



AMERICA'S ESSENTIAL HOSPITALS

Patient Harm Series III: VTE Measures and Prevention

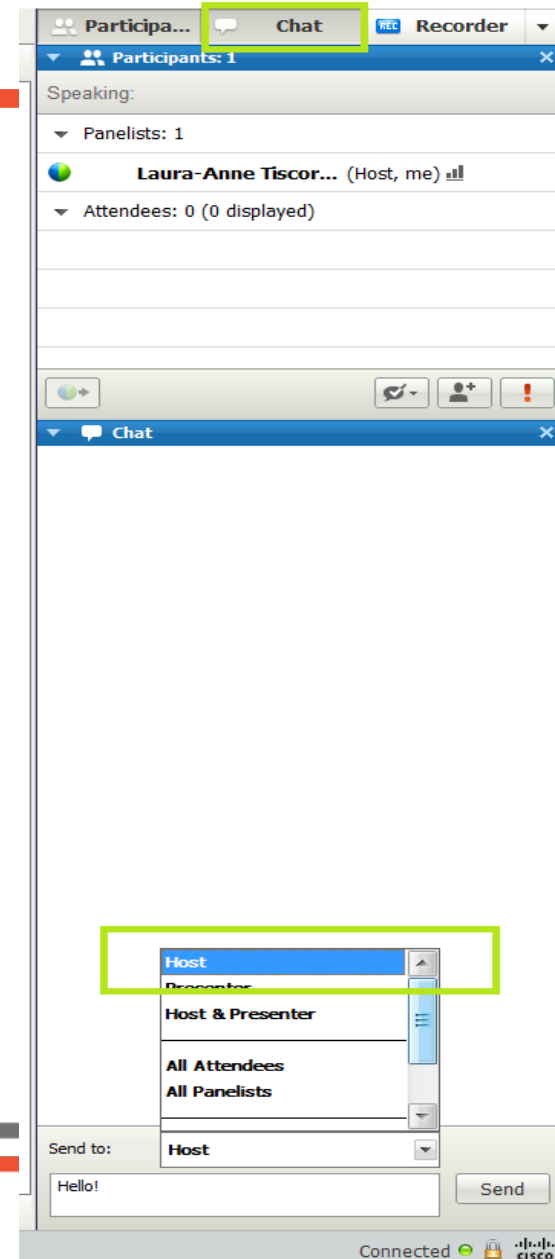
Essential Hospitals Engagement Network

May 8, 2014



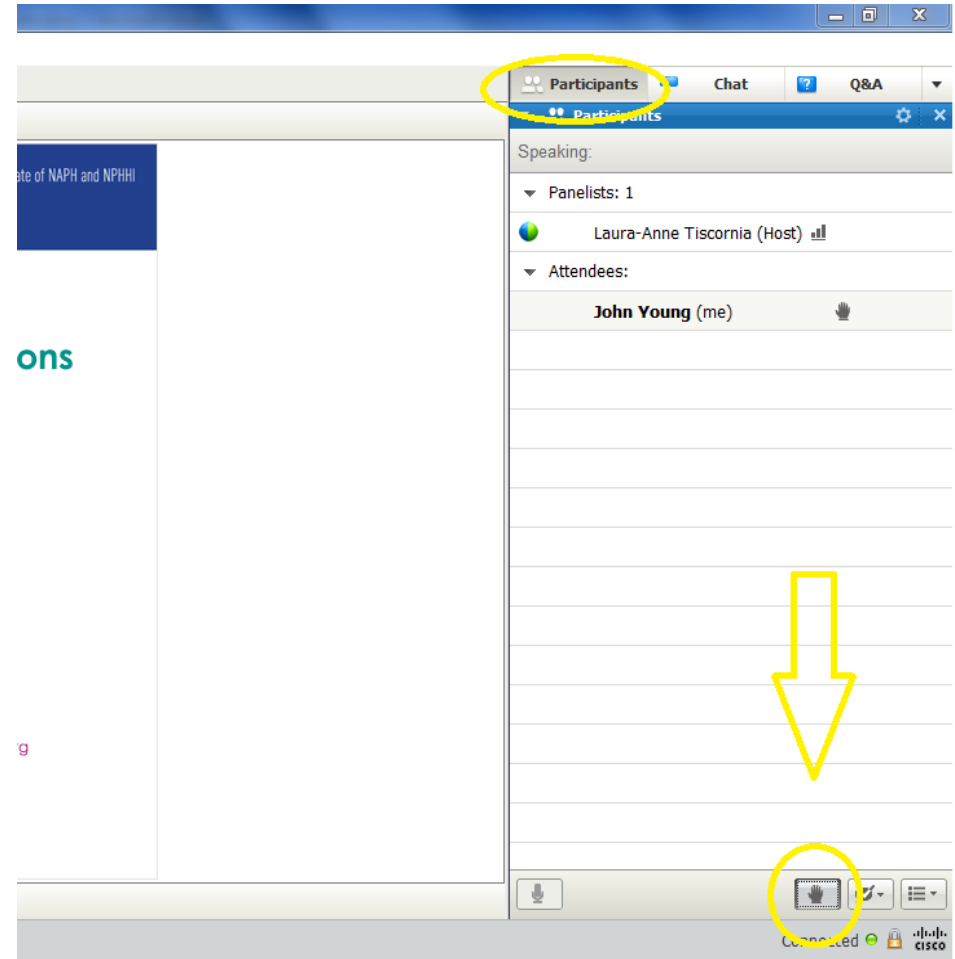
CHAT FEATURE

The chat tool is available to ask questions or comments at any time during this event.

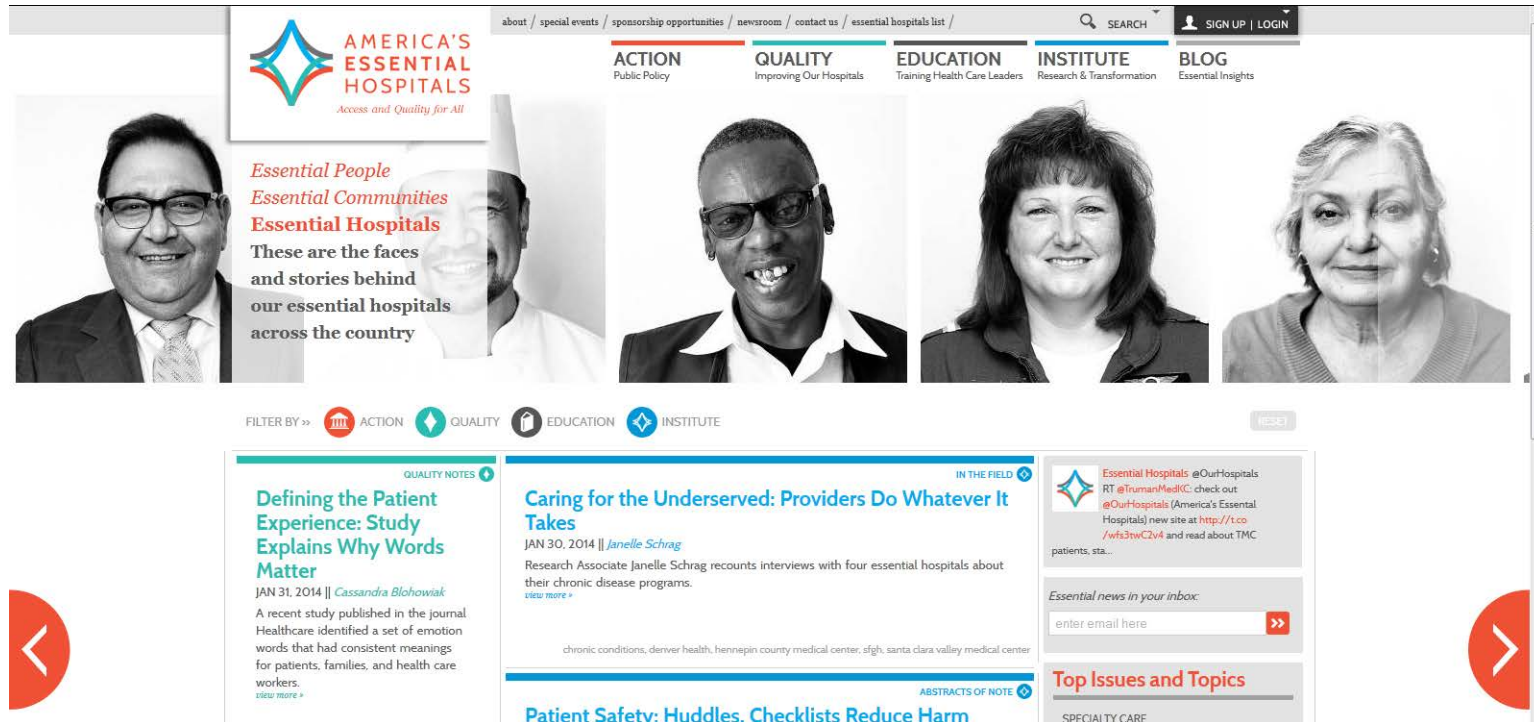


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If you wish to speak telephonically, please “raise your hand.” We will call your name, when your phone line is unmuted.



