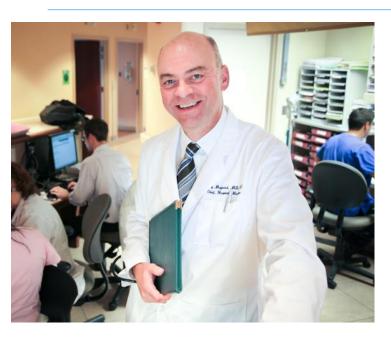


Preventing Hospital-Associated Venous Thromboembolism: Practical Strategies That Work

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CDC Webinar

Conflict of Interest



Executive Committee -Mariner VTE Prevention trial of extended duration prophylaxis in medical patients

AHRQ DVT Prevention Guide author

Abbreviations – Terms

- VTE venous thromboembolism
- VTE-P VTE Prevention / prophylaxis
- HA VTE hospital-associated VTE
- CDS Clinical decision support
- IPCD intermittent pneumatic compression devices
- SCD sequential compression devices
- GCS graduated compression stockings
- Extended duration prophylaxis beyond hospital stay
- LMWH low-molecular weight heparin
- UFH unfractionated heparin
- LDUH low dose unfractionated heparin
- PAH Pulmonary artery hypertension
- AT8 ACCP Anticoagulation / DVT Prevention guidelines (2008)
- AT9 ACCP Anticoagulation / DVT Prevention guidelines (2102)

A Major Source of Mortality and Morbidity

- 350,000 to 650,000 with VTE per year
- 100,000 to > 200,000 deaths per year
- About half are hospital related.
- VTE is primary cause of fatality in half-
 - More than HIV, MVAs, Breast CA <u>combined</u>
 - Equals 1 jumbo jet crash / day
- 10% of hospital deaths
 - PE among top sources of preventable hospital related death
- Huge costs and morbidity (recurrence, postthrombotic syndrome, chronic PAH, anticoag)

QI Framework and Strategies that Work

- UC San Diego and Univ. of California VTEP Collaborative
- SHM / AHRQ improvement guides and Collaborative
- Experience, mentoring other hospitals via UCSD CIIS
- Johns Hopkins experience
- Systematic reviews

Kahn SR, Morrison DR, Cohen JM, Emed J, Tagalakis V, Roussin A, Geerts W. Interventions for implementation of thromboprophylaxis in hospitalized medical and surgical patients at risk for venous thromboembolism (Review). *Cochrane Database of Systematic Reviews* 2013, Issue 7. Art. No.: CD008201. DOI: 10.1002/14651858.CD008201.pub2.

Streiff MB, Carolan HT, Hobson DB, Kraus PS, Holzmueller CG, Demski R, et al. Lessons from the Johns Hopkins Multi-Disciplinary Venous Thromboembolism (VTE) Prevention Collaborative. *BMJ* 2012; Jun 19;344:e3935.

Kakkar AK, Davidson BL, Haas SK. Compliance with recommended prophylaxis for venous thromboembolism: improving the use and rate of uptake of clinical practice guidelines. *J Thromb Haemost*. 2004;2:221–227.

Tooher R, Middleton P, Pham C, et al. A systematic review of strategies to improve prophylaxis for venous thromboembolism in hospitals. *Ann Surg* 2005; 241:397–415.

Maynard G, Stein J. Designing and Implementing Effective VTE Prevention Protocols: Lessons from Collaboratives. *J Thromb Thrombolysis* 2010 Feb:29(2):159-166.

Strategies to Reduce HA VTE

- Centralized steering group for institution wide approach
- Review and distill the evidence / best practices
- Standardize Create a VTE Prevention Protocol
- Embed protocol guidance into order sets, hard stops for use on admission, transfer, and post op – Provide seamless CDS
- Go beyond core measures / SCIP better measures
- Active day-to-day surveillance, in addition to monthly / quarterly
- Multiple mutually reinforcing interventions to reinforce protocol
- Active vs passive interventions
- Address adherence / administration of prophylaxis
- Address other failure modes / contributing factors to HA VTE

Review Evidence, identify best practice

Guidelines ATP, ATS, ACP1, AAOS, ASCO, ACOG, NICE

Regulatory TJC/NQF/SCIP/C MS

Local standards

Other guidance Literature, meta-analyses Define Local Best Practice Standards and Expectations

Policies

Summarize

Translate

Protocols
VTE and bleeding
risk assessment
linked to risk
appropriate
prophylaxis
options



Design Multi-faceted Interventions

Order sets Embedded VTE prevention protocol

Education

Increase delivery of ordered prophylaxis

Checklists

Alerts

Reduced use of central venous catheters

Enhance mobility and activity

Audit and feedback

Care pathways



Ensure reliable delivery of best practice -_ Implement, monitor, revise, and refine,

Engage Explain why important

Educate Share evidence, include "Just in time" education



Evaluate Regularly assess, revise and refine as needed Spread and Sustain







Analyze Care Delivery Survey Previous / Ongoing Efforts Environment, baseline, failure modes, barriers to implementation Continue analysis of care delivery Local barriers, impact of interventions

Track Metrics

Key Metrics:

Prevalence of appropriate VTE Prophylaxis Incidence of Hospital-Associated VTE

Establish Foundation:

Institutional Support and Centralized, Empowered VTE Prevention Steering Team

The Essential First Intervention



- 1) a standardized VTE risk assessment, linked to...
- 2) a menu of appropriate prophylaxis options, plus...
- 3) a list of contraindications to pharmacologic VTE prophylaxis

Challenges:

Make it easy to use ("automatic")

Make sure it captures almost all patients

Trade-off between guidance and ease of use /
efficiency

Protocol

Characteristics of the hypothetical ideal protocol

Trade-offs and prioritization of characteristics often needed

- Accurately detects all patients at risk for DVT.
- Reliably <u>excludes</u> patients who would be unlikely to develop DVT, minimizing inappropriate overprophylaxis in those of lower risk.
- Provides actionable recommendations for permutations of VTE and bleeding risk.
- Simple to use in routine clinical practice
- Identifies patients that should have a combination of mechanical and anticoagulant prophylaxis.
- Lends itself to automation or dynamic ongoing reevaluations.
- Integration results in convincing decreases in hospitalassociated VTE without any increase in bleeding.

