



North Dakota Cardiac System of Care STEMI, NSTEMI, & Acute Coronary Syndrome Guide

Tertiary Hospital One Call:

<u>Altru Health System – Grand Forks</u>

Phone: 701-780-5206 or 1-855-425-8781 Fax: 701-780-1097

CHI St. Alexius Health - Bismarck

Phone: 701-530-7699 or 1-877-735-7699 Fax: 701-530-7005

Essentia Health System - Fargo

Phone: 701-364-CALL (2255) or 844-865-CALL (2255) Fax: 701-364-8405

Sanford Health System-Bismarck

Phone: 1-855-550-1225 Fax: 701-323-5751

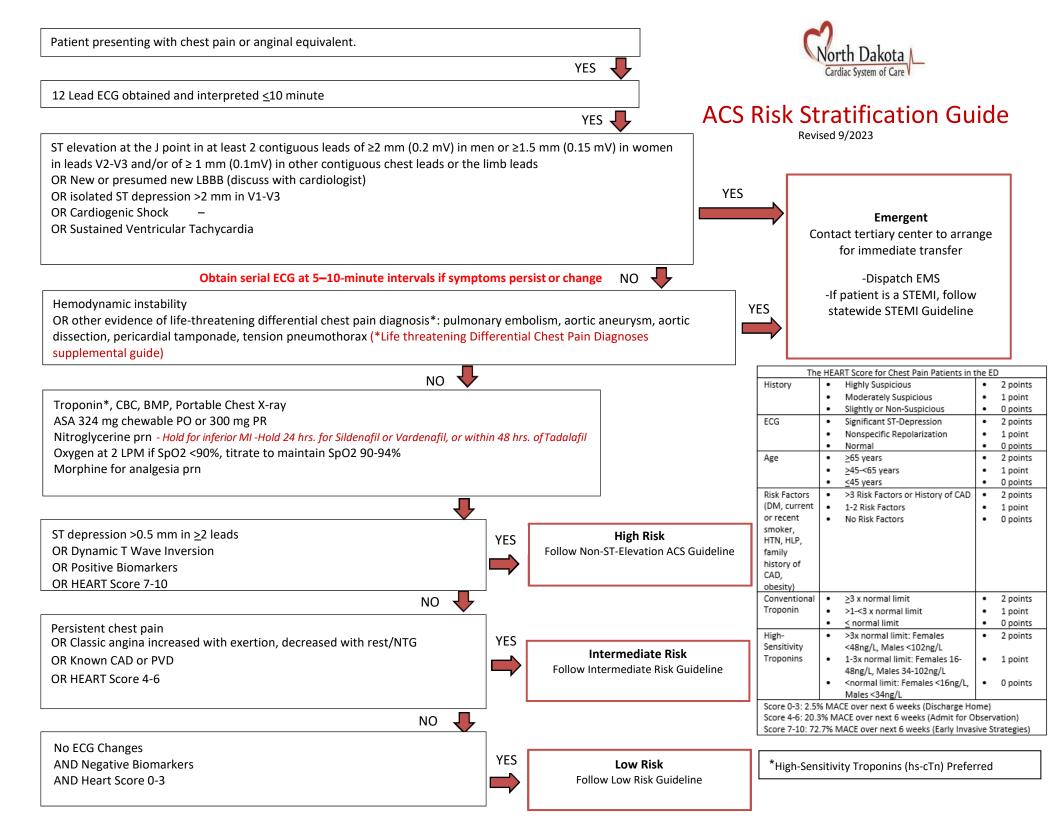
Sanford Health System- Fargo

Phone: 701-234-6304 or 1-877-647-1225 Fax: 701-234-7203

Trinity Health System - Minot

Phone: 701-857-3000 or 1-800-223-1596 Fax: 701-857-3260

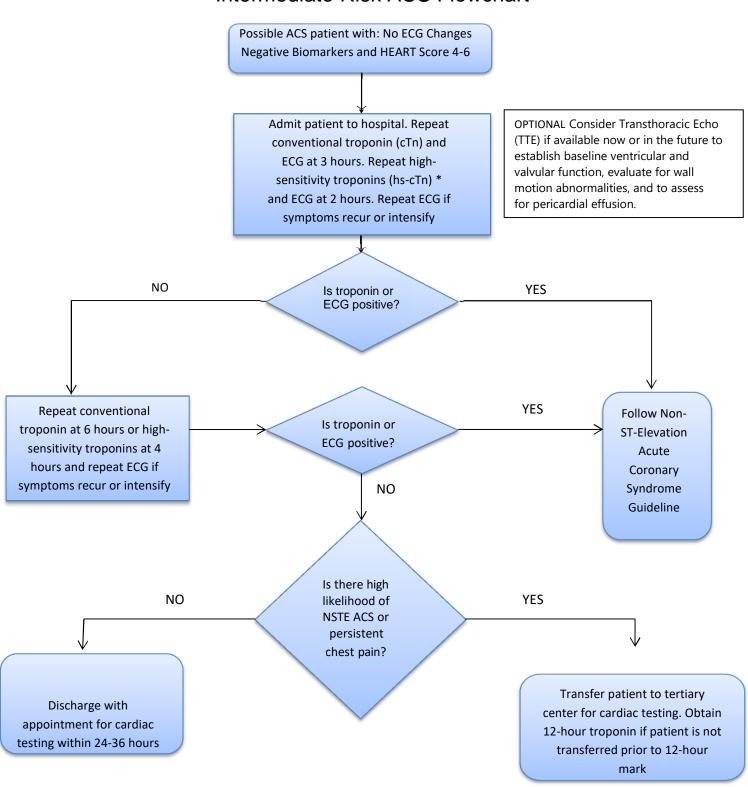
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Revised 9/2023

Intermediate Risk ACS Flowchart

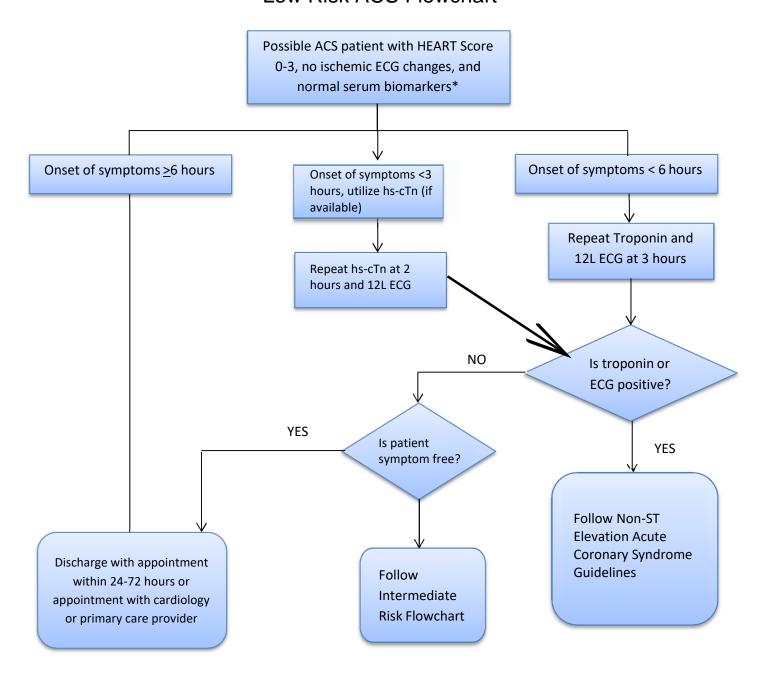


^{*}High-Sensitivity Troponins (hs-cTn) Preferred See reference for Cardiac Testing pg. 9-10



Revised 9/2023

Low Risk ACS Flowchart



^{*}High-Sensitivity Troponins (hs-cTn) Preferred







ND STEMI Inter-Hospital Transfer Guideline

(ST-Segment Elevation Myocardial Infarction)

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Ideal STEMI Treatment Goals:

- First Medical Contact-to-First ECG time <10 minutes unless pre-hospital ECG obtained
- All eligible patients receive Reperfusion (PCI or fibrinolysis) therapy
- Fibrinolytic-eligible patients with **Door-to-Needle** time < 30 minutes
- Reperfusion eligible patients transferred to a PCI receiving center with referring center
 Door in- Door out time (Length of Stay) < 45 minutes
- Referring Center ED **Door-to-PCI device time** ≤ 100 minutes (includes transport time)
- All STEMI patients without a contraindication receiving aspirin before ED discharge
- Upon Transfer Fax the following documents to the accepting facility: 12 L ECG, ED Record, Lab Results, Current Medication Record, ND STEMI documentation

Patients with a contraindication to transfer or PCI/Medical Therapy Option:

