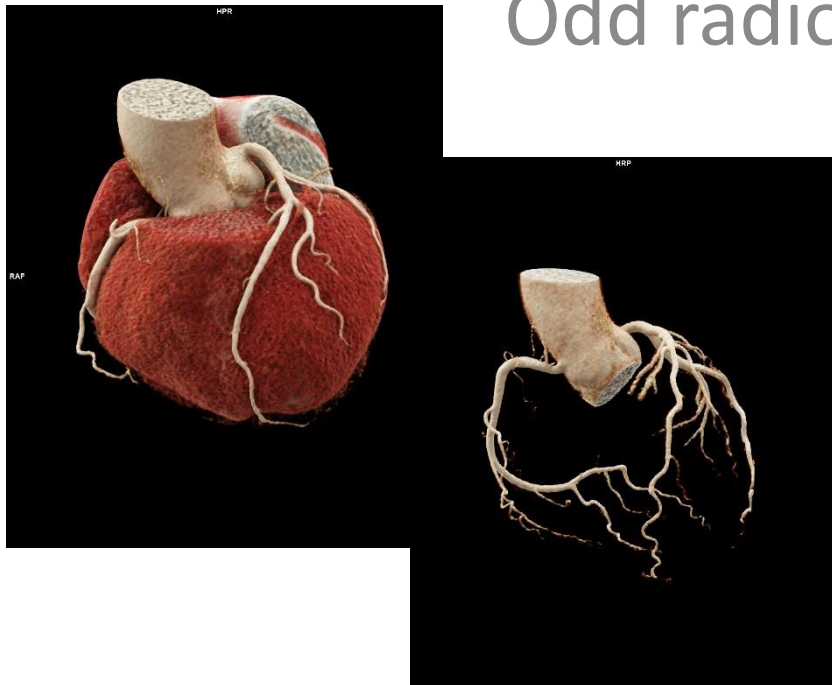


Akútna bolesť na hrudi a CTCA

Maroš Daxner

Odd rádiológie SÚSCCH





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

[Clinical Practice Guideline: Full Text](#)

Writing Committee Members, Martha Gulati, Phillip D. Levy, Debabrata Mukherjee, Ezra Amsterdam, ... [SEE ALL AUTHORS](#) ▾

J Am Coll Cardiol. 2021 Nov, 78 (22) e187–e285

2020 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation: The Task Force for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation of the European Society of Cardiology (ESC)

Jean-Philippe Collet , Holger Thiele , Emanuele Barbato, Olivier Barthélémy, Johann Bauersachs, Deepak L Bhatt, Paul Dendale, Maria Dorobantu, Thor Edvardsen, Thierry Folliguet ... [Show more](#)

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it

Robert A Byrne ✉, Xavier Rossello, J J Coughlan, Emanuele Barbato, Colin Berry, Alaide Chieffo, Marc J Claeys, Gheorghe-Andrei Dan, Marc R Dweck, Mary Galbraith ...

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CLASS (STRENGTH) OF RECOMMENDATION	
CLASS 1 (STRONG)	Benefit >>> Risk
Suggested phrases for writing recommendations: <ul style="list-style-type: none"> • Is recommended • Is indicated/useful/effective/beneficial • Should be performed/administered/other • Comparative-Effectiveness Phrases†: <ul style="list-style-type: none"> – Treatment/strategy A is recommended/indicated in preference to treatment B – Treatment A should be chosen over treatment B 	
CLASS 2a (MODERATE)	Benefit >> Risk
Suggested phrases for writing recommendations: <ul style="list-style-type: none"> • Is reasonable • Can be useful/effective/beneficial • Comparative-Effectiveness Phrases†: <ul style="list-style-type: none"> – Treatment/strategy A is probably recommended/indicated in preference to treatment B – It is reasonable to choose treatment A over treatment B 	
CLASS 2b (WEAK)	Benefit ≥ Risk
Suggested phrases for writing recommendations: <ul style="list-style-type: none"> • May/might be reasonable • May/might be considered • Usefulness/effectiveness is unknown/unclear/uncertain or not well-established 	
CLASS 3: No Benefit (MODERATE) (Generally, LOE A or B use only)	Benefit = Risk
Suggested phrases for writing recommendations: <ul style="list-style-type: none"> • Is not recommended • Is not indicated/useful/effective/beneficial • Should not be performed/administered/other 	
Class 3: Harm (STRONG)	Risk > Benefit
Suggested phrases for writing recommendations: <ul style="list-style-type: none"> • Potentially harmful • Causes harm • Associated with excess morbidity/mortality • Should not be performed/administered/other 	

LEVEL (QUALITY) OF EVIDENCE‡	
LEVEL A	
<ul style="list-style-type: none"> • High-quality evidence‡ from more than 1 RCT • Meta-analyses of high-quality RCTs • One or more RCTs corroborated by high-quality registry studies 	
LEVEL B-R	(Randomized)
<ul style="list-style-type: none"> • Moderate-quality evidence‡ from 1 or more RCTs • Meta-analyses of moderate-quality RCTs 	
LEVEL B-NR	(Nonrandomized)
<ul style="list-style-type: none"> • Moderate-quality evidence‡ from 1 or more well-designed, well-executed nonrandomized studies, observational studies, or registry studies • Meta-analyses of such studies 	
LEVEL C-LD	(Limited Data)
<ul style="list-style-type: none"> • Randomized or nonrandomized observational or registry studies with limitations of design or execution • Meta-analyses of such studies • Physiological or mechanistic studies in human subjects 	
LEVEL C-E0	(Expert Opinion)
<ul style="list-style-type: none"> • Consensus of expert opinion based on clinical experience 	

COR and LOE are determined independently (any COR may be paired with any LOE).

A recommendation with LOE C does not imply that the recommendation is weak. Many important clinical questions addressed in guidelines do not lend themselves to clinical trials. Although RCTs are unavailable, there may be a very clear clinical consensus that a particular test or therapy is useful or effective.



* The outcome or result of the intervention should be specified (an improved clinical outcome or increased diagnostic accuracy or incremental prognostic information).

† For comparative-effectiveness recommendations (COR 1 and 2a; LOE A and B only), studies that support the use of comparator verbs should involve direct comparisons of the treatments or strategies being evaluated.

‡ The method of assessing quality is evolving, including the application of standardized, widely-used, and preferably validated evidence grading tools; and for systematic reviews, the incorporation of an Evidence Review Committee.

COR indicates Class of Recommendation; E0, expert opinion; LD, limited data; LOE, Level of Evidence; NR, nonrandomized; R, randomized; and RCT, randomized controlled trial.

2020 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation: The Task Force for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation of the European Society of Cardiology (ESC) ^{FREE}

Jean-Philippe Collet , Holger Thiele , Emanuele Barbato, Olivier Barthelemy, Johann Bauersachs, Deepak L Bhatt, Paul Dendale, Maria Dorobantu, Thor Edvardsen, Thierry Folliguet ... Show more

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[Author Notes](#)

