

Do Risk Factors Change in Medical Patients While Receiving VTE Prophylaxis?

A retrospective, cross-sectional study assessing VTE and Bleeding Risk



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Background

- An estimated 75-80% of non-surgical medical patients receive VTE prophylaxis during hospital admission^{1,2}
- Current guidelines recommend pharmacological prophylaxis for patients at increased risk of thrombosis and low risk of bleed, and recommend risk stratification through the use of externally validated risk-assessment models (RAMs) such as the IMPROVE models³
- Risk factors may change during the course of admission and re-assessment of risk categories represents an opportunity to discontinue unnecessary or unsafe drug therapy

Objectives

- To describe whether risk categories for thrombosis and bleed change during hospital admission to warrant discontinuation of prophylaxis using the IMPROVE VTE and Bleed RAMs, respectively
- Primary outcome:** the no. of patients who moved from high to low risk of VTE, or who moved from low to high risk of bleed during hospital admission (→)
- Secondary outcomes:** the no. of patients who:
 - Remained at low risk of VTE throughout admission
 - Remained at high risk of bleeding throughout admission
 - Remained or changed to low risk of VTE and continued prophylaxis until discharge
 - Remained or changed to high risk of bleed and continued prophylaxis until discharge

Methods

- Design:** Cross-sectional, retrospective chart review
- Population:** Non-surgical medical patients who received prophylactic doses of dalteparin or UFH while admitted in the Fraser Health Authority (FH) between April 1-30, 2017
- Sample:** Systematic random sampling of 200 patients with proportionate representation from all acute FH sites
- Sample size calculation:** Convenient sample size of 200 patients calculated to yield a 95% CI of ± 6.7%, assuming a conservative estimate of 50% for the no. of patients whose risk factors changed during admission