- Documentation of contraindication or Patient refusal to transfer for PCI or medical treatment
- Aspirin within 24 hours of hospital arrival, and aspirin at discharge
- Beta blocker at discharge
- High intensity statin at discharge
- P2Y12 (Plavix or Brilinta) at discharge
- STEMI patients who smoke receive smoking cessation counseling at discharge
- Scheduled Cardiology Consultation within 1-2 weeks at discharge
- Cardiac Rehabilitation referral at discharge

ND STEMI Guideline (ST-Segment Elevation Myocardial Infarction)

Revised 9/2023







Diagnostic Criteria for STEMI

- ST elevation at the J point in at least 2 contiguous leads of ≥2 mm (0.2 mV) in men or ≥1.5 mm (0.15 mV) in women in leads V2–V3 and/or of ≥1 mm (0.1 mV) in other contiguous chest leads or the limb leads.
- New or presumably new LBBB at presentation occurs infrequently, may interfere with ST-elevation analysis, and should not be considered diagnostic of acute
 myocardial infarction (MI) in isolation. If doubt persists, immediate referral for invasive angiography may be necessary. Consult with Cardiology.
- ECG demonstrates evidence of ST depression suspect of a Posterior MI consult with PCI receiving center
- If initial ECG is not diagnostic but suspicion is high for STEMI obtain serial ECG at 5–10-minute intervals



ACTIVATE STEMI ALERT at Receiving PCI Hospital

STANDARD ORDERS & LABS							
□ Apply Continuous Cardiac Monitor							
□ Insert (2) peripheral IV large bore Saline lock							
□ Troponin (hs-cTn preferred)							
□ВМР	□ Magnesium						
□ CBC	□ Glucose						
□INR	□aPTT						
-Do not delay transfer awaiting results							

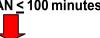
CONSIDER: Estimated transfer time in minutes to PCI facility take into account first medical contact time: Air:and/or Ground:						
ASSESS: SymptomOnset Date:Time:						
Code Status □ Full Code □ DNR -If patient wishes to remain DNR Status, consult receiving facility MD prior to initiation of transfer						
REVIEW: Thrombolytic Contraindications Page 7						

Choose One Reperfusion Pathway

OPTIONAL MEDICATION

- □ Nitroglycerin IV or 0.4 mg SL
- -Hold for inferior MI
- -Evaluate if erectile dysfunction or pulmonary hypertension medications taken in the past 24 hours including: Sildenafil (Viagra, Revatio), Vardenafil (Levitra, Staxyn), or Avanafil (Stendra), Tadalafil (Cialis, Adcirca). -Hold nitrates for 48 hours following the last dose.
- ☐ Analgesia as needed
- □ **Ondansetron** (Zofran) 4 mg POor IV
- □ Metoprolol Tartrate □ 25 mg PO CONTRAINDICATION FOR METOPROLOL TARTATE: <u>Do not give</u> if any of the following: signs of heart failure or shock, heart rate less than 60 or more than 110, systolic blood pressure less than 100-, second- or third-degree heart block, severe asthma, or reactive airway disease

Time of First Medical Contact to PCI arrival Expected to be LESS THAN ≤ 100 minutes



PRIMARY PCI Direct to CATH LAB for Emergent PCI

- □ **Aspirin** 324 mg chewed or 300mg rectally
- □ Ticagrelor (Brilinta) 180 mg PO preferred OR
 - □ Clopidogrel (Plavix) 600 mg PO
 - -DO NOT give both Plavix & Brilinta
- □ **Heparin IV Bolus** (60 Units/kg, max 4,000 Units)
- □ **Transport** patient <u>directly</u> to Cath Lab for PCI (Percutaneous Coronary Intervention)
 - Goal Arrival to Departure < 30 minutes unless awaiting air transport
- □ Oxygen as needed to keep SpO2 90-94%

-DO NOT give Thrombolytics TNKase, rPA, or TPA

Time of First Medical Contact to PCI arrival anticipated to be GREATER THAN > 100 minutes



THROMBOLYTIC Therapy

- □ **Aspirin** 324 mg chewed or 300mg rectally
- □ **Tenecteplase IV (TNKase**) per attached protocol
 - Facility Arrival to lytic administration goal LESS THAN < 30 minutes
- □ Plavix 300 mgPO