9

	Hierarchy of Daliability	Predicted Prophylaxis rate	
Level	Hierarchy of Reliability		
1	No protocol* ("State of Nature")	40%	
2	Decision support exists but not linked to order writing, or prompts within orders but no decision support	f 50%	
3	Protocol well-integrated (into orders at point-of-care)	65-85%	
4	Protocol enhanced	90%	
	(by other QI / high reliability strategies)		
5	Oversights identified and addressed in	95+%	
	——————————————————————————————————————	San Diego TH SCIENCES	

Protocol

- Local Standards of best practice
- Written out
- Algorithmic decision trees can be useful
- Include operational definitions*
- Must have enough detail to be measurable and make judgments re:

Is this case meeting our standard of care?

- Examples requiring operational definitions*
 - High INR
 - Low platelet counts
 - Impaired mobility
 - "Low Risk"

Prompt - Not a protocol - No CDS offered

DVT PROPHYLAXIS ORDERS

- Anti thromboembolism Stockings
- Sequential Compression Devices
- UFH 5000 units SubQ q 12 hours
- UFH 5000 units SubQ q 8 hours
- LMWH (Enoxaparin) 40 mg SubQ q day
- LMWH (Enoxaparin) 30 mg SubQ q 12 hours
- No Prophylaxis, Ambulate

Over 20 different VTE risk assessment models

- No consensus on what is best in clinical practice
- Individualized point-based scoring (quantitative) models
 - Generally more rigorously validated in determining risk, but not in clinical practice
 - Examples:
 - Caprini
 - IMPROVE

Padua

- Grouping or "bucket" models
 - Generally not as well validated in predicting risk, but easier to implement, more published / unpublished success stories in reducing HA VTE

Examples:

- NICE / NHS guidelines, Australia / New Zealand working group model
- Classic "3 bucket" model
- Updated "3 bucket" grouping model

Each Risk Factor I	Represents 1 Poin	t	Each Risk Factor Represents 2 Points	
☐ Age 41-60 years ☐ Acu	te myocardial infarction	on	☐ Age 61-74 years ☐ Central venous access	ŝ
☐ Swollen legs (current) ☐ Con	gestive heart failure (<1 month)	☐ Arthroscopic surgery ☐ Major surgery (>45 mir	nutes)
	dical patient currently		☐ Malignancy (present or previous)	
	ory of inflammatory b		a Laparoscopic surgery (*45 minutes)	total:
0 , 1	ory of prior major sur	, , ,	,	
☐ Sepsis (<1 month) ☐ Abn	ormal pulmonary fund	ction (COPD)	☐ Immobilizing plaster cast (<1 month)	
 Serious lung disease including p 	,)	Each Risk Factor Represents 3 Points	
Oral contraceptives or hormone Pregnancy or postpartum (<1 m History of unexplained stillborn is abortion (≥ 3), premature birth with Other risk factors	onth) nfant, recurrent spont toxemia or growth-re	stricted infant Subtotal:	□ Age 75 years or older □ Family history of throm □ History of DVT/PE □ Positive Prothrombin 202 □ Positive Factor V Leiden □ Positive Lupus anticoagu □ Elevated serum homocysteine □ Heparin-induced thrombocytopenia (HIT) (Do not use heparin or any low molecular weight hepa □ Elevated anticardiolipin antibodies □ Other congenital or acquired thrombophilia Sub	210A ulant
☐ Stroke (<1 month)	☐ Multiple trauma	(<1 month)	If yes: Type	
 Elective major lower extremity a 	rthroplasty		* most frequently missed risk factor	
☐ Hip, pelvis or leg fracture (<1 month) ☐ Acute spinal cord injury (paralysis) (<1 month)			TOTAL RISK FACTOR SCORE:	
Caprini Score Risk	VTE Incidence	Recommend	nded Prophylaxis	

Caprini Score	Risk	VTE Incidence	Recommended Prophylaxis
0 - 2	very low - low	< 1.5% ¹	Early ambulation, IPC
3 - 4	moderate	3% ¹	LMWH; UFH; or IPC. If high bleeding risk, IPC until bleeding risk diminishes.
5-8	high	6% ¹	LMWH + IPC; or UFH + IPC. If high bleeding risk, IPC until bleeding risk diminishes.
>8	very high	6.5 - 18.3%	LMWH + IPC; or UFH + IPC. If high bleeding risk, IPC until bleeding risk diminishes. Consider extended duration prophylaxis.

^{*} Abdominal or pelvic surgery for cancer should receive extended VTE prophylaxis with LMWH x 30 days. 1

IPC = intermittent pneumatic compression LMWH = low molecular weight heparin UFH = unfractionated heparin

 Gould MK, Garcia DA, Wren SM, et.al. Prevention of VTE in nonorthopedic surgical patients: antithrombotic therapy and prevention of thrombosis, 9th ed: Americal College of Chest Physicians Evidence-Based Clinical Practice Guidelines. Chest. 2012; 141(2)(Suppl): e227S-e277S.

Caprini Model

- Validated in predicting risk
- Can be difficult to use reliably
- Only 1
 published
 success in
 clinical practice
 published after
 30 years of use.
- Works best in centers with advanced CDS to make it easier / more automated

