# Anticoagulation Safety: Reducing Adverse Drug Events

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#### Disclosure

I have nothing to disclose.



#### **Objectives**

- Review reasons for errors in relation to prescribing and administering anticoagulants
- Identify opportunities for improvement in the medication use process for safer anticoagulation use



#### **Statistics**

- Incidence of Disease Requiring Anticoagulation
  - Atrial fibrillation: 6.1 million Americans
  - Venous thromboembolism: 183 per 100,000 patient-years
- In 2016, over 11 million prescriptions for oral anticoagulants were filled in the US.
- Anticoagulants account for 17.6% of ED visits annually.



#### **Agents Used for Anticoagulation**

#### Parenteral Agents

Factor Xa Inhibitors	Direct Thrombin Inhibitors
Heparin	Heparin
Enoxaparin (Lovenox <sup>®</sup> )	Argatroban
Fondaparinux (Arixtra®)	Bivalirudin (Angiomax <sup>®</sup> )
Dalteparin (Fragmin <sup>®</sup> )	

Oral Agents

Vitamin K Antagonists	Factor Xa Inhibitors	Direct Thrombin Inhibitors
Warfarin (Coumadin <sup>®</sup> )	Rivaroxaban (Xarelto <sup>®</sup> )	Dabigatran (Pradaxa®)
	Apixaban (Eliquis®)	
	Edoxaban (Savaysa <sup>®</sup> )	

#### Question

"What's the usual dose for Xarelto®?"



#### **Complexities of Dosing Agents**

- Based on weight and renal function
  - Fondaparinux for VTE prophylaxis contraindicated if body weight < 50 kg or CrCl < 30 mL/min</li>
- Based on indication and renal function

Drug	Dose	Renal Function
Rivaroxaban for	20 mg with dinner	CrCl > 50 mL/min
atrial fibrillation	15 mg with dinner	CrCl 15-50 mL/min
Rivaroxaban for VTE treatment	15 mg BID x 21 days followed by 20 mg daily	Not recommended if CrCl < 30 mL/min



#### **Parenteral Anticoagulants**

- Potential for medication error in transcription, administration, monitoring and titration of continuous infusion agents
- Heparin
  - Selection of appropriate dosing nomogram
  - Drawing of blood samples for activated partial thromboplastin time (aPTT) or anti-Factor Xa levels
  - Dose adjustment based on interpretation of blood sample levels

Example	Titration	Nomogram
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PTT (sec)	BOLUS DOSE	STOP INFUSION	RATE CHANGE	REPEAT PTT
Less than 39 <sup>∓</sup>	Re-bolus with 5000 units IV	No	Increase infusion rate by 200 units/hr (2mL/hr)	6 hours
39 - 54	Re-bolus with 2500 units IV	No	Increase infusion rate by 100 units/hr (1mL/hr)	6 hours
55 - 75 Therapeutic Range	None	No	No Change	6 hours until therapeutic times 2 values then every 12 hours
76 - 100	None	No	Decrease infusion rate by 100 units/hr (1 mL/hr)	6 hours
101 - 115	None	Hold infusion for 30 minutes	Decrease infusion rate by 100 units/hr (1mL/hr)	6 hours after restarting infusion
116 - 139	None	Hold infusion for 1 hour	Decrease infusion rate by 200 units/hr (2mL/hr)	6 hours after restarting infusion
		Hold infusion and	*If PTT less than 39 re-bolus with 5000 units IV bolus and decrease rate by 200 units/hr (2 mL/hr)	
Greater than 139	None recheck PT hours	recheck PTT in 2 hours	*If PTT 39-54 re-bolus with 2500 units IV bolus and decrease rate by 200 units/hr (2 mL/hr)	6 hours after restarting infusion
		with PTT results	*If PTT therapeutic (55-75) decrease rate by 300 units/hr (3mL/hr)	
			* If PTT still greater than 75 contact prescriber	

 How would you adjust the rate if the patient's current PTT is 50 seconds?while the previous aPTT was 140 seconds?