-If patient > 75 yrs. consult with cardiologist and reduce dosage to 75 mg PO

- □ **Heparin IV Bolus** (60 Units/kg, max 4,000 Units)
- □ **Heparin IV Drip** (12 Units/kg/hr., max 1,000 Units/hr.)
- ☐ **Transport** patient urgently <u>directly</u> to PCI capable hospital
 - Goal Arrival to Departure < 45 minutes unless awaiting air transport







	ND STEINI	(51-Segment	Elevation	wyocardiai iiii	arction) Guid		; 		
Tened	cteplase (TNKase) I	Dosing	Weight:	lb. kg	Height:	in.		Age:	yrs.
Patient weight (kg)	TNK (mg)	TNK (mL)							
Less than 60 kg	30 mg	6 mL	Allergies:						
60 or more but less than 70	35 mg	7 mL	Medication		Dose	Time S	Start	Time Stop	RN (Initials)
70 or more but less than 80	40 mg	8 mL	Aspirin (81 m	ng chew x 4)	324 mg				
80 or more but less than 90	45 mg	9 mL							
90 or more kg	50 mg	10 mL	Ticagrelor (Brili (PPCI therapy		180 mg	i			
ABSOLUTE CONTRAINDICATIONS FOR FIBRINOLYSIS (TNK) IN STEMI		-Do not give tion Plavix together	eagrelor (Brilinta) and						
Any prior intracranial hemorrhage Known structural cerebral vascular lesion (e.g.,		Clopidogrel (PPCI therapy		600 mg					
arteriovenous malformation) 3. Known malignant intracranial neoplasm (primary or metastatic) 4. Ischemic stroke within 3 months except acute ischemic		Clopidogrel (Lytic therapy If >75 y/o, gir	dose	300 mg 75mg					
stroke within 4.5 hours 5. Suspected aortic dis	section		Heparin IV B		Units				
6. Active bleeding or ble 7. Significant closed-he	eeding diathesis (e		PCI Dose 60 U	/kg, max 4000 Units					
Significant closed-ne Chest Pain/Symptor			Lytic Dose 60 t Heparin IV Ir	J/kg, max 4000 Units nfusion	Units/hr.				
RELATIVE CONTRAII (TNK) IN STEMI	NDICATIONS FOR	R FIBRINOLYSIS:	Lytic Dose 12 U/hr.	U/kg/hr. max 1000					
 History of chronic, se Severe uncontrolled (SBP more than 180 	hypertension on pi	resentation	•	e (TNKase) IV agrelor (Brilinta) with	mg (= mL)				
History of prior ischer	mic stroke more tha	an 3 months,	Nitroglycerin		0.4				
dementia, or known int contraindications			-Erectile Dysfu past 24 hrs. \Box	Inction Medication within	mg 0.4 mg				
5. Maior augus (within last 2 weeks)		•		0.4 mg					
Recent internal bleeding (within last 2-4 weeks)		Nitroglycerin IV Infusion		mcg/min					
0. Ctrantakingga (aniatranlaga, prior aypagura (mara than F		Morphine Su	Ilfate IV	mg	<u> </u>				
days ago) or prior allergic reaction to these agents		Fentanyl IV Ondansetron	(Zofran) PO	mcg 4 mg				_	
10. Active peptic ulcer		Ondanselion	(Zoliali) i O	Ting					
		Ondansetron (Zofran) IV		4 mg					
		Metoprolol Tartrate 25 mg or 50 mg PO		mg					
Notes:			<u>PU</u>						
RN Name (Print):		RN Signature:		RN Initials:	Date:		Т	ime:	
Data Elements in the ND State STEMI Registry			☐ Call Report w	hen patient leaves yo				n update	
1. Initial Sympton	om Onset Time	e			departure tim				
Date:	Time:			 Copy ECG, ED physician and Nurses documentation and send with patient – DO NOT delay transport 					
2. FMC EMS Ag	ency:				•				-
3. Referring Ho	spital Arrival ((Door-in)			ork to referring Hospita Order, Notes, Medica				
Date:	Time:			·					
4. Referring Hos	spital 1st ECG	Time:	_	D. C. A.					
5. STEMI ECG T	ime:			Patient Name:					
6. STEMI Activa	tion (STEMI R	eceiving contacte	ed)						
Date:	Time:								
7. Referring Hos	spital Departu	re (Door-out)							
Date:	Time:								
8. Transfer EMS	Agency Nam	e:							