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Published: 25 August 2023

Definition

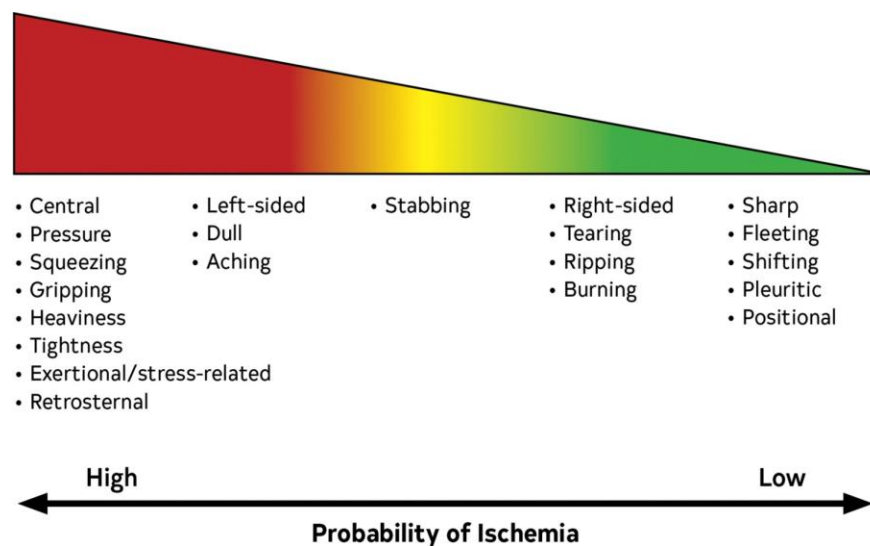
Wording to use

Class	Definition	Wording to use
Class I	Evidence and/or general agreement that a given treatment or procedure is beneficial, useful, effective.	Is recommended or is indicated
Class II	Conflicting evidence and/or a divergence of opinion about the usefulness/efficacy of the given treatment or procedure.	
Class IIa	Weight of evidence/opinion is in favour of usefulness/efficacy.	Should be considered
Class IIb	Usefulness/efficacy is less well established by evidence/opinion.	May be considered
Class III	Evidence or general agreement that the given treatment or procedure is not useful/effective, and in some cases may be harmful.	Is not recommended

Level of evidence A	Data derived from multiple randomized clinical trials or meta-analyses.
Level of evidence B	Data derived from a single randomized clinical trial or large non-randomized studies.
Level of evidence C	Consensus of opinion of the experts and/or small studies, retrospective studies, registries.

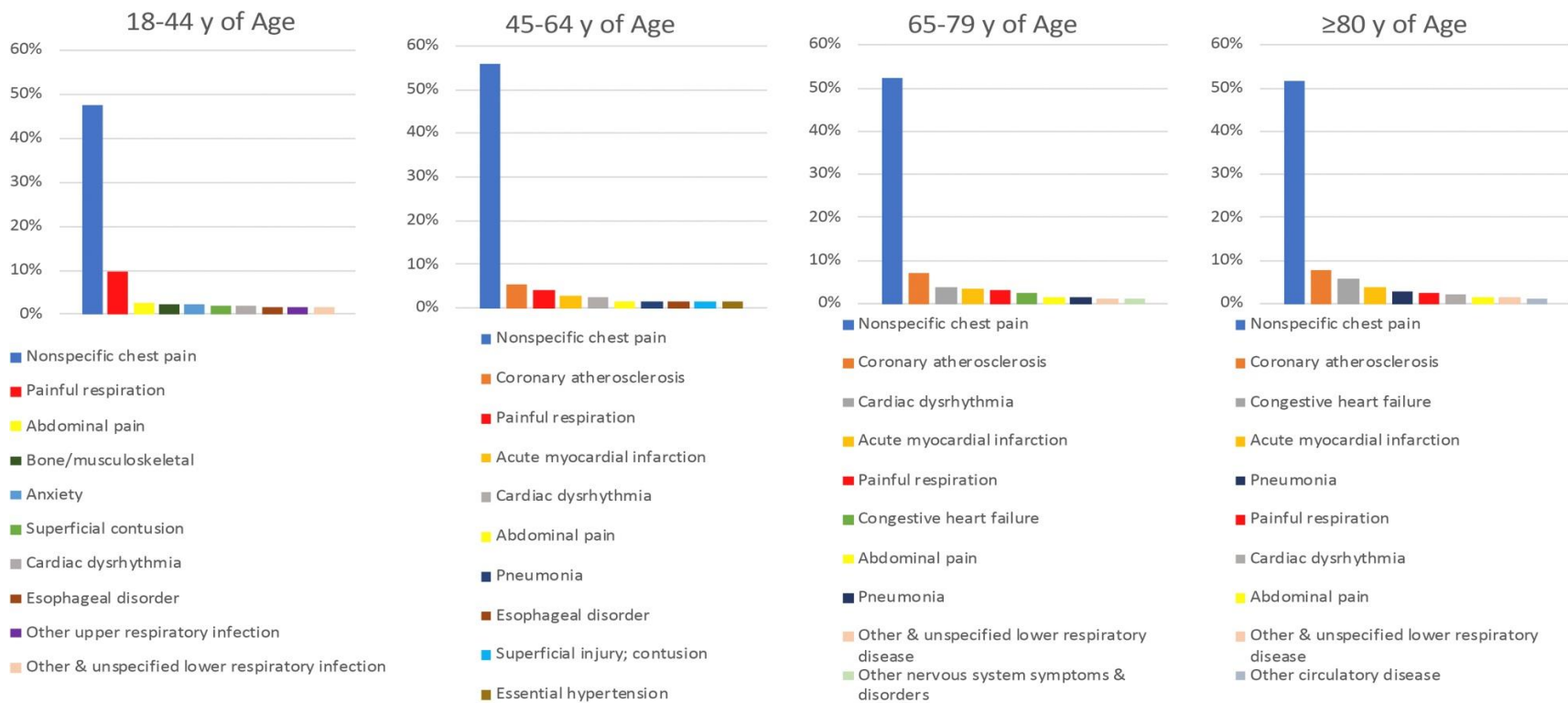


Hlavným príznakom u pacientov s podozrením na AKS je akútna bolesť/diskomfort na hrudníku



“kardiálna” “pravdepodobne kardiálna,” a “nekardiálna”
“typická” a “atypická”

