



FLINT

McLaren Cardiac and Thoracic Surgery Michigan Heart Valve Institute
Phone (810)-342-2590 Fax (810)-342-2591

LAAC Device: WATCHMAN™ Pre Procedure Assessment & Eligibility Form

PATIENT INFORMATION

Name: _____ DOB: _____ Sex: M F Home Phone: _____
Address: _____ Alt/Cell Phone: _____
Referring Physician: _____ PCP: _____ Cardiologist: _____
Allergies: NKA Latex: YES NO Contrast dye: Yes No Nickel: Yes No

INSURANCE

Insurance #1: _____ Contract# : _____
Insurance #2: _____ Contract# : _____
Authorization #: _____

MEDICATION ASSESSMENT

Current Alzheimer: None Aricept Cognex Exelon Namenda Razadyne
Current Anticoagulation: None Coumadin Eliquis (Apixaban)
 Pradaxa (Dabigatran) Savayesa (Edoxaban) Xarelto (Rivaroxaban)
Current Antiplatelet: None Plavix (Clopidogrel) Brilinta (Ticagrelor) Effient (Prasugrel)
 Ticlid (Ticlopidine) Zontivity (Vorapaxar) ASA Dose: _____
Current use of NSAIDS (OTC or RX): Name: _____ Dose _____ Frequency _____
Other Medications: See attached medication list

RISK SCORING

CHA2DS2VASc Score: _____	<input type="checkbox"/> CHF (1)	<input type="checkbox"/> HTN (1)	<input type="checkbox"/> DM (1)	<input type="checkbox"/> Stroke, TIA, or Thrombembolism (2)
Stroke Risk: _____	<input type="checkbox"/> 65-74 (1)	<input type="checkbox"/> ≥ 75 (2)	<input type="checkbox"/> Female (1)	<input type="checkbox"/> Prior MI, PAD or Aortic Plaque (1)
If yes for CHF: NYHA classification is <input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III <input type="checkbox"/> IV				
HAS-BLED Score: _____	<input type="checkbox"/> HTN (1)	<input type="checkbox"/> Abnormal Renal/liver function (1pt ea.)	<input type="checkbox"/> Stroke (1)	<input type="checkbox"/> Current drugs (Antiplatelet or NSAIDs) or alcohol use (1pt ea.)
Bleed risk: _____	<input type="checkbox"/> Labile INR (1)	<input type="checkbox"/> Bleeding History or disposition (1)	<input type="checkbox"/> Age > 65 (1)	

RATIONALE: Patient has an appropriate rationale to seek a non-pharmacologic alternative to warfarin:

<input type="checkbox"/> History of bleeding (eg. Intracerebral, subdural, GI, retro-peritoneal)	<input type="checkbox"/> Severe renal failure
<input type="checkbox"/> Documented poor compliance with anticoagulant therapy	<input type="checkbox"/> High risk of recurrent falls
<input type="checkbox"/> Inability or significant difficulty maintaining therapeutic range	<input type="checkbox"/> Cognitive impairment
<input type="checkbox"/> Occupation/ lifestyle with increased bleeding risk	<input type="checkbox"/> Intolerance oral anticoagulation
<input type="checkbox"/> Need for prolonged dual anti platelet therapy	<input type="checkbox"/> Other situations for which anticoagulation is inappropriate
<input type="checkbox"/> Increased bleeding risk (eg. Thrombocytopenia, cancer, or risk or tumor associated bleeding in case of systemic anticoagulation)	



680

PT.

MR.#/P

DR.



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PROCEDURAL ELIGIBILITY

Patient has non Valvular Atrial Fibrillation (NVAf)	<input type="checkbox"/> YES <input type="checkbox"/> NO
Patient has an increased risk for stroke and systemic embolism based on CHA2DS2VASc score ≥ 3 and can be recommended for anticoagulation therapy	<input type="checkbox"/> YES <input type="checkbox"/> NO
Patient is suitable for short-term Coumadin (Warfarin) therapy but deemed unable to take long term anticoagulation therapy	<input type="checkbox"/> YES <input type="checkbox"/> NO
Patient is currently on anticoagulation therapy (AC), and it is currently on hold (If AC is on hold by an outside provider, a note from that provider is required)	<input type="checkbox"/> YES <input type="checkbox"/> NO
Patient is able to take Plavix and aspirin	<input type="checkbox"/> YES <input type="checkbox"/> NO
Therapeutic options and risks/benefits have been discussed with the patient including but not limited to estimated stroke risk and estimated bleed risk	<input type="checkbox"/> YES <input type="checkbox"/> NO

PROCEDURAL EXCLUSIONS:

Current Thrombus	<input type="checkbox"/> YES <input type="checkbox"/> NO	Prior atrial septal repair or device	<input type="checkbox"/> YES <input type="checkbox"/> NO
Prior Mechanical Valve	<input type="checkbox"/> YES <input type="checkbox"/> NO	Other anticoagulation indication such as DVT	<input type="checkbox"/> YES <input type="checkbox"/> NO
Women pregnant or planning to become	<input type="checkbox"/> YES <input type="checkbox"/> NO		

DECISION

Shared decision documented with independent, non-implanting physician: _____
(physician name)

Based on the above assessment, I believe this patient to be a candidate for the Watchman Device.

Physician Signature

Physician Printed Name

Date/ Time



PT.

MR./P

DR.