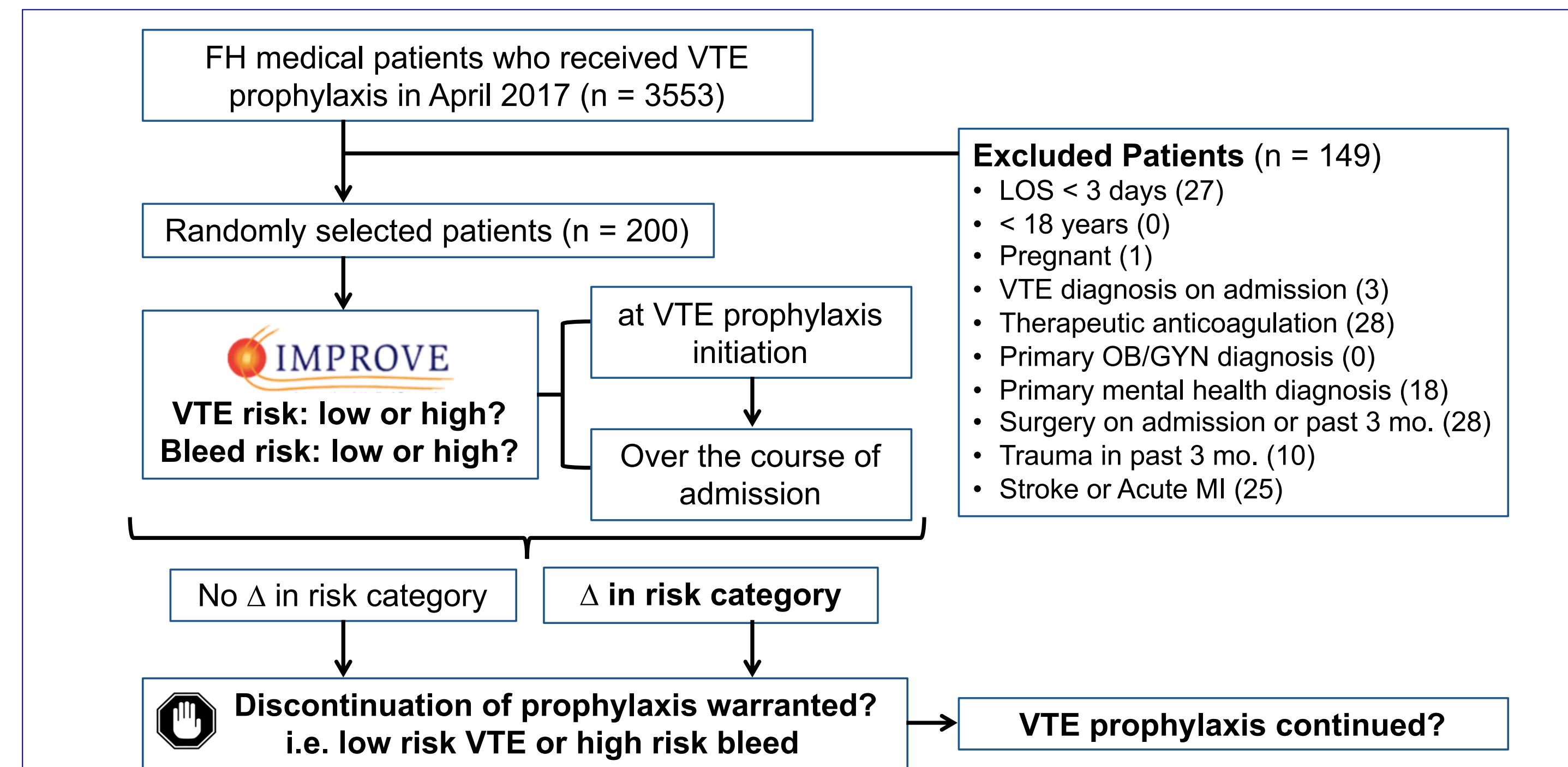


Figure 1: Study Flow Diagram

VTE Risk Factor	Score	Bleed Risk Factor	Score
Previous VTE	3	Gastroduodenal ulcer	4.5
Thrombophilia	2	Bleeding in prior 3 months	4
Age > 60 years	1	Platelets < 50 x 10 ⁹ /mL	4
Active cancer	2	Hepatic failure: INR > 1.5	2.5
Immobility	1	GFR 30-59 mL/min	1
ICU/CCU stay	1	GFR < 30 mL/min	2.5
Lower limb paralysis	2	Central venous catheter	2
		Rheumatic disease	2
		Active cancer	2
		Age > 40 years	1.5
		Male sex	1
Total Score	Associated Risk	Category	
< 2	< 1%	low risk	
≥ 2	≥ 1%	high risk	

Total Score	Associated Risk	Category
< 7	< 2%	low risk
≥ 7	> 3%	high risk

Table 1: IMPROVE Risk Assessment Models

Male, no. (%)	97 (49)
Age, mean years ± SD	70 ± 16
VTE prophylaxis agent used, no. (%)	
▫ dalteparin	150 (75)
▫ heparin	40 (20)
▫ both dalteparin and heparin used	10 (5.0)
VTE prophylaxis duration, mean days ± SD	10.7 ± 12.4
Length of hospital stay, mean days ± SD	12.5 ± 12.4
IMPROVE VTE score at prophylaxis initiation, mean ± S.D.	1.5 ± 1.1
IMPROVE Bleed score at prophylaxis initiation, mean ± S.D.	4.9 ± 2.2

Table 2: Patient Characteristics (n = 200)

