Incidence of DVT in high risk patients at NYGH

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Introduction

Post operative venous thromboembolism (VTE) events

Deep Venous Thrombosis (DVT)
Pulmonary Embolism (PE)



 leading causes of morbidity and mortality in surgical patients

- Pulmonary embolism
 - may cause sudden death
 - may independently reduce survival for up to 3 months after diagnosis
 - Those who live may develop pulmonary hypertension

- Deep venous thrombosis result in
 - venous hypertension
 - lead to debilitating swelling and chronic pain

- Prevention of these events requires
 - diligent prophylaxis which must be considered for all surgical patients

Early recognition and treatment of VTE is crucial

- New Yangon General Hospital
 - 200 bedded hospital opened since 1984
 - Yet no standardized risk-stratified prophylaxis protocol for VTE till 2016

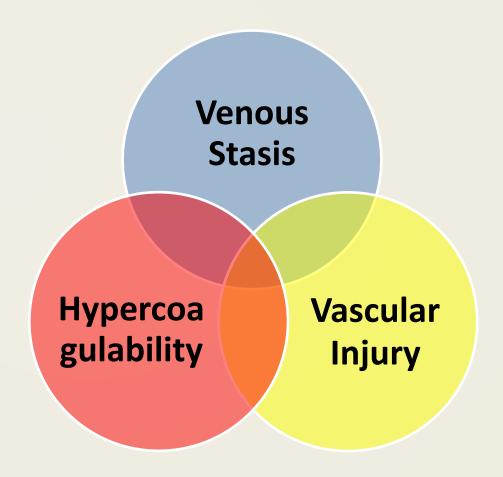
Plan to detect VTE incidence and reduce VTE complications

- In screening of VTE,
 - important to detect the risk factors for VTE in patients.

 a lot of risk factors assessment that link to development of VTE

Rudolph Virchow

• link the development of VTE to the presence of at least 1 of these 3 conditions:



- In 1992
 - the Thromboembolic Risk Factors (THRIFT)
 Consensus Group identified acquired risk factors for VTE

- Sixteen years later
 - the most recent update of the American College of Chest Physicians (ACCP) guidelines for VTE prophylaxis revealed the same risk factors for VTE

Selected acquired risk factors for VTE

ACCP 2008	THRIFT 1992
 Increasing age Immobility, paresis Previous VTE Cancer and/or its treatment Trauma (major or lower limb) Obesity Central venous catheters Inflammatory bowel disease Nephrotic syndrome Pregnancy and postpartum Estrogen therapy or estrogen containing oral contraceptive Acute medical illness 	 Increasing age Immobility (> 4 d), limb paralysis Previous VTE malignancy Surgery (pelvis, hips, legs) Trauma (pelvis, hips, legs) Obesity Varicose veins Heart failure Recent myocardial infarct Inflammatory bowel diseases Nephrotic syndrome Pregnancy High dose estrogen therapy infection

decided to use the Caprini risk stratification method

Caparini Risk Scoring Model

- adaptable to individual patient's risk factors
- less likely to underestimate the hazards of VTE
- Has been well validated
- The most up-to-date model

Methods

- study design
 - hospital based, prospective and interventional study
- Study duration
 - from October 2016 to September 2017 of one year duration

- Study population
 - all surgical patients admitted to NYGH
 - Total number of 2119 patients

Caprini risk scoring model

more than 30 risk factors in the Caprini model

categorized as

low (score 1-2) moderate (score 3-4) high (score 5 and more)

 All patients admitted to our hospital was assessed with Caprini score .