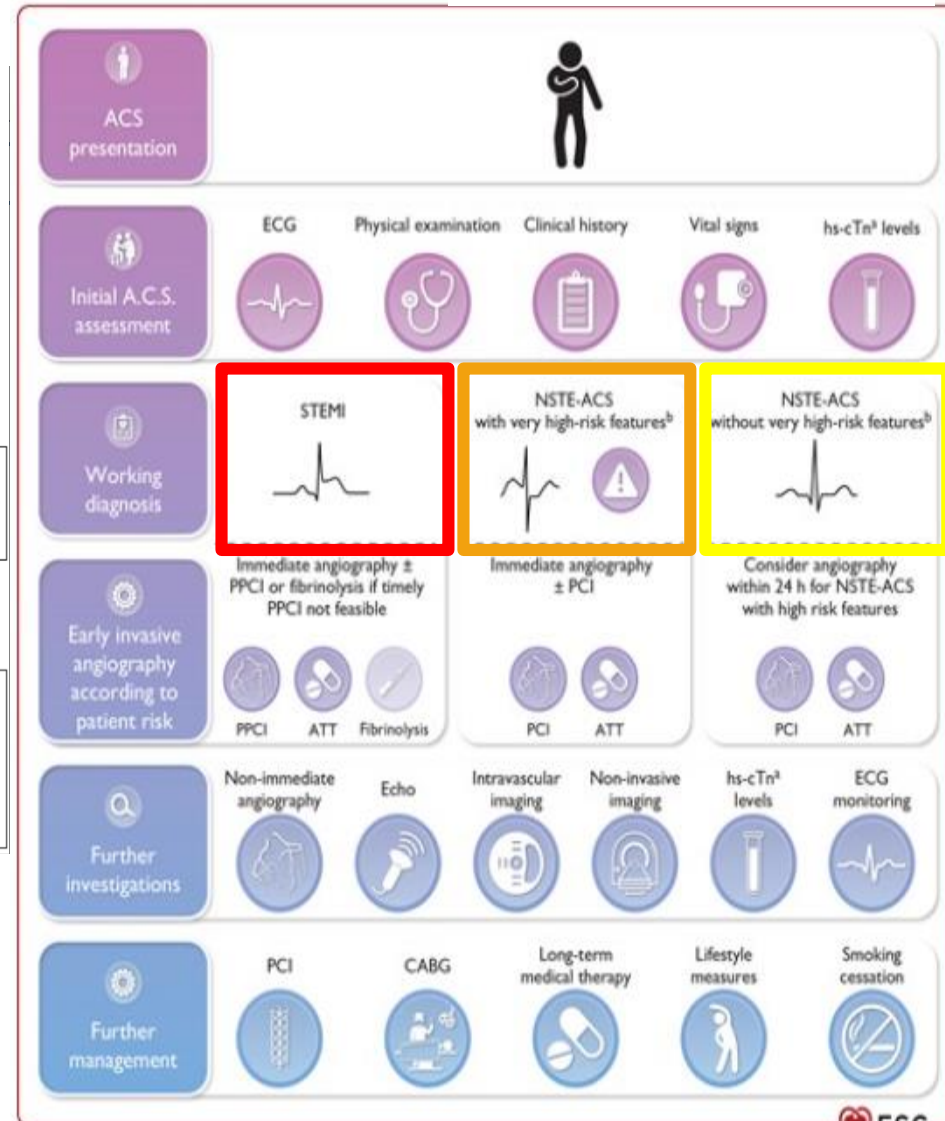
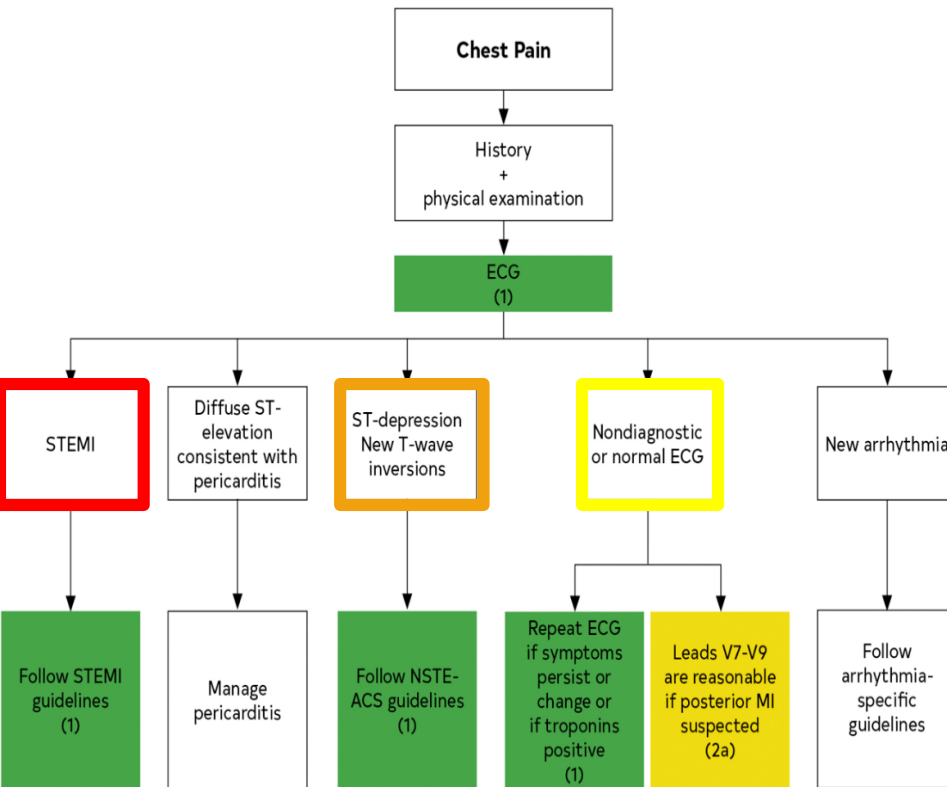




Po úrazoch druhá najčastejšia príčina príchodu UP u dospelých v US viac ako 6,5 milióna návštev, čo je 4,7 % zo všetkých návštev na UP. Zo všetkých pacientov na UP s bolesťou na hrudníku iba 5,1 % má akútny koronárny syndróm

COR	LOE	Recommendations
1	B-NR	1. Women who present with chest pain are at risk for underdiagnosis, and potential cardiac causes should always be considered (1-7).
1	B-NR	2. In women presenting with chest pain, it is recommended to obtain a history that emphasizes accompanying symptoms that are more common in women with ACS (1-7).

ISCHEMIA trial - Ženy - menšia pravdepodobnosť, že budú mať včasnú a primeranú starostlivosť.
- sú symptomatickejšie ako muži s výskytom prodromálnych symptómov pri stredne ťažkej a ťažkej ischémii

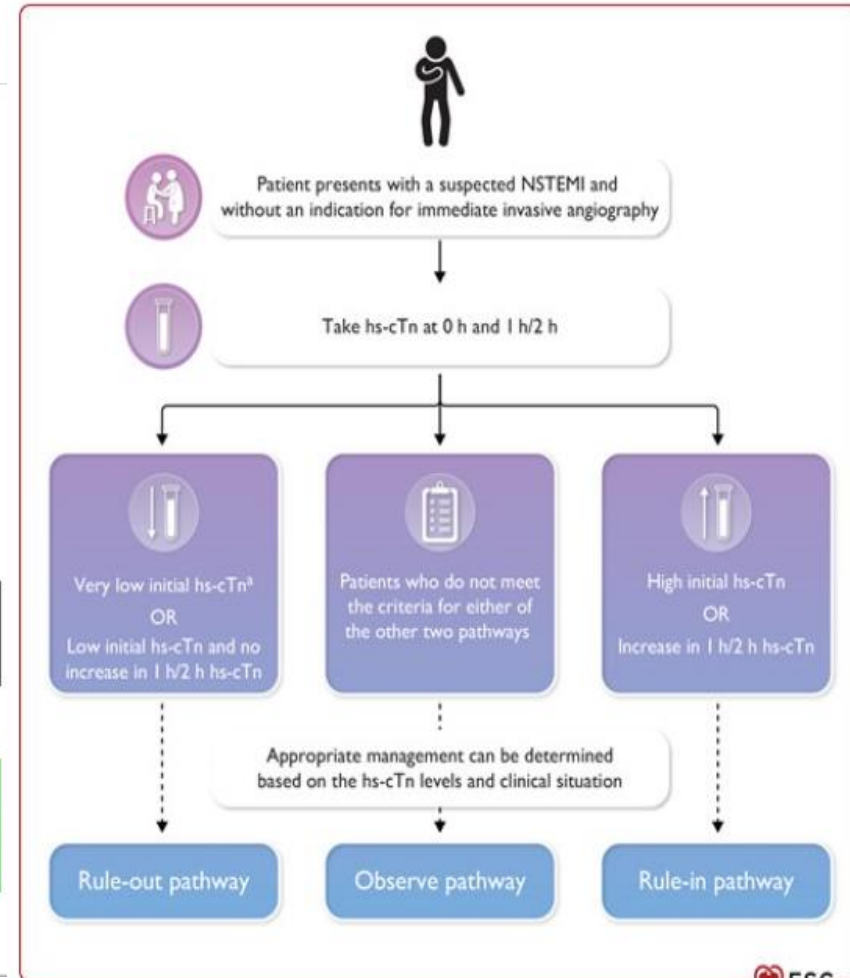
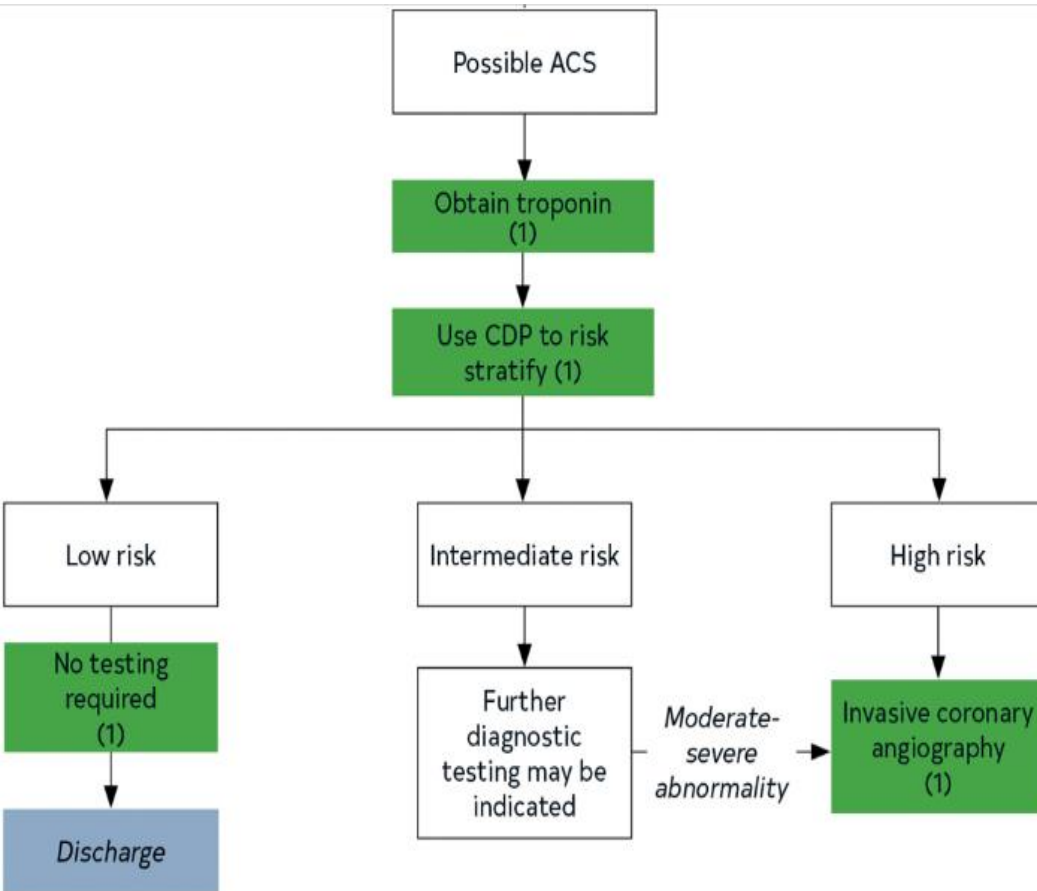