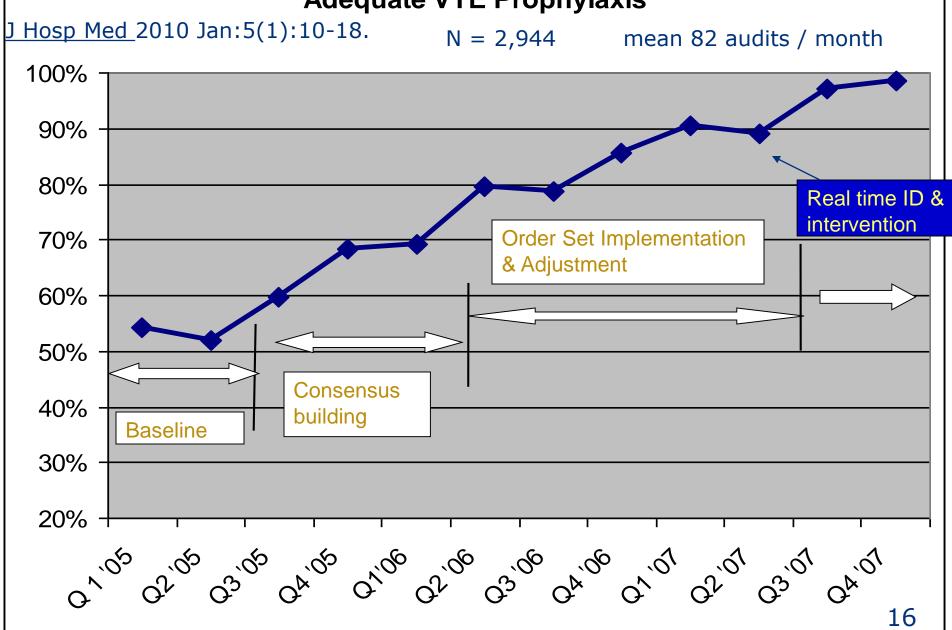
Risk Assessment

- Classic "3 bucket" model derived from AT8

Low Risk: Minor surgery in mobile patients. Medical patients who are fully mobile. Observation patients with expected hospital stay < 48 hours.	No prophylaxis, reassess periodically, ambulate.
Moderate Risk: Most general, thoracic, open gynecologic or urologic surgery patients. Medical patients, impaired mobility from baseline or acutely ill.	UFH or LMWH prophylaxis*
High Risk: Hip or knee arthroplasty, hip fracture surgery. multiple major trauma, spinal cord injury or major spinal surgery, Abdominal-pelvic surgery for cancer.	IPCD <u>AND</u> LMWH or other anticoagulant*

^{*}For those at moderate or high risk and contraindications to anticoagulation, use IPCD.

Percent of Randomly Sampled Inpatients with Adequate VTE Prophylaxis



UCSD results

Hospital Acquired VTE by Year				
	2005	2006	2007	2008
Patients at Risk	9,720	9,923	11,207	
Cases w/ any VTE Risk for HA VTE Odds Ratio (95% CI)	1.0	138 1 in 73 1.03 (0.81, 1.32)	92 1 in 122 0.61# (0.46, 0.80)	80
Cases with PE Risk for PE Odds Ratio (95% CI)	21 1 in 463 1.0	22 1 in 451 1.02 (0.54, 1.96)	15 1 in 747 0.62 (0.30, 1.26)	12
Cases with DVT (and no PE) Risk for DVT Odds Ratio (95% CI)	1.0	116 1 in 85 1.03 (0.79, 1.96)	77 1 in 146 0.61* (0.45, 0.82)	68
Cases w/ Preventable VTE Risk for Preventable VTE Odds Ratio (95% CI)	44 1 in 221 1.0	21 1 in 473 0.47# (0.26, 0.80)	7 1 in 1,601 0.14* (0.05, 0.31)	6
	# p < 0.01 *p < 0.001)10 1 F(1) - 10	10	17

J Hosp Med 2010 Jan:5(1):10-18.

Updated Model – More c/w AT9 guidelines

Updated "3 bucket" model, now in use at authors' site (UC San Diego)

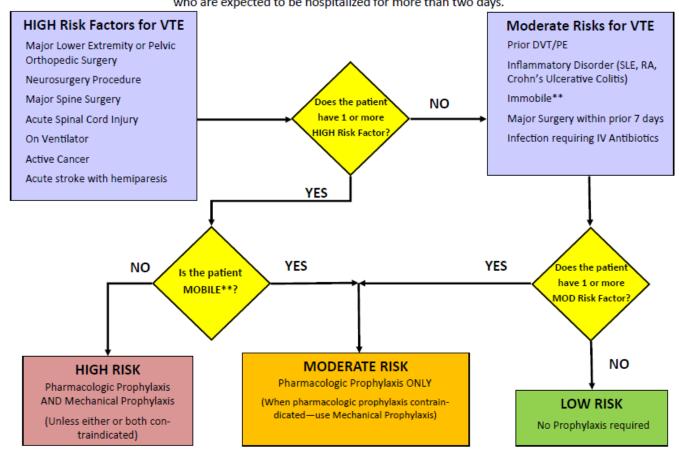
Low Risk: Observation status, expected LOS < 48 hours. Minor ambulatory surgery unless multiple strong risk factors. Medical patients ambulatory in hall and not Moderate or High Risk. Ambulatory cancer patients admitted for short chemotherapy infusion.	No prophylaxis, reassess periodically, ambulate.
Moderate Risk (Most general medical / surgical patients): Most general, thoracic, open gynecologic or urologic surgery patients. Active cancer or past VTE / known thrombophilia in medical patient with LOS > 48 hours. Medical patient with decrease in usual ambulation AND VTE risk factors (MI, Stroke, CHF, PNA, active inflammation / infection, dehydration, age > 65)	UFH or LMWH prophylaxis*
High Risk: Hip or knee arthroplasty, hip fracture surgery, multiple major trauma, spinal cord injury or major neurosurgery, abdominal-pelvic surgery for cancer	IPCD <u>AND</u> LMWH or other anticoagulant*

^{*}For those at moderate or high VTE risk and contraindications to anticoagulation, use IPCD alone until bleeding risk subsides.

UC Davis Medical Center

3 bucket model algorithm assoc. w/ reduction in HA VTE

VTE Risk Assessment Tool—for patients ADMITTED (not OBS or same day surgery) to UCDMC and who are expected to be hospitalized for more than two days.



^{**}Minimum Criteria for MOBILITY = Walks occasionally for short distances, with or without assistance. May spend most of the day in bed.