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AGENDA

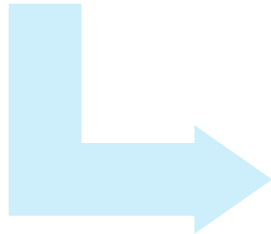
- Partnership for Patients and 2014
- AHRQ PSI-12 and Postoperative VTE after TKA (Banafsheh Sadeghi, MD, PhD)
- VTE prevention system (Anneliese Schleyer, MD, MHA/Ellen F. Robinson, PT)
- Q & A
- Upcoming events



2014 PARTNERSHIP FOR PATIENTS

Partnership for Patients (PfP)

- CMS-funded
- Reduce nine hospital-acquired conditions by 40 percent
- Reduce readmissions by 20 percent



Hospital Engagement Networks (HENs)

- 27 contracted organizations
- 3,700 U.S. hospitals



Essential Hospitals Engagement Network (EHEN)

- 22 hospitals nationwide
- Only essential hospital-focused HEN
- Special focus on health equity

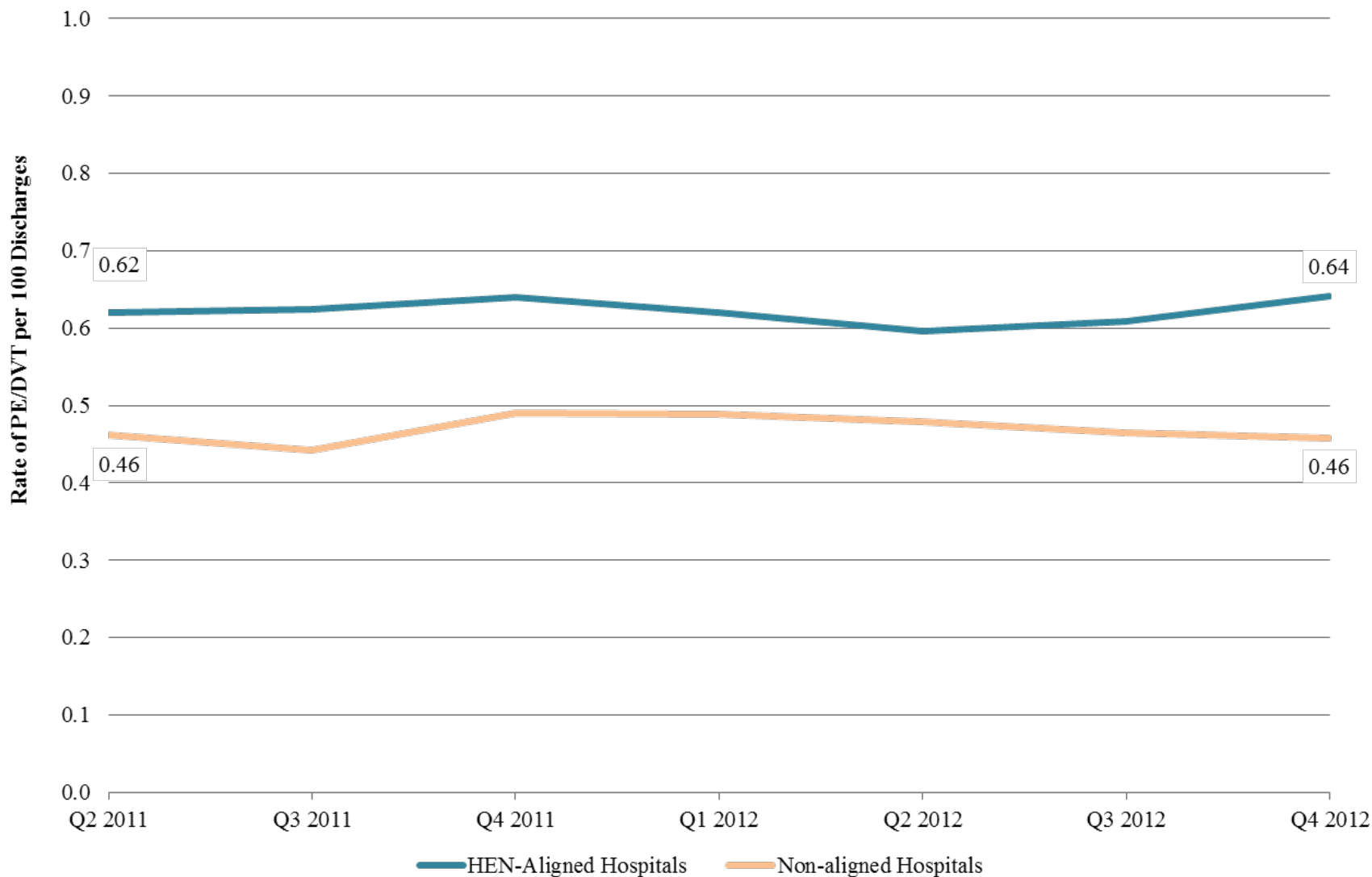


MEMBERS-ONLY VTE RESOURCES

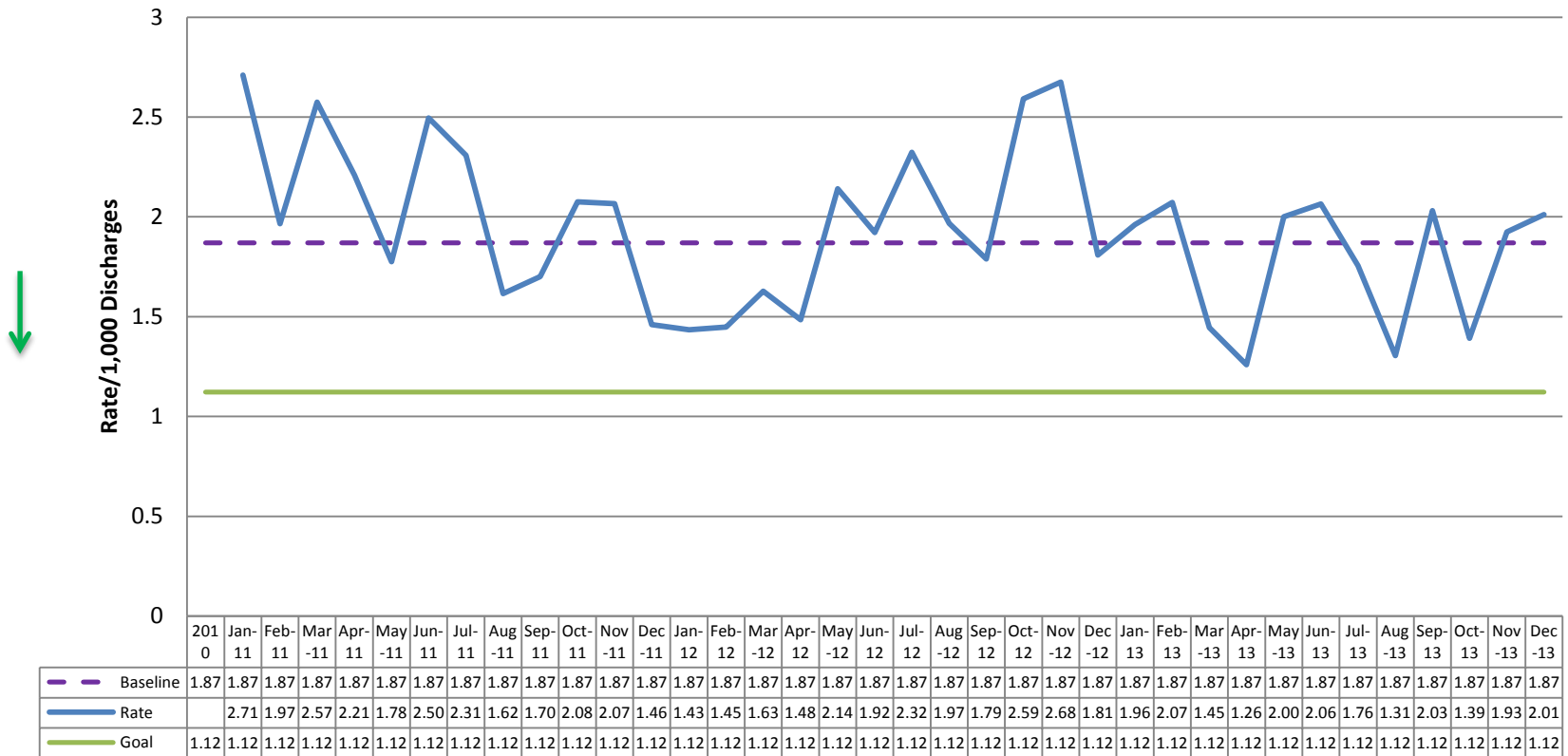
- VTE tab includes
 - » Resources
 - » Links
 - » Discussion thread
 - » Recordings of past webinars
- <http://essentialhospitals.org/groups/ehen/venous-thromboembolism/>



Post-Operative Pulmonary Embolism or DVT, Medicare FFS (PSI 12)



EHEN VTE: PE/DVT RATE (ALL DISCHARGES) [UHC-MODIFIED AHRQ PSI-12]



SPEAKER INFORMATION



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Overview AHRQ Quality Indicators: PSIs

- The AHRQ Patient Safety Indicators (PSIs)
- In hospital complications and adverse events following surgeries, procedures, and childbirth.
- Comprehensive literature review, analysis of ICD-9-CM codes



AHRQ Quality Indicators: PSIs

- Improve the safety of inpatient care
- Public reporting and pay-for-performance
- Identify potentially avoidable events
- Incidence of adverse events and in hospital complications
- using administrative data
 - hospital avoidable adverse events
 - regional level avoidable adverse event