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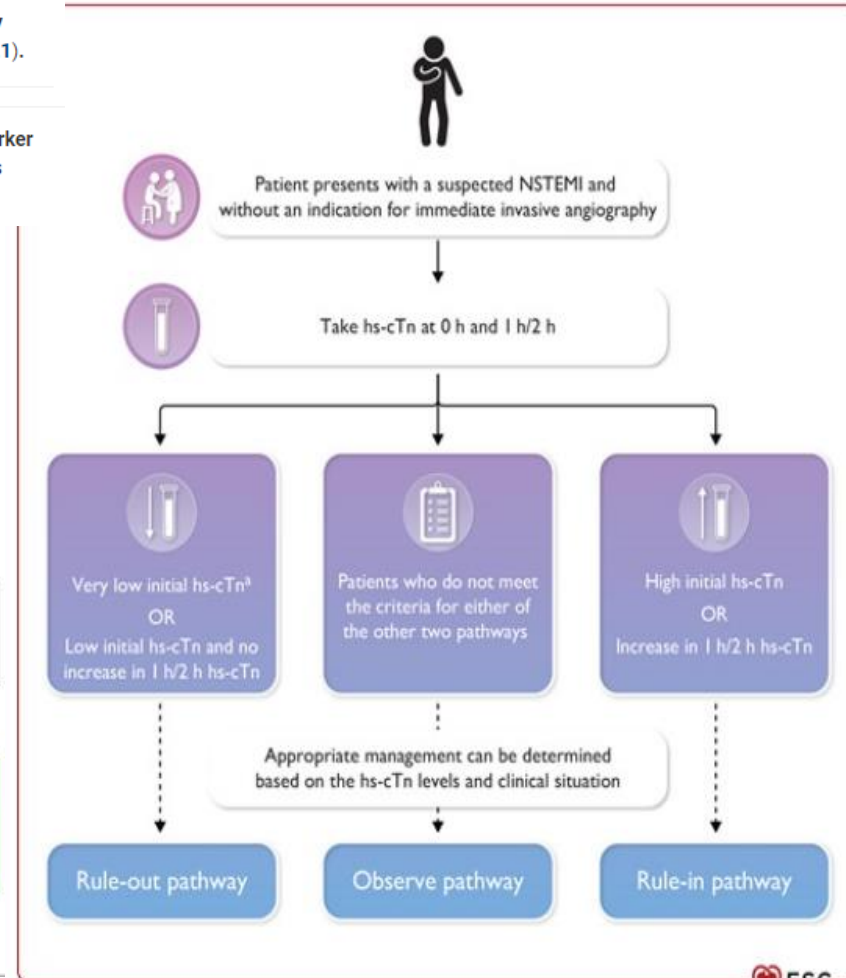
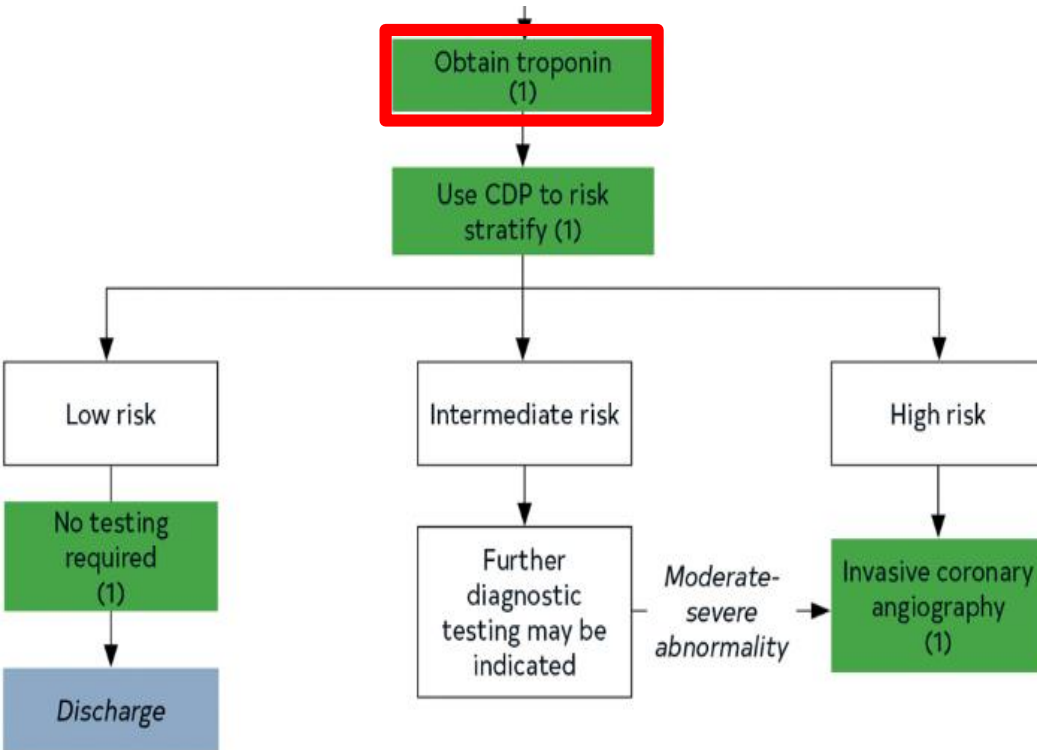
American Heart Association



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1 B-NR 1. In patients presenting with acute chest pain, serial cTn I or T levels are useful to identify abnormal values and a rising or falling pattern indicative of acute myocardial injury (1-21).

1 B-NR 2. In patients presenting with acute chest pain, high-sensitivity cTn is the preferred biomarker because it enables more rapid detection or exclusion of myocardial injury and increases diagnostic accuracy (17,21-25).