Effective Implementation / CDS Principles

- 1. Keep it simple for the end user
 - a. Some adjustments can be done behind the scenes (pharmacy adjustment of dose or periop timing, for example)
 - b. Minimize calculations / clicks, automate process for them
 - c. Streamline options, offer only preferred choices
- Don't interrupt the work flow
 - a. Integrate risk assessment in admit / transfer / post op process
 - b. Keep VTE risk assessment, bleeding risk assessment, and ordering of risk-appropriate prophylaxis together as a unified process.
- Design reliability into the process
 - **a.** Forcing functions / hard stop for VTEP
 - b. Present **preferred risk appropriate prophylaxis as the default** option once risk level chosen
 - **c. Scheduling** and **redundant** checks for highest risk patients
 - **Standardization** for services / groups of patients (discourage over-customization at provider level)

Effective Implementation / CDS Principles

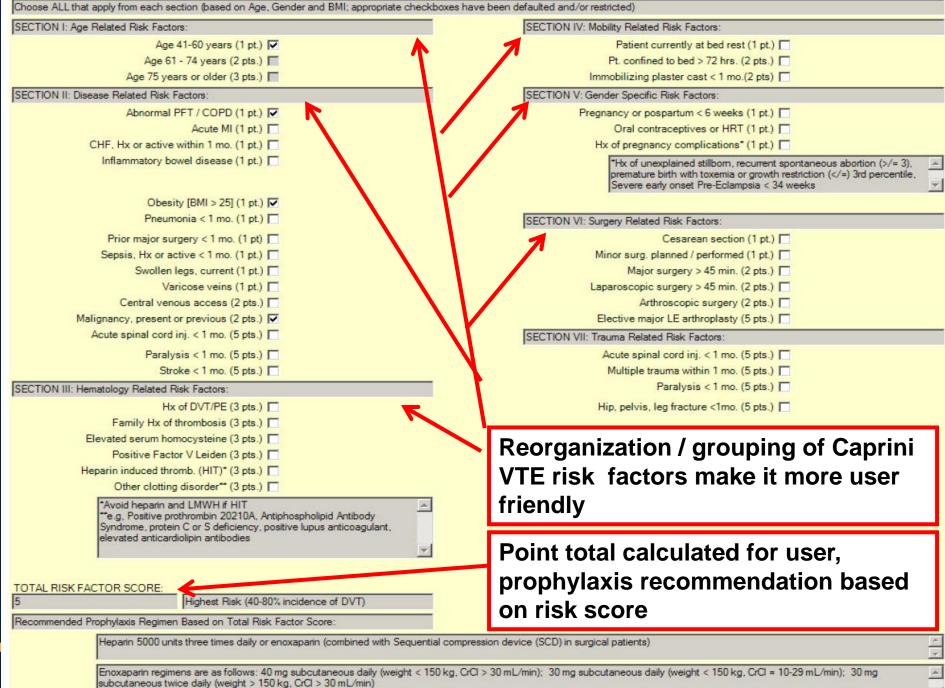
- 4. Pilot interventions on a small scale
 - a. Engage medical staff groups, look for barriers and special needs
 - b. Use case histories or real patient scenarios to simulate use of the order set
- 5. Monitor use of the protocol. Build measurement and monitoring into order set and documentation tools
 - a. Capture VTE risk, declaration of contraindications, what is ordered
 - b. Ambulation, IPCD adherence
 - c. Audits order sets being used? Completed properly?
 - d. Learn for variation from protocol