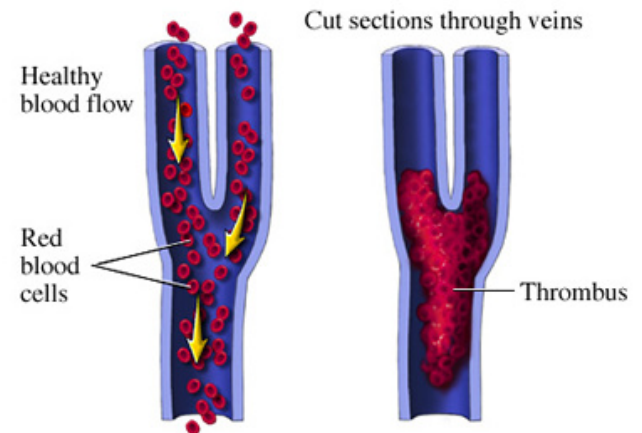
www.qualityindicators.ahrq.gov/psi_download.htm

Postoperative Venous Thromboembolism (PSI #12)

Patient Safety Indicators #12 (PSI #12)

Numerator

- Discharges, among cases meeting the inclusion and exclusion rules for the denominator, with a secondary ICD-9-CM diagnosis code for deep vein thrombosis or a secondary ICD-9-CM diagnosis code for pulmonary embolism.
- ICD-9-CM Deep vein thrombosis diagnosis codes¹:
45111, 45119, 4512, 45181, 4519, 45340, 4538, 4539
- ICD-9-CM Pulmonary embolism diagnosis codes¹:
4151, 41513, 41511, 41519



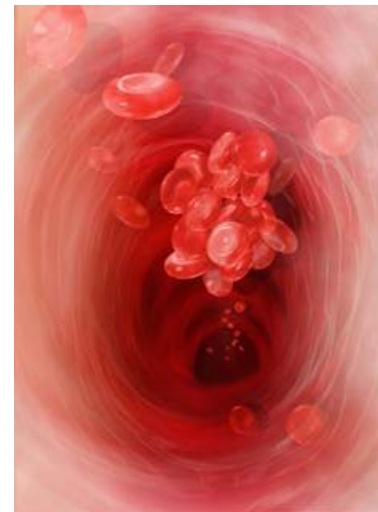
Postoperative Venous Thromboembolism (PSI #12)

Denominator

- Surgical discharges, for patients ages 18 years and older, with any-listed ICD-9-CM procedure codes for an operating room procedure. Surgical discharges are defined by specific DRG or MS-DRG codes.

Exclude cases:

- with a principal ICD-9-CM diagnosis code (or secondary diagnosis present on admission) for deep vein thrombosis (see above)
- with a principal ICD-9-CM diagnosis code (or secondary diagnosis present on admission) for pulmonary embolism (see above)
- where the only operating room procedure is interruption of vena cava
- where a procedure for interruption of vena cava occurs before or on the same day as the first operating room procedure†
- MDC 14 (pregnancy, childbirth, and puerperium)
- with missing gender (SEX=missing), age (AGE=missing), quarter (DQTR=missing), year (YEAR=missing), or principal diagnosis (DX1=missing)



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AHRQ Mission

AHRQ's mission is to produce evidence to make health care safer, higher quality, more accessible, equitable, and affordable, and to work with HHS and other partners to make sure

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Funding for Implementing PCOR

Questions Are The Answer

Treatment Options

ORIGINAL RESEARCH

Mechanical and Suboptimal Pharmacologic Prophylaxis and Delayed Mobilization but Not Morbid Obesity Are Associated With Venous Thromboembolism After Total Knee Arthroplasty: A Case-Control Study

Banafsheh Sadeghi, MD, PhD^{1*}, Patrick S. Romano, MD, MPH¹, Gregory Maynard, MD, MS², Amy L. Strater, MPH, MBA³, Laurie Hensley, MHA³, Julie Cerese, RN, MSN³, Richard H. White, MD, FACP¹

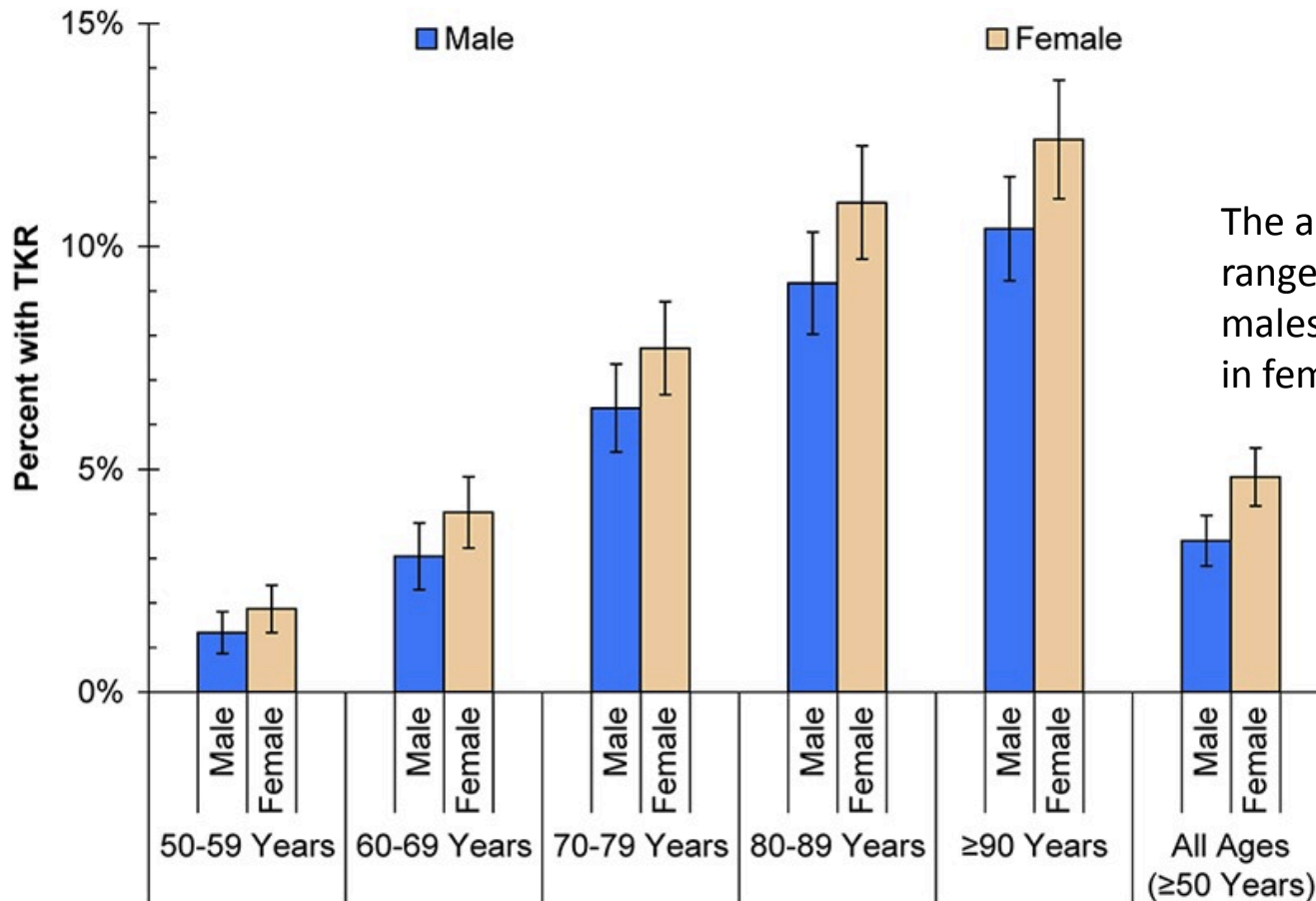
¹School of Medicine, Department of Internal Medicine, Division of General Medicine; ²School of Medicine, Division of Hospital Medicine; ³University HealthSystem Consortium, Chicago, Illinois.



Sadeghi B, Romano PS, Maynard G, Strater AL, Hensley L, Cerese J, White RH.

J Hosp Med. 2012 Nov-Dec;7(9):665-71.
PMID: 23042665

Total Knee Replacement (TKA)



The annual incidence of TKA ranged from 1.6% to 11.9% in males and from 2.0% to 10.9% in females.



J Bone Joint Surg Am. 2013 Mar 6;95(5):385-92. doi: 10.2106/JBJS.L.00206.

Estimating the burden of total knee replacement in the United States.

[Weinstein AM1, Rome BN, Reichmann WM, Collins JE, Burbine SA, Thornhill TS, Wright J, Katz JN, Losina E.](#)



Possible TKA Complications

- **Infection.** Major or deep infections may require more surgery and removal of the prosthesis.
- **VTE.** Blood clots in the leg veins are the most common complication of knee replacement surgery. PE can be life-threatening.
- **Implant problems.** Implant surfaces may wear down and the components may loosen. Additionally, although an average of 115° of motion is generally anticipated after surgery, scarring of the knee can occasionally occur, and motion may be more limited.
- **Continued pain.** A small number of patients continue to have pain after a knee replacement. This complication is rare.
- **Neurovascular injury.** While rare, injury to the nerves or blood vessels around the knee can occur during surgery.

VTE Prophylaxis after TKA

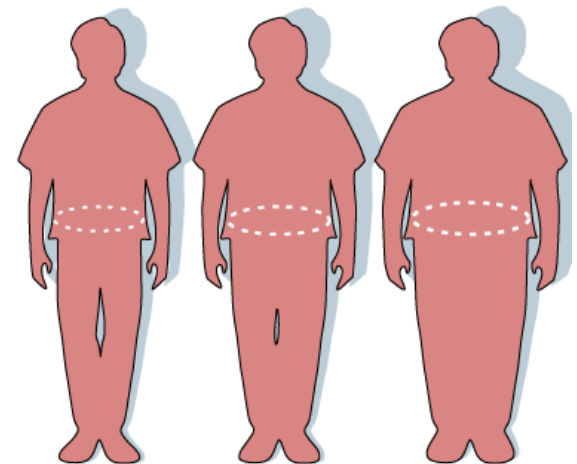
- Most guidelines from North America recommend the use of postoperative low-molecular-weight heparin (LMWH), fondaparinux, or warfarin for at least 10 days after TKA.