Deep Vein Thrombosis (DVT)

Prophylaxis Orders (For use in Elective General Surgery Patients)

Thrombosis Risk Factor Assessment (Choose all that apply)	CPI No. SEX M F VISIT No.
Age 41-60 years	Age 61-74 years Central venous access Arthroscopic surgery Major surgery (>45 minutes) Major surgery Major surgery (>45 minutes) Laparoscopic surgery (>45 minutes) Patient confined to bed (>72 hours) Immobilizing plaster cast (<1 month) Each Risk Factor Represents 3 Points Age 75 years or older Family History of thrombosis* History of DVTPE Positive Frothrombin 20210A Positive Factor V Leiden Positive Luous anticoaculant Elevated serum homocysteine Heparin-induced thrombocytopenia (HIT) (Do not use heparin or any low molecular weight heparin)
Stroke (<1 month)	□ Elevated anticardiolipin antibodies □ Other congenital or acquired thrombophilis if yes: Type • most frequently missed risk faotor TOTAL RISK FACTOR SCORE:
FACTORS ASSOCIATED WITH	INCREASED BLEEDING

BIRTHDATE

NAME

Active Bleed, Ingestion of Oral Anticoagulants, Administration of glycoprotein libitilis inhibitors, History of heparin induced thrombocytopenia GLINICAL CONSIDERATIONS FOR THE USE OF SEQUENTIAL COMPRESSION DEVICES (SCO)

Patients with Severe Peripheral Arterial Disease, CHF, Acute Superficial DVT

Total Risk Factor Score	Rick Level	Prophylaxic Regimen	
0	VERY LOW	☐ Early ambulation	
1-2	LOW	☐ Sequential Compression Device (SCD)	
3.4	MODERATE	Choose QNE of the following medications +/- compression devices: □ Sequential Compression Device (SCD) - Optional □ Heparin 5000 units 9Q TID □ Enoxaparin/Lovenox: □ 40mg SQ daily (WT < 150kg, CrCl > 30mL/min) □ 30mg SQ daily (WT < 150kg, CrCl = 10-29mL/min) □ 30mg SQ BID (WT > 150kg, CrCl > 30mL/min) (Please refer to Dosing Guidelines on the back of this form)	
5 or more	HIGH	(Please refer to Dosing Guidelines on the back of this form) Choose QNE of the following medications PLUS compression devices: □ Sequential Compression Device (SCD) □ Heparin 5000 units 9Q TID (Preferred with Epidurals) □ Enoxaparin/Lovenox (Preferred): □ 48mg 9Q daily (WT < 150kg, CrCl > 30mL/min) □ 30mg 9Q daily (WT < 150kg, CrCl = 10-29mL/min) □ 30mg 9Q BID (WT > 150kg, CrCl > 30mL/min) (Please refer to Dosing Guidelines on the back of this form)	

□ Ambulatory Surgery - No orders for venous thromboembolic prophylaxis required □ VTE Prophylaxis Contraindicated, Reason: VTE Real Actor Assessment To					Caprini, MD, MS, FACS, RVT actor Assessment Tool
Physician Signature Dr. #			Date	Time	
Processed By:	Date/Time:				
	White-Medical Record Yellow-MIS Pink-Pharma	ıcy N	University of Wichigan Health System	DVT Propi	hylaxis Regimen

UMHS ENOXAPARIN DOSING GUIDELINES

- o MUST wait 24 hours before starting Enoxaparin if patient has epidural catheter
- D/C Enoxaparin 10-12 hours prior to removing epidural catheter
- May restart Enoxaparin 24 hours after epidural catheter has been removed.

NON-PREGNANT PATIENTS

Body weight < 150kg, CrCl > 30mL/min: Enoxaparin 40mg \$Q daily Body weight < 150kg, CrCl = 10-29mL/min: Enoxaparin 30mg SQ daily Body weight > 150kg, CrCl > 30mL/min: Enoxaparin 30mg SQ BID

PREGNANT PATIENTS

Prevention of DVT:*

Maternal body weight (start of therapy) < 75 kg:

Recommend 30 mg SQ once dally until 20 weeks Recommend 30 mg SQ BID after 20 weeks

Maternal body weight (start of therapy) ≥ 75 kg:

Recommend 40 mg SQ once dally until 20 weeks

Recommend 40 mg SQ BID after 20 weeks

*Walt 12 hours before regional anesthesia

MONITORING RECOMMENDATIONS

- Patients who are obese (actual body weight > 150 kg)
- Patients who are pregnant
- Patients with renal insufficiency (creatinine clearance < 30 ml/min)

Indication	Desired Level (Draw 4 hours after the 4 th dose)	Recommendations for Dose Alteration		
		Anti-factor Xa Level (units/ml)	Dose Adjustment	Repeat Anti-factor Xa To Be Obtainted
		< 0.2	Increase by 25 %	4 hours after 4th dose
		0.2 to 0.5	No change	Repeat in 1 week, then monthly thereafter
		0.6 to 1	Decrease by 20 %	4 hours after 4 th dose
Prevention of DVT/PE	0.2 to 0.5 units/ml	>1	Hold for 3 hours, then decrease next dose by 30%	4 hours after 4th dose

Ideal Body Weight

IBW, men = 50 kg + 2.3 (Inches > 5 feet)

IBW, women = 45.5 kg + 2.3 (Inches > 5 feet)

Each risk factor represent one point					
	Age 41-60 years	For Wo	men only		
	Swollen legs (current)		Oral contraceptives o	r hormone	
	Varicose vein		replacement therapy		
	Obesity (BMI >25)		Pregnancy or postpar	tum (within 1	
	Minor surgery planned		month)		
	Sepsis (within 1 month)		History of unexplaine		
	Acute myocardial infarction		infant, recurrent spor		
	Congestive heart failure (within 1 month)		abortion (≥3), premat		
	Medical patient currently at bed		toxemia or growth re	stricted infant	
	History of inflammatory bowel disease				
	History of prior major surgery (within 1 month)				
	Abnormal pulmonary function (COPD)				
	Serious lung disease including pneumonia				
	(within 1 month)				
	Other risk factors				
			Subtotal		

Each risk factor represent 2 points					
	Age 61- 74years				
	Central venous access				
	☐ Major surgery (>45 minutes)				
	Malignancy (present or previous)				
	laparoscopic surgery (>45 minutes)				
	Patient confined to bed (> 72 hours)				
	Immobilization plaster cast (<1 month)	Subtotal			

Each risk factor represent 3 points				
 □ Age 75 or older □ Family history of thrombosis □ History of DVT/PE 				
☐ Congenital or acquired thrombophi	lia Subtotal			
Each risk factor represent 5 points				
☐ Stroke (within 1 month) ☐ Hip, pelvis or leg fracture (within 1	month)			
☐ Acute spinal cord injury (paralysis)				

 In this study, we only emphasized in high risk score patients for chemoprophylaxis

- For low and moderate risk score patients
 - encouraged early ambulation after post operative period

 no intermittent pneumatic compression device (for mechanical prophylaxis) in operation theatre

Chemoprophylaxis

all patients with Caprini high score (5 or more)

- low molecular weight heparin(Enoxaprin)
- subcutaneously once a day dose adjusted to patient's body weight

- Duration of prophylaxis
 - at least 5 post operative days or until patient can ambulate

started prophylaxis post operative 12 – 24 hours

- DVT was confirmed by Duplex Ultrasound.
- Any suspected leg swelling and pain in post operative period
 - checked by Duplex Ultrasound to detect DVT

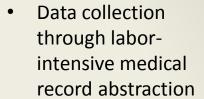
- Association between the incidence of DVT and background characteristics
 - calculated by Fisher's exact test
 - p value 0.05 was statistically significant

Results

Total admitted number **2119 patients**



High risk
(Caparini score 5 and more)
73 patients



incomplete and lost data



Drop out **5 patients**



Recorded for Chemoprophylaxis
68 patients

Background characteristics of high risk patients (N = 68)

Age	Numbers (%)		
<=40	7 (10.3)		
41-60	24 (35.3)		
>60	37 (54.4)		
Sex			
Male	37 (54.4)		
Female	31 (45.6)		
Pathology			
Benign	13 (19.1)		
Malignant	55 (80.9)		

Background characteristics of high risk patients continued

BMI	Numbers (%)		
<= 25	50 (73.5)		
> 25	18 (26.5)		
Operation			
Major	55 (80.9)		
Minor	4 (5.9)		
Observed	9 (13.2)		
Duration of Enoxaparin			
≤ 5 days	27 (39.7)		
> 5days	41 (60.3)		

Clinical characteristics of high risk patients

DVT	Numbers (%)		
Yes	7 (10.3)		
No	61 (89.7)		
Complication of Enoxa			
Yes	4 (5.9)		
No	64 (94.1)		
Viral infection			
В	4 (5.9)		
С	3 (4.4)		
Non	61 (89.7)		
Sepsis			
Yes	6 (8.8)		
No	62 (91.2)		

- overall incidence of DVT in the study period
 - 7 out of 2119 admitted patients i.e.; 0.3%

- The incidence of DVT in high risk patients
 - 7 out of 73 patients i.e.; **9%**

Association between occurrence of DVT and background characteristics

	Yes	No	P value (Fisher's exact test)
Age (years)			0.606
<=40	1 (14.3)	6 (85.7)	
41-60	3 (12.5)	21 (87.5)	
>60	3 (8.1)	34 (91.9)	
Sex			1.000
Male	4 (10.8)	33 (89.2)	
Female	3 (9.7)	28 (90.3)	
BMI (kg/m2)			1.000
≤25	5 (10.0)	45 (90.0)	
>25	2 (11.1)	16 (88.9)	
Pathology			0.598
Benign	1 (7.7)	12 (92.3)	
Malignant	6 (10.9)	49 (89.1)	

Association between occurrence of DVT and background characteristics

		Yes	No	P value (Fisher's exact test)
	Operation			0.023
	Major	3 (5.5)	52 (94.5)	
	Minor	1 (25.0)	3 (75.0)	
	Observed	3 (33.3)	6 (66.7)	
	Duration of Enoxa			0.105
	≤ 5days	5 (18.5)	22 (81.5)	
	>5days	2 (4.9)	39 (95.1)	
	Sepsis			0.507
	Yes	0 (0.0)	6 (100.0)	
	No	7 (11.3)	55 (88.7)	

Discussion

 Reviewing the hospital data analysis, there was no DVT patients in low and moderate risk patients

 data were collected only for in-patients not extended to follow-up period

- In this study period,
 - had not found documented PE despite there were cases of unexplained death.
 - But autopsy result did not show PE in unexplained death
- So there was no PE cases in this study

 We intended to strict adherence to risk stratification guideline in our hospital but...

- 5 patients drop-out data
- 8 high risk patients did not included in the prophylaxis regimen
- 4 patients who suffered DVT did not get DVT chemoprophylaxis properly
- 23% of high risk patients did not receive the recommended prophylaxis

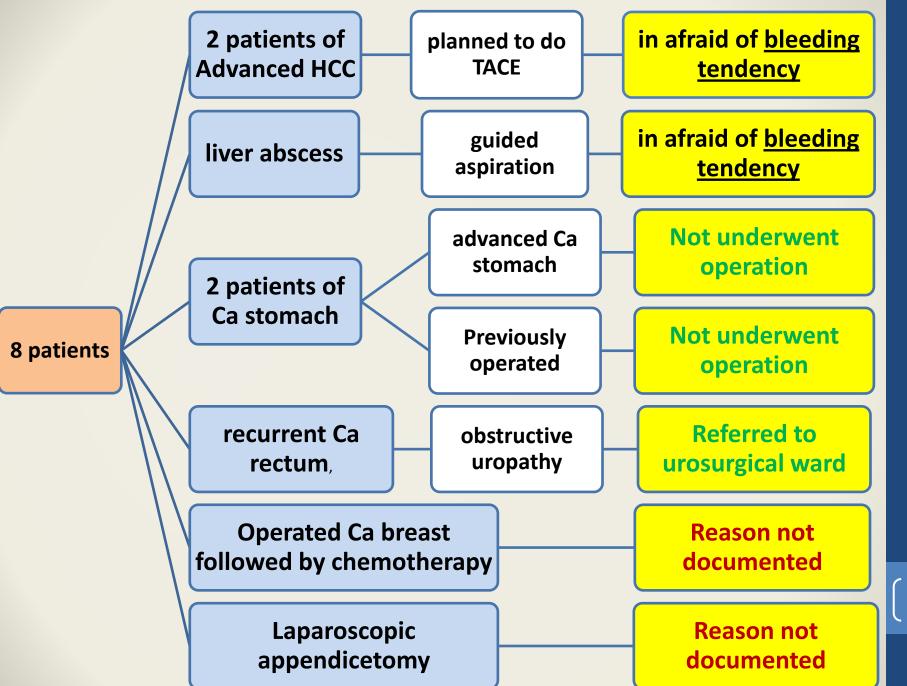
 This indicate that there was a leakage in our practice of DVT prophylaxis.