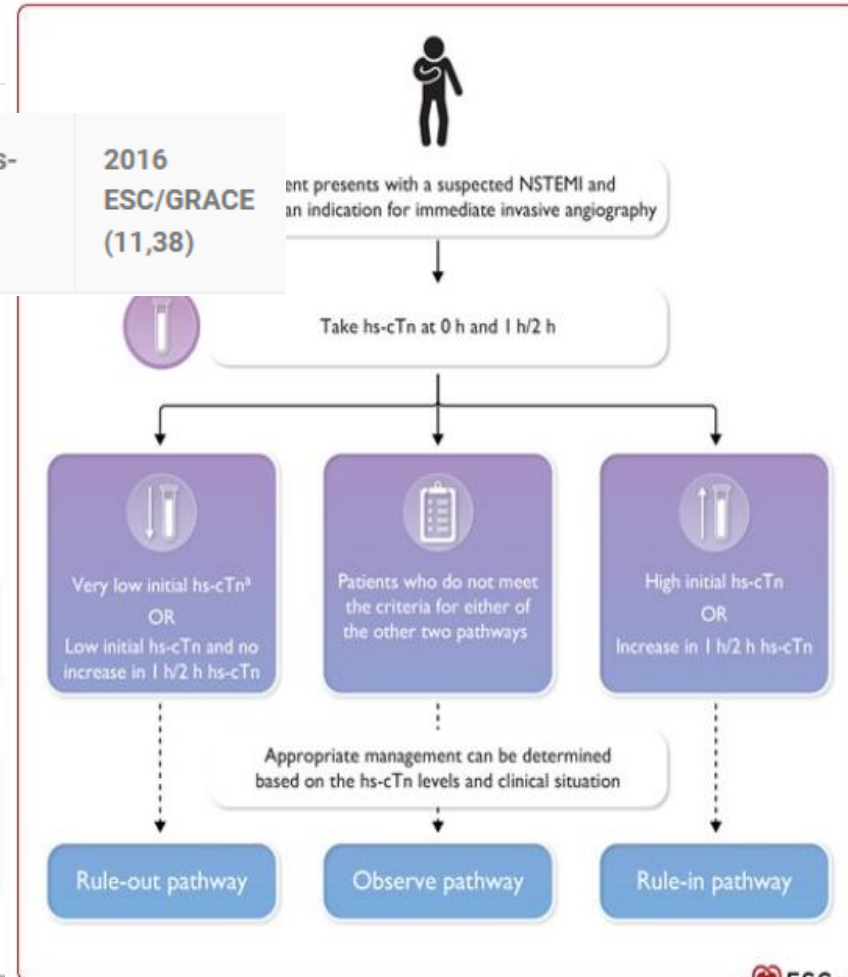
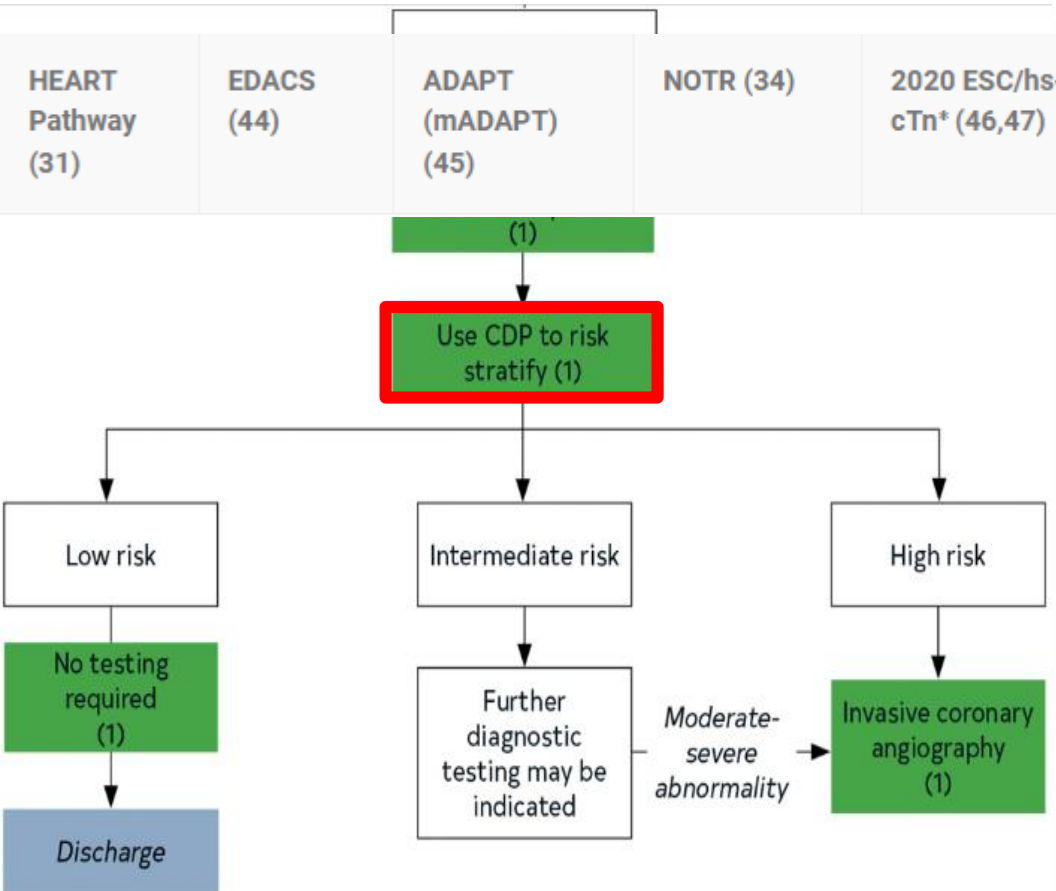




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Približne 582 000 výsledkov (0,23 sekundy)

University of Washington
http://depts.washington.edu/uw...

UW Medicine ED Chest Pain Pathway

11. 2. 2018 — Tool to help ED physicians decide on pathway for chest pain patients, University of Washington Medical Center.

UWMC ED Chest Pain Pathway
Flowchart of this pathway on OCCAM

STEMI or Sudden Death No

HEART Score for Early Discharge in Acute Chest Pain*

History
Posterior chest pain, pressure, radiation to jaw/left shoulder/arms, duration 5-15 min, initiated by exertion/indignation, perspiration, nausea/vomiting, radiation on nitroglycerin, patient recognizes symptoms, Lower risk features of chest pain include: well localized, sharp, non-exertional, no diaphoresis, no nausea or vomiting, and reproducible with palpation.

EKG
+1 point: No ST depression but LBBB, LVH, repolarization changes (see algorithm)
+2 points: ST depression/elevation not due to LBBB, LVH, or digoxin

Age
≤45
45-60
>60

Risk factors
+1PTV = hypercholesterolemia + DM + obesity (BMI >30 kg/m²)
+ smoking (current, or smoking cessation <3 mo) + positive family history (parent or sibling with CVD before age 65) + atherosclerotic disease, prior MI, PVD/CAD, CVA/TIA, or peripheral arterial disease

Initial troponin
Less than 0.04 ng/mL (normal)
0.04 to 0.08 ng/mL (1-2x normal)
Greater than 0.08 ng/mL (> 2x normal)

4 points
HEART Pathway Score

Intermediate risk
12-16.5% 6-week MACE
Additional risk stratification recommended
Next step depends on GRACE score

Additional Risk Stratification

Highly suspicious
EKG
+1 point: No ST depression but LBBB, LVH, repolarization changes (see algorithm)
+2 points: ST depression/elevation not due to LBBB, LVH, or digoxin