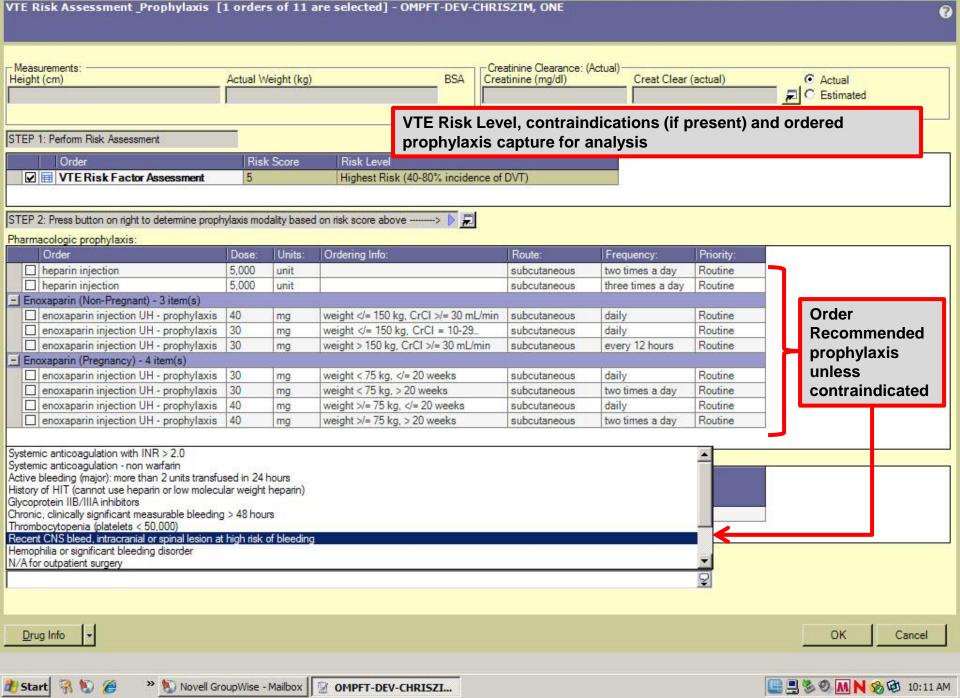


Key Strategies Implementing Caprini Model

Courtesy Marc Moote, PA-C

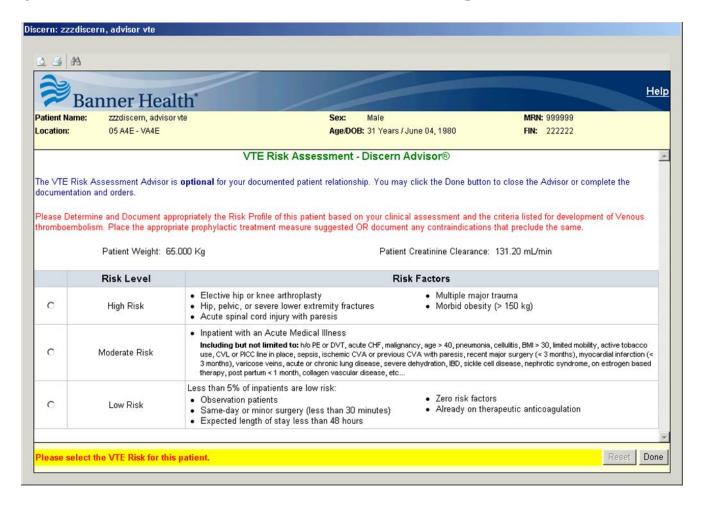
- Scope: <u>ALL</u> adult inpatients
- Standardized VTE Protocol Caprini model
- Mandatory risk assessment with CPOE hard-stop
- Clinical decision support to drive clinical practice
- Required documentation of contraindications
- Data feedback to services regarding performance
- VTE prophylaxis included as peer review (OPPE) indicator for many services
- Review of EVERY VTE event that occurs in the health system for preventability



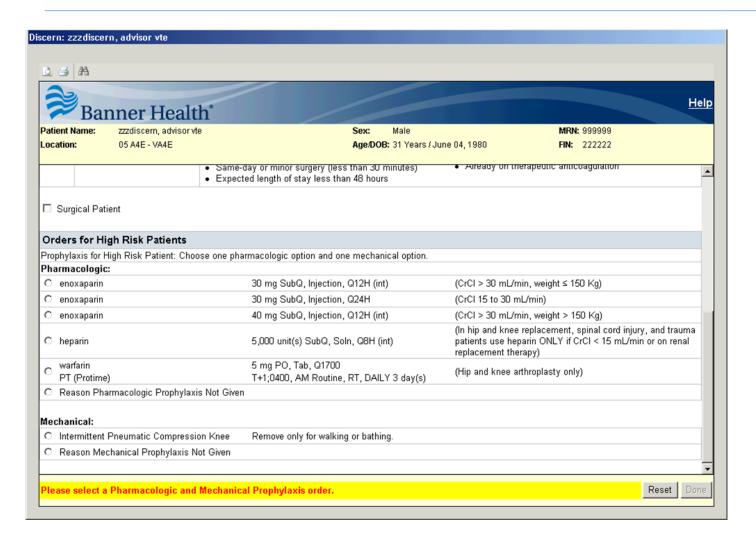


Classic 3 bucket model implementation

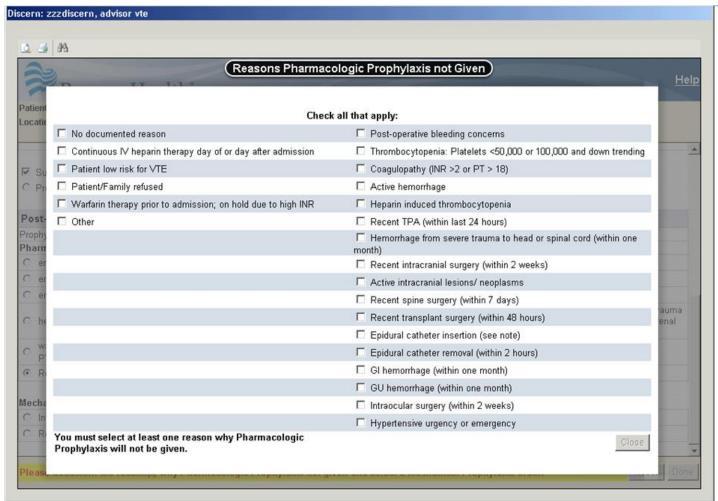
Courtesy Dr. Lori Porter, Good Samaritan Regional Medical Center

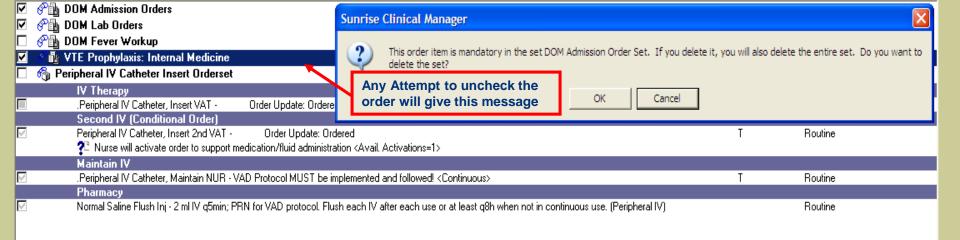


Risk-appropriate prophylaxis options appear after risk level chosen. High Risk requires dual prophylaxis



Contraindications captured if pharmacologic prophylaxis not ordered for a patient at risk of DVT.





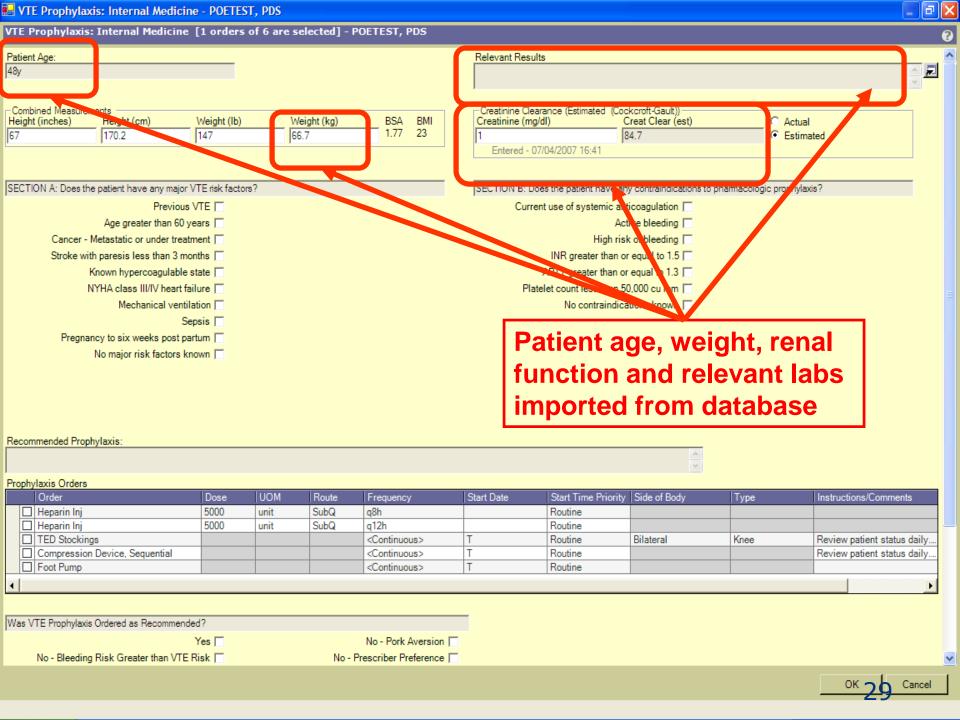
Johns Hopkins Medicine DVT Prevention Order Set Example Courtesy Dr. Michael Streiff

