



Government of **Western Australia**
Department of **Health**

WA Adult Anticoagulation Medication Chart

For Chart Version 05/22

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Overview

This presentation will provide an overview of:

- The layout of the WA Anticoagulation Medication Chart (WA AMC)
- The management of anticoagulants using the chart:
 - Low Molecular Weight Heparins (i.e. enoxaparin)
 - Unfractionated heparin (UFH)
 - Warfarin
 - Direct oral anticoagulants (DOACs)

Anticoagulants – High Risk Medications

- Anticoagulants are consistently identified as causing preventable harm to patients.
- Top 10 medication categories involved in confirmed medication-related clinical incidents (July 2021 - June 2022)

1. Opioid analgesics
2. Antimicrobials
3. Insulins
4. Anticoagulants
5. Antipsychotics
6. Vaccines
7. Antihypertensives
8. Non-opioid analgesics
9. Medication for anxiety and sleep disorders
10. Antiepileptics

- When used in error or omitted, they can cause life-threatening or fatal bleeding or thrombosis.

Those most commonly prescribed anticoagulants are:

- unfractionated heparin
- low-molecular weight heparin (LMWH)
 - enoxaparin sodium (Clexane[®])
 - dalteparin sodium (Fragmin[®]) and
- warfarin.


Direct oral anticoagulants are also available and are being prescribed more frequently:

- dabigatran (Pradaxa[®])
- rivaroxaban (Xarelto[®])
- apixaban (Eliquis[®]).

Factors that increase the potential for error and harm include:

- Low margin for error
 - over-dose → bleeding
 - under-dose or omission → thrombosis
- Wide variation in individual patient response
 - multiple indications
 - wide range and complexity of dosage
 - frequent dose adjustment/monitoring
 - interaction with other medicines, herbals, over-the-counter products, food and alcohol.

Benefits of the WA Anticoagulant Medication Chart

- Provides one chart for all anticoagulant prescriptions to reduce the risk of duplicate prescribing.
 - Point of care guidelines for initiation, monitoring and reversal of anticoagulants.
 - Enables the effective achievement of therapeutic levels.
 - Minimise the risk of bleeding events due to supra-therapeutic levels.
 - To achieve this the chart includes:
 - Optimal dosing guidelines and monitoring requirements
 - important information required for dosing including test results, weight and renal function
- 

Importance of Cross-Referencing Anticoagulant Chart with WA HMC

- The main WA Hospital medication chart (WA HMC) **MUST** be annotated (cross-referenced) to identify when the anticoagulation chart is in use to reduce the risk of duplicated orders or dose omissions.

Front of WA HMC →

Hospital name.....
 Hospital Provider number.....
 Ward..... Team.....

Medication chart number of

Additional charts Variable dose
 IV fluid BGL/insulin Acute pain Other
 Palliative care Chemotherapy Anticoagulation

Inside WA HMC →

Venous Thromboembolism (VTE) risk assessment / Anticoagulation		Risk Assessment completed by (name)	Date/Time	Continue Y / N
<input type="checkbox"/> VTE risk considered (refer guidelines)	<input type="checkbox"/> Bleeding risk considered			
Pharmacological Prophylaxis: <input type="checkbox"/> Indicated* <input type="checkbox"/> Not Indicated <input type="checkbox"/> Contraindicated <small>*Consider surgical and anaesthetic implications prior to prescribing</small>				
Mechanical Prophylaxis: <input type="checkbox"/> GCS <input type="checkbox"/> IPC <input type="checkbox"/> VFP <input type="checkbox"/> Not Indicated <input type="checkbox"/> Contraindicated				
Key: GCS – Graduated Compression Stockings; IPC – Intermittent Pneumatic Compression; VFP – Venous Foot Pumps		If risk changes document VTE prophylaxis requirements on new chart		

**Warfarin/
Anticoagulant
in use**
Refer to
Anticoagulation Chart for
administration details

The front page

Facility/Service: **XXX**

Ward/Unit: _____

Consultant: _____

WA Anticoagulation Chart

ATTACH PATIENT IDENTIFICATION LABEL HERE AND OVERLEAF

URMN: _____
 Family Name: _____
 Given Name: _____
 Address: _____
 DOB: _____ Gender: M F

Attach ADR Sticker

Patient weight _____ kg Date weighed ____/____/____
 Height _____ cm

Bleeding Risk considered before prescribing anticoagulants _____ assessed by (prescriber) _____ Date: ____/____/____

ONCE ONLY AND TELEPHONE (Prescriber to sign with 24 hours of order)

Date prescribed	Medicine (print generic name)	Route	Dose	Date/Time of dose	Nurse	Prescriber	Given by	Time Given
					N1	N2	Sign	Print Name

REGULAR DOSE ORDERS - PROPHYLACTIC DOSES Check platelets and coagulation profile before commencing (Subcutaneous unfractionated heparin and direct oral anticoagulants - DOACs)

YEAR 20____ DAY AND MONTH →

Date	Medicine (Print generic name)	Route	Dose AND Frequency (NDI) enter times →	Pharmacy	Dispense	Compliance	Request	Dispense

REGULAR DOSE ORDERS - THERAPEUTIC DOSES Check platelets and coagulation profile before commencing (Subcutaneous low molecular weight heparins and direct oral anticoagulants - DOACs)

YEAR 20____ DAY AND MONTH →

Date	Medicine (Print generic name)	Route	Dose AND Frequency (NDI) enter times →	Pharmacy	Dispense	Compliance	Request	Dispense

WARFARIN VARIABLE DOSE ORDERS

YEAR 20____ DAY AND MONTH →

Date	Medicine	Route	Dose Time	INR Result	Prescriber	Telephone order N1/N2	Given by

Warfarin Discharge Plan Dose _____ mg Target INR _____ Duration _____ next INR due ____/____/____ Prescriber _____

ANTICOAGULANT DISCHARGE PLANNING

Warfarin DOAC _____ LMWH _____ Patient has booklet Patient education completed

Signature: _____ Designation: _____ Date: _____

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- Bleeding risk considered
- Once only and telephone
- Regular dose prophylactic doses
- Regular dose orders Treatment doses
- Variable dose orders - warfarin

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The back page

Treatment recommendations do not cover all clinical scenarios and do not replace the need for clinical judgement					
RECOMMENDATIONS FOR DIRECT ORAL ANTICOAGULANTS					
Direct Oral Anticoagulant Agents (DOACs) – Apixaban, Dabigatran, Rivaroxaban (also known as NOACs) • Prescribe with care in elderly (>75 years), underweight (<50kg), overweight (>150kg) and patients with renal impairment (CrCl < 50mL/min). • Prior to DOAC initiation: Record: FBC, Coagulation status (INR, aPTT and PT), renal and liver function. Check for drug interactions prior to prescribing. • If the patient is on warfarin: Discontinue warfarin and start DOAC when INR is 2.0 or less • Refer to local prescribing guidelines for further information.					
Apixaban (Eliquis®)	Dabigatran (Pradaxa®) Idarucizumab is the reversal agent for dabigatran Refer to local hospital guidelines.	Rivaroxaban (Xarelto®) (Use with caution if CrCl 15-29mL/min)			
Treatment of DVT/PE: • CrCl >25 mL/min: 10mg twice daily for first 7 days, then 5mg twice daily thereafter		Treatment and Prevention of DVT/PE: • CrCl ≥ 15 mL/min: 15mg twice daily for 3 weeks, then 20mg once daily • Seek specialist advice if CrCl 15-29mL/min			
Non-Valvular Atrial Fibrillation (therapeutic dose): 5mg twice daily Reduce to 2.5mg twice daily IF at least 2 of the following risks: <input type="checkbox"/> SCr ≥ 133 micromol/L <input type="checkbox"/> Age ≥ 80 years, <input type="checkbox"/> Weight ≤ 60 kg	Non-Valvular Atrial Fibrillation (therapeutic dose): • CrCl ≥ 50 mL/min: 150mg twice daily • CrCl 30-49 mL/min or ≥ 75years: 110mg twice daily	Non-Valvular Atrial Fibrillation (therapeutic dose): • CrCl ≥ 50 mL/min: 20mg once daily • CrCl 30-49 mL/min: 15mg once daily • CrCl 15-29 mL/min: seek specialist advice			
VTE prophylaxis: Total Hip or Knee Replacement • CrCl > 25mL/min: 2.5mg twice daily Hip: up to 38 days Knee: up to 14 days	VTE prophylaxis: Total Hip or Knee Replacement • CrCl > 50 mL/min: 220mg (2 x 110 mg) once daily • CrCl 30-50 mL/min: 150mg (2 x 75 mg) once daily Hip: up to 35 days Knee: up to 10 days	VTE prophylaxis: Total Hip or Knee Replacement • CrCl ≥ 15 mL/min: 10mg once daily Hip: up to 35 days Knee: up to 14 days			
RECOMMENDATIONS FOR WARFARIN Warfarin brands are NOT equivalent and cannot be used interchangeably.					
TARGET INR RANGE					
2.0-3.0	• Therapy for DVT or PE • Preventing DVT: high risk patients e.g. hip or knee surgery • Preventing systemic embolism: AF valvular heart disease, post MI, bioprosthetic heart valves (first 3 months)				
2.0-3.0	• Aortic bileaflet mechanical heart valve – if no other risk factors				
2.5-3.5	• Starr-Edwards mechanical heart valves. Mitral bileaflet mechanical heart valve or aortic if risk factors for thromboembolic event including AF, previous thromboembolism, LV dysfunction, hypercoagulable condition.				
(ADULT) DOSING FOR WARFARIN NAÏVE PATIENTS (TARGET INR 2-3)		DOSING WITH ONGOING WARFARIN THERAPY			
Consider if bridging with heparin is indicated. Refer to WATAG or local warfarin guidelines for further information. Record baseline FBC, coagulation status (INR, aPTT and PT) and liver function. • Suggested initial dosing of 5mg daily for first 2 days, modify dosing for day 3 based on day 3 INR. • For younger patients (< 60 years) consider 7-10mg on day 1 and day 2. • Consider smaller starting doses when the patient is elderly, has low body weight or abnormal liver function, is at high bleeding risk or has severe chronic renal impairment. • Consider dose modification in the presence of interacting drugs. • Discontinue heparin after a minimum of 5 days therapy and INR is 2.0 or greater.		• Patients being re-initiated on warfarin post surgery/ intervention should be restarted on the dose prescribed prior to intervention and check INR day 3. • In acutely ill patients with ongoing warfarin therapy: daily monitoring of INR may be appropriate. • Monitor INR more frequently when any change in treatment involves drugs known to interact with warfarin.			
REVERSING WARFARIN OVER-TREATMENT (bleeding risk increases exponentially as INR rises above 3. Monitor closely INR ≥ 6)					
Clinical Setting		Management			
INR	Bleeding	Warfarin	Vitamin K (seek advice if cardiac valve replacement)	Prothrombinex VF	Comments
Greater than therapeutic range but <4.5	Absent	Reduce dose or omit next dose			Resume warfarin at reduced dose when INR approaches therapeutic range. If INR <10% above therapeutic level, dose reduction may not be necessary.
4.5 – 10	Absent (Low risk)	Stop			Measure INR in 24 hours. Resume warfarin at reduced dose when INR approaches the therapeutic range.
	Absent (High Risk)*	Stop	Consider 1–2 mg (oral) ¹ Or 0.5–1mg IV ²		Measure INR within 24 hours. Resume warfarin at reduced dose when INR approaches the therapeutic range.
>10	Absent (Low risk)	Stop	3–5mg (oral) ¹ Or IV ²		Measure INR in 12-24 hours. Resume warfarin at reduced dose when INR approaches the therapeutic range.
	Absent (High Risk)*	Stop	3–5mg IV ²	Consider 15-30 Units/kg ^{3,4} See weight based nomogram	Measure INR in 12-24 hours. Resume warfarin at reduced dose when INR approaches the therapeutic range. Close monitoring over the following week.
Clinically significant bleeding where warfarin is a contributing factor. e.g. Intracranial or massive haemorrhage		Stop	5–10 mg (IV) ²	25–50 Units/kg ^{3,4} doses may be appropriate as per warfarin reversal guidelines. See weight based nomogram	Only add Fresh Frozen Plasma (FFP) if critical organ bleeding (150-300mL) or if Prothrombinex VF is unavailable (FFP 15mL/kg). If required seek consultation with a haematologist / specialist.
Notes ¹ undiluted paediatric IV formulation ² at a rate of 3mL/min. 500 Units of factor IX in 1 vial of Prothrombinex VF ³ undiluted as slow IV bolus over at least 30 seconds ⁴ available from transfusion service For reversal prior to a procedure – Refer to hospital guidelines or seek specialist advice. Seek advice with Vitamin K in cardiac valve replacement.					
*High Bleeding Risk One or more ⇨ • Recent surgery / trauma / bleed • Advanced age • Renal Failure • Hypertension • Alcohol abuse • Active GI bleed • Antiplatelet therapy • Other relevant co-morbidity					

• Recommendations for direct oral anticoagulants

• Recommendations for warfarin

• Updated Warfarin Reversal Guidelines

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The middle pages

-prescribing and administering IV heparin

Attach Patient Sticker

REASON FOR NURSES NOT ADMINISTERING Codes MUST be circled	
Absent (A)	Refused - notify Doctor (R)
Fasting (F)	Not Available (N)
Vomiting (V)	Obtain supply or contact doctor
On Leave (L)	Self Administering (S)
	Withheld (W)
	Enter reason in clinical notes

RECOMMENDATIONS FOR INTRAVENOUS UNFRACTIONATED HEPARIN

Standard dilution: 50 units / mL : dilute 25,000 units of unfractionated heparin in 500mL of 0.9% sodium chloride or 5% glucose

Target aPTT: VTE/ACS: xx + xx seconds or as otherwise specified by consultant.
Target aPTT and dose nomograms are HOSPITAL SPECIFIC – consult Pathology Laboratory for correct aPTT ranges.

Monitoring: Measure baseline aPTT prior to commencing treatment, then within 6 hours of every rate change, otherwise daily.
Measure platelets at baseline and at least twice weekly.
Contact haematologist in all suspected cases of Heparin Induced Thrombocytopenia (HIT).

Reversing heparin treatment: Seek specialist or senior colleague advice. Protamine reversal should be used for cases of major bleeding or where required prior to emergency surgery. For a high aPTT without bleeding follow nomogram (page 3).
As a guide: Estimate heparin dose received in last hour. Administer 1 mg protamine sulphate per 100 units of heparin (maximum 50mg) as a slow IV push (over 10 minutes). Monitor aPTT after bolus then as required.

INTRAVENOUS PRESCRIPTION ORDER
Prescriber to complete. A new prescription is required if the order (total dose, fluid or volume) is changed

Target aPTT: Indication: VTE Acute Coronary Syndrome (ACS) Other(specify) Weight: kg

Date	Drug	Total dose (units)	Fluid	Volume (mL)	Signature	Print Name	Contact
	HEPARIN	25,000 units	0.9% SODIUM CHLORIDE	500 mL			

INITIAL BOLUS DOSE AND INITIAL INFUSION RATE Prescriber to complete ORDER

Date	Baseline aPTT	Baseline Platelets	Date/Time of dose	Initial Bolus (units)	Initial Infusion Rate (mL/hour)	Prescriber Signature	Print Name	Nurse Signature	Time

MAINTENANCE INFUSION RATE CHANGES AND BOLUS DOSES

Prescriber to complete order Prescriber to be contacted following each aPTT test
 Nursing staff to adjust dose based on nomogram using _____ kg column

Date	Prescriber Signature	Print Name	Contact	Pharmacy

aPTT test

Date	Time Taken	aPTT	Time	IV Bolus (units)	Bolus (Sign)	Hold (mins)	Time Stopped	Hold (Sign)	Time Started	New Rate (mL/hour)	Rate (Sign)	Prescriber (Sign)	Platelets

INFUSION CEASED: Date: / / Time: Prescriber Signature: Print Name:

INFUSION BAG CHANGES Nursing staff to document each new bag. Infusion should only be interrupted when indicated by aPTT

Date	Time Commenced	Checked	Given	Time Completed	Volume Infused (mL)	Date	Time Commenced	Checked	Given	Time Completed	Volume Infused (mL)

- Recommendation for IV unfractionated heparin
- Intravenous prescription order
- Initial bolus and infusion rate
- Maintenance infusion rate and bolus dose
- Infusion bag changes

The middle pages-dosing recommendations

Treatment recommendations do NOT cover all clinical scenarios and do not replace the need for clinical judgement.

INFUSION NOMOGRAM FOR INTRAVENOUS UNFRACTIONATED HEPARIN USE

- This nomogram (weight-based guide) is only valid when using an unfractionated heparin concentration of 25,000 units in 50mL and **not** 10,000 units in 50mL.
- Fluid Restricted Patients: A dilution of 25,000 units of unfractionated heparin in 50mL sodium chloride 0.9% infusion with associated nomogram is available for patients requiring severe fluid restrictions. Please contact your pharmacist for advice. If required, strike out nomogram below and staple Fluid Restricted Nomogram over page 3 of this chart.

INITIAL ORDER: Prescriber should complete order (initial bolus and initial infusion rate) on page 2. See below for recommended dose for Venous Thromboembolism (VTE) or Acute Coronary Syndrome (ACS).

- It is important that a bolus dose of unfractionated heparin is prescribed and administered on initiating an unfractionated heparin infusion to ensure that the therapeutic range is reached within the first 24 hours of therapy.

MAINTENANCE: Prescriber to indicate on page 2 whether nurse should maintain infusion rate based on nomogram as indicated OR whether the prescriber is to be contacted following each aPTT test.

IT IS RECOMMENDED THAT ALL BOLUS DOSES BE DRAWN UP FROM SEPARATE AMPOULES INTO A SYRINGE FOR ADMINISTRATION.

