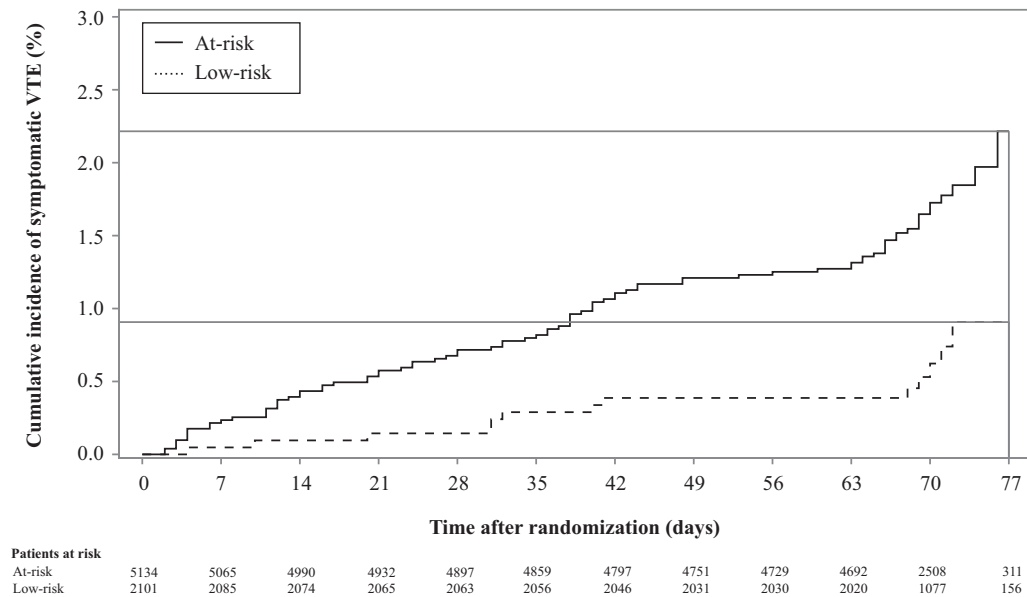
the incorporation of biomarker may enhance the performance of clinical RAM, it remains uncertain whether this refinement is clinically meaningful to warrant the additional complexity and expense. Future studies are therefore required to evaluate the practicability of IMPROVEDD score in the “real-world” setting.

Limitations

VTE risk factors encompassed in the analyses were not all inclusive. First, other previously described risk factors (e.g., myocardial infarction, chronic obstructive pulmonary disease, chronic kidney disease, and dyslipidemia) and

Table 7 Risk stratification by the IMPROVEDD VTE risk category

Risk category	Event rate (95% CI)	Odds ratio (95% CI)	p-Value
At 42 d			
At-risk (≥ 2 points)	1.40% (1.08–1.72%)	2.97 (1.53–5.77)	0.0002
Low-risk (0–1 points)	0.48% (0.18–0.77%)	Reference	
At 77 d			
At-risk (≥ 2 points)	1.64% (1.29–1.98%)	2.67 (1.49–4.80)	0.0002
Low-risk (0–1 points)	0.62% (0.28–0.95%)	Reference	

**Fig. 3** Kaplan–Meier curves for symptomatic VTE stratified by the IMPROVEDD risk category.

biomarkers (e.g., prothrombin fragment 1 + 2, soluble P-selectin, clotting factor VIII, and thrombin generation potential) may offer additional refinement in risk stratification among acutely ill, hospitalized medical patients. Second, small numbers of events in certain score categories may preclude an accurate calibration of the IMPROVEDD model. Also the lack of patients who were immobilized ≥ 7 days as a risk factor in the IMPROVE model may have altered the model characteristics. Nonetheless, the cutoff of an IMPROVEDD score of ≥ 2 demonstrated excellent discriminatory capacity in this population. Third, asymptomatic DVT was not included as a component of the outcome in the analysis. Fourth, it should be noted that D-dimer measurement could be influenced by the analytical methods and reporting standards from different laboratories.²⁵ Finally, the data were derived from a population that agreed to participate in a clinical trial. The results are applicable to patients who received primary prophylaxis with either standard-duration enoxaparin or extended-duration betrixaban and may not be generalizable to other settings.

Conclusion

Baseline D-dimer level demonstrated a robust, incremental prognostic value to the IMPROVE RAM in VTE risk stratification for medical patients. Strategies to improve VTE risk stratification should consider incorporation of D-dimer measurement into standard RAMs. An IMPROVEDD VTE risk score of two or greater identifies a subset of hospitalized, medically ill patients receiving thromboprophylaxis at a sustained, heightened risk for symptomatic VTE through 77 days. This population may potentially benefit from an extended course of thromboprophylaxis. Independent validation and impact analysis should be undertaken before employing the IMPROVEDD VTE risk score in clinical practice.

Clinical Trial Registration

URL: <http://www.clinicaltrials.gov>. Unique identifier: NCT01583218.

Conflict of Interest

None declared.

Funding

The study was funded by Portola Pharmaceuticals, Inc.