### **Examples of Anticoagulation Medication Errors**



#### **Examples of Anticoagulation Errors**

- Transcription/Documentation
  - Reading handwriting
- Prescribing
  - Two prescribers entering orders for same patient
- Dispensing
  - Similar looking packaging
  - Mix-ups in medication names
- Administration
  - Titration of continuous infusion medications
  - Double dosing due to poor timing/scheduling



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#### **Classic Examples of High Risk** Errors

Combin 4~ Pis



#### The Joint Commission Sentinel Events

- From 1997-2007, 32 sentinel events reported related to anticoagulants
  - Heparin: 21
  - Warfarin: 6
  - Enoxaparin: 3
- Most events occurred in hospital settings
- 28/34 patients died



#### **The Joint Commission**

- National Patient Safety Goal (NPSG) 03.05.01
  - Reduce the likelihood of patient harm associated with the use of anticoagulation therapy
- Elements of Performance
  - Use approved protocols for initiation and maintenance of anticoagulation therapy
  - Before starting a patient on warfarin, assess the patient's baseline coagulation status and titrate warfarin based on INR.
  - Provide education to the patient and family.
  - Evaluate anticoagulation safe practices.



# Strategies for Reducing the Risk of Anticoagulation Errors



#### **ISMP Risk Assessment**

- Patient Information
- Drug Information
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#### **Patient Information**

- Readily available in the patient's medical record
  - Diagnosis
  - Allergies, height, weight
  - Laboratory values (PTT, INR, platelet count)
  - Concomitant diseases/conditions
  - History of heparin induced thrombocytopenia (HIT)
- Used to make monitor and manage the effects of anticoagulants
  - Goal INR for warfarin patients
  - Guidelines/protocols for unfractionated heparin use
  - Bridging parenteral and oral anticoagulation
  - Assessment of subtherapeutic INRs
  - 4T score if concerned about HIT

#### **Focus on Warfarin**

2. Baseline INR obtained

5. INR obtained prior to initiating warfarin therapy

- 6. INR on admission for patients on warfarin as an outpatient
- 7. Defined frequency for INR monitoring during hospitalization
- 9. Blood specimens for INR drawn at standard times each day
- 11-12. Access to inpatient and outpatient labs as necessary
- 19. Indication for anticoagulation is documented
- 20. Monitor INR levels to adjust warfarin dosing
- 24. Bridging for patients with active thrombosis
- 25. If discharged with subtherapeutic INR, close monitoring to determine when to discontinue parenteral therapy



#### Strategies for Implementation of "Patient Information" Recommendations

- Electronic medical records
  - Complete with demographic, laboratory and other information necessary for making appropriate decisions
- Policies, Guidelines and Dosing Protocols
  - Anticoagulation policy
  - Oral anticoagulant dosing support
  - Heparin infusion protocols
- Heparin Induced Thrombocytopenia
  - Alerts regarding platelet drops
  - HIT antibody testing
  - Argatroban infusion protocol

#### Warfarin in Computerized Physician Order Entry (CPOE) System

- Require all warfarin orders to be entered with indication and goal INR
- Allows all providers and pharmacists to be aware of goal for patient and evaluate daily INRs appropriately

Indication:	Goal INR:
1 ATRIAL FIBRILLATION 2 DVT/PE TREATMENT 3 CARDIAC THROMBUS 4 HYPERCOAGULABLE STATE 5 LVAD ANTICOAGULATION 6 MECHANICAL HEART VALVE	1         INR BETWEEN 2 AND 3           2         INR BETWEEN 2.5 TO 3.5           3         INR BETWEEN 2 TO 2.5           4         INR BETWEEN 2.5 TO 3           5         INR BETWEEN 1.8 TO 2.2 (LVAD)



#### **Direct Oral Anticoagulants in CPOE**

- Currently listed apixaban, dabigatran and rivaroxaban with indications and indication-specific dosing
- Default doses and frequencies
  - 15 mg vs. 20 mg
  - With meals vs. standard administration times
  - Daily vs. BID

rivaroxaban [XARELTO]

Please select an indication:

- 1. non-valvular atrial fibrillation
- 2. venous thromboembolism treatment initial 3 weeks
- 3. venous thromboembolism treatment beyond 3 weeks
- 4. orthosurgery vte prophylaxis

rivaroxaban [ XARELTO ] Weight=5 KG on Tue Jun 05 14:05 Information: 20mg Daily if Crcl>50mL/min (Cockroft-Gault formula); 15mg Daily if Crcl 15-50mL/min; Contraindicated if Crcl<15mL/min, on dialysis, or moderate to severe hepatic impairment (Child-Pugh Class B or C).