VTE Prophylaxis after TKA

Thromboprophylaxis reduces the risk of developing asymptomatic VTE by more than 60%.

Pharmacologic prophylaxis using LMWH, fondaparinux, or warfarin alone is recommended by the ACCP and other organizations, with use of mechanical pneumatic compression, low-dose unfractionated heparin, or aspirin as alternative options.



Study Hypothesis and Goals:

1. Use of standard pharmacological thromboprophylaxis drug is associated with lower risk of acute VTE compared with mechanical prophylaxis only.
2. Among patients given LMWH/fondaparinux, excessive obesity ($\text{BMI} > 35$) is associated with a higher risk of developing VTE.
3. Delayed ambulation after TKA is associated with higher risk for VTE.



Methods: Partnership

- Univ. of California Davis (UCD)
 - 631-bed hospital serves
 - key referral center for 33 counties and 6 million residents.
 - Northern California's only level I trauma center
 - UC Davis Medical Center ranks among the top 50 hospitals in America according to annual U.S. News & World Report survey



- University HealthSystem Consortium (UHC)
 - UHC is an alliance of 120 academic medical centers and 299 of their affiliated hospitals.



Methods: Study Design

- Retrospective case-control study.
- Fifteen volunteer hospitals nationwide abstracted medical records up to 40.

Cases:

- Having one or more secondary diagnosis codes for VTE as defined by AHRQ PSI-12 coupled with a POA flag of “no”

OR

- Were readmitted with principal diagnosis of VTE within 90 days of date of surgery



Controls:

- Patients who did not develop acute VTE during the index hospitalization or within 90 days of surgery.

Methods: Inclusion & Exclusion Criteria

- Inclusion:
 - Admission between Oct. 1, 2008 and March 31, 2010;
 - ICD-9-CM procedure code of 81.54 or 81.55;
 - Age of 40 years or more;
- Exclusion:
 - Pregnancy related principal diagnosis;
 - Inferior vena cava interruption on or before the date of the first operating room procedure.



Data Collection: Chart Abstraction Tool

A chart abstraction tool was constructed and personnel at each site were taught how to obtain the desired information.



UHC University HealthSystem Consortium	
VTE Prophylaxis in Total Knee Replacement (TKR) Benchmarking Project Patient Level Data Collection Form	
BENCHMARKING & IMPROVEMENT SERVICES	
Methods: Retrospective review of up to 40 cases undergoing total knee replacement (TKR) surgery and discharged between October 1, 2008, and March 31, 2010, and meeting specific enrollment criteria for each of 2 patient groups described below.	
Inclusion Criteria for all patients: <ul style="list-style-type: none">• Adult patients ≥ 40 years of age• Total Knee Replacement (TKR) procedure (unilateral, bilateral, or revision not for infection) performed during index admission	
Group 1a: Patients who developed acute VTE after TKR surgery during the index admission and/or	
Group 1b: Patients who developed acute VTE within 90 days of discharge from the index admission (NOTE: If the number of eligible cases per site exceeds 20, Group 1 cases will be selected via a randomization scheme constructed by UHC.)	
Group 2: A sample of control cases that did not develop acute VTE after TKR surgery during the index admission or within 90 days of discharge from the index admission	
Exclusion Criteria for all patients: <ul style="list-style-type: none">• Had another hip or knee surgery within 90 days prior to the indexed TKR• Patients with principal ICD-9-CM diagnosis code for DVT/PE for the index admission• Patients with secondary ICD-9-CM diagnosis code for DVT/PE that is POA=yes for the index admission• A primary diagnosis or reason for admission related to pregnancy, childbirth or puerperium (MDC 14)	
A. ADMINISTRATIVE	
1. Encounter number: (CDB) (required) <input type="text"/>	
2. Age: ___ years (CDB) (required)	
3. Hospital admission date (CDB) (required)	
Date: / / (mm/dd/yyyy)	

Data Collection: Chart Abstraction Tool

Data elements:

- Age
- Gender
- Height
- Weight
- Type of TKA
- Use of pharmacologic and mechanical prophylaxis
- Ambulation status

Types of prophylaxis categories:

1. LMWH/fondaparinux with or without mechanical prophylaxis;
2. Warfarin alone, with or without mechanical prophylaxis;
3. LMWH/fondaparinux and warfarin with or without mechanical prophylaxis;
4. Mechanical prophylaxis (Pneumatic compression devices, graduated compression stockings, or foot pump) alone (without any pharmacological prophylaxis but with or without aspirin);
5. Aspirin only, without any other pharmacologic or mechanical prophylaxis.

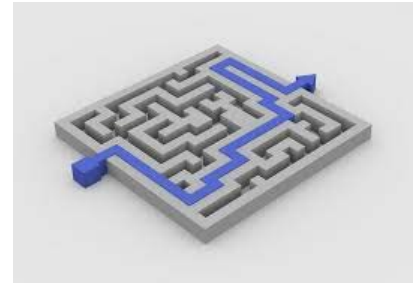
FDA-approval status:

- FDA-approved
- Pharmacologic prophylaxis
- Other prophylaxis



Outcome of Study

- Principal outcome:
 - Validated symptomatic objectively confirmed VTE
 - Patients with VTE on the day of surgery or the day after surgery were not included in the principal analysis.



- Statistical analysis:
 - Bivariate analysis: student t-test, chi-square, unadjusted Odds Ratio (OR)
 - Multivariate analysis: logistic regression and two-way interactions adjusted for correlated data, using SAS-PC 9.2



Results:

Total of 593 TKA record were abstracted

All patients underwent TKA on the day of admission or the day after:

- 114 cases (44 PE, 68 DVT, 2 both)
- 463 controls
- 16 cases (12 PE and 4 DVT) excluded.

Result: Bivariate Analysis

- Age in cases significantly higher than controls: 65.5 ± 10.4 vs. 63.5 ± 10.4 ($P < 0.05$)
- Cases underwent more bilateral: 23% vs. 7% ($P < 0.001$)
- Cases had marginally higher BMI than controls: 34.6 ± 8.0 vs. 33.3 ± 7.1 ($P = 0.07$)
 - Among PE cases, there were more morbidly obese cases compared with controls: 30% vs. 16% ($P = 0.01$)



Result: Bivariate Analysis

- Fewer VTE cases began ambulation on or before the second post-op day compared with controls: 47% vs. 73% ($P < 0.001$).
- All patients received at least 1 type of pharmacological or mechanical prophylaxis within 24 hrs. after TKA.
- Controls had marginally higher odds of receiving FDA-approved pharmacologic prophylaxis than cases 61% vs. 48% ($P = 0.07$).



Results: Multivariable Analysis

Predictive Factor	Odds Ratio (95% CI)	P value
Age	1.02 (0.99 – 1.05)	0.20
Gender (ref: male)	1.7 (0.9 – 2.9)	0.90
Ambulation (ref: no ambulation) <ul style="list-style-type: none">• Taking steps day 1 or 2• Taking steps after day 2	0.3 (0.1 – 0.9) 0.7 (0.2 – 2.1)	0.005 0.56
Type of TKA (ref: unilateral TKA) <ul style="list-style-type: none">• Bilateral TKR	4.2 (1.9 – 9.1)	0.004
Recommended pharmacologic prophylaxis (ref: only mechanical)	0.5 (0.3 – 0.8)	0.01
BMI ≥ 35 (ref: BMI < 35)	0.9 (0.5 – 1.6)	0.66

Results: Prophylaxis

- FDA-approved pharmacologic prophylaxis:
 - LMWH, fondaparinux, or warfarin
- Odds Ratio = 0.5; 95% CI: 0.3 – 0.8, P-value= 0.01
- Risk of VTE across the BMI levels



Results: Ambulation

- Early mobilization in the first 2 days after:
 - Odds Ratio = 0.3, 95% CI: 0.1 – 0.9, P-value = 0.005
- Mobilization at or after day 3



Results: Bilateral vs. Unilateral

- Bilateral simultaneous TKA strongly associated with VTE:
 - Odds Ratio = 4.2, 95% CI: 1.9 – 9.1, P-value <0.001
 - The effect still existed after adjusting for obesity or time of mobilization.