Non-ST-Elevation Acute Coronary Syndrome Guideline

Diagnostic Criteria

- New >0.5 mm ST segment depression or new >2 mm anterior T-wave inversion and/or positive biomarkers
- If patient experiences persistent or worsening symptoms obtain serial ECGs at 15–30-minute intervals to monitor for new onset ST elevation



- Contact PCI Center to arrange for transfer of patient
 - Dispatch EMS service once transfer is confirmed



ACC/AHA Guideline Based Treatment

Standard orders and labs

- Assess vital signs stat, repeat per unit routine
- Continuous cardiac monitoring (telemetry)
- Insert 1-2 large bore peripheral saline lock IV(s)
- Obtain following labs: CBC, BMP, PT/INR, PTT, Troponin I at 3 and 6 hours or hs-cTn at 2 and 4 hours (if stay is extended)
- Oxygen at 2 LPM if SpO2<90%, titrate to maintain SpO2 90-94%

Standard Medications-Discuss with accepting provider prior to administration

- Aspirin 324 mg (chewable non-enteric coated 81 mg x 4) orally stat x 1 or if patient is unable to swallow give: Aspirin 300 mg rectally
- Ticagrelor (Brilinta) 180 mg orally stat x 1 OR
 Clopidogrel (Plavix) 300 mg orally stat x 1 (do not give both Ticagrelor and Clopidogrel)

Heparin-Adjust dose according to weight-based protocol if patient stay is extended

- Heparin 60 units/kg IV bolus (max bolus 4000 units)
- Heparin IV drip 12 units/kg/hr. (max 1000 units/hr.)



Optional Labs

BNP, HCG

Optional Medications

- Nitroglycerine 0.4 mg SL every 5 minutes x 3 as needed for chest discomfort
- Nitroglycerine IV continuous infusion as needed for chest pain
- -Hold Nitro if recent phosphodiesterase inhibitor, 24 h of sildenafil or vardenafil, or within 48 h of tadalafil.
- For severe uncontrolled pain, consider use of **Morphine** or other narcotic analgesic of choice IV as needed.
- Ondansetron (Zofran) 4 mg IV as needed for nausea/vomiting x 1
- Metoprolol Tartrate (Lopressor) 25 mg orally x 1
- -Hold Beta Blocker if Signs of heart failure or shock, SBP less than 110, Heart rate less than 60 bpm or heart block, severe asthma, or reactive airway disease



- Transfer patient to PCI center for possible early invasive strategy
- Send with or fax the following documents to accepting facility:12L ECG, ED record, lab results, current medication record, EMS record

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Revised 9/2023

Cardiac Testing

Revised 9/2023

Diagnostic Testing may include:

- Exercise ECG Coronary Computed Tomography Angiography
- Echocardiography/Stress Echocardiography Invasive Coronary Angiography
- Stress Nuclear (PET or SPECT) Myocardial Perfusion Imaging
- Cardiovascular Magnetic Resonance Imaging

Anatomic Testing may include:

- Coronary Computed Tomography Angiography
- Invasive Coronary Angiography

For further direction on choosing proper provocative testing, see attached algorithms from the 2021 AHA/ACC/ASE/CHEST/SAEM/SC CT/SCMR Chest Pain Guideline and cardiac testing algorithms:

- Patients With Suspected ACS at Intermediate Risk With No Known CAD
- Patients With Suspected ACS at Intermediate Risk With Known CAD