Normal
Non-specific repolarization disturbance
Significant ST depression

Age
≤45
45-60
>60

Risk factors
+ HTN + hypercholesterolemia + DM + obesity (BMI >30 kg/m²)
+ smoking (current, or smoking cessation <3 mo) + positive family history (parent or sibling with CVD before age 65) + atherosclerotic disease, prior MI, PVD/CAD, CVA/TIA, or peripheral arterial disease

1-2 risk factors
≥3 risk factors or history of atherosclerotic disease

Initial troponin
Less than 0.04 ng/mL (normal)
0.04 to 0.08 ng/mL (1-2x normal)
Greater than 0.08 ng/mL (> 2x normal)

5 points
HEART Pathway Score

Intermediate risk
12-16.5% 6-week MACE
Additional risk stratification recommended
Next step depends on GRACE score

Additional Risk Stratification

Slightly suspicious

Moderately suspicious

Highly suspicious

Normal
Non-specific repolarization disturbance
Significant ST depression

Age
≤45
45-60
>60

No known risk factors
1-2 risk factors
≥3 risk factors or history of atherosclerotic disease

Initial troponin
Less than 0.04 ng/mL (normal)
0.04 to 0.08 ng/mL (1-2x normal)
Greater than 0.08 ng/mL (> 2x normal)

Low/Intermediate Risk – Additional risk stratification appropriate

Initial Assessment: Intermediate Risk (HEART 4 pts, contributions: +2 History, +1 Age, +1 Additional) → Additional risk stratification recommended.
46 yr old male, w/ typical chest pain, no diabetes, dyslipidaemia, hypertensive, non smoker → 36% risk of obstructive CAD,
75 BPM, 105 mmHg, no diuretic, no renal failure, STEMI absent, normal troponin, no arrest at admission → 4% risk of in-hospital event (by GRACE 2.0)
Since risk of in-hospital event is 0-6%, expedited diagnostic testing is recommended.

High likelihood CAD No

Iodine contrast allergy No

Atrial Fibrillation No

Can walk on treadmill No

Bronchospastic lung disease No

Heart block No

Left bundle branch block No

Recommended Test:
Coronary CTA

Alternative Tests:
Vasodilator SPECT (NM Cardiac Stress Perfusion)

Dobutamine Stress Echocardiogram

Cardiology Consult

Initial Assessment: Intermediate Risk (HEART 5 pts, contributions: +2 History, +2 Age, +1 Troponin) → Additional risk stratification recommended.
70 yr old male, w/ typical chest pain, no diabetes, no dyslipidaemia, hypertensive, smoker → 72% risk of obstructive CAD,
85 BPM, 135 mmHg, no diuretic, no renal failure, STEMI absent, normal troponin, no arrest at admission → 6% risk of in-hospital event (by GRACE 2.0)
Since risk of in-hospital event is 0-6%, expedited diagnostic testing is recommended.

High likelihood CAD Yes

Iodine contrast allergy No

Atrial Fibrillation No

Can walk on treadmill No

Bronchospastic lung disease No

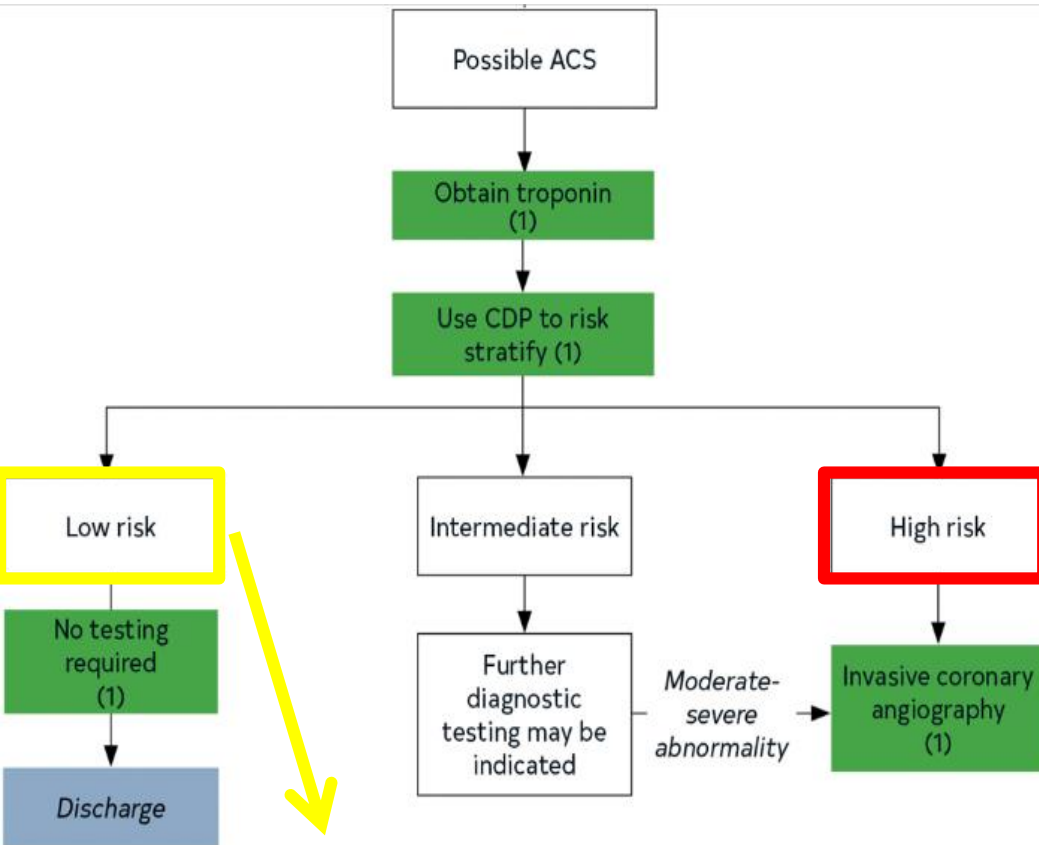
Heart block No

Left bundle branch block No

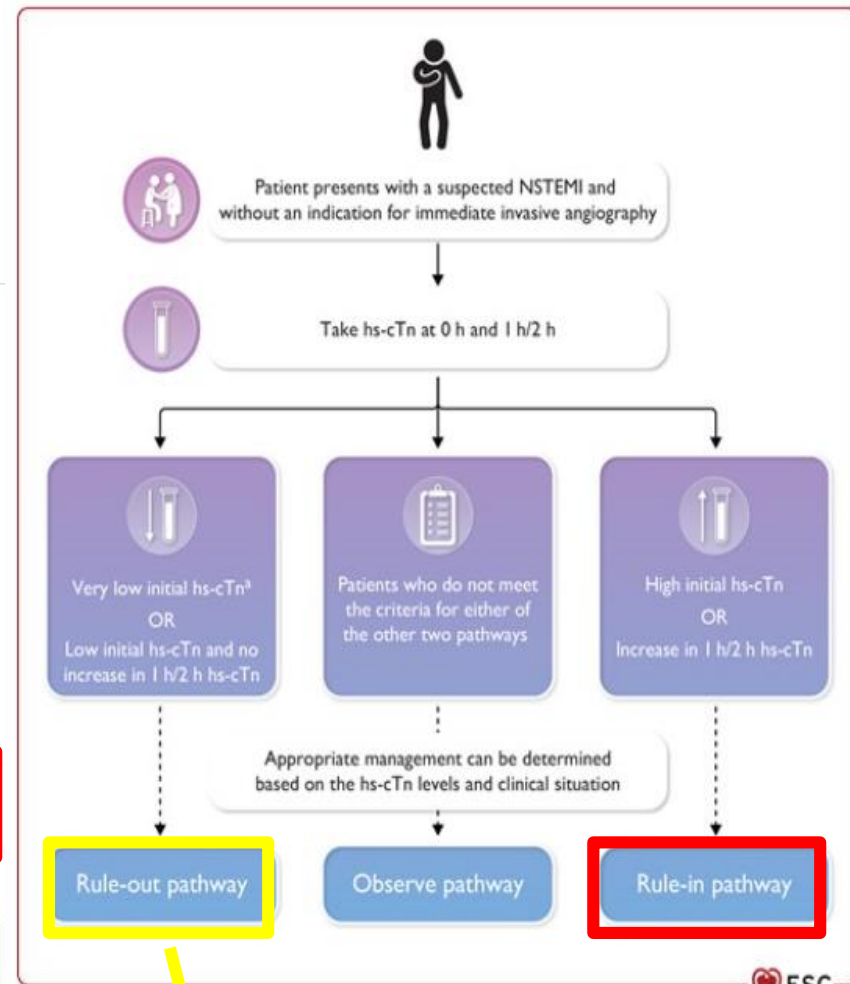
Recommended Test:
Vasodilator SPECT (NM Cardiac Stress Perfusion)