Study limitations

- No record of readmission to other hospitals
- Effect of hospital volume and specialization
- Data collection by employees and no double collection
- All teaching hospitals
- Inherent case-control limitations



Conclusion



- Actionable opportunities to improve care to prevent VTE persist despite 100% compliance with existing TJC process measures
- FDA approved pharmacologic prophylaxis
- Early mobilization
- Bilateral vs. unilateral
- AHRQ PSI#12
- Further research studies needed

Evidence-Based Clinical Practice Guidelines

Guideline	Accepted VTE Prophylaxis	Defined Risk
<p>American College of Chest Physicians: Antithrombotic Therapy and Prevention of Thrombosis, 9th ed.</p> <p>Guyatt GH, Akl EA, Crowther M. Executive summary: Evidence-Based Clinical Practice Guidelines. <i>Chest</i>. 2012;141(2 suppl):7S-47S.</p> <p>Geerts WH, Bergquist D, Pineo GF, et al. Prevention of venous thromboembolism. <i>Chest</i>. 2008;133(6):381S-453S.</p>	<ul style="list-style-type: none"> One of the following for 10-14 days <ul style="list-style-type: none"> LMWH started ≥ 12 h before or after surgery Low-dose unfractionated heparin Factor Xa inhibitor (fondaparinux, apixaban, rivaroxaban) Warfarin Dabigatran Recommends against the use of aspirin alone for any group In patients with elevated bleeding risk: <ul style="list-style-type: none"> Intermittent pneumatic compression device (18 hours daily) 	<p>Total knee surgeries are considered high risk for VTE, regardless of age, activity level, or comorbidities</p>
<p>American Academy of Orthopedic Surgeons</p> <p>http://www.aaos.org/research/guidelines/VTE/VTE_guideline.asp</p>	<p>Pharmacologic prophylaxis as above</p> <p>Mechanical prophylaxis for patients with elevated bleeding risk:</p> <ul style="list-style-type: none"> Pneumatic compression devices Foot and leg pumps 	<p>Total knee surgeries are considered standard risk for PE. Bleeding risk defined as:</p> <ul style="list-style-type: none"> History of a bleeding disorder History of recent gastrointestinal bleed History of recent hemorrhagic stroke
<p>Surgical Care Improvement Project</p>	<ul style="list-style-type: none"> LMWH Factor Xa inhibitor (fondaparinux) Warfarin Intermittent pneumatic compression devices Venous foot pump 	<p>Orthopedic surgeries with surgical time > 60 min and LOS > 3 days</p>

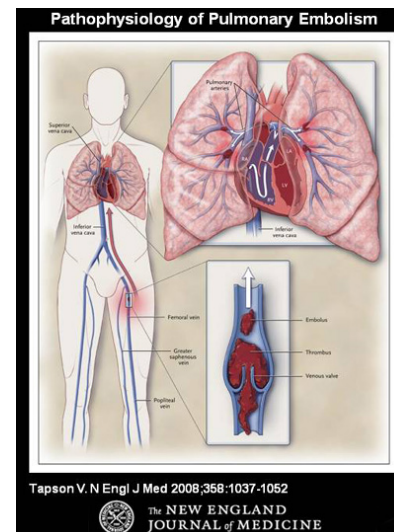
Q & A



Banafsheh Sadeghi, M.D., Ph.D.
Assistant Adjunct Professor, School of Medicine
UC Davis Department of Internal Medicine
Sacramento, CA

Today's Objectives

- Describe Multidisciplinary VTE Team
- Discuss innovative approaches that have reduced post-operative DVT/PE
- Outline systematic event review methods
- Provide overview of the tools developed for engaging front line staff



Harborview Medical Center



Key Centers of Excellence

- Level I adult and pediatric trauma and burn care
- Neurosciences Institute
- Orthopaedic reconstruction and rehabilitation
- Comprehensive Eye Institute and vision science center
- Vascular
- Behavioral health
- AIDS/STD

2012 Statistics

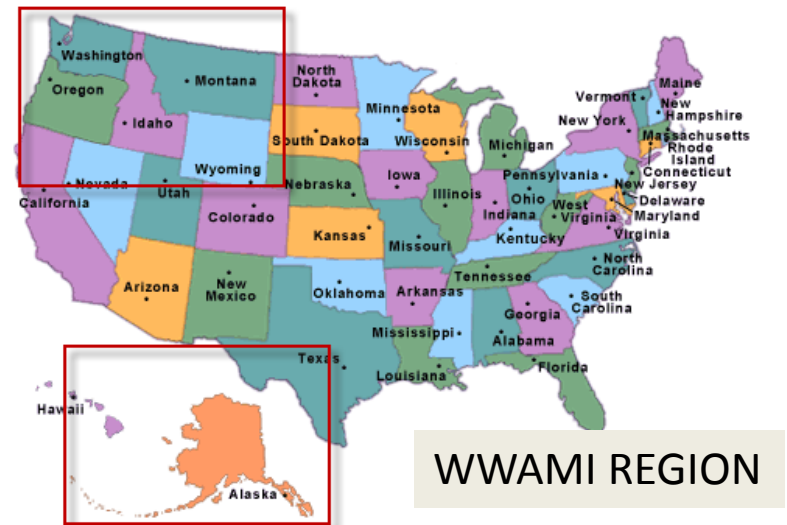
Licensed beds	413
Employees	4,684
Physicians	1,243
Admissions	19,094
Emergency Department visits	62,432
Clinic visits	247,246
Surgery cases	14,872

Charity Care

Provided \$210 million in charity care in fiscal year 2012.



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VTE Team Members

- Physician Co-Chair – Surgeon and Hospitalist
- Pharmacy – clinical, safety, administration
- Nursing – trauma, clinical education
- Quality Improvement – analyst, programmer
- Input from Orthopedic, Neurology, Neurosurgery physicians, EMAR IT analysts, UWMC pharmacy team, discharge planning

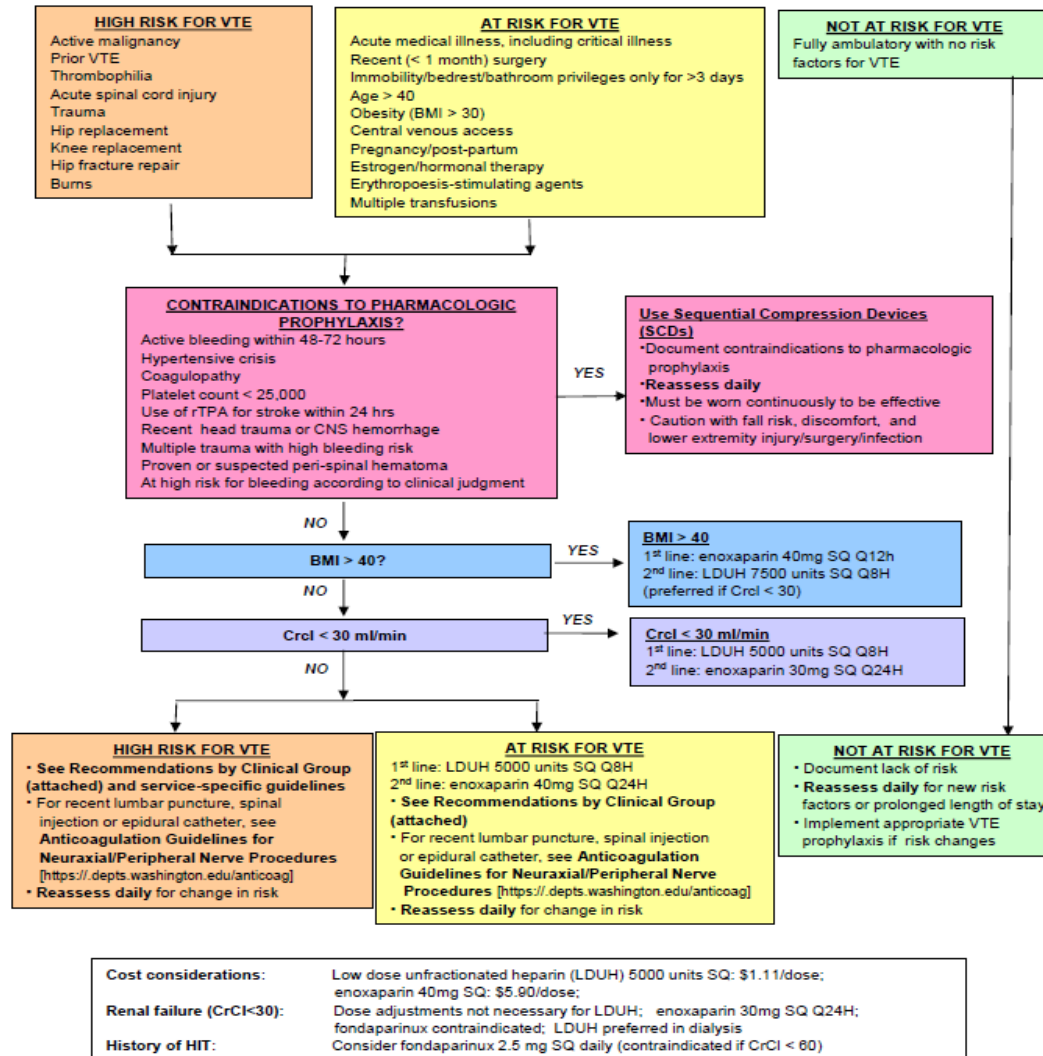
UW Medicine Guidelines for Prevention of VTE

Evidence based following national guidelines

UW Medicine

GUIDELINES FOR PREVENTION OF VENOUS THROMBOEMBOLISM (VTE) IN HOSPITALIZED PATIENTS

UW Medicine Recommended Practices based on Antithrombotic Therapy and Prevention of Thrombosis, 9th Edition, American College of Chest Physicians Evidence Based Clinical Practice Guidelines; Chest 2012 (suppl 2).



<http://uwmcacc.org>

THESE GUIDELINES ARE NOT INTENDED TO SUPERCEDE CLINICAL JUDGEMENT

UW Medicine VTE Prophylaxis Taskforce
J Cuschieri MD, R Dumitru PharmD, P Kritek MD, A Schleyer MD, A Wittkowsky PharmD
November 2013

VTE Case Reviews

*Hospital Acquired VTE

- Clinical information for quality review imbedded complex electronic medical record (EMR - Cerner)
- Time consuming, not systematic
- Created an electronic tool to allow for efficient monthly team review



- For all HAC VTE
- Team reviews following elements for guideline adherence
 - Risk Assessment performed
 - Chemical Prophylaxis timing, drug selection, dosing
 - Mechanical Prophylaxis utilized if chemical not appropriate
 - Charted Contraindications

UW Medicine

MRN	Name	Age	Admit Dt Tm	Disch Dt Tm	BMI	Notes

Patient presents with right lower extremity swelling. Deep vein thrombosis (non-occlusive) in the right mid femoral vein. Deep vein thrombosis (occlusive) in the right posterior tibial veins. Could not rule out non-occlusive thrombus in the right peroneal veins due to vessel depth and poor visualization. No deep vein thrombosis in the right common femoral, proximal profunda femoris, popliteal, or left lower extremity veins.