Venous Thromboembolism (DVT/PE) Bolus and Initial Rate Requirements

		Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
Bolus Dose	80 units/kg	Units	3200	3600	4000	4400	4800	5200	5600	6000	6400	6800	7200	7200
Initial Rate	18 units/kg/hour	Rate (mL/hour)	14	16	18	20	22	23	25	27	29	31	32	32

Acute Coronary Syndrome Bolus and Initial Rate Requirements

		Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
Bolus Dose	60 units/kg	Units	2400	2800	3000	3300	3600	4000	4000	4000	4000	4000	4000	4000
Initial Rate	12 units/kg/hour	Rate (mL/hour)	10	11	12	13	14	15	17	19	20	20	20	20

Nomogram for modifying rate of administration for Venous Thromboembolism and Acute Coronary Syndrome

MAINTENANCE ORDER		Weight Based Rate For Maintenance Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
aPTT	Dose Adjustment Use weight column on nomogram and row for aPTT range for mL/hour conversion of units/kg/hour	Rate Change (mL/hour)	This rate equals recommended change in units/hour for a 50 units/kg/hour infusion. Remeasure aPTT within 6 hours of each rate change.											
≤ Kk	Bolus dose as per indication (VTE OR ACS listed above) Then increase 3 units/kg/hour		+2	+3	+3	+3	+4	+4	+4	+5	+5	+5	+5	+5
L-Nm	Increase 2 units/kg/hour For VTE consider 40 units/kg bolus dose		+2	+2	+2	+2	+2	+3	+3	+3	+3	+3	+4	+4
Nm-Pp	No Change		Remeasure aPTT within 24 hours (or next morning)											
Qq-Rr	Reduce 1 units/kg/hour		-1	-1	-1	-1	-1	-1	-1	-2	-2	-2	-2	-2
Ss-Tt	Hold 30 minutes Then reduce 2 units/kg/hour		-2	-2	-2	-2	-2	-3	-3	-3	-3	-3	-4	-4
> Zz	Contact doctor Hold 60 minutes Then reduce 3 units/kg/hour		-2	-3	-3	-3	-4	-4	-4	-5	-5	-5	-5	-6

Slight variances of aPTT ranges may occur due to changes in laboratory reagents used. Please check with your Pathology Laboratory.

RECOMMENDATIONS FOR UNFRACTIONATED SUBCUTANEOUS HEPARIN

Dosing	VTE prophylaxis: 5000 units bid (0600 & 1800) High Risk Thromboembolism: 5000 units tds (0600,1200,1800)
Withholding subcutaneous UFH	Withhold subcutaneous heparin a minimum of 6 to 8 hours prior to intervention Interventional (surgical) procedure: may commence prophylactic doses 2 hours after procedure.
Monitoring	Full blood count: Measure platelets at baseline and at least twice weekly. Medical review if platelets less than 50 x 10 ⁹ /L

RECOMMENDATIONS FOR LOW MOLECULAR WEIGHT HEPARIN (LMWH)

Preferred administration times for twice daily dosing are 0600 and 1800 hr. Daily thromboprophylaxis should be given in the evening.

Enoxaparin Dosage and Frequency (Seek specialist advice in patients weighing < 50kg and > 120kg)

INDICATION	Normal renal function	Impaired renal function (CrCl < 30mL/min)
VTE prophylaxis	40mg once daily	20mg once daily or consider alternative
DVT/PE treatment	1.5mg/kg once daily OR 1 mg/kg twice daily	1mg/kg once daily or consider alternative
Acute Coronary Syndrome/Cardiac Valves	1mg/kg twice daily	1mg/kg once daily or consider alternative

Dalteparin is commonly used for VTE treatment in cancer patients: dose 200 Units/kg daily subcutaneously for 30 days, then 150 Units/kg daily for 5 months. Total daily dose should not exceed 18,000 Units. Dose adjustment is required for renal impairment and thrombocytopenia. See prescribing guidelines.

Monitoring	Baseline full blood count and U&Es. Measure platelets at baseline and at least twice weekly. Medical review if platelets less than 50 x 10 ⁹ /L Seek specialist advice for monitoring anti-Xa, dose modification or alternative therapeutic options. Consider anti-Xa levels for patients on high doses, and in obese, pregnant, renal impairment and frail elderly patients.
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Reversing Overtreatment	Seek specialist advice as prothrombin only partially neutralises low molecular heparin. Only consider prothrombin if LMWH has been given within the last 12 hours. Check hospital guidelines for more detailed advice on prothrombin use. As a guide: Give 1mg prothrombin sulfate per 1mg enoxaparin (maximum 50mg as a single dose). Administer initial dose (up to 50mg) by slow IV push (over 10 minutes) and remaining dose by intravenous infusion (maximum infusion rate 5mg/minute). Reassess the patient and the aPTT in 2-4 hours and consider a repeat dose if the patient is still bleeding or the aPTT remains prolonged.
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- Infusion nomogram for intravenous unfractionated heparin use
- Venous Thromboembolism (VTE) bolus and initial rate
- Acute Coronary Syndromes (ACS) bolus and initial rate
- Nomogram for rate change
- Recommendations for unfractionated subcutaneous heparin
- Recommendations for LMWH

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Prescribing anticoagulant agents

When prescribing anticoagulant agents, it is important to first check for:

- co-existing conditions,
 - past history of anticoagulant related adverse events and
 - concomitant therapy
- These may influence the decision to prescribe a particular anticoagulant or indicate a need for closer monitoring and/or dose adjustment.
 - The Bleeding Risk considered before prescribing anticoagulants prompt is on the front of the anticoagulant chart.

Bleeding Risk considered before prescribing anticoagulants Completed by (prescriber) _____ Date: _/ _/ _
Please refer to Local Venous Thromboembolism Guidelines for Bleeding Risk Assessment. Caution should be considered for patients on Dual Antiplatelet Therapy (DAPT)

- The prescriber MUST complete this section.
- Please refer to local Venous Thromboembolism guidelines for bleeding risk assessment and the Warfarin and Heparin guidelines associated with this chart for further information

Regular dose orders

DATE AND MONTH for each separate order
Ensure bleeding AND VTE risk is reassessed

REGULAR DOSE ORDERS - PROPHYLACTIC DOSES Check platelets and coagulation profile before commencing (Subcutaneous unfractionated and low molecular weight heparins and direct oral anticoagulants - DOACs)

YEAR 20__ DAY AND MONTH →

Date: _____ Medicine (First generic name): _____

CrCl mL/min: _____ Route: _____ Dose AND Frequency NOW enter times → _____

Indication: **VTE Prophylaxis** Pharmacy: _____ Creatinine: _____

Prescriber Sign: _____ Print Name: _____ Contact No.: _____ Platelets: _____

REGULAR DOSE ORDERS - THERAPEUTIC DOSES Check platelets and coagulation profile before commencing (Subcutaneous low molecular weight heparins and direct oral anticoagulants - DOACs)

YEAR 20__ DAY AND MONTH →

Date: _____ Medicine (First generic name): _____

CrCl mL/min: _____ Route: _____ Dose AND Frequency NOW enter times → _____

Indication: **Therapeutic** Pharmacy: _____ Creatinine: _____

Prescriber Sign: _____ Print Name: _____ Contact No.: _____ Platelets: _____

Pharmaceutical review: _____

WARFARIN OR DOAC DRUG INTERACTIONS (Pharmacy: Indicate drug and expected interaction) Details: _____

Sign: _____ Date: _____

Continual Discharge: YES / NO _____

Dispense: YES / NO _____

Duration: _____ days / Qry: _____

Pharmacist: _____ Date: _____

Record creatinine and platelets results

Calculate and record Creatinine Clearance

Please document the indication here e.g. DVT

- Subcutaneous unfractionated heparin
- Subcutaneous enoxaparin or dalteparin dosing based on indication and the patient's renal function and weight.
- Direct oral anticoagulant (eg. Rivaroxaban, apixaban and dabigatran are to be prescribed in this section of the chart depending on indication).

Example of Correct Use of Regular Dose Order Section

When changing the anticoagulant agent or the indication, the day and month must be carried in the corresponding column across the order as shown below:

REGULAR DOSE ORDERS - PROPHYLACTIC DOSES					Check platelets and coagulation profile before commencing																				
(Subcutaneous unfractionated and low molecular weight heparins and direct oral anticoagulants - DOACs)																									
YEAR 20 <u>22</u>		DAY AND MONTH →			4/8	5/8	6/8	7/8																	
Date	4/8	Medicine (Print generic name)	Heparin		0600	ZA	ZA	ZA	ZA	/										Continue at Discharge: YES / NO					
CrCl mL/min	68	Route	subcut		1800	MN	MN	MN	MN												Disperse YES / NO				
Dose AND Frequency NOW enter times →			5000 units BD		Ceased 7/8/22										Duration: _____ days. Qty: _____										
Indication: VTE Prophylaxis		Pharmacy			A.B 4/8		Creatinine																		
Prescriber Sign		A.Medic		Print Name		A.Medic		Contact No.		1234		Platelets													
YEAR 20 <u>22</u>		DAY AND MONTH →																							
Date	8/8	Medicine (Print generic name)	Enoxaparin		1800	X	X	X	X	8/8	9/8	10/8	11/8	/						Continue at Discharge: YES / NO					
CrCl mL/min	66	Route	subcut		40mg daily	TN	TN	TN	TN	Disperse YES / NO															
Dose AND Frequency NOW enter times →			40mg daily		Ceased 11/8/22										Duration: _____ days. Qty: _____										
Indication: VTE Prophylaxis		Pharmacy			A.B 8/8		Creatinine																		
Prescriber Sign		A.Medic		Print Name		A.Medic		Contact No.		1234		Platelets													
REGULAR DOSE ORDERS - THERAPEUTIC DOSES					Check platelets and coagulation profile before commencing																				
(Subcutaneous low molecular weight heparins and direct oral anticoagulants - DOACs)																									
YEAR 20 <u>22</u>		DAY AND MONTH →																							
Date	12/8	Medicine (Print generic name)	Enoxaparin		0600	X	X	X	X	X	X	X	X	12/8	13/8	14/8	15/8	/						Continue at Discharge: YES / NO	
CrCl mL/min	66	Route	subcut		1800	X	X	X	X	X	X	X	ST	ST	ST	ST	Disperse YES / NO								
Dose AND Frequency NOW enter times →			80mg BD												Duration: _____ days. Qty: _____										
Indication: DVT Therapeutic		Pharmacy			A.B 12/8		Creatinine																		
Prescriber Sign		A.Medic		Print Name		A.Medic		Contact No.		1234		Platelets													

Example of Correct Use of Regular Dose Order Section

If the anticoagulant is the same and there is no change in indication, you can continue the prescription order as shown below:

REGULAR DOSE ORDERS - <u>PROPHYLACTIC DOSES</u>					Check platelets and coagulation profile before commencing (Subcutaneous unfractionated and low molecular weight heparins and direct oral anticoagulants - DOACs)														
YEAR 20 <u>22</u>		DAY AND MONTH →			4/8	5/8	6/8	7/8	8/8	9/8	10/8	11/8	12/8	13/8	14/8	15/8	Continue at Discharge: YES / NO	Dispense YES / NO	Duration: _____ days, Qty: _____
Date 4/8	Medicine (Print generic name) Enoxaparin			1800															
CrCl mL/min 28	Route subcut	Dose AND Frequency NOW enter times → 20mg daily			AD	CT	CT	CT	PL	PL	PL	AD	PL	ZA	CT		ZA		
Indication: VTE Prophylaxis		Pharmacy A.B 4/8		Creatinine	153										156				
Prescriber Sign <i>A.Medic</i>	Print Name A.Medic	Contact No. pager 1234		Platelets		177			178										
YEAR 20 <u>22</u>		DAY AND MONTH →														Continue at Discharge: YES / NO	Dispense YES / NO	Duration: _____ days, Qty: _____	
Date 16/8	Medicine (Print generic name) Enoxaparin			1800		16/8	17/8	18/8	19/8	20/8	21/8	22/8	23/8	24/8	25/8				26/8
CrCl mL/min 28	Route subcut	Dose AND Frequency NOW enter times → 20mg daily			ZA	AD	ZA	AD	CT	KF	KF	KF	KF	AD	MN	MN			
Indication: VTE Prophylaxis		Pharmacy A.B 16/8		Creatinine	160										154				
Prescriber Sign <i>A.Medic</i>	Print Name A.Medic	Contact No. pager 1234		Platelets	201									201					

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Recommendations for Low Molecular Weight Heparin (LMWH)

- Dosing of LMWH (i.e. enoxaparin and dalteparin) is based on the indication, risk of bleeding risk and modifying factors (e.g., renal function and patient weight).
- Dose modification of these drugs is required when the creatinine clearance (CrCl) is less than 30mL/min.

RECOMMENDATIONS FOR UNFRACTIONATED SUBCUTANEOUS HEPARIN		
Dosing	<i>VTE prophylaxis:</i> 5000 units bd (0600 & 1800) <i>High Risk Thromboembolism:</i> 5000 units tds (0600,1200,1800)	
Withholding subcutaneous UFH	<ul style="list-style-type: none"> • Withhold subcutaneous heparin a minimum of 6 to 8 hours prior to intervention • Interventional (surgical) procedure: may commence prophylactic doses 2 hours after procedure. 	
Monitoring	<ul style="list-style-type: none"> • Full blood count: Measure platelets at baseline and at least twice weekly. Medical review if platelets less than 50 x 10⁹/L 	
RECOMMENDATIONS FOR LOW MOLECULAR WEIGHT HEPARIN (LMWH)		
Preferred administration times for twice daily dosing are 0600 and 1800 hr. Daily thromboprophylaxis should be given in the evening.		
Enoxaparin Dosage and Frequency (Seek specialist advice in patients weighing < 40kg and > 120kg)		
INDICATION	Normal renal function	Impaired renal function (CrCl<30mL/min)
<i>VTE prophylaxis</i>	40mg once daily	20mg once daily or consider alternative
<i>DVT/PE treatment</i>	1.5mg/kg once daily OR 1 mg/kg twice daily	1mg/kg once daily or consider alternative
<i>Acute Coronary Syndrome/Cardiac Valves</i>	1mg/kg twice daily	1mg/kg once daily or consider alternative
Dalteparin is commonly used for VTE treatment in cancer patients: dose 200 Units/kg daily subcutaneously for 30 days, then 150 Units/kg daily for 5 months. Total daily dose should not exceed 18,000 Units. Dose adjustment is required for renal impairment and thrombocytopenia. See prescribing guidelines.		
Monitoring	<ul style="list-style-type: none"> • Baseline full blood count and U&Es. Measure platelets at baseline and at least twice weekly. Medical review if platelets less than 50 x 10⁹/L • Seek specialist advice for monitoring anti-Xa, dose modification or alternative therapeutic options. • Consider anti-Xa levels for patients on high doses, and in obese, pregnant, renal impairment and frail elderly patients. 	
Reversing Overtreatment	<ul style="list-style-type: none"> • Seek specialist advice as protamine only partially neutralises low molecular heparin. Only consider protamine if LMWH has been given within the last 12 hours. • Check hospital guidelines for more detailed advice on protamine use. As a guide: Give 1mg protamine sulfate per 1mg enoxaparin (maximum 50mg as a single dose). • Administer initial dose (up to 50mg) by slow IV push (over 10 minutes) and remaining dose by intravenous infusion (maximum infusion rate 5mg/minute). Reassess the patient and the APTT in 2-4 hours and consider a repeat dose if the patient is still bleeding or the aPTT remains prolonged. 	

Recommendations for low molecular weight heparin (LMWH)

- Routine monitoring of residual anti-Xa activity as a measure of LMWH therapy is not required.
- However, in the case of patients at high risk of bleeding, obese patients, patients on high doses, pregnant, renal impairment and frail elderly patients, anti-factor Xa monitoring may be appropriate.
- While the risk of heparin induced thrombocytopenia (HIT) is lower with LMWH than unfractionated heparin, screening for HIT with a platelet count at day 5 of therapy is recommended.

Prescribing Intravenous Unfractionated Heparin (UFH)

- **Initial order** – prescriber should complete order (initial bolus and initial infusion rate) on page 2 of chart.
- **Maintenance** – prescriber to indicate whether nurse should maintain infusion rate based on nomogram as indicated OR whether prescriber is to be contacted
- **It is important, especially for serious pulmonary embolism (PE), that a bolus dose of UFH is prescribed and administered on initiating UFH infusion to ensure that the therapeutic range is reached within the first 24 hours of therapy**

Heparin Infusion Nomogram

Venous Thromboembolism (DVT/PE) Bolus and Initial Rate Requirements													
		Weight Based Guide For Initial Dose											
	Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
Bolus Dose	80 units/kg	Units	3200	3600	4000	4400	4800	5200	5600	6000	6400	6800	
Initial Rate	18 units/kg/hour	Rate (mL/hour)	14	16	18	20	22	23	25	27	29	31	
Acute Coronary Syndrome Bolus and Initial Rate Requirements													
		Weight Based Guide For Initial Dose											
	Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
Bolus Dose	60 units/kg	Units	2400	2800	3000	3300	3600	4000	4000	4000	4000	4000	4000
Initial Rate	12 units/kg/hour	Rate (mL/hour)	10	11	12	13	14	15	17	19	20	20	20
Nomogram for modifying rate of administration for Venous Thromboembolism and Acute Coronary Syndrome													
MAINTENANCE ORDER		Weight Based Rate For Maintenance Dose											
	Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
aPTT	Dose Adjustment Use weight column on nomogram and row for aPTT range for mL/hour conversion of unit/kg/hour	Rate Change (mL/hour) This rate equals recommended change in units/hour for a 50 unit/mL dilution. Remeasure aPTT within 6 hours of each rate change.											
≤ Kk	Bolus dose as per indication (VTE OR ACS listed above) Then increase 3 units/kg/hour	+2	+3	+3	+3	+4	+4	+4	+5	+5	+5	+5	+6
Ll-Mm	Increase 2 units/kg/hour For VTE consider 40 units/kg bolus dose	+2	+2	+2	+2	+2	+3	+3	+3	+3	+3	+3	+3
Nn-Pp	No Change	Remeasure aPTT within 24 hours (or next morning)											
Qq-Rr	Reduce 1 unit/kg/hour	-1	-1	-1	-1	-1	-1	-1	-2	-2	-2	-2	-2
Ss-Tt	Hold 30 minutes Then reduce 2 units/kg/hour	-2	-2	-2	-2	-2	-3	-3	-3	-3	-3	-3	-3
> Zz	• Contact doctor • Hold 60 minutes • Then reduce 3 units/kg/hour	-2	-3	-3	-3	-4	-4	-4	-5	-5	-5	-5	-6
Slight variances of aPTT ranges may occur due to changes in laboratory reagents used. Please check with your Pathology Laboratory.													