Alternative Tests:
Dobutamine Stress Echocardiogram

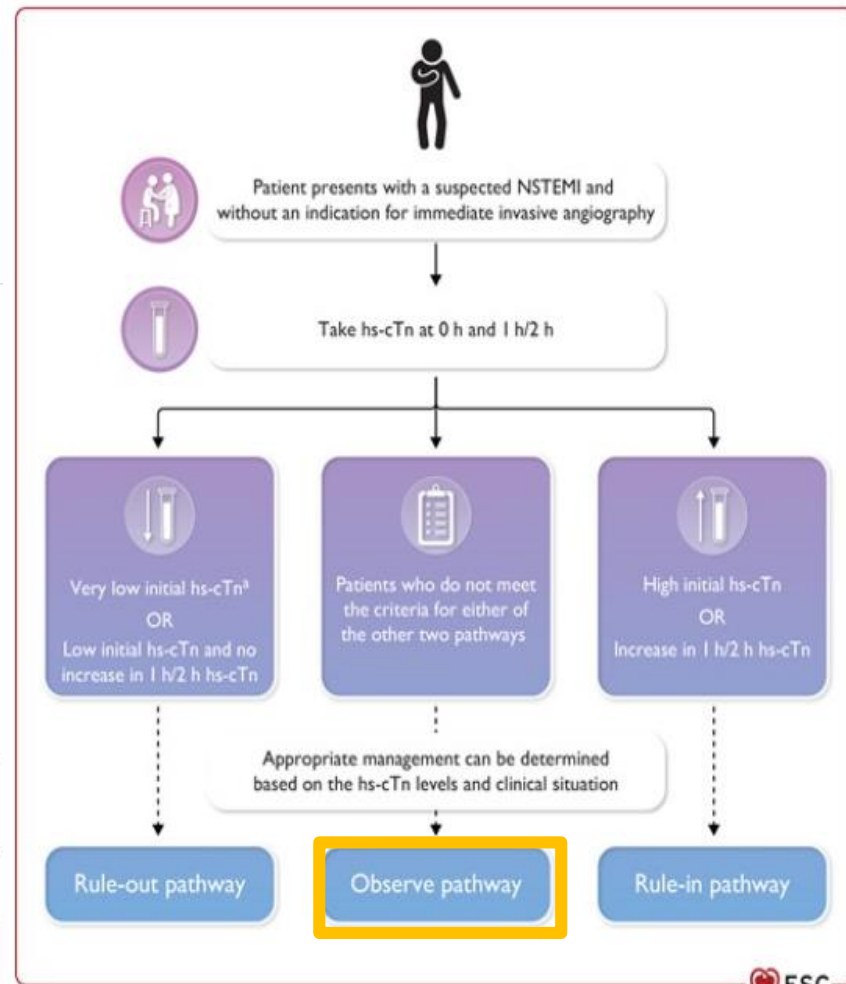
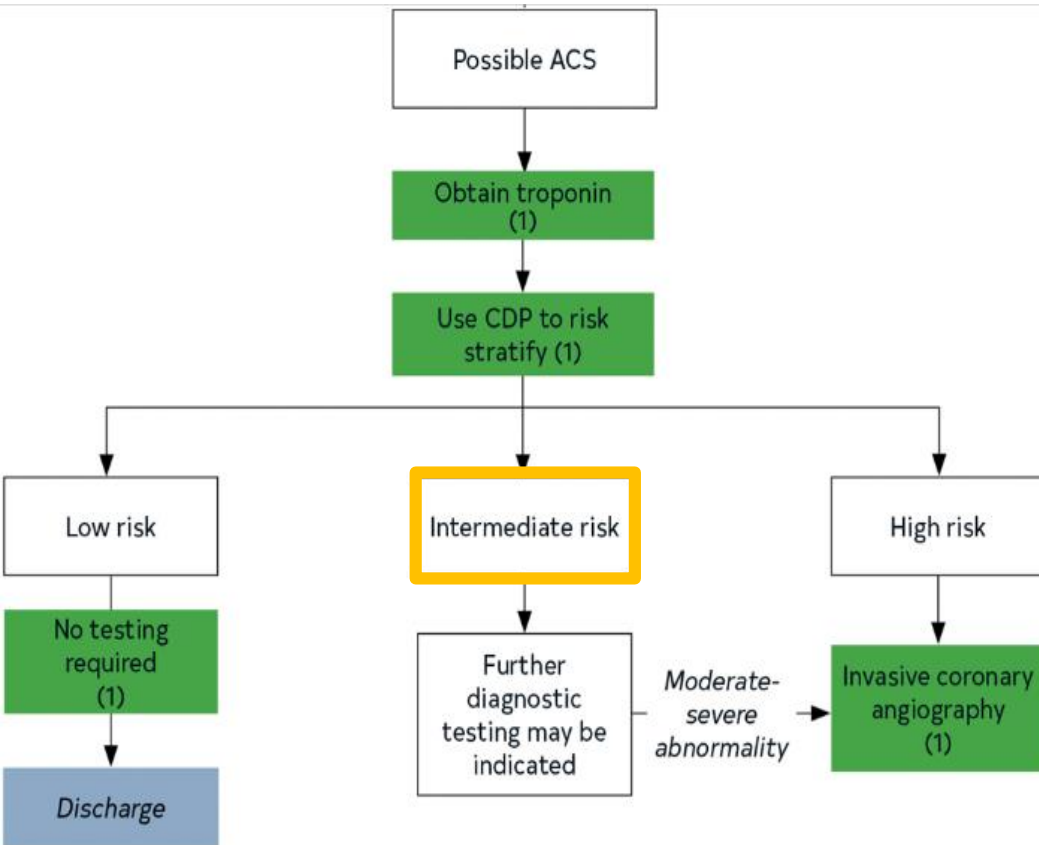
Cardiology Consult



probability of MACE within 30 days is $\leq 1\%$

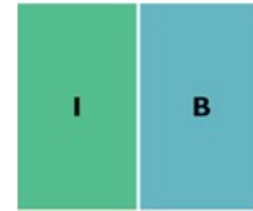


The NPV for MI in patients assigned to the 'rule-out' pathway has exceeded 99% elective non-invasive or invasive imaging may be appropriate according to clinical and risk assessment



Observe pathway

cTn at 3 h (\pm echocardiography) - odporúčané



- vysoký stupeň klinického podozrenia pre AKS (napr. relevantné zvýšenie 3h cTn) – ICA
- nízka až stredná pravdepodobnosť pre AKS - podľa klinického posúdenia - neinvazívne zobrazovacie metódy - CTCA