Team review utilizing pertinent data points

- [illegible]

* *J Hosp Med* 2014 Jan; 9(1) 48-53

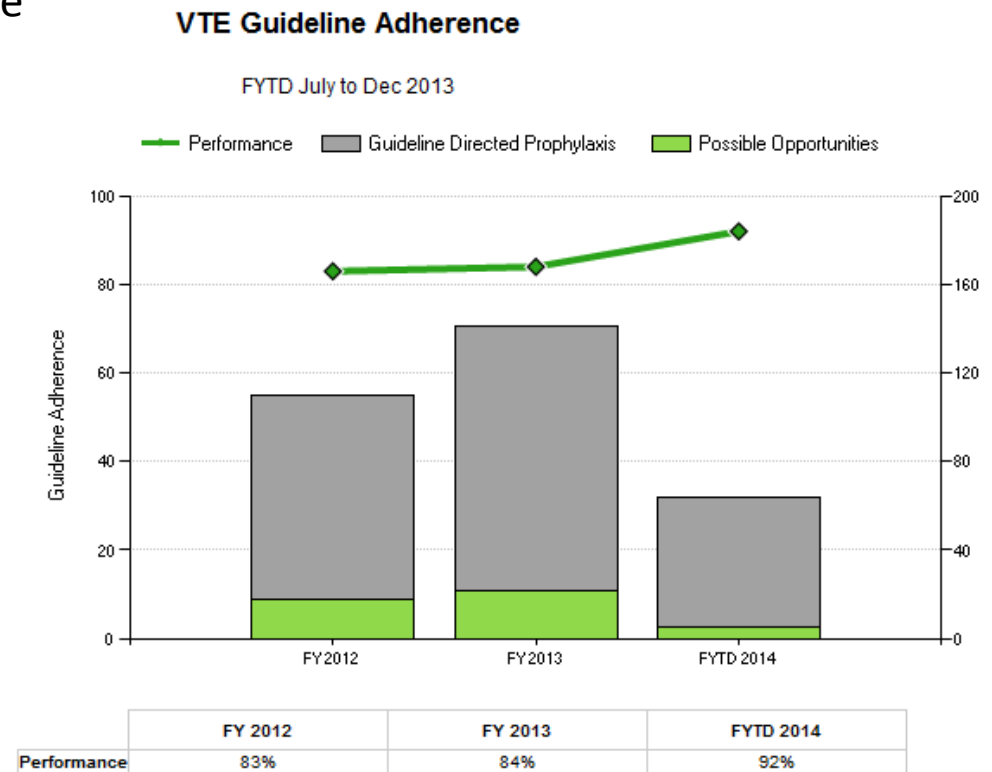
From Retrospective to “Real Time”

Room	Admit Date	Age	LOS	Reason	INR	SCD	Drug Dose	Drug Ordered
			3	LEFT INTERTROCHANTERIC FEMUR FRACTURE; LEFT PARASYMPHYSEAL PELVIC FRACTURE	1.1 2013-12-09 15:35	OFF	Not Given: Pt still on PNC enoxaparin 2013-12-11 21:00	
			9	LUMBAR 2 BURST FRACTURE	1.1 2013-12-03 11:40	OFF	Not Done: hold per MD order, capping PNC's in am enoxaparin 2013-12-11 21:00	
			1	Ulcer of other part of foot		OFF		Not Ordered
			4	GUN SHOT WOUND TO LEFT HAND	1.0 2013-12-08 16:30	None Applied		Not Ordered
			2	Acquired spondylolisthesis		ON		Not Ordered
			1	Spondylolisthesis of lumbar region		ON		Not Ordered
			2	Hallux valgus (acquired)		ON	40 mg enoxaparin 2013-12-11 09:00	
			11	LEFT HUMERUS FRACTURE	1.1 2013-12-12 07:26	OFF	5,000 units heparin 2013-12-05 17:00	
			1	SUBARACHNOID HEMORRHAGE	0.9 2013-12-12 06:22	None Applied		Not Ordered
			1	Spinal stenosis, lumbar region, without neurogenic claudication	1.2 2013-12-11 22:00	ON		Not Ordered
			1	GASTROINTESTINAL BLEED	1.7 2013-12-12 04:45	None Applied		warfarin 2013-12-11 19:39
			2	POLYTRAUMA	1.0 2013-12-12 04:12	None Applied		Not Ordered
			6	Gunshot Wound to the Left Chest and Neck	1.0 2013-12-12 06:05	None Applied	Not Given: Patient Refused heparin 2013-12-12 01:00	

Intervention can be related to resident, nursing or patient education

Guideline Directed Prophylaxis

Cases that do not adhere to guidelines per VTE team review sent for secondary review to each medical service and results reported through Medical QI Committee

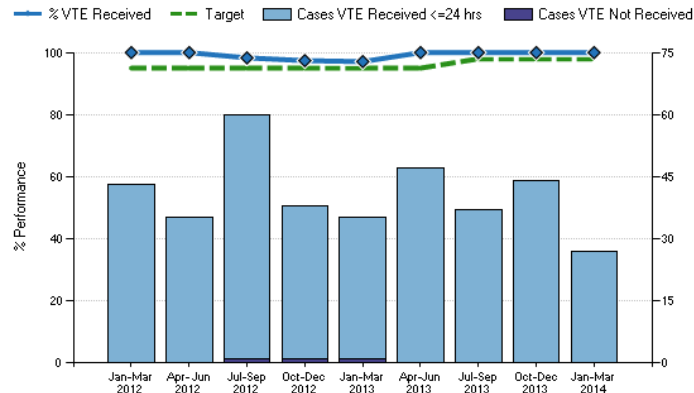


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SCIP/VTE Prophylaxis Core Measures

HMC Core Measures SCIP Appropriate Venous Thromboembolism Prophylaxis Received within 24 hrs

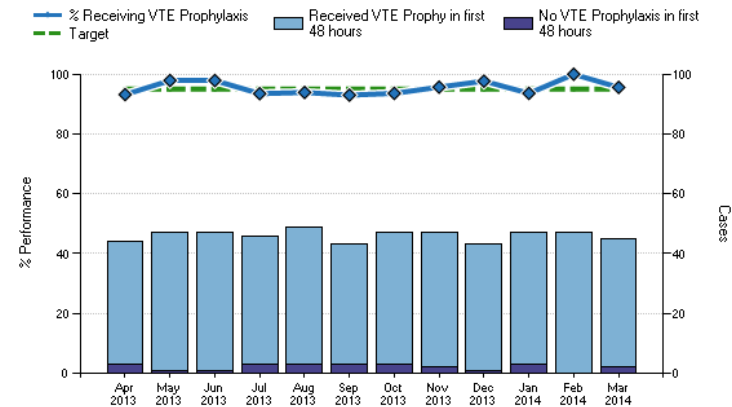
Q1 2014: No Fallouts
VBP Threshold: 94.9 VBP Benchmark: 99.8%



	Jan-Mar 2012	Apr-Jun 2012	Jul-Sep 2012	Oct-Dec 2012	Jan-Mar 2013	Apr-Jun 2013	Jul-Sep 2013	Oct-Dec 2013	Jan-Mar 2014
% VTE Received	100%	100%	98.3%	97.4%	97.1%	100%	100%	100%	100%
Cases VTE Received <=24 hrs	43	35	59	37	34	47	37	44	27
Cases VTE Not Received	0	0	1	1	1	0	0	0	0
Eligible Cases	43	35	60	38	35	47	37	44	27
Target	95%	95%	95%	95%	95%	95%	98%	98%	98%

Venous Thromboembolism - Prophylaxis Acute Care and ICU

March 2014: SCDs ordered but not placed - 2W, 7E



	Apr 2013	May 2013	Jun 2013	Jul 2013	Aug 2013	Sep 2013	Oct 2013	Nov 2013	Dec 2013	Jan 2014	Feb 2014	Mar 2014
% Receiving VTE Prophylaxis	93.2%	97.9%	97.9%	93.6%	93.9%	93.0%	93.6%	95.7%	97.7%	93.6%	100%	95.6%
Received VTE Prophylaxis in first 48 hours	41	46	46	43	46	40	44	45	42	44	47	43
No VTE Prophylaxis in first 48 hours	3	1	1	3	3	3	3	2	1	3	0	2
Total Patients (sampled)	44	47	47	46	49	43	47	47	43	47	47	45
Target	95%	95%	95%	95%	95%	95%	95%	95%	95%	95%	95%	95%

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Expanding to “All Patients”

- Take findings from Case Reviews and implement system change for all hospitalized patients
- Areas for ongoing improvement
 - VTE Risk Assessment
 - Chemical prophylaxis dosing
 - Held chemical doses
 - SCD utilization



Ongoing Improvement – VTE Risk Assessment

CPOE VTE Prophylaxis order set April 2013
Allowed Risk Assessment to be captured

Ongoing Monthly Monitoring (March 2014)
93% of inpatients had risk assessed

Educated Epilepsy to assess risk

Medical and Surgical Teams educating
“high risk” versus “at risk”

RX VTE Prophylaxis (Planned Pending)

Medications

Pharmacologic Prophylaxis

☐ See evidence link for Anticoagulation Guidelines for Neuraxial or Peripheral Nerve Procedures.