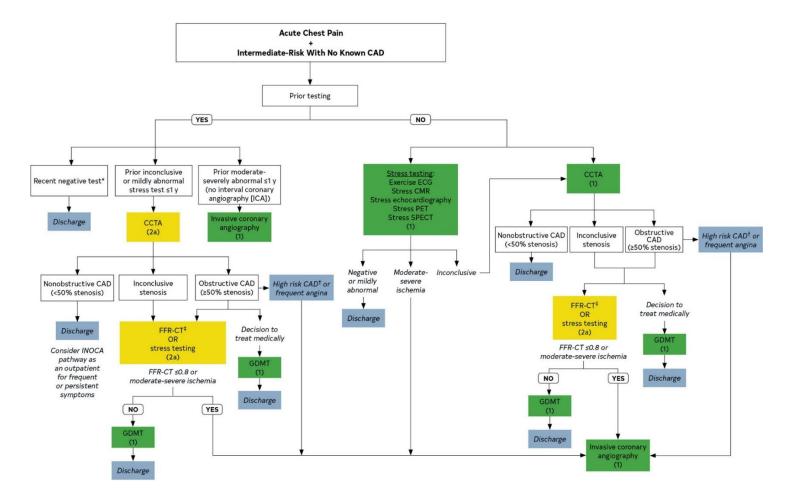


Figure 9. Evaluation Algorithm for Patients With Suspected ACS at Intermediate Risk With No Known CAD Test choice should be guided by local availability and expertise. *Recent negative test: normal CCTA ≤2 years (no plaque/no stenosis) OR negative stress test ≤1 year, given adequate stress. †High-risk CAD means left main stenosis ≥ 50%; anatomically significant 3-vessel disease (≥70% stenosis). ‡For FFR-CT, turnaround times may impact prompt clinical care decisions. However, the use of FFR-CT does not require additional testing, as would be the case when adding stress testing. CAD indicates coronary artery disease; CCTA, coronary CT angiography; CMR, cardiovascular magnetic resonance imaging; CT, computed tomography; FFR-CT, fractional flow reserve with CT; GDMT, guideline-directed medical therapy; ICA, invasive coronary angiography; INOCA, ischemia and no obstructive coronary artery disease; PET, positron emission tomography; and SPECT, single-photon emission CT.

Cardiac Testing

Revised 9/2023

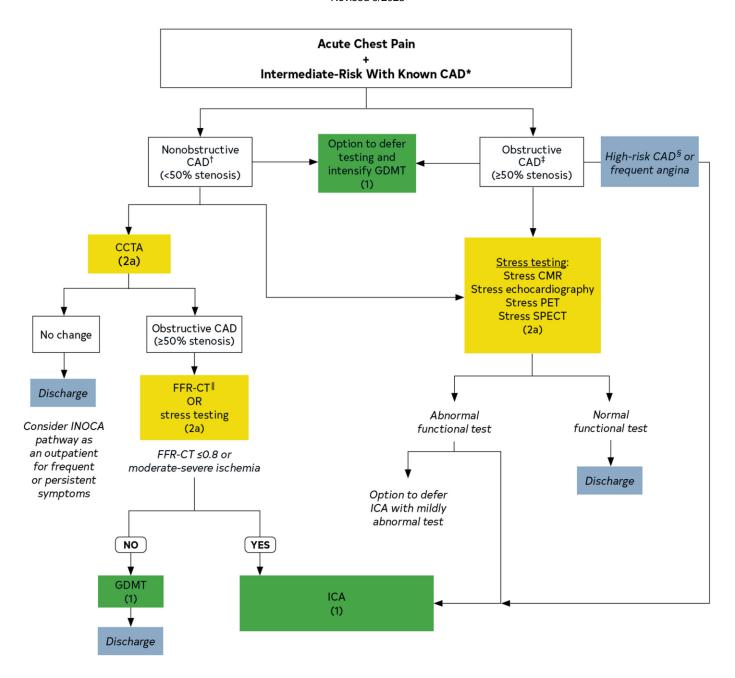


Figure 10. Evaluation Algorithm for Patients With Suspected ACS at Intermediate Risk With Known CAD

Test choice should be guided by local availability and expertise. *Known CAD is prior MI, revascularization, known obstructive or nonobstructive CAD on invasive or CCTA. †If extensive plaque is present a high-quality CCTA is unlikely to be achieved, and stress testing is preferred ‡Obstructive CAD includes prior coronary artery bypass graft/percutaneous coronary intervention. §High-risk CAD means left main stenosis ≥50%; anatomically significant 3-vessel disease (≥70% stenosis). ∥FFR-CT turnaround times may impact prompt clinical care decisions. ACS indicates acute coronary syndrome; CAD, coronary artery disease; CCTA, coronary CT angiography; CMR, cardiovascular magnetic resonance; CT, computed tomography; FFR-CT, fractional flow reserve with CT; GDMT, guideline-directed medical therapy; ICA, invasive coronary angiography; INOCA, ischemia and no obstructive coronary artery disease; PET, positron emission tomography; and SPECT, single-photon emission CT.

Reference: 2021 AHA/ACC/ASE/CHEST/SAEM/SCCT/SCMR Guideline for the Evaluation and Diagnosis of Chest Pain: A Report of the American College of Cardiology/American Heart Association Joint Committee on Clinical Practice Guidelines J Am Coll Cardiol. Oct 28, 2021. Epublished DOI: 10.1016/j.jacc.2021.07.053