- In other studies
 - there is evidence that prophylaxis measures are often under used with at-risk patients receiving inappropriate or no prophylaxis.

 A large multinational study revealed that only 59% of surgical patients received evidence based VTE prophylaxis

- Based on Cassidy practice of prophylaxis,
 - among patients stratification to the high risk category,
 - 89% received appropriate pharmacologic prophylaxis and duration.
 - 10% of patients did not receive the recommended prophylaxis.

 He also advice to collect data through the electronic inpatient medical record system. Audit to 8 high risk patients who did not receive chemoprophylaxis



 Those 8 patients did not suffer DVT in their hospitalized period

6 DVT patients with malignant origin

Did not received DVT prophylasxis

4 patients

6 patients

received DVT prophylasxis

2 patient

Operated Ca stomach last 1 year ago presenting with intestinal obstruction

Operated recurrent Ca rectum last 6 months ago presenting with obstructive uropathy

Emergency appendicectomy

biopsy result of carcimatosis peritonei from ovarian tumor

Operated Ca rectum last 2 years ago with recurrent tumor

A case of retroperitoneal liposarcoma and debulking surgery

DVT on 8th POD

Multifocal HCC undergone liver section DVT on 5th POD

DVT patient with benign pathology

- History of previous DVT with bilateral hydrocele with right sided atrophic testis
- He underwent right orchidectomy with DVT prophylaxis.
- In the period of chemoprophylaxis, he encountered the complication of LMWH at post operative day 9th, bleeding from wound site.
- So we stopped chemoprophylaxis.

- After stopping of chemoprophylaxis, he noticed leg swelling and pain and confirmed DVT by Duplex scan.
- Then we gave therapeutic regime of LMWH followed by life long warfarin

After auditing the data and found that 17
 patients with high risk Caprini score were not get
 proper DVT prophylaxis.

Among them, 4 patients were suffered DVT;
 24%.

- Despite guideline from multiple sources for VTE prophylaxis regimes
 - PE and DVT remains significant problems among hospitalized patients in United States

 High risk patients tend to receive insufficient prophylaxis and low risk patients may be over treated

 Caprini suggest that the solution to this problem is standardize risk assessment and commensurate prophylaxis Complication of low molecular weighted heparin History of DVT with right orchidectomy for atrophic testis .

Complication at 9th
day of
chemoprophylaxis

4 patients

Retroperitoneal tumor patient underwent debulking surgery biopsy result of Non Hodgkin Lymphoma

Complication at 3rd day of chemoprophylaxis

Bleeding

from operated wounds

Ca rectum with liver secondary performed ARR, splenectomy, RFA

Complication at 6th day of chemoprophylaxis

Ca stomach, performed TG, OJ,JJ, splenectomy

Complication at 5th day of chemoprophylaxis

44

Conclusion

 first standardizes risk stratification prophylaxis protocol for DVT in the surgical ward of NYGH

 overall incidence of DVT patients during one year period is 0.3%

incidence of DVT in high risk patients is 9%

 So strict adherence of risk stratification and standardized prophylaxis guideline is very important

 to reduce the incidence of DVT formation in surgical patients.

- After auditing the prophylaxis protocol, 23% of high risk patients actually did not received proper chemoprophylaxis.
- Because of improper chemoprophylaxis, 4 patients suffered DVT; 24%.
- That shows there was a leakage in labour intensive medical record system.

 need to change our labor intensive data recording system to electronic prophylaxis recommendation system in future.

Limitations

- single center based study
- only detected the symptomatic DVT, not asymptomatic DVT
- 3. only used chemoprophylaxis, not used other mechanical prophylaxis for DVT
- 4. detected in general surgical patients, not included trauma, orthopedic, vascular patients.

Take home message

Once VTE occurs

21.5% of patients will have a recurrent VTE within 5 years

2.6% incidence of PE

Prevention is important in our patients.

Thank You