References

- Jha AK, Larizgoitia I, Audera-Lopez C, Prasopa-Plaizier N, Waters H, Bates DW. The global burden of unsafe medical care: analytic modeling of observational studies. *BMJ Qual Saf* 2013;22(10):809–815
- Kahn SR, Lim W, Dunn AS, et al. Prevention of VTE in nonsurgical patients: Antithrombotic Therapy and Prevention of Thrombosis, 9th ed: American College of Chest Physicians Evidence-Based Clinical Practice Guidelines. *Chest* 2012;141(2, Suppl):e195S–e226S
- Qaseem A, Chou R, Humphrey LL, Starkey M, Shekelle P; Clinical Guidelines Committee of the American College of Physicians. Venous thromboembolism prophylaxis in hospitalized patients: a clinical practice guideline from the American College of Physicians. *Ann Intern Med* 2011;155(09):625–632
- ISTH Steering Committee for World Thrombosis Day. Venous thromboembolism: A Call for risk assessment in all hospitalised patients. *Thromb Haemost* 2016;116(05):777–779
- Flanders SA, Greene MT, Grant P, et al. Hospital performance for pharmacologic venous thromboembolism prophylaxis and rate of venous thromboembolism: a cohort study. *JAMA Intern Med* 2014;174(10):1577–1584
- Greene MT, Spyropoulos AC, Chopra V, et al. Validation of risk assessment models of venous thromboembolism in hospitalized medical patients. *Am J Med* 2016;129(09):1001.e9–1001.e18
- Desjardins L, Bara L, Boutitie F, et al. Correlation of plasma coagulation parameters with thromboprophylaxis, patient characteristics, and outcome in the MEDENOX study. *Arch Pathol Lab Med* 2004;128(05):519–526
- Fan J, Li X, Cheng Y, Yao C, Zhong N; Investigators Group. Measurement of D-dimer as aid in risk evaluation of VTE in elderly patients hospitalized for acute illness: a prospective, multicenter study in China. *Clin Invest Med* 2011;34(02):E96–E104
- Cohen AT, Harrington R, Goldhaber SZ, et al. The design and rationale for the Acute Medically Ill Venous Thromboembolism Prevention with Extended Duration Betrixaban (APEX) study. *Am Heart J* 2014;167(03):335–341
- Mahan CE, Liu Y, Turpie AG, et al. External validation of a risk assessment model for venous thromboembolism in the hospitalised acutely-ill medical patient (VTE-VALOURR). *Thromb Haemost* 2014;112(04):692–699
- Rosenberg D, Eichorn A, Alarcon M, McCullagh L, McGinn T, Spyropoulos AC. External validation of the risk assessment model of the International Medical Prevention Registry on Venous Thromboembolism (IMPROVE) for medical patients in a tertiary health system. *J Am Heart Assoc* 2014;3(06):e001152
- Spyropoulos AC, Anderson FA Jr, FitzGerald G, et al; IMPROVE Investigators. Predictive and associative models to identify hospitalized medical patients at risk for VTE. *Chest* 2011;140(03):706–714
- Cohen AT, Harrington RA, Goldhaber SZ, et al; APEX Investigators. Extended thromboprophylaxis with betrixaban in acutely ill medical patients. *N Engl J Med* 2016;375(06):534–544
- Pencina MJ, D'Agostino RB Sr, D'Agostino RB Jr, Vasan RS. Evaluating the added predictive ability of a new marker: from area under the ROC curve to reclassification and beyond. *Stat Med* 2008;27(02):157–172, discussion 207–212
- Vickers AJ, Elkin EB. Decision curve analysis: a novel method for evaluating prediction models. *Med Decis Making* 2006;26(06):565–574
- Lester W, Freemantle N, Begaj I, Ray D, Wood J, Pagano D. Fatal venous thromboembolism associated with hospital admission: a cohort study to assess the impact of a national risk assessment target. *Heart* 2013;99(23):1734–1739
- Roberts LN, Porter G, Barker RD, et al. Comprehensive VTE prevention program incorporating mandatory risk assessment reduces the incidence of hospital-associated thrombosis. *Chest* 2013;144(04):1276–1281
- Catterick D, Hunt BJ. Impact of the national venous thromboembolism risk assessment tool in secondary care in England: retrospective population-based database study. *Blood Coagul Fibrinolysis* 2014;25(06):571–576
- Barbar S, Noventa F, Rossetto V, et al. A risk assessment model for the identification of hospitalized medical patients at risk for venous thromboembolism: the Padua Prediction Score. *J Thromb Haemost* 2010;8(11):2450–2457
- Kucher N, Koo S, Quiroz R, et al. Electronic alerts to prevent venous thromboembolism among hospitalized patients. *N Engl J Med* 2005;352(10):969–977
- Woller SC, Stevens SM, Jones JP, et al. Derivation and validation of a simple model to identify venous thromboembolism risk in medical patients. *Am J Med* 2011;124(10):947–954.e2
- Pabinger I, Thaler J, Ay C. Biomarkers for prediction of venous thromboembolism in cancer. *Blood* 2013;122(12):2011–2018
- Eichinger S, Heinze G, Jandek LM, Kyrle PA. Risk assessment of recurrence in patients with unprovoked deep vein thrombosis or pulmonary embolism: the Vienna prediction model. *Circulation* 2010;121(14):1630–1636
- Raskob GE, Spyropoulos AC, Zrubek J, et al. The MARINER trial of rivaroxaban after hospital discharge for medical patients at high risk of VTE. Design, rationale, and clinical implications. *Thromb Haemost* 2016;115(06):1240–1248
- Lippi G, Tripodi A, Simundic AM, Favaloro EJ. International survey on D-dimer test reporting: a call for standardization. *Semin Thromb Hemost* 2015;41(03):287–293