#### **Future State**

Enhanced decision support for prescribers

rivaroxaban (XARELTO) tablet Atrial Fibrillation

rivaroxaban (XARELTO) tablet VTE Prophylaxis Post Hip Replacement

rivaroxaban (XARELTO) tablet VTE Prophylaxis Post Knee Replacement

rivaroxaban (XARELTO) tablet VTE Treatment Beyond 3 Weeks

rivaroxaban (XARELTO) tablet VTE Treatment Initial 3 Weeks

#### **Future State**

Key is to ensure that "it's easy to do the right thing"



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#### **Drug Information**

- Essential patient information used to monitor effects and adjust therapy
  - Disease-specific guidelines for antithrombotic therapy
  - Reviewed by practitioners prior to use
  - Reviewed every 3 years
  - Guidelines for periprocedural management (cessation, resumption) and management when using neuraxial analgesia
  - Weight-based protocol for unfractionated heparin
  - Alerts for practitioners regarding serious drug interactions
- Essential drug information readily available to guide management of adverse reactions
  - Protocols and guidelines for managing supratherapeutic INRs, life-threatening bleeding
  - Management of HIT



#### Strategies for Implementation of "Drug Information" Recommendations

- Antithrombotic Reversal Guidelines
- Periprocedural Management of Antithrombotic Therapy Guidelines
- Heparin and Heparin Induced Thrombocytopenia (HIT) Order Sets

#### **Antithrombotic Reversal Guidelines**

Oral Direct Thrombin Inhibitor

Dabigatran

(Pradaxa®) Half-life CrCl > 80: 14-17 hours CrCl 50-79: 16.6 hours CrCl 30-49: 18.7 hours CrCl < 30: 27.5 hours



#### **Continuous Anticoagulant Infusion Order Set**

- Key Points to Cover
  - Indications
  - Dosing
  - Titration
  - Laboratory testing
- Key to ensure that all parties who will use the order set find it user-friendly
  - Prescribers for ordering
  - Nursing for titrating



#### **Heparin Order Set**

- **Baseline** labs
- Duplicate anticoagulants
- Patient weight
- Clear indication for dosing and boluses
- Titration algorithm

ATRIAL FIBR	ILLATION/ACS HEPARIN TRI ORDERS - ADULT	EATMENT		
- DATE & TIME ALL ORDERS -				
Drug Allergies:				
1. STAT CBC & PTT	prior to initiating heparin			
2. Daily CBC x 14 da	ays			
<ol> <li>Orders for the fol dabigatran, edox</li> </ol>	lowing anticoagulants should be aban, enoxaparin, fondaparinux, h	discontinued by o neparin (subcutar	ordering prescriber: apixaban, argat eous), rivaroxaban.	roban, bivalirudin,
4. wirnen Actual Bo	dy Weight for Bolus and Infusion	Dosing <u>(must co</u>	mplete):kg	
5. Initial Heparin	Bolus (Choose one of the follow	ing orders):		
Heparin Bolus	60 units/kg IV push (maximum i	nitial bolus of 4,0	00 units - refer to table)	
□ NO INITIAL E	BOLUS			
🗆 Alternate initia	I bolus dose of units/I	kg IV push x	kg = units (round	to nearest 500 units)
<ol> <li><u>Heparin Infusio</u> Initial Heparin Inf 12 units/kg/hr Alternate initial</li> </ol>	Heparin Infusion (standard concentration 25,000 units in 250 mL) Initial Heparin Infusion Rate (Choose <u>one</u> of the following orders): 12 units/kg/hr (maximum initial infusion rate of 1,000 units/hr - refer to table) Atemate initial infusion of units/kg/hr second to perfect to units/hr - refer to table)			
7. Subsequent He	parin Bolus			· · · · ·
INO BOLUSES	should be given during heparin ti	tration		
8. PTT 6 hours after	heparin infusion is started.			
9. Heparin dosing to	be titrated according to PTT as	per Protocol belo	w:	
PTT (seconds)	BOLUS DOSE	STOP INFUSION	RATE CHANGE	REPEAT PTT
Less than 39 *	If ordered, re-bolus 5,000 units IV	No	Increase rate by 200 units/hr (2 mL/hr)	6 hours
39 - 54	If ordered, re-bolus 2,500 units IV	No	Increase rate by 100 units/hr (1 mL/hr)	6 hours
55 - 75 Therapeutic Range *	None	No	No change	6 hours until therapeutic for two values, then every 12 hours
76 - 100	None	No	Decrease infusion rate by 100 units/hr (1 mL/hr)	6 hours
101 - 115	None	Hold infusion for 30 minutes	Decrease infusion rate by 200 units/hr (2 mL/hr)	6 hours after restarting infusion
116 - 139	None	Hold infusion for 1 hour	Decrease infusion rate by 200 units/hr (2 mL/hr)	6 hours after restarting infusion
140 - 180	None	Hold infusion for 1 hour	Decrease infusion rate by 300 units/hr (3 mL/hr)	6 hours after restarting infusion
Greater than 180:				
Appropriately timed and collected sample	None	HOLD heparin infusion. Repeat PTT hourly until less than 101	When PTT less than 101, decrease rate by 400 units/hr (4 mL/hr). Call MD if two consecutive PTTs are greater than 180	6 hours after restarting infusion

If suspected improperly timed sample, or potentially contaminated sample, repeat PTT

DOCTOR'S ORDER

If patient's PTT is less than 39 seconds for 12 hours or greater, call prescriber. PTT levels between 57 and 74 correspond to anti-Xa levels between 0.2 and 0.4 units/ml.



