

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 ³ , active CA or recent hx of CA MHV: Mitral valve without additional stroke risk factors ¹ ; 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 ⁴ , antiphospholipid antibodies, associated with vena cava filter, associated with active CA with high VTE risk ⁵ 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
Minimal	Do not interrupt ²	Do not interrupt ²	Do not interrupt ²
Low/Moderate/High	-Interrupt -Do not bridge	-Interrupt -Do not bridge	-Interrupt -Bridging suggested ^{6,7} (2022 CHEST) -Consider bridging in AF (if recent stroke/TIA or mechanical valve) (2023 ACC/AHA/ACCP)

¹AF, prior stroke/TIA, prior valve thrombosis, rheumatic heart disease, hypertension, diabetes, congestive heart failure, age ≥ 75 years. ²Interruption 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); ³heterozygous factor V Leiden or prothrombin gene mutation; ⁴eg, deficiency of protein C, protein S or antithrombin, homozygous factor V Leiden or prothrombin gene mutation or double heterozygous for each mutation, multiple thrombophilias; ⁵pancreatic cancer, myeloproliferative disorders, primary brain cancer, gastric cancer, and esophageal cancer; ⁶Address any reversible patient risk factors such as high INR or aspirin use, and consider bleed history before bridging; ⁷Bridging not suggested for colonoscopies with anticipated polypectomy; ⁸Consider delaying procedure, if possible, in high thrombotic risk patients with recent thromboembolism (within 3 months).

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 -Foot/hand 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 GI 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) -Neuraxial 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

Patient/ procedure 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 bridging agent
CrCl \geq 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 UFH: at least 24 hours following procedure ^e ; 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 UFH or LMWH: 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

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

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

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

Joglar, et al. 2023 ACC/AHA/ACCP/HRS Guideline for the Diagnosis and Management of Atrial Fibrillation Circulation. 2023;148:e00-e00. DOI: 10.1161/CIR.0000000000001193

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