Initial dose will vary depending on the indication – VTE or ACS

Maintenance order will depend on patients weight and aPTT level

aPTT ranges in above nomogram are an EXAMPLE ONLY to illustrate use of chart in following slides. Please check with your Pathology Laboratory for aPTT ranges for your hospital

Intravenous infusions

Eg: for patient with Venous Thromboembolism

INTRAVENOUS PRESCRIPTION ORDER								
Prescriber to complete. A new prescription is required if the order (total dose, fluid or volume) is changed)								
Target aPTT: 70-95		Indication: <input checked="" type="checkbox"/> VTE <input type="checkbox"/> Acute Coronary Syndrome (ACS) <input type="checkbox"/> Other (specify)					Weight: 74 kg	
Date	Drug	Total dose (units)	Fluid	Volume (mL)	Signature	Print Name	Contact	
31/8	HEPARIN	25,000 units	0.9% SODIUM CHLORIDE	500 mL	<i>A. Doctor</i>	A.Doctor	4025	
INITIAL BOLUS DOSE AND INITIAL INFUSION RATE Prescriber to complete ORDER								
Date	Baseline aPTT	Date/Time of dose	Initial Bolus (units)	Initial Infusion Rate (mL/hour)	Prescriber		Nurse	
					Signature	Print Name	Time	N1/N2
31/8	42	31/8/22 0200	6000 units	27mL/hr	<i>A. Doctor</i>	A.Doctor	1430	SR DA
MAINTENANCE INFUSION RATE CHANGES AND BOLUS DOSES								
Prescriber to complete order <input type="checkbox"/> Prescriber to be contacted following each aPTT test								
<input checked="" type="checkbox"/> Nursing staff to adjust dose based on nomogram using <u>75</u> kg column								
Date 31/8/22	Prescriber signature <i>A. Doctor</i>		Print Name A.Doctor		Contact 4025	Pharmacy P.Harmacist		

Heparin Infusion Nomogram use for VTE

Venous Thromboembolism (DVT/PE) Bolus and Initial Rate Requirements														
Bolus Dose 80 units/kg Initial Rate 18 units/kg/hour		Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
		Units	3200	3600	4000	4400	4800	5200	5600	6000	6400	6800	7200	7200
		Rate (mL/hour)	14	16	18	20	22	23	25	27	29	31	32	32
Acute Coronary Syndrome Bolus and Initial Rate Requirements														
Bolus Dose 60 units/kg Initial Rate 12 units/kg/hour		Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
		Units	2400	2800	3000	3300	3600	4000	4000	4000	4000	4000	4000	4000
		Rate (mL/hour)	10	11	12	13	14	15	17	19	20	20	20	20
Nomogram for modifying rate of administration for Venous Thromboembolism and Acute Coronary Syndrome														
MAINTENANCE ORDER		Weight Based Rate For Maintenance Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
aPTT	Dose Adjustment Use weight column on nomogram and row for aPTT range for mL/hour conversion of unit/kg/hour	Rate Change (mL/hour) This rate equals recommended change in units/hour for a 50 unit/mL dilution. Remeasure aPTT within 6 hours of each rate change.												
≤ Kk	Bolus dose as per indication (VTE OR ACS listed above) Then increase 3 units/kg/hour	+2	+3	+3	+3	+4	+4	+4	+5	+5	+5	+5	+6	
Li-Mm	Increase 2 units/kg/hour For VTE consider 40 units/kg bolus dose	+2	+2	+2	+2	+2	+3	+3	+3	+3	+3	+4	+4	
Nn-Pp	No Change	Remeasure aPTT within 24 hours (or next morning)												
Qq-Rr	Reduce 1 unit/kg/hour	-1	-1	-1	-1	-1	-1	-1	-2	-2	-2	-2	-2	
Ss-Tt	Hold 30 minutes Then reduce 2 units/kg/hour	-2	-2	-2	-2	-2	-3	-3	-3	-3	-3	-4	-4	
> Zz	• Contact doctor • Hold 60 minutes • Then reduce 3 units/kg/hour	-2	-3	-3	-3	-4	-4	-4	-5	-5	-5	-5	-6	

Slight variances of aPTT ranges may occur due to changes in laboratory reagents used. Please check with your Pathology Laboratory.

aPTT ranges in above nomogram are an EXAMPLE ONLY to illustrate use of chart in following slides. Please check with your Pathology Laboratory for aPTT ranges for your hospital

Maintaining the infusion regimen using the weight-based nomogram and weight-based guide

aPTT test			Bolus and infusion rate administration									
Date	Time Taken	aPTT	Time	IV bolus (units)	Bolus (Sign)	Hold (minutes)	Time stopped	Hold (Sign)	Time started	New Rate (mL / hour)	Rate (Sign)	Prescriber Sign
31/8			0800	6000	AL MC				0800	27	KC JK	27 + 3
31/8	1400	90							1430	27	KF MC	
1/9	1400	62	1430	3000	DA SW				1430	30	DA SW	30 - 1
1/9	2000	85							2030	30	KW SU	
2/9	2000	109							2030	29	CP MR	
3/9	0400	125				60 minutes	0430		0530	24	CP MR	
INFUSION CEASED:			Date	Time	Prescriber signature			Print Name		Contact	Pharmacy	

1. Contact Doctor
2. Withhold infusion for 60 minutes
3. Reduce rate by 3 units/kg/hour, which is 5mL/hour as per nomogram= 24mL/hour

(mL) | (mL)

Maintenance regimen IV Heparin Continuous infusion – should only be stopped when indicated by nomogram or as directed by the prescriber.

- aPTT should be checked:
 - within 6 hours of every rate change or
 - within 24 hours (next morning) – when aPTT within target range
- There should be a prompt dose adjustment to each aPTT measurement
- The infusion should be continuous– only stop when indicated by aPTT (nomogram)
- **Prescriber should always be contacted for EXTREME aPTT levels**
- In all cases the prescriber should frequently check the aPTT result and subsequent infusion rate changes
- It is recommended that bolus doses be drawn up (as prescribed) from a separate ampoule into a syringe for administration.

Fluid Restricted Patients

- Renal failure and heart failure
- 25,000 units in 50mL nomogram available
- Watch rate changes
- 10 x difference to normal nomograms
- Print and staple to WA Anticoagulation Chart

Heparin Infusions

- Important to make sure correct dilution used
- Standard dilution **25,000 units in 500mL** on WA Anticoagulation Chart
- Fluid Restricted Patients **25,000 units in 50mL**
- *** Not all sites will require a fluid restricted nomogram**
 - check local guidelines
- Different nomograms required
 - 10x rate errors
- Monitoring and rate adjustment important for safe management

Treatment recommendations do NOT cover all clinical scenarios and do not replace the need for clinical judgement.

Infusion Nomogram for Intravenous Unfractionated Heparin For FLUID RESTRICTED PATIENTS 25,000 units in 50 mL

Patients requiring fluid restrictions (e.g. patient with heart failure or severe renal impairment) may require a more concentrated dilution of unfractionated heparin than the standard dilution used in the WA Anticoagulation Medication Chart – 25,000 units in 500mL of sodium chloride 0.9% (50units/mL).

Print a copy of the FLUID RESTRICTED nomogram and ATTACH to Anticoagulation Chart over existing page 3 – put a line through the original nomogram on the WA Anticoagulation Medication Chart.

This nomogram (weight-based guides) is ONLY valid when using an unfractionated heparin concentration of 25,000 units in 50mL and STANDARD aPTT targets.

INITIAL ORDER : Prescriber should complete order (initial bolus and initial infusion rate) on page 2. See below for recommended dose for Venous Thromboembolism (VTE) or Acute Coronary Syndrome (ACS).

- It is important that a bolus dose of unfractionated heparin is prescribed and administered on initiating an unfractionated heparin infusion to ensure that the therapeutic range is reached within the first 24 hours of therapy.

MAINTENANCE : Prescriber to indicate on page 2 whether nurse should maintain infusion rate based on nomogram as indicated OR whether the prescriber is to be contacted following each aPTT test.