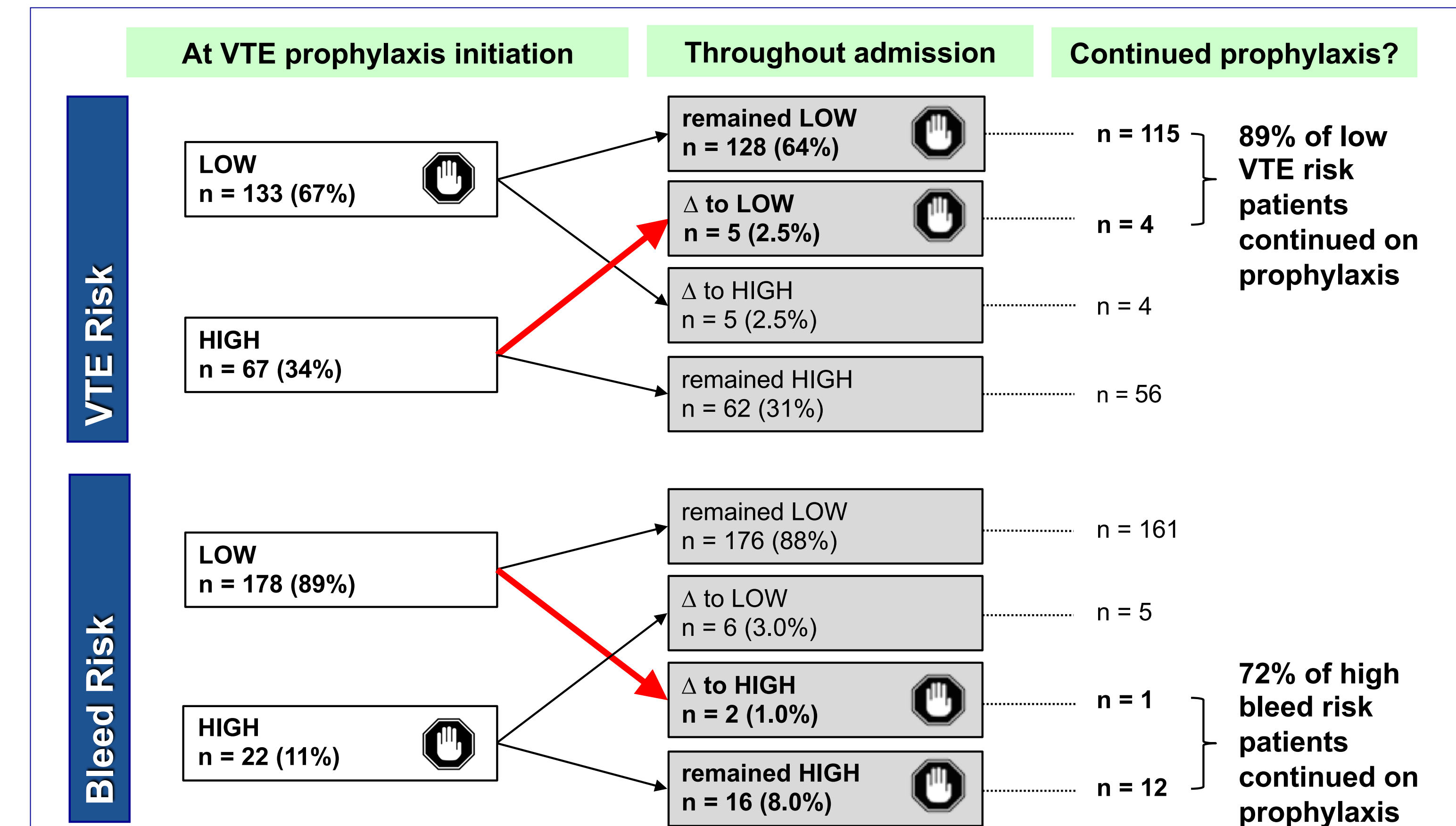


Figure 2: VTE and Bleeding Risk Assessments

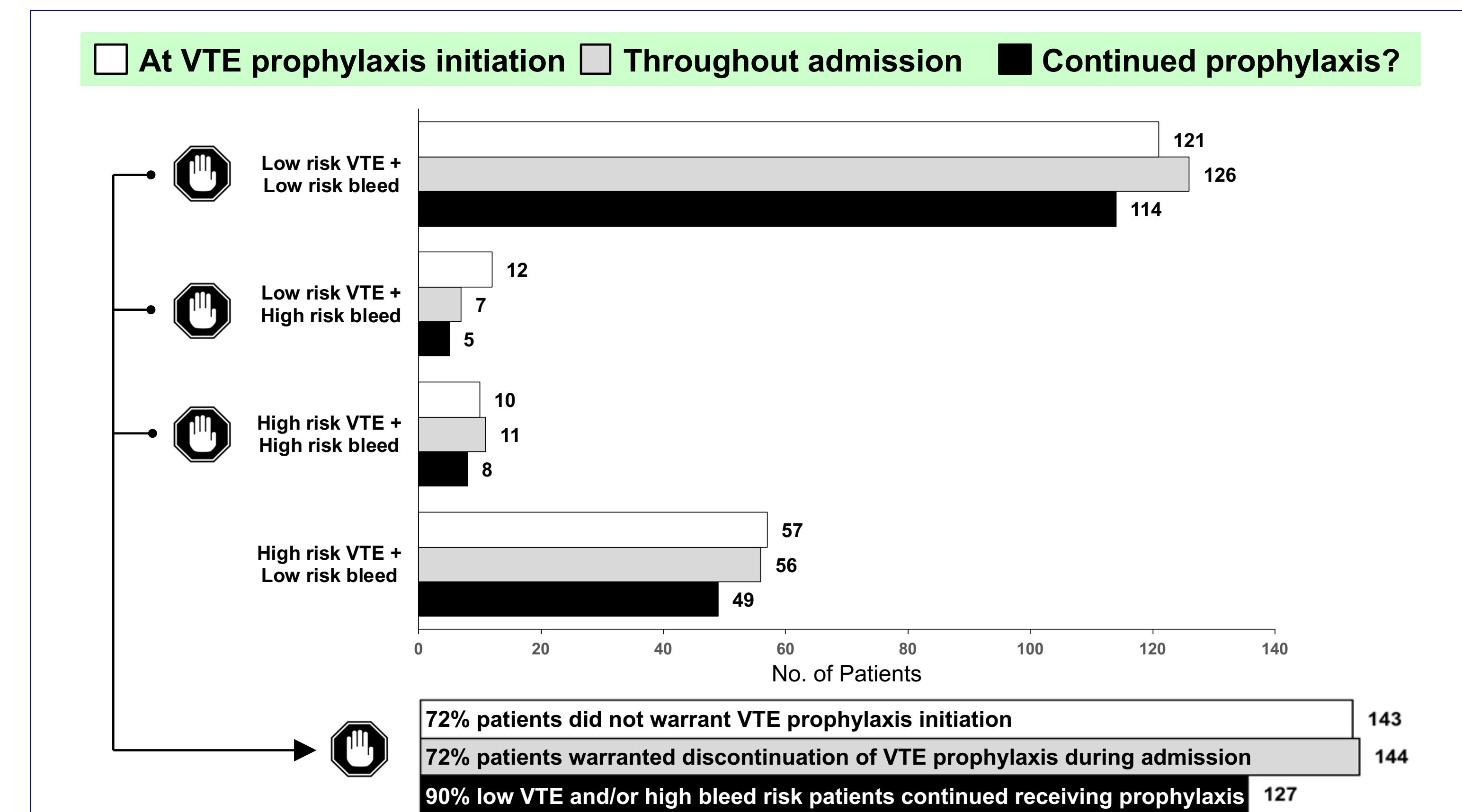


Figure 3: Combined VTE and Bleed Risk Categories Warranting Discontinuation of VTE prophylaxis

Conclusions

- For the majority of patients, VTE and bleed risk categories did not change throughout admission
- The majority of VTE prophylaxis given to medical patients was unnecessary or unsafe and was continued throughout admission
- Future efforts to minimize inappropriate VTE prophylaxis should focus on risk assessment prior to VTE prophylaxis initiation