#### Argatroban Order Set

#### • Step 1: Diagnosis

Scoring System for the Pretest Probability for the presence of HIT A score (from 0 to 2) is determined for each of the below categories, resulting in the potential	Individual Point Score	Section Points
1. Thrombocytopenia		
Platelet count decrease greater than 50% and nadir greater than or equal to 20.000	2	
Platelet count decrease 30-50% or nadir between 10.000-19.000	1	
Platelet count decrease less than 30% or nadir less than 10,000	0	
2. Timing of platelet count fall		
Clear onset between 5 to 10 days <b>or</b> platelet count fall less than or equal to 1 day if prior heparin exposure within the last 30 days	2	
Consistent with fall at 5 to 10 days but not clear (e.g. missing platelet counts) <b>or</b> onset after day 10 or fall less than or equal to 1 day with prior heparin exposure within the last 30-100 days	1	
Platelet count fall at less than 4 days without recent exposure	0	
3. Thrombosis or other sequelae		
Confirmed new thrombosis, skin necrosis, <b>or</b> acute systemic reaction post-IV unfractionated heparin bolus	2	
Progressive or recurrent thrombosis, non-necrotizing (erythematous) skin lesions, or suspected thrombosis which has not been proven	1	
None	0	
4. Other causes for thrombocytopenia present		
None apparent	2	
Possible:	1	
Definite:	0	
	Total Score	
	(1+2+3+4)	
	Total Score	Pretest Probability
	0-3	Low
	4-5	Intermediate
	6-8	High



#### **Argatroban Order Set (cont.)**

Step 2: Determine next steps based on 4T score

Diagnosis of HIT based on laboratory results				
DTI=Direct Thrombin Ir	DTI=Direct Thrombin Inhibitor LMWH=Low Molecular Weight Heparin SRA=Serotonin Release Assay			
Pretest Probability	Heparin Induced Platelet Antibody			
,,	Positive (Greater than or equal to 0.4 optical density)	Negative (Less than 0.4 optical density)		
High	HIT confirmed	HIT possible		
Stop all heparin and LMWH products	Continue DTI	<ul> <li>Send SRA for confirmation</li> </ul>		
Start DTI (see orders page 2)		<ul> <li>Consider other causes of thrombocytopenia</li> </ul>		
Intermediate	HIT possible	<ul> <li>Use clinical judgement regarding continued</li> </ul>		
Stop all heparin and LMWH products	Continue DTI	use of heparin and LMWH products		
Start DTI (see orders page 2)	<ul> <li>Send SRA for confirmation</li> </ul>	<ul> <li>Use of DTI specific to patient risk vs. benefit</li> </ul>		
Low	HIT possible	HIT unlikely		
	<ul> <li>Send SRA for confirmation</li> </ul>	<ul> <li>Consider restarting / continuation of heparin</li> </ul>		
	<ul> <li>Consider other causes of thrombocytopenia</li> </ul>	or LMWH if necessary		
	<ul> <li>Use clinical judgement regarding continued use of heparin and LMWH products</li> <li>Use of DTI specific to patient risk vs. benefit</li> </ul>	<ul> <li>Use of DTI probably not necessary</li> </ul>		