IT IS RECOMMENDED FOR SAFETY THAT

- All bolus doses be drawn up from separate ampoules into a syringe for administration
- A syringe driver is used to administer the infusion due to the very low infusion rates required

Venous Thromboembolism (DVT/PE) Bolus and Initial Rate Requirements														
Bolus Dose	80 units/kg	Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
		Units	3200	3600	4000	4400	4800	5200	5600	6000	6400	6800	7200	7200
Initial Rate	18 units/kg/hour	Rate (mL/hour)	1.4	1.6	1.8	2	2.2	2.3	2.5	2.7	2.9	3.1	3.2	3.2

Acute Coronary Syndrome Bolus and Initial Rate Requirements														
Bolus Dose	60 units/kg	Weight Based Guide For Initial Dose												
		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
		Units	2400	2800	3000	3300	3600	4000	4000	4000	4000	4000	4000	4000
Initial Rate	12 units/kg/hour	Rate (mL/hour)	1	1.1	1.2	1.3	1.4	1.5	1.7	1.9	2	2	2	2

Nomogram for modifying rate of administration for Venous Thromboembolism and Acute Coronary Syndrome

MAINTENANCE ORDER		Weight Based Rate For Maintenance Dose												
Use weight column on nomogram and row for aPTT range for mL/hour conversion of units/kg/hour		Weight	≤40kg	45kg	50kg	55kg	60kg	65kg	70kg	75kg	80kg	85kg	90kg	≥95kg
aPTT	Dose Adjustment	Rate Change (mL/hour)	This rate equals recommended change in units/hour for a 50 unit/mL dilution. Remeasure aPTT within 6 hours of each rate change											
< Kk	Bolus dose as per indication (VTE OR ACS listed above) Then increase 3 units/kg/hour	+0.2 +0.3 +0.3 +0.3 +0.4 +0.4 +0.4 +0.5 +0.5 +0.5 +0.5 +0.5 +0.6												
Ll-Mm	Increase 2 units/kg/hour For VTE consider 40units/kg bolus dose	+0.2 +0.2 +0.2 +0.2 +0.2 +0.3 +0.3 +0.3 +0.3 +0.3 +0.4 +0.4												
Nll-Pp	No Change	Remeasure aPTT within 24 hours (or next morning)												
Qq-Rr	Reduce 1 unit/kg/hour	-0.1 -0.1 -0.1 -0.1 -0.1 -0.1 -0.1 -0.2 -0.2 -0.2 -0.2 -0.2 -0.2												
Ss-Tt	Hold 30 minutes Then reduce 2 units/kg/hour	-0.2 -0.2 -0.2 -0.2 -0.2 -0.3 -0.3 -0.3 -0.3 -0.3 -0.4 -0.4												
> Zz	Contact doctor Hold 60 minutes Then reduce 3 units/kg/hour	-0.2 -0.3 -0.3 -0.3 -0.4 -0.4 -0.4 -0.5 -0.5 -0.5 -0.5 -0.5												

Slight variances of aPTT ranges may occur due to changes in laboratory reagents used. Please check with your Pathology Laboratory

Please note: Each hospital is required to check with their Pathology laboratory should determine its own therapeutic target range for heparin against a gold standard test (eg residual anti-Xa activity). Because of this hospitals should not use a WA Anticoagulation Chart from another hospital as ranges will change from hospital to hospital

Reported Heparin Infusion Issues

- Wrong rate due to using the incorrect nomogram
- Be aware that ICU may have a different dilution they use for renal perfusion, if this is the case then a new prescription on the Anticoagulation Chart must be initiated and a new infusion solution must be used.
- Accidentally pushing through a large volume when not required. (often occurs when 'pushing' through volume of infusion bag rather than drawing up into a syringe for a push).
- Not monitoring aPTT and changing rate in accordance with aPTT results has led to subtherapeutic and supratherapeutic heparin management
- Not administering a bolus dose when required by nomogram for low aPTT values resulting in subtherapeutic heparin management

Warfarin

- The following is to be documented:
 - INR results
 - daily warfarin dose & prescriber’s initials prior to 1600hrs according to the most recent INR
 - indication & target INR range
 - brand of warfarin to be used
 - initials of administering and checking nurses/midwives

WARFARIN and DOAC DRUG INTERACTIONS (Pharmacy: Indicate drug and expected interaction) Details: <i>Ciprofloxacin increasing INR</i>											Sign <i>RF</i>		
											Date 13/9/22		
WARFARIN VARIABLE DOSE ORDERS													
Year 20 <u>22</u>		DAY AND MONTH →			12/9								Continue at Discharge YES / NO <input type="checkbox"/> Take as Directed Dispense YES / NO Date / / Marevan 5mg qty <u>3</u> mg qty <u>1</u> mg qty <u> </u> OR Coumadin 5mg qty <u>2</u> mg qty <u>1</u> mg qty <u> </u>
Dose at admission: Dose _____ mg <input checked="" type="checkbox"/> Not applicable				INR Result		1.1							
Brand: <input checked="" type="checkbox"/> Marevan® or <input type="checkbox"/> Coumadin®				DOSE		5	mg	mg	mg	mg	mg	mg	
Date	Medicine		Dose Time	Prescriber		AP							
12/9/22	WARFARIN			Telephone order N1/N2		/	/	/	/	/	/	/	
Indication		Route	16:00 hr	Given by		SW							
AF		ORAL		Contact No.		4152							
Target INR		Pharmacy		Prescriber sign		Print name		Contact No.		Given by			
2-3		<i>RF</i>		<i>A. Prescriber</i>		<i>A. Prescriber</i>		4152		<i>SW</i>			

Best practice when initiating warfarin

- Consider if the benefits of anticoagulation outweigh the risks for each patient
- Measure baseline INR prior to starting therapy.
- For the majority of patients > 60 years a starting dose of 5mg for day 1 and day 2 is recommended, with dose modification tailored to INR on Day 3.
- For younger patients (< 60 years) consider 7-10mg on day 1 and day 2
- Consider smaller starting doses for high risk patients (elderly, low body weight, abnormal liver function or is at high bleeding risk)
- Consider dose modification in the presence of interacting drugs
- Warfarin doses should be modified based on the INR result.

Warfarin dosing nomogram

- This warfarin dosing nomogram can be found in the Guidelines for Anticoagulation Using Warfarin

Day	INR	Suggested dose
1	1-1.4	5 mg
2	No INR	5 mg
3	<1.8	5 mg
	≥1.8	1 mg
4&5	<1.5	7 mg
	1.5-1.9	5 mg
	2.0-2.5	4 mg
	2.6-3.5	3 mg
	3.6-4.0	2 mg
	4.1-4.5	1 mg
	>4.5	See treatment reversal
6 onwards	Measure on alternate days until stable (daily if drug interaction or high bleeding risk)	As for day 4&5 or per clinical judgement

Bridging with heparin

- Bridging with heparin is recommended for patients at high risk of thrombotic events.
- Acute treatment of venous thromboembolism (DVT or PE) should be treated with heparin (unfractionated or low molecular weight) for at least of 5 days and INR is > 2
- No heparin cover is required for patients at low risk of thrombosis

Ongoing warfarin therapy:

- Brand substitution is not allowed
- Marevan[®] is the preferred brand for initiation
- In acutely ill patients daily monitoring of INR may be appropriate.
- Monitor INR more frequently when any change in treatment involves drugs known to interact with warfarin.
- Patients being re-initiated on warfarin post surgery/ procedure should be restarted on the dose prescribed prior to the intervention and check INR on day 3

Warfarin discharge planning

If patient is on warfarin, doctor to complete warfarin discharge plan prior to discharge