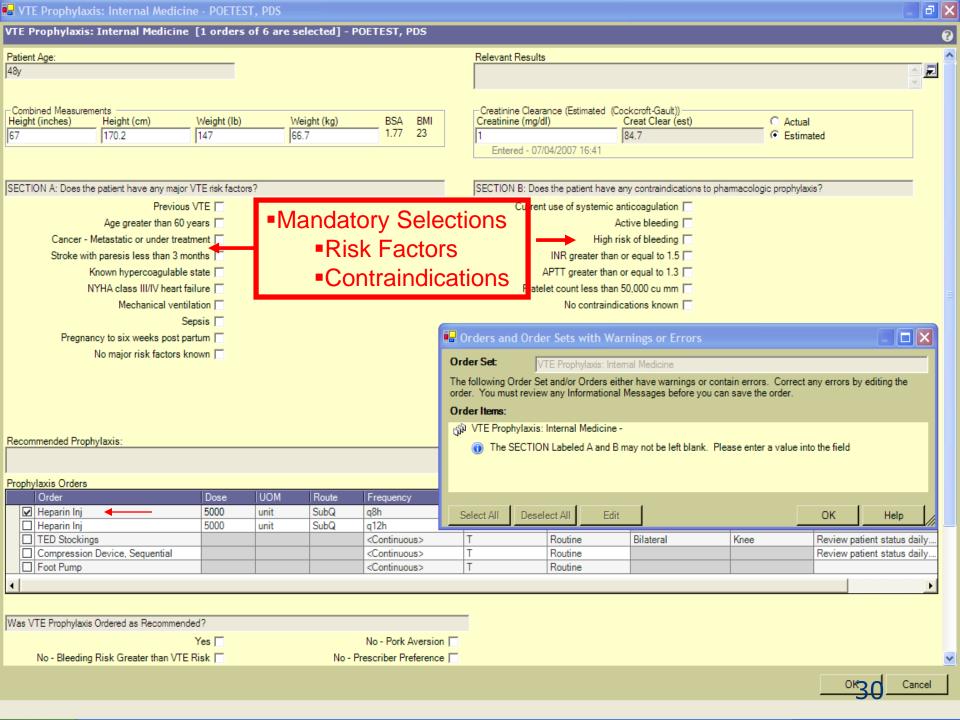
- Embedded in Medicine Admission Orders
- Hard Stop to use (vs delete entire order set)

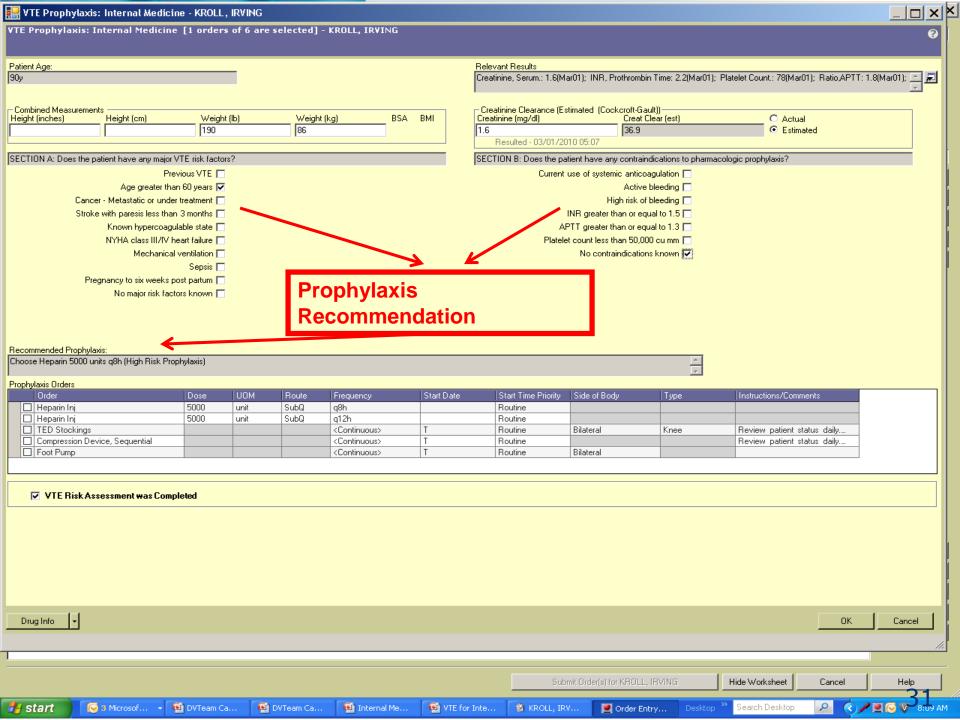
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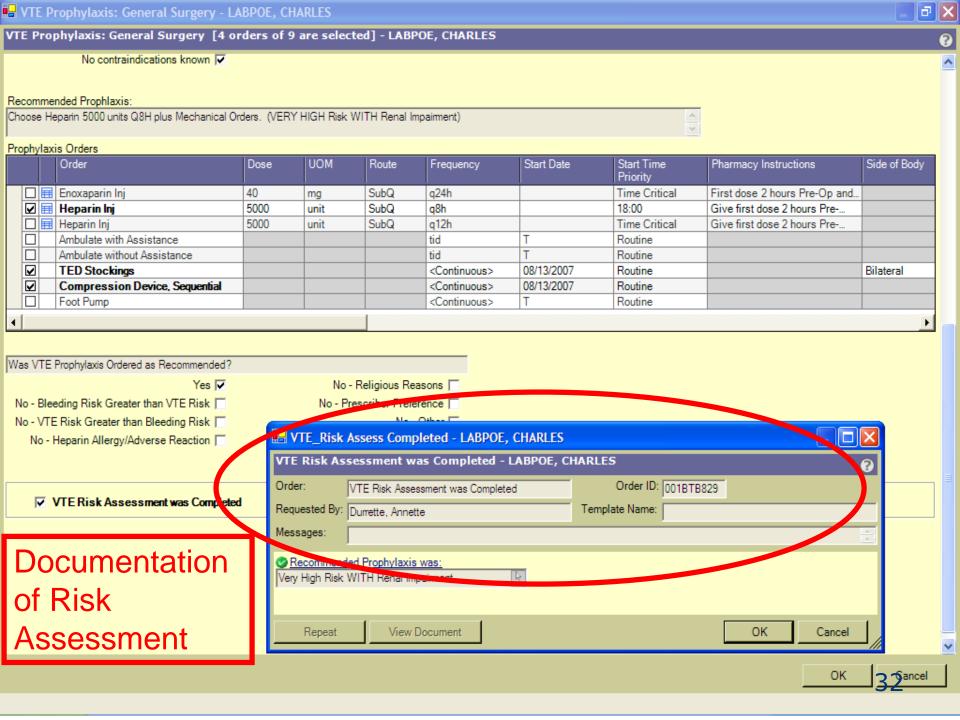
Cancel

Help









TJC and SCIP Measures

- Relatively low bar
- Do not drive rapid cycle QI
- Looks only at set points in hospitalization
 - Does not address patients who "fall off" protocol
- TJC measures: any prophylaxis = adequate prophylaxis

Go Beyond Core Measures to achieve better results

- Judge adequacy of prophylaxis by adherence to your protocol
- HA VTE = readmitted cases with new VTE + those not present on admission
- Monitor for lapses in care on a day-to-day basis

Outcomes measure for HA VTE and Preventable VTE

- Real time capture using imaging system, and concurrent review of cases to see if they are HA or community acquired, preventable / not preventable. Not practical for most, but may be gold standard.
- Improved methodology using administrative data
 - Captures readmitted patients as well as those with POA = No
 - Captures UE DVT, but tracks them separately
 - Higher bar for 'preventable'
 - Audits to validate coding
- Administrative coding Caveats

Need to address all common failures in process

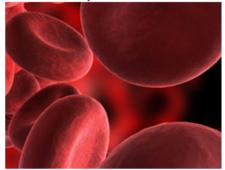
- No protocol / standardized order sets
- Order sets / prompts for VTE P in place, but no guidance
- Order sets with guidance in place but bypassed
- Order sets with guidance in place and used, but used incorrectly
- Patient gets placed on right prophylaxis, but VTE / bleeding risk changes and adjustment not made.
- Prophylaxis gets missed / changed on transfer / peri-op setting
- Correct prophylaxis ordered, but not administered, or patient refuses.
- Patient not mobilized optimally
- Preventable risk factors (central line) not optimally managed
- Patient had indication for extended duration prophylaxis, but did not get it

Strategies for VTE Prevention Beyond order sets

- Good protocol driven order set is well integrated
- Assessing administration / adherence
 - (not just orders)
- Alert Systems
 - Electronic alerts (E-alerts)
 - Human alerts
- Raising situational awareness (eg checklists)
- Audit and feedback
- Measure-vention
- Increase activity
- Optimize central lines
- Focus on extended duration for select populations

What is a blood clot?

- Clumps of thickened blood that blocks blood flow
- Blood clots most often form in your legs, arms, and groin but could move to your lungs, heart or brain
- Blood clots can be dangerous and deadly



Why am I at risk in the hospital?

- You are not moving around well *
- You recently had surgery or an injury
- Your disease may increase your chance of getting a clot

To prevent a blood clot from happening during your hospital stay, your doctor may ask you to take a medication or wear a leg device.

If your doctor asks you to take a medication....

- The medication is a blood thinner
- This medication is a small injection into fatty tissue just below the skin
- It may be given more than once a day
- You will likely not need the medication once you leave the hospital





If your doctor asks you to wear a leg device...

- Sleeves will be placed on your legs that will squeeze your legs off and on during the day
- This light squeeze will increase the flow of blood in your legs to prevent clots from forming
- These sleeves should be removed before you are out of bed and walking because they can cause you to trip and fall
- Be sure you to ask for the sleeves to be put back on when you are back in bed

^{*}If you are able to walk, this may decrease your risk. Please ask your nurse for help before getting out of bed.

MEASURE-VENTION

Daily <u>measure</u>ment drives concurrent inter<u>vention</u> (i.e., same as Level 5 in Hierarchy)

Identify suboptimal prophylaxis in real time

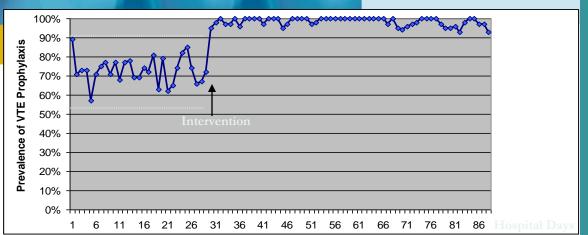
- Ongoing assessment
- Use for real-time intervention

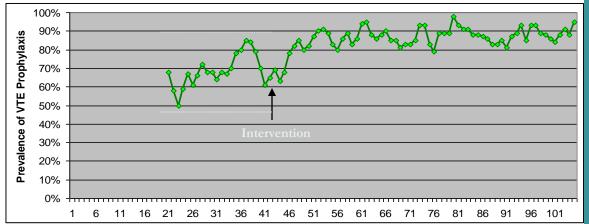
28 Patients – Measure-vention

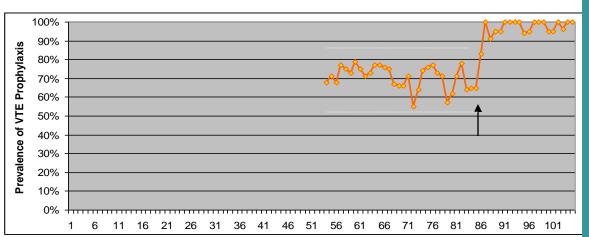
20 on anticoagulation

4 on mechanical prophylaxis with lab contraindication 3 on Nothing 1 mechanical

	יוווון אינט ל	9	± mcchamca					
							Orders	Orders
						Lab	state	state LOW
BED_LABEL	Service	VTE Risk Category		Dose	SCD	Contra	contra	VTE Risk
2250A	Medicine Thornton	LOW	warfarin (COUMADIN) tablet 3 mg	3 mg EVERY EVENING Oral	Υ	N	N	Υ
2250B	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 30 mg	30 mg DAILY Subcutaneous	Υ	N	N	N
2251	Medicine Thornton	MODERATE	heparin injection 5,000 Units	5000 Units EVERY 12 HOURS Su	Υ	N	N	N
2252	Cardiothoracic Surgery	MODERATE/HIGH	No Anticoag Med	No Anticoag Dose	Υ	Υ	N	Υ
2253	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	Υ	N	N
2254	Medicine Thornton	MODERATE	heparin injection 5,000 Units	5000 Units EVERY 8 HOURS Sub	Υ	N	N	N
2255	Medicine Thornton	MODERATE	heparin injection 5,000 Units	5000 Units EVERY 12 HOURS Su	Υ	N	N	N
2256A	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	N	N	N
2256B	Pulmonary Vascular Medicine	MODERATE/HIGH	enoxaparin (LOVENOX) injection 50 mg	50 mg EVERY 12 HOURS Subcut	Υ	Υ	N	N
2257A	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	N	N	N
2257B	Gynecology	MODERATE/HIGH	No Anticoag Med	No Anticoag Dose	Υ	Υ	N	N
2258	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 30 mg	30 mg DAILY Subcutaneous	Υ	N	N	Υ
2259	Medicine Thornton	MODERATE	No Anticoag Med	No Anticoag Dose	Υ	N	N	N
2260	Pulmonary/Critical Care	LOW	No Anticoag Med	No Anticoag Dose	N	N	N	Υ
2261	Medicine Thornton	MODERATE/HIGH	No Anticoag Med	No Anticoag Dose	Υ	Υ	N	N
2262A	Medicine Thornton	LOW	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	N	N	Υ
2262B	Medicine Thornton	MODERATE	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	N	N	N
2263	Medicine Thornton	MODERATE/HIGH	No Anticoag Med	No Anticoag Dose	Υ	Υ	N	N
2264	Pulmonary Vascular Medicine	MODERATE	warfarin (COUMADIN) tablet 5 mg	5 mg EVERY EVENING Oral	Υ	Υ	N	Υ
2265	Pulmonary Vascular Medicine	LOW	heparin injection 5,000 Units	5000 Units EVERY 8 HOURS Sub	Υ	N	N	Υ
2265	Pulmonary Vascular Medicine	LOW	warfarin (COUMADIN) tablet 10 mg	10 mg EVERY EVENING Oral	Υ	N	N	Υ
2266	Medicine Thornton	MODERATE	heparin injection 5,000 Units	5000 Units EVERY 8 HOURS Sub	Υ	N	N	N
2267	Pulmonary Vascular Medicine	HIGH	enoxaparin (LOVENOX) injection 100 mg	100 mg EVERY 12 HOURS Subcu	Υ	Υ	N	Υ
2268	Cardiothoracic Surgery	LOW	enoxaparin (LOVENOX) injection 40 mg	40 mg DAILY Subcutaneous	Υ	N	N	Υ
2269	Cardiothoracic Surgery	No Risk Category	No Anticoag Med	No Anticoag Dose	N	N	N	N
2270	Cardiothoracic Surgery	No Risk Category	No Anticoag Med	No Anticoag Dose	N	N	N	N
		MAGDEDATE	hanarin injection E 000 Units	5000 Units EVERY 12 HOURS Su	v	N	NI	N
2271	Medicine Thornton	MODERATE	heparin injection 5,000 Units	3000 UIIILS EVENT 12 HOURS SU	T	IN	N	







Effect of Situational Awareness on Prevalence of VTE Prophylaxis by Nursing Unit

Hospital A, 1st Nursing Unit

<u>Baseline</u> <u>Post-Intervention</u>

UCL: 93% 104%

Mean: 73% 99% (p < 0.01)

LCL: 53% 93%

Hospital A, 2nd Nursing Unit

Baseline Post-Intervention
UCL: 90% 102%

Mean: 68% 87% (p < 0.01)

LCL: 46% 72%

Hospital B, 1st Nursing Unit

Baseline Post-Intervention

UCL: 89% 108%

Mean: 71% 98% (p < 0.01)

LCL: 53% 88%

UCL = Upper Control Limit LCL = Lower Control Limit

Patient Enemy #1: Bed

Complications Associated with Hospital Beds:

- Aspiration pneumonia
- Deep Vein Thrombosis
- Delirium
- Pulmonary Emboli
- Pressure Ulcers
- Ileus, Bowel Paralysis

PICC Lines

- Increasing use
- Symptomatic VTE associated with PICC during hospitalization
 3.0 -7.8%
- Significant CLABSI burden
- Occlusion complications / lytics

Practices to Reduce PICC complications

- Minimize exposure to PICCs
 - Maximize midline / PIV
 - Remove asap
- Size matters smaller PICCs = fewer DVT
- Smallest number of lumens
- Proper flushing
- Following all infection control practices
- Fewer attempts to place PICC
- Appropriately sized catheter in proper position
- Appropriate DVT prophylaxis probably helps some, but not as much as for leg DVT
- Special catheters?

Questions / Answers / Comments?

- Coming Spring 2015 Major Revision / Update AHRQ DVT Prevention Guide
- Questions on this webinar series? Contact Cynthia Sayers at 404-498-0020.