#### Argatroban Order Set (cont.)

#### Step 3: Dosing of Argatroban

DISCONTINUE ALL HEPARIN PRODUCTS (including flushes and LMWH products) ADD heparin to patient allergy list
Obtain specimen: ☐ Heparin Induced Platelet Antibody (ELISA) ☐ Serotonin Release Assay (SRA) (if indicated, refer to table on Page 1)
Indication for direct thrombin inhibitor use <ul> <li>Anticoagulation for prophylaxis/treatment of thrombosis in patients with suspected/documented HIT</li> <li>Anticoagulation for prophylaxis/treatment of thrombosis in patients refractory or allergic to heparin</li> </ul>
Baseline CBC, Basic Metabolic Panel, LFTs, PTT, INR
Daily CBC, Basic Metabolic Panel, PTT, INR while patient on argatroban or bivalirudin therapy
Patient weight: kg (calculations are based on total body weight)
Order 🗆 STAT 📋 Begin at (date and time)
<ul> <li>□ BIVALIRUDIN 250 mg/250 mL D5W</li> <li>□ Standard dose: mg/kg/hr (normal renal function, recommended dose is 0.15 mg/kg/hr)</li> <li>□ Adjusted dose: Renal impairment (*CrCl less than 60 mL/min)</li> <li>□ CrCl 30-59 mL/min: 0.08 mg/kg/hr</li> <li>□ CrCl less than 30 mL/min: 0.05 mg/kg/hr</li> <li>□ CVVH: 0.05 mg/kg/hr</li> <li>□ Hemodialysis: 0.02 mg/kg/hr</li> </ul>
*CrCl = [(140-age) x body weight (kg) / (72 x SCr)] [x 0.85 for women]
<ul> <li>ARGATROBAN 125 mg/125 mL D5W</li> <li>Standard dose mcg/kg/min (recommended dose is 1 mcg/kg/min, max start dose is 2 mcg/kg/min)</li> <li>Adjusted dose: mcg/kg/min (recommended dose is 0.5 mcg/kg/min in patients with moderate hepatic failure (Child Pugh Score of 6 or greater; refer to table on reverse page for calculations), heart failure, multiple system organ failure/critically ill, severe anasarca, the early post cardiac surgery period)</li> </ul>



#### **Transitioning to Warfarin**





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## **Communication of Drug Orders and Other Drug Information**

- Computer order entry system
  - Interfaced with laboratory system
  - Alerts regarding abnormal laboratory values
  - Duplicate anticoagulation order alerts
- Dosing Standardization
  - Ordering and charting of heparin boluses
  - Rounding of doses



#### Strategies for Implementation of "Communication of Drug Orders and Other Drug Information"

Enoxaparin Dose (mg)	Round To (mg)	Administer (mL)
75-84	80	0.8
85-94	90	0.9
95-104	100	1
105	105	0.7
106-111	111	0.74
112-114	114	0.76
114-124	120	0.8
125-129	129	0.86
130-135	135	0.9
136-141	141	0.94



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#### Drug Storage, Stock, Standardization and Distribution

- Standard concentrations for unfractionated heparin
- Premixed solutions used whenever possible
  - Intravenous heparin
  - Glycoprotein IIb/IIIa inhibitors
- Thrombolytic therapy prepared by the pharmacy or trained practitioners
- Standard administration times for antithrombotics
- Ready access to reversal agents or antidotes



#### Strategies for Implementation of "Drug Storage, Stock, Standardization and Distribution"







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# Medication Device Acquisition, Use and Monitoring

- Laboratory reagents used to test aPTT
  - Re-establishment of therapeutic range required when reagents are changed
  - Updates to protocols must occur simultaneously
- Use of smart infusion pumps
  - Ensure accuracy of pump libraries
  - Annual updates and testing
  - Minimal deviation allowed to minimize risk of errors
  - Involvement of multidisciplinary team



### Impact of a Pharmacist on Anticoagulation Safety



#### **Clinical Pharmacist Anticoagulation Services**

- Pharmacy-driven warfarin dosing
- Pharmacist monitoring of argatroban
  - Appropriate prescribing
  - Appropriate dose titration
- Automatic renal dose adjustment policies



#### **Inpatient Warfarin Management**

- Study conducted by the Clinical Pharmacist Anticoagulation Service (CPAS) at Kaiser Permanente
- Evaluated the impact of 24/7 warfarin dosing by pharmacists in an inpatient setting
- Included over 6,000 patients in final analysis
- Results
  - Pharmacist-dosing led to significantly increased percent of time in therapeutic range and less time outside of therapeutic range
  - Reduced risk of hemorrhagic and thromboembolic complications



#### **Inpatient Argatroban Management**

- Study conducted by pharmacists at the Medical University of South Carolina
- Evaluated the impact of pharmacist-driven dosing and titration of argatroban infusions
- Final analysis included 50 patients
- Results
  - Therapeutic PTT values were achieved significantly more quickly in the pharmacist arm
  - Lower rate of medication errors
  - No difference in adverse events



#### Conclusion

- Anticoagulants are high risk medication with a narrow therapeutic window that require appropriate dosing and monitoring
- Requires the involvement of all healthcare professionals to ensure appropriate use
- Strategies to minimize ADRs include multidisciplinary involvement in patient care, identification of weakness in the medication use process and optimization of electronic methods.



## Anticoagulation Safety: Reducing Adverse Drug Events

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