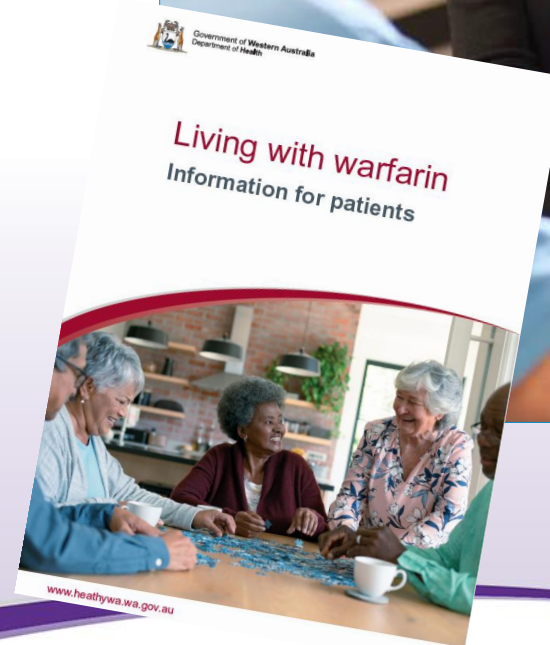
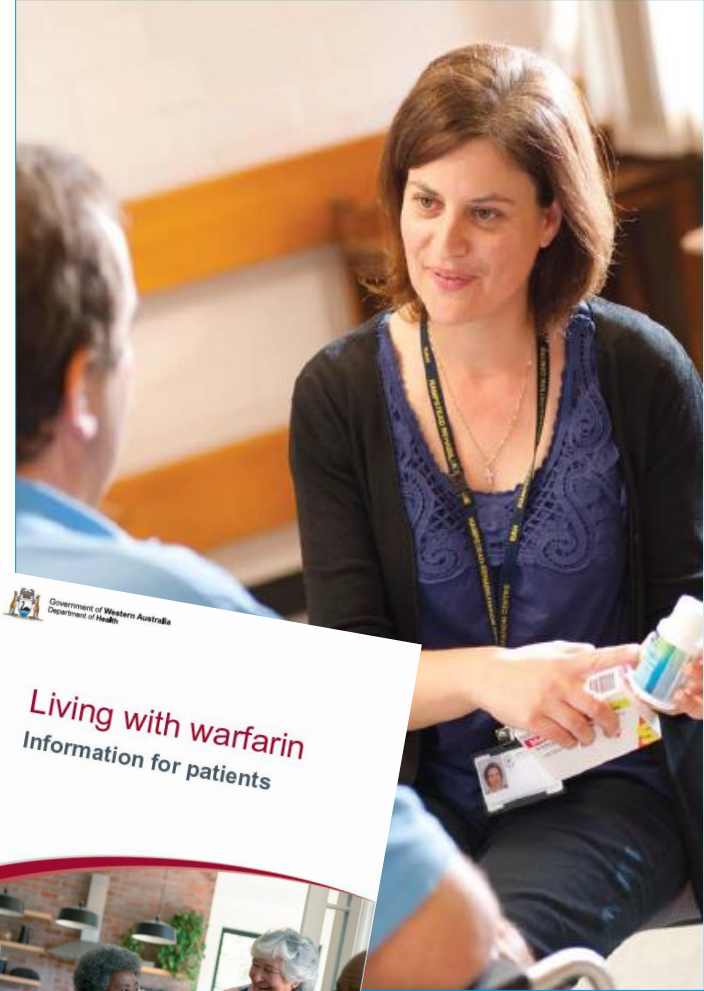
WARFARIN VARIABLE DOSE ORDERS												
YEAR 20__		DAY AND MONTH →										
Dose at admission: Dose _____ mg <input type="checkbox"/> Not applicable						INR Result						
Brand: <input type="checkbox"/> Marevan® or <input type="checkbox"/> Coumadin®												
Date	Medicine WARFARIN					Dose Time 16:00 hr	DOSE					YES / NO YES / NO Dispense 3mg 1mg OR Coumadin Qty: 5mg 2mg 1mg
Indication			Route ORAL		Prescriber							
Target INR		Pharmacy			Telephone order N1/N2							
Prescriber Sign			Print Name		Contact No.		Given by					
Warfarin Discharge Plan		Dose _____ mg	Target INR _____		Duration _____	next INR due ___/___/___		Prescriber _____				
ANTICOAGULANT DISCHARGE PLANNING <input type="checkbox"/> Patient has booklet <input type="checkbox"/> Patient education completed												
<input type="checkbox"/> Warfarin <input type="checkbox"/> DOAC _____ <input type="checkbox"/> LMWH <input type="checkbox"/> Patient given treatment plan <input type="checkbox"/> Duration _____ <input type="checkbox"/> GP informed <input type="checkbox"/> GP faxed chart												
Signature: _____			Designation: _____			Date: _____						

Patient Information Warfarin

- Engage the patient and family in self-management of warfarin
 - highlight the importance of identifying & reporting signs of bleeding
 - provide verbal counselling and education booklets
 - highlight the importance of:
 - regular INR monitoring
 - Medicines and food/alcohol that interfere with the way warfarin works.

Medication safety resources

[Medication safety resources \(health.wa.gov.au\)](http://health.wa.gov.au)



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Direct Oral Anticoagulants

- Direct Oral Anticoagulants (DOACs) are to be prescribed on the WA AMC.
- Prescribe in the Regular Dose Order section (either prophylaxis or treatment depending on indication)
- Prescribe with care in patients with poor renal function and elderly, underweight (<50kg) or overweight (>150kg) patients.
- Idarucizumab is the reversal agent for dabigatran
 - Refer to local hospital guidelines
- No Specific Reversal Agents for the other DOACs – Contact Haematology for advice if serious bleeding occurs.

Recommendations for DOACs

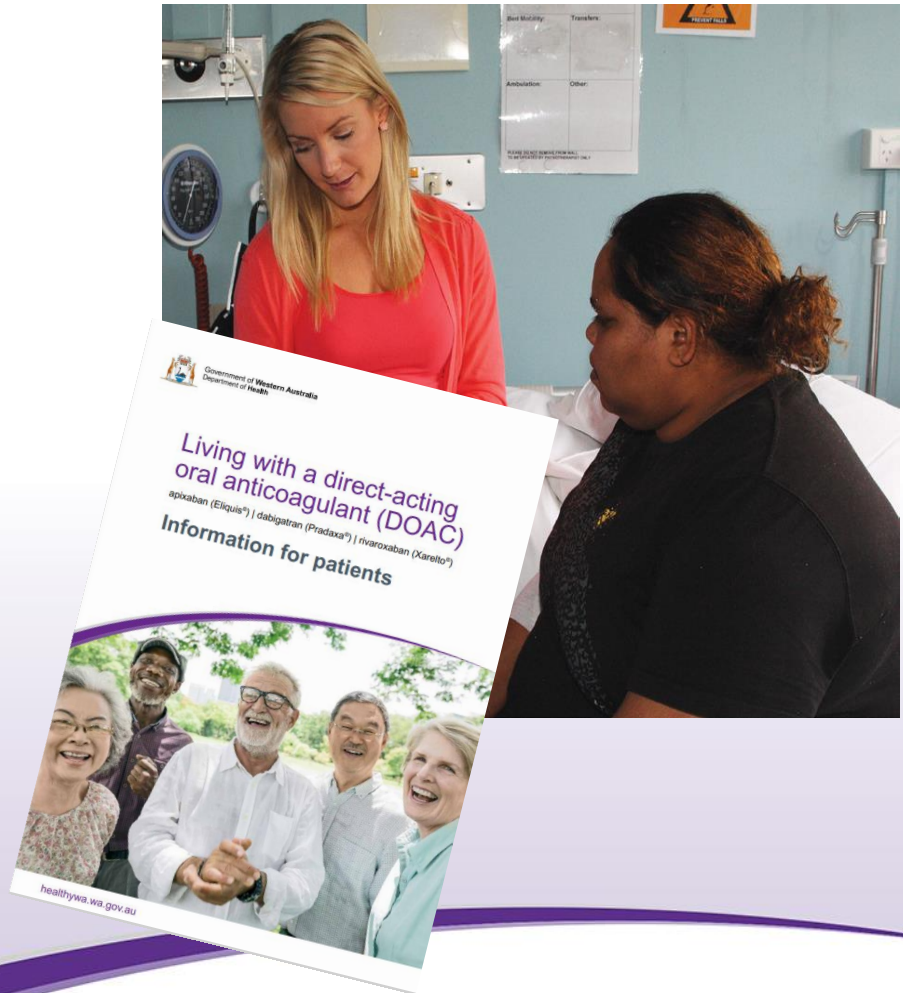
Page 4 of the WAAMC has recommendations for DOACs

Treatment recommendations do not cover all clinical scenarios and do not replace the need for clinical judgement		
RECOMMENDATIONS FOR DIRECT ORAL ANTICOAGULANTS		
Direct Oral Anticoagulant Agents (DOACs) – Apixaban, Dabigatran, Rivaroxaban (also known as NOACs) <ul style="list-style-type: none"> • Prescribe with care in elderly (>75 years), underweight (<50kg), overweight (>150kg) and patients with renal impairment (CrCl < 50mL/min). • Prior to DOAC initiation: Record: FBC, Coagulation status (INR, aPTT and PT), renal and liver function. Check for drug interactions prior to prescribing. • If the patient is on warfarin: Discontinue warfarin and start DOAC when INR is 2.0 or less • Refer to local prescribing guidelines for further information. 		
Apixaban (Eliquis®)	Dabigatran (Pradaxa®) Idarucizumab is the reversal agent for dabigatran Refer to local hospital guidelines.	Rivaroxaban (Xarelto®) (Use with caution if CrCL 15-29mL/min)
Treatment of DVT/PE: <ul style="list-style-type: none"> • CrCl >25 mL/min: 10mg twice daily for first 7 days, then 5mg twice daily thereafter 		Treatment and Prevention of DVT/PE: <ul style="list-style-type: none"> • CrCl ≥ 15 mL/min: 15mg twice daily for 3 weeks, then 20mg once daily • Seek specialist advice if CrCl 15-29mL/min
Non-Valvular Atrial Fibrillation (therapeutic dose): 5mg twice daily Reduce to 2.5mg twice daily IF at least 2 of the following risks: <input type="checkbox"/> SCr ≥ 133 micromol/L <input type="checkbox"/> Age ≥ 80 years, <input type="checkbox"/> Weight ≤ 60 kg	Non-Valvular Atrial Fibrillation (therapeutic dose): <ul style="list-style-type: none"> • CrCl ≥ 50 mL/min: 150mg twice daily • CrCl 30-49 mL/min or ≥ 75years: 110mg twice daily 	Non-Valvular Atrial Fibrillation (therapeutic dose): <ul style="list-style-type: none"> • CrCl ≥ 50 mL/min: 20mg once daily • CrCl 30-49 mL/min: 15mg once daily • CrCl 15-29 mL/min: seek specialist advice
VTE prophylaxis: Total Hip or Knee Replacement <ul style="list-style-type: none"> • CrCl > 25mL/min: 2.5mg twice daily Hip: up to 38 days Knee: up to 14 days 	VTE prophylaxis: Total Hip or Knee Replacement <ul style="list-style-type: none"> • CrCl > 50 mL/min: 220mg (2 x 110 mg) once daily • CrCl 30-50 mL/min: 150mg (2 x 75 mg) once daily Hip: up to 35 days Knee: up to 10 days 	VTE prophylaxis: Total Hip or Knee Replacement <ul style="list-style-type: none"> • CrCl ≥ 15 mL/min: 10mg once daily Hip: up to 35 days Knee: up to 14 days
		Prevention of cardiovascular events in chronic stable CAD/PVD (in combination with aspirin): <ul style="list-style-type: none"> • CrCl ≥ 15mL/min: 2.5 mg twice daily