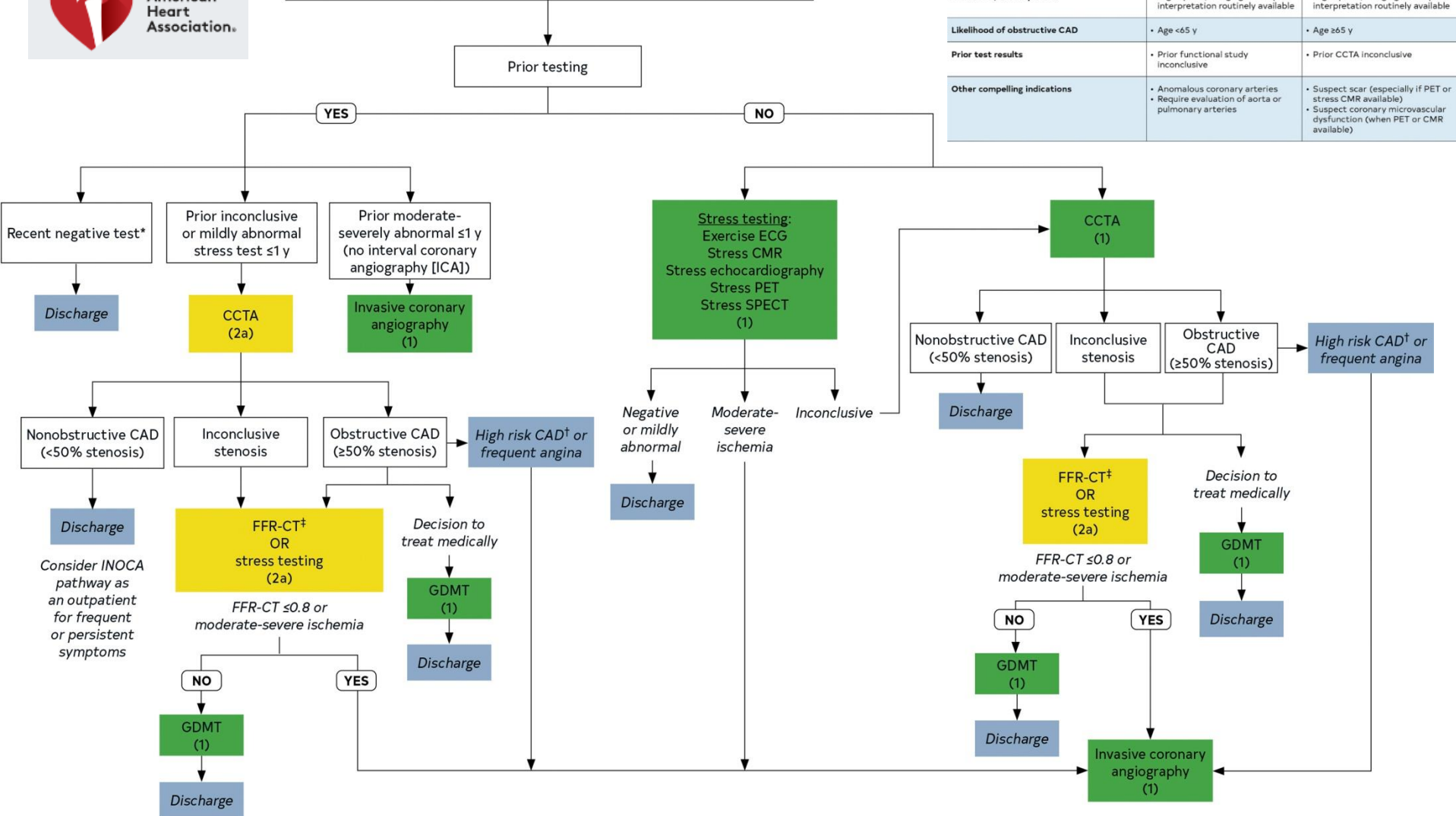


CTCA

- neobštrukčné AS postihnutie koronárnych aa - prepustenie
- Obštrukčné AS postihnutie koronárnych aa – zväžiť revaskularizáciu



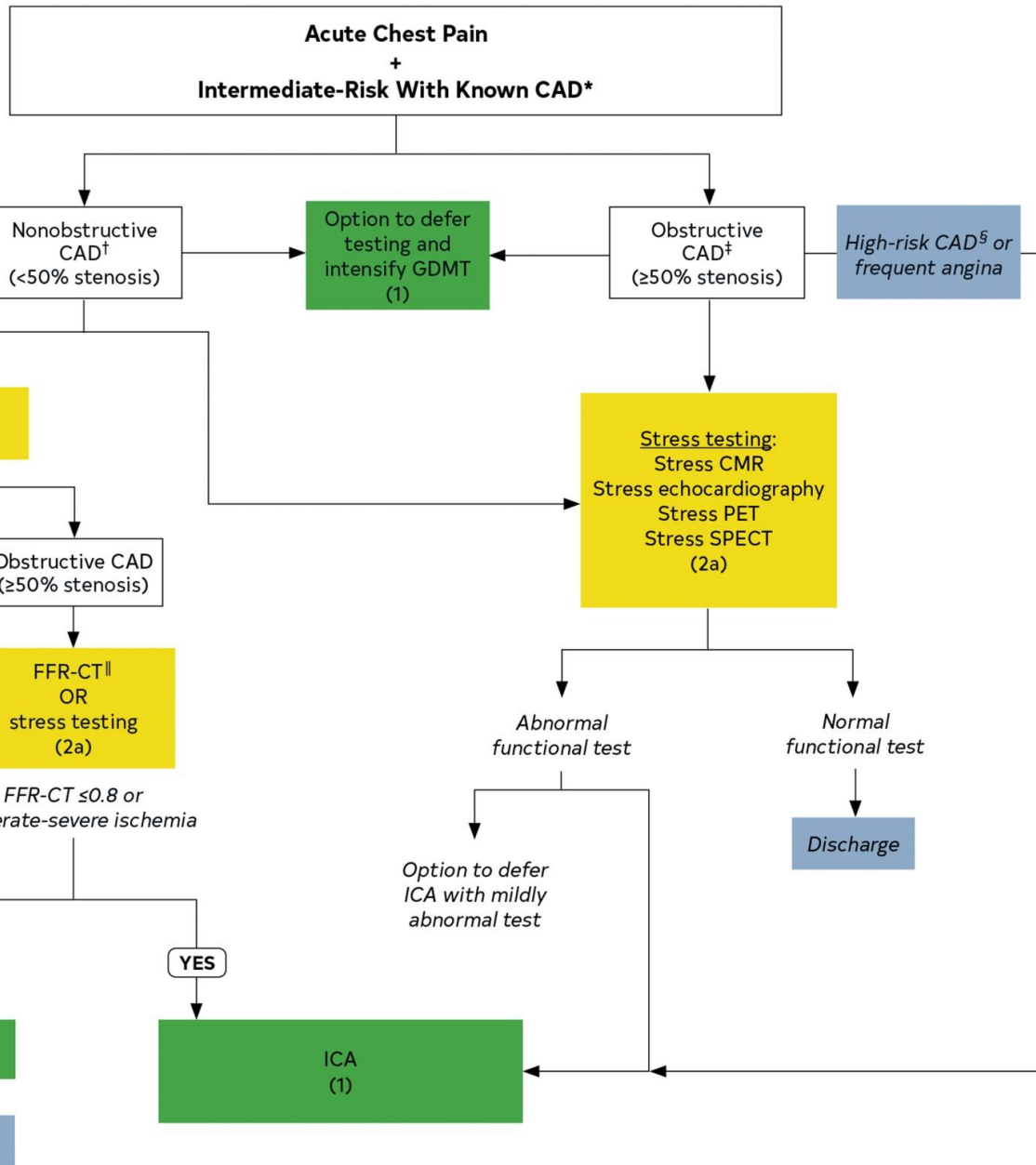
**Acute Chest Pain
+
Intermediate-Risk With No Known CAD**



	Favors use of CCTA	Favors use of stress imaging
Goal	<ul style="list-style-type: none"> Rule out obstructive CAD Detect nonobstructive CAD 	<ul style="list-style-type: none"> Ischemia-guided management
Availability and expertise	<ul style="list-style-type: none"> High-quality imaging and expert interpretation routