Periprocedural Management (Warfarin) v1.2

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Procedure Bleed Risk (see table below)	Low thromboembolic risk	Moderate thromboembolic risk	High thromboembolic risk ¹⁰	
	AF: CHA ₂ DS ₂ -VASc score 1-4 or CHADS ₂ score of 0-2 (and no prior stroke or TIA) VTE: VTE >12 months ago MHV: Bileaflet aortic valve prosthesis without major risk factors for stroke ¹	AF: CHA ₂ DS ₂ -VASc 5-6 or CHADS ₂ score 3 or 4 VTE: VTE within past 3-12 months, recurrent VTE, non-severe thrombophilia ⁴ , recurrent VTE, active CA or recent hx of CA ⁵ MHV: Mechanical mitral valve without additional stroke risk factors ^{1,2,11} ; Bileaflet aortic valve prosthesis with major risk factors for stroke ¹	AF: CHA ₂ DS ₂ -VASc score ≥ 7 or CHADS ₂ score 5 or 6, recent stroke or TIA (< 3 months), or rheumatic valvular heart disease VTE: VTE < 3 months, severe thrombophilia ⁶ , antiphospholipid antibodies, associated with vena cava filter, associated with vena cava filter, associated with active CA with high VTE risk ⁷ MHV: Mitral valve with major risk factors for stroke ¹ ; caged-ball or tilting disc mitral/aortic valve prosthesis; recent (<3 months) stroke or TIA, or other high risk stroke risk factors ²	
Minimal	Do not interrupt ³	Do not interrupt ³	Do not interrupt ³	
Low/Moderate/High	-Interrupt ⁷ -Do not bridge	-Interrupt ⁷ -Do not bridge	-Interrupt ⁷ -Bridging suggested ^{8,9}	

Imultiple prior strokes, prior perioperative stroke, or prior valve thrombosis; 2 trial fibrillation, prior stroke or TIA, HTN, Diabetes, CHF, or age>75; 3 Interruption may be appropriate if there is increased concern for bleeding due to patient factors (eg. dental extraction in a patient with poor dentition, a screening colonoscopy in a patient with history of polyps that may require resection, or coronary angiography with a femoral (instead of radial) access; 4 heterozygous factor V Leiden or prothrombin gene mutation; 5 within 5 years if history of cancer, excluding non-melanoma skin cancer; 6 eg. deficiency of protein C, protein S or antithrombin, homozygous factor V Leiden or prothrombin gene mutation or double heterozygous for each mutation, multiple thrombophilias; 7 shorter interruption periods may be acceptable for low/moderate bleed risk procedures; 8 Address any reversible patient risk factors such as high INR or aspirin use, and consider bleed history before bridging; 9 finding not suggested for colonoscopies with anticipated polypectomy; 10 Consider delaying procedure, if possible, in high thrombotic risk patients with recent thromboembolism (within 3 months). 11 Based on MAQI consensus: mechanical mitral valves without stroke risk factors are not listed in 2022 CHEST guidelines thrombotic risk table.

Estimate Procedure Bleed Risk (examples)					
Minimal	Low/Moderate		High		
-Minor dermatologic procedures -Ophthalmologic (cataract) procedures -Minor dental procedures -Pacemaker or cardioverter-defibrillator device implantation	-Arthroscopy -Cutaneous/lymph node biopsies -Foothand surgery -Coronary angiography -GI endoscopy ± biopsy -Colonoscopy ± biopsy	-Abdominal hysterectomy -Laparoscopic cholecystectomy -Abdominal hernia repair -Hemorrhoidal surgery -Bronchoscopy ± biopsy	-Major surgery with extensive tissue injury -Cancer surgery, especially, solid tumor resection -Major orthopedic surgery, including shoulder replacement surgery -Reconstructive plastic surgery -Major thoracic surgery -Urologic or Gl surgery, especially anastomosis surgery -Transurethral prostate resection, bladder resection, or tumor ablation	-Colonic polyp resection -Bowel resection -Percutaneous endoscopic gastrostomy placement, endoscopic -Retrograde cholangiopancreatography -Surgery in highly vascular organs (kidneys, liver, spleen) -Cardiac, intracranial, or spinal surgery -Any major operation (procedure duration > 45 minutes) -Neuraxula anesthesia -Epidural injections	

Stopping warfarin

INR result (5-7 days before procedure)	Supratherapeutic	Therapeutic	Subtherapeutic
When to start holding warfarin	At least 5 days before	5 days before	3-4 days before

Bridging

Bridging							
Patient/ proce- dure factors	Bridging agent	When to start bridging agent prior to procedure	When to stop bridging agent prior to procedure	When to restart anticoagulants following procedure d	When to stop bridg- ing agent		
CrCl ≥30	LMWH	Start LMWH when INR goes below therapeutic range or after omitting 2-3 doses of warfarin (if INR not checked)	24 hours prior to the procedure. a	Warfarin: within 24 hours LMWH: at least 24 hours following low/ moderate risk procedure; at least 48-72 hours in high bleed risk procedures	When INR becomes therapeutic		
	UFH	Start UFH when INR goes below therapeutic range or after omitting 2-3 doses of warfarin (if INR not checked)	At least 4 hours prior to procedure and if aPTT is in normal range. ^b	Warfarin: within 24 hours <u>UFH</u> : at least 24 hours following proce- dure ² ; after 48-72 hours in high bleed risk procedures	When INR becomes therapeutic		
CrCl <30	UFH (recommended over LMWH) ^c	Start UFH when INR goes below therapeutic range or after omitting 2-3 doses of warfarin (if INR not checked)	At least 4 hours prior to procedure and if aPTT is in normal range. ^b	Warfarin: within 24 hours <u>UFH or LMWH</u> : at least 24 hours following procedure; after 48-72 hours in high bleed risk procedures	When INR becomes therapeutic		
Heparin allergy or recent HIT	Follow local protocol	Follow local protocol	Follow local protocol	Follow local protocol	Follow local protocol		

Doherty et al. 2017 ACC Expert Consensus Decision Pathway for Periprocedural Management of Anticoagulation in Patients With Nonvalvular Atrial Fibrillation. DOI: 10.1016/j.jacc.2016.11.024
Douketis et al. Perioperative Management of Antithrombotic Therapy: An American College of Chest Physicians Clinical Practice Guideline, Chest, Volume 162, Issue 5, 2022, Pages e207-e243, ISSN 0012-3692

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^a Half the total daily dose of LMWH the day prior to the procedure is suggested.

^bIf aPTT is not in normal range, delay procedure and recheck aPTT every 2 hours until in normal range.