☐ heparin 5,000 units, Subcutaneous, Q8 Hours, Injection

☐ enoxaparin 40 mg, Subcutaneous, QHS, Injection

☐ enoxaparin dosing for Severe Renal Failure (CrCl < 30)

☐ enoxaparin 30 mg, Subcutaneous, QHS, Injection

☐ enoxaparin dosing for Obesity (BMI > 40)

☐ enoxaparin 40 mg, Subcutaneous, Q12 Hours, Injection

☐ Trauma, Acute Spinal Cord Injury, Total Hip Replacement, Total Knee Replacement, Hip Fracture Repair or Burns.

☐ enoxaparin 30 mg, Subcutaneous, Q12 Hours, Injection

VTE Prophylaxis

☐ GENERAL GUIDELINES: Review Reference Text

1. Risk for Venous Thromboembolism (VTE): Not at Risk, At Risk or High Risk

2. Contraindications to pharmacologic and mechanical prophylaxis

☐ At Risk - VTE Prophylaxis (most patients)

☐ At Risk - VTE Prophylaxis DELAYED Pharmacologic Pro...

☐ High Risk - VTE Prophylaxis

☐ No Risk - VTE Prophylaxis

March 2014	LOS > 1 day			
Service at time of	No Risk Assessed	VTE Risk Assessment - At Risk	VTE Risk Assessment - High Risk	VTE Risk Assessment - No Risk
H EPILEPSY	0	9	0	6
H MED H1 A	2	25	0	1
H MED H2 A	0	6	0	0
H MED H3 A	0	13	2	0
H SICU A	0	71	9	0
H SICU B	3	71	21	1
H SURGERY I	1	31	3	0
H SURGERY II	1	34	6	0
Grand Total	91	1045	145	18
% Distribution	7%	80%	11%	1%

HIGH RISK FOR VTE

Active malignancy
Prior VTE
Thrombophilia
Acute spinal cord injury
Trauma
Hip replacement
Knee replacement
Hip fracture repair
Burns

Ongoing Improvement – Enoxaparin Dosing

Prophylactic enoxaparin doses (6 months admissions)

30mg and 40mg; q12h and q24h

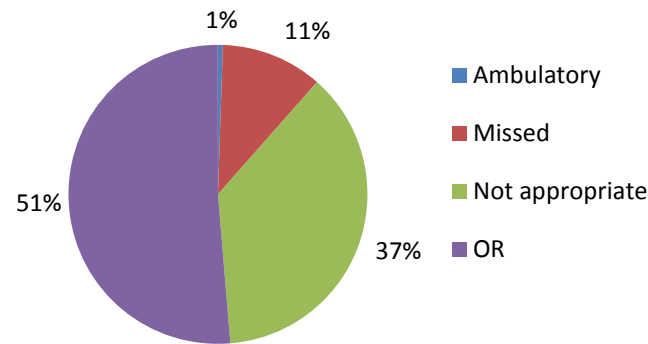
- Analysis to compare Risk Assessment to drug dosing by a review of on patient clinical factors/disease process
- 47% were accurately “Risk Assessed”
 - Reasons related to Trauma patients “at risk” (*high risk*)
 - Patients with history of/active cancer “at risk” (*high risk*)
- 85% received the correct dosing
 - Reasons related to BMI > 40 - given 30 mg dosing
 - CCRL > or < 30 given 30 vs. 40 mg dosing
 - Trauma patients given Q24 vs. Q12 dosing

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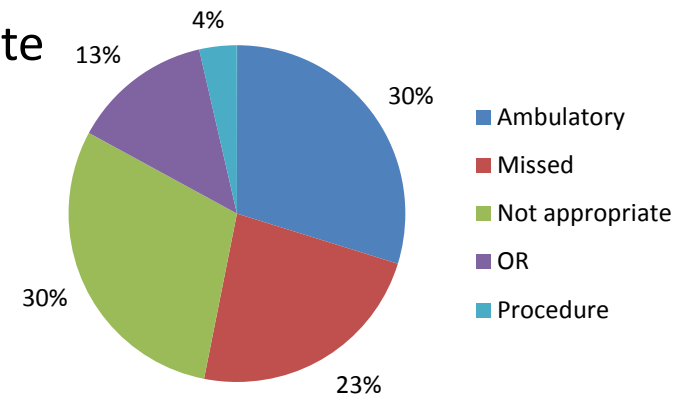
Ongoing Improvement – Held Doses

3 months of chemical prophylaxis reviewed

- Enoxaparin – 88% given – 12% Held
- Majority related to OR



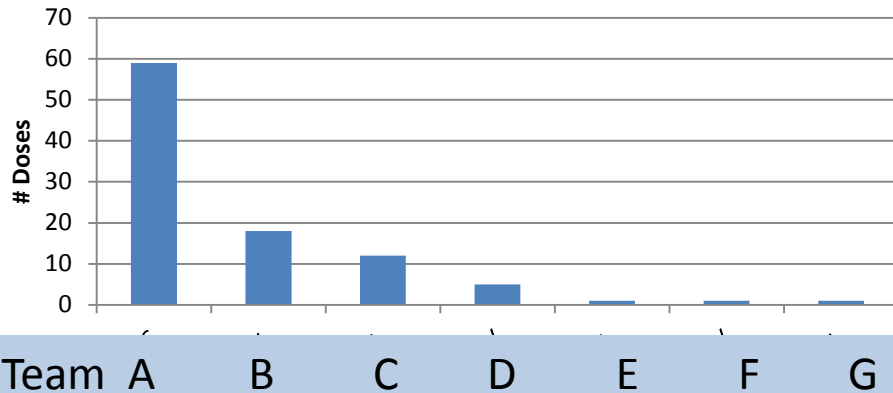
- Heparin – 85% given – 15% held
- Majority related to Ambulatory or Not appropriate



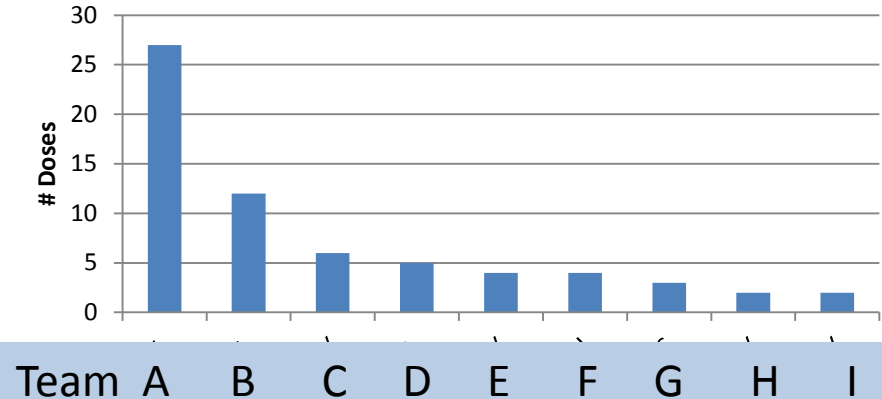
Held for OR by Service

Variation for Holding doses: May be appropriate for certain cases
Working to standardize practice across services

Enoxaparin



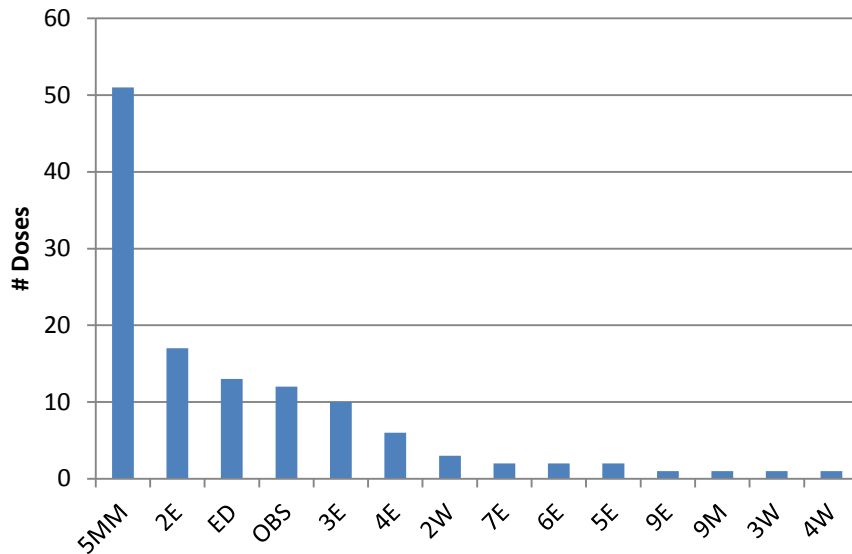
Heparin



Held Doses: “Ambulating” and “Refused” By Nursing Unit

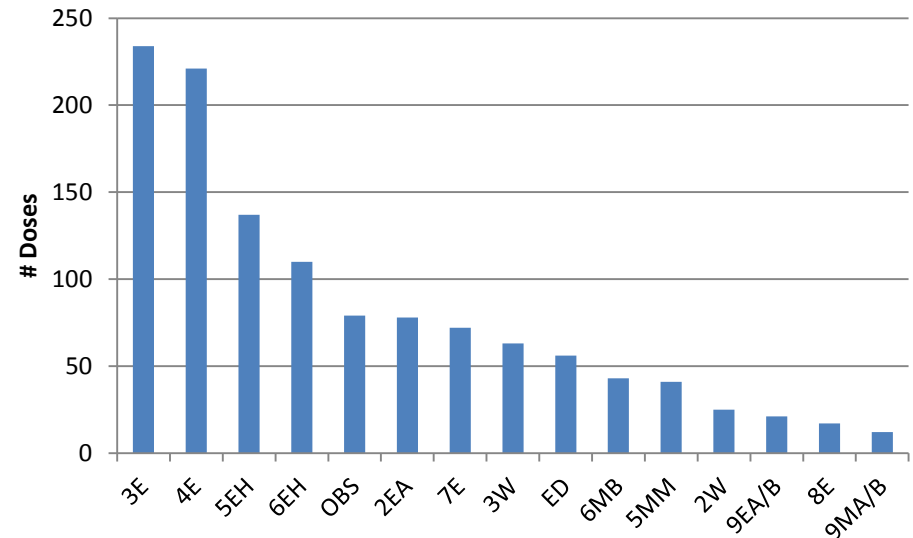
“Ambulating”