Patient Information

Direct Oral Anticoagulant Agents (DOACs)

- Engage the patient and family in self-management of NOACs
 - Including
 - Dabigatran
 - Apixaban
 - Rivaroxaban
 - highlight the importance of identifying & reporting signs of bleeding
 - provide verbal counselling and education booklets



Medication safety resources

[Medication safety resources \(health.wa.gov.au\)](http://health.wa.gov.au)

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Anticoagulant discharge planning

- This section should be completed for any patient that is being discharged on an anticoagulant.
- This should be used as a prompt to ensure all aspects of discharge planning are completed and handed over to the patient's GP

WARFARIN VARIABLE DOSE ORDERS												
YEAR 20__				DAY AND MONTH →								
Dose at admission: Dose _____ mg <input type="checkbox"/> Not applicable				INR Result								
Brand: <input type="checkbox"/> Marevan® or <input type="checkbox"/> Coumadin®												
Date	Medicine WARFARIN			DOSE				Continue at Discharge YES/NO <input type="checkbox"/> Take as Directed Dispense YES/NO Marevan Qty: 5mg _____ 3mg _____ 1mg _____ OR Coumadin Qty: 5mg _____ 2mg _____ 1mg _____				
Indication	Route ORAL	Dose Time 16:00 hr		Prescriber								
Target INR	Pharmacy			Telephone order N1/N2								
Prescriber Sign	Print Name	Contact No.		Given by								
Warfarin Discharge Plan				Dose _____ mg	Target INR _____	Duration _____	next INR due ____/____/____		Prescriber _____			
ANTICOAGULANT DISCHARGE PLANNING <input type="checkbox"/> Patient has booklet <input type="checkbox"/> Patient education completed <input type="checkbox"/> Warfarin <input type="checkbox"/> DOAC _____ <input type="checkbox"/> LMWH <input type="checkbox"/> Patient given treatment plan <input type="checkbox"/> Duration _____ <input type="checkbox"/> GP informed <input type="checkbox"/> GP faxed chart Signature: _____ Designation: _____ Date: _____												

Minimising Risks with Anticoagulants

- Careful prescribing

- Use Standardised abbreviations- write “Units”

Date 5/12	Medication (Print Generic Name) Heparin	Tick if Slow release
Route S/C	Dose 5000U	Frequency & enter times tds

Mistaken for
50 000 units

Date 1/11	Medication (Print Generic Name) Clexate	Tick if Slow release
Route S/C	Dose 40mg	Frequency & enter times QD
Pharmacy/Additional Information		
Inclusion		
Dose Calculation (e.g. mg/kg per DOS)		

Once daily or
twice daily ???

- Brand specification for warfarin

- Marevan[®] preferred unless patient previously stabilised on Coumadin[®]
 - If not available on ward, ensure staff are familiar with ordering medications to ensure correct brand is supplied for patient

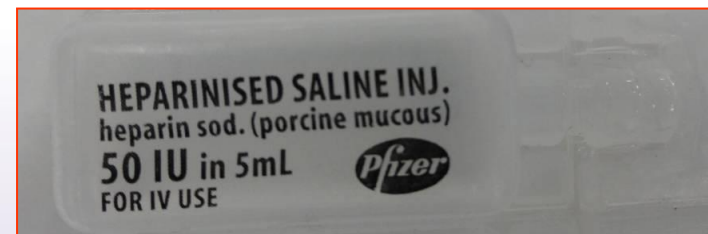
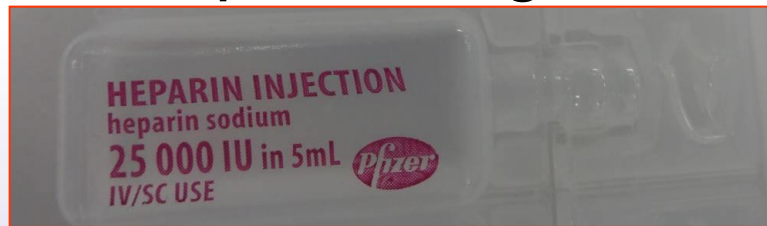
Minimising Risks with Anticoagulants

Choosing the correct product for administration

- Correct brand and strength of warfarin chosen



- Multiple strengths of heparin available



- Confusion with other medications



Adverse Effects of Anticoagulants

- The major side effect of anticoagulants is bleeding
- All symptoms must be followed up and appropriate action implemented according to the severity of the bleed
- Bleeds may be:
 - minor
 - major
 - critical

Adverse Effects of Anticoagulants

• **Minor bleeds:**

- bleeding from gums after brushing teeth
- bruising easily
- nose bleeds
- prolonged bleeding from cuts/wounds
- excessive menstrual or vaginal bleeding

Major bleeds:

- blood in stools (melena):
 - bright red blood-stained stools
 - black tarry stools
 - rectal bleeding
- vomiting blood (hematemesis)
 - may have a 'coffee ground' appearance
- passing blood in urine (hematuria):
 - bright red urine
 - dark brown, rusty coloured urine
- coughing up blood (hemoptysis)
 - pink or blood-streaked sputum
- painful, swollen, hot joints
- patient feeling tired and looking pale (anaemia)

Intracranial Haemorrhage

- An intra-cerebral bleed is a clinically critical bleed
- Symptoms may include:
 - sudden, severe headache
 - change in vision, speech
 - difficulty in walking, dizziness
 - confusion
 - weakness or numbness in one arm/leg or side of face.

Reversal of Heparin Over-treatment

Unfractionated heparin

Reversing heparin treatment	<ul style="list-style-type: none">• Seek specialist or senior colleague advice. Protamine reversal should be used for cases of major bleeding or where required prior to emergency surgery. For a high aPTT without bleeding follow nomogram (page 3).• As a guide: Estimate heparin dose received in last hour. Administer 1 mg protamine sulphate per 100 units of heparin (maximum 50mg) as a slow IV push (over 10 minutes). Monitor aPTT after bolus then as required.
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Information found on page 2 of chart

Low molecular weight heparins (e.g. enoxaparin and dalteparin)

Reversing Overtreatment	<ul style="list-style-type: none">• Seek specialist advice as protamine only partially neutralises low molecular heparin. Only consider protamine if LMWH has been given within the last 12 hours.• Check hospital guidelines for more detailed advice on protamine use. As a guide: Give 1mg protamine sulfate per 1mg enoxaparin (maximum 50mg as a single dose).• Administer initial dose (up to 50mg) by slow IV push (over 10 minutes) and remaining dose by intravenous infusion (maximum infusion rate 5mg/minute). Reassess the patient and the APTT in 2-4 hours and consider a repeat dose if the patient is still bleeding or the aPTT remains prolonged.
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Information found on page 3 of chart

Safe management of anticoagulants Pre and Post Invasive Procedures



- A protocol for withholding or resuming anticoagulants pre and post invasive procedures should be readily accessible to staff.
- Consideration should be made based on agent half life, surgery type, bleeding risk and thrombotic risk
- For more information refer to local guidelines

Summary

Anticoagulants are high risk medications

Anticoagulants

- have complex dosing regimens
- require monitoring for safe management
- The WA Anticoagulant Medication Chart is designed to enable safe and appropriate dose selection and monitoring.

Add Local Data/Information Here

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Risk Register

- Medication Safety

Medicines and Technology Unit,

Patient Safety and Clinical Quality Directorate.

WA Department of Health

DoH.medicinesandtechnologyunit@health.wa.gov.au

- Local Risk Register

– Contact: _____

WA Anticoagulation Steering Group

The Quality Improvement and Change Management Unit would like to acknowledge the contribution of the WA Anticoagulation Steering Group members to the revision of the WA Anticoagulation Medication Chart in 2022.

- Dr Dominic Pepperrell
- Dr Tony Ryan
- Dr Mark Newman
- Dr Justin Yeung
- Ms Michaela Walters
- Ms Barbara O'Callaghan
- Dr Tony Calogero
- Mr David Lui
- Ms Tandy-Sue Copeland
- Ms Ann Berwick
- Ms Cindy Tan
- Mr David McKnight
- Dr Rosslyn de Wet