Heparin



“Patient Refused”

Heparin



Ongoing Improvement - SCDs

Service	Admit Dt	Admit Reason	SCD Order Reason (may be blank)	Order By	Last Chart	Chart By	Chart Dt
H SURGERY II	12/5/2013 1:23:00 PM	pleural effusion; other pneumothorax; ot...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started.		OFF Legs - Bilaterally		12/12/2013 12:30:00 AM
H SURGERY I	12/7/2013 5:10:00 PM	pancreas injury nos- clo; inj liv no op w...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started.		OFF Legs - Bilaterally		12/12/2013 12:15:00 AM
H THORACIC SURGERY	11/28/2013 2:33:00 PM	atrial fibrillation; occlu,sten carotid ...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started. Do not place on RLE!!!		OFF Legs - Bilaterally		12/12/2013 12:10:00 AM
H SURGERY II	12/7/2013 6:17:00 PM	swelling of limb; nb subarachnoid hemorr...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started.		OFF Legs - Bilaterally		12/11/2013 8:15:00 PM
H SURGERY II	12/7/2013 6:23:00 PM	traumatic subdural hem; other and unspec...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started.		OFF Legs - Bilaterally		12/11/2013 9:35:00 PM
H ORTHOPEDICS	12/3/2013 11:04:00 AM	joint pain-upper arm; pain in limb; fx l...	Contraindication to pharmacologic VTE prophylaxis. Continue until pharmacologic prophylaxis is started.		OFF Legs - Bilaterally		12/11/2013 8:16:00 PM

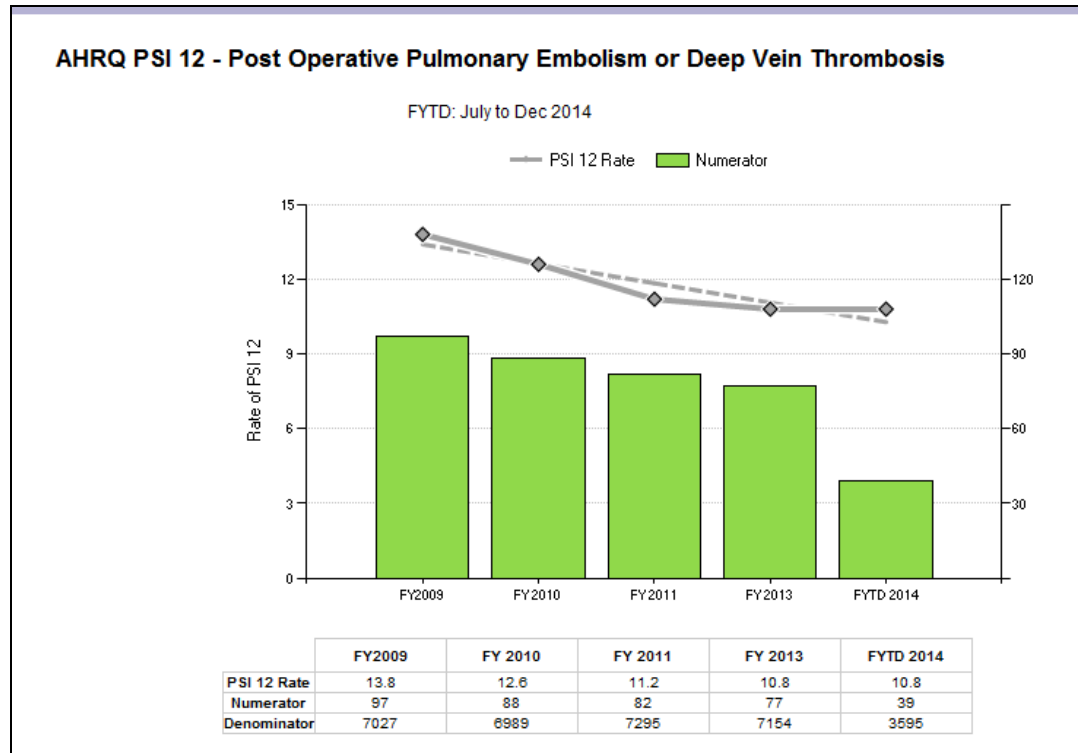
Daily review – 30 to 35 patients per day had SCD orders with the devices charted as “off” sometime in the previous 24 hours

Allows for real time education interventions with nursing staff

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Results: Rate of VTE – PSI 12



Focus on VTE prevention has resulted in a decrease in VTE events

VTE 6 -Incidence of Potentially Preventable VTE



Hospital Quality Measures Report

Welcome Ellen Robinson!

Wednesday, April 30, 2014

[Report Resources](#)

Harborview Medical Center

Jul - Sep 2013 (Q3)

VTE-6 Incidence of Potentially-Preventable Venous Thromboembolism %

Definition: The percentage of patients diagnosed with confirmed VTE during hospitalization (not POA) who did not receive VTE prophylaxis between hospital admission and the day before the VTE diagnostic testing order date.

Denominator: Patients who developed confirmed VTE during hospitalization. Excludes patients less than 18 years of age; who have a LOS greater than 120 days; with comfort measures only; enrolled in clinical trials; with a principal diagnosis code of VTE; with VTE POA; with reasons for not administering mechanical and pharmacologic prophylaxis; without VTE confirmed by diagnostic testing.

Numerator: Patients who received no VTE prophylaxis prior to the VTE diagnostic test order date.

Target: The Joint Commission target upper limit

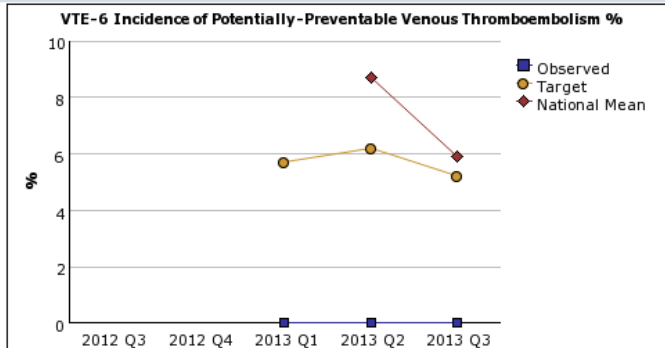
National Mean: The mean of all comparison group observed rates obtained from the Core Measure National Comparison Group File.

See Also: http://www.jointcommission.org/performance_measurement.aspx

Jul - Sep 2013 (Q3)

Oct 2012 - Sep 2013 (recent year)

	Relative Performance	Numerator (n)	Denom (n)	Observed	UHC Median-TJC Method	Target		
Current Quarter		0	24	0.0	0.0	5.2		
UHC Population								
	Value	%tile	10th	25th	50th	75th	90th	Mean
Cases (denom.)	24	79	2	4	11	21	30	15
Observed	0.0	0	0.0	0.0	0.0	9.8	22.2	7.5
Target	5.2							
National Mean	5.9							



	Relative Performance	Numerator (n)	Denom (n)	Observed	UHC Median-TJC Method	Target		
Recent Year		0	58	0.0	0.0	5.2		
UHC Population								
	Value	%tile	10th	25th	50th	75th	90th	Mean
Cases (denom.)	58.0	66	10	15	39	67	84	47
Observed	0.0	0	0.0	2.0	5.6	11.1	20.0	8.4
Target	5.2							

Recent Year UHC Top 10 in This Metric	Observed	Denom
The Ohio State University Wexner Medical Center	0.0	77
Mayo Clinic Hospital - Rochester	0.0	75
UC Irvine Medical Center	0.0	67
UMass Memorial Health Care	0.0	58
Harborview Medical Center	0.0	58
West Virginia University Hospitals, Inc.	0.0	53
Beaumont Hospital, Troy	0.0	31
Cedars-Sinai Medical Center	1.2	85
The Cleveland Clinic Foundation	1.8	168
Rush University Medical Center	1.8	56

* To qualify, facilities must meet the metric target and have an annual volume of at least 30 cases. Fewer than 10 facilities will be listed if less than 10 facilities meet these criteria.

Conclusion

- Systematic Processes have resulted in a decrease in VTE events
- Success of the project can be attributed to physician leadership, communicated expectations, and front line staff engagement
- Questions and contact:
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Q & A



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UPCOMING EVENTS

- Webinar: Antimicrobial Stewardship and *C. difficile* Infection
June 5, 2-3 pm ET



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VITAL2014, America's Essential Hospitals' annual conference, is coming to San Antonio! Plan now to join us Wednesday, June 25, through Friday, June 27, at the Westin Riverwalk for the premier national event for hospital and health system professionals. Together, we will support our shared mission of ensuring high-quality health care for vulnerable patients.

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THANK YOU FOR ATTENDING

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<http://essentialhospitals.org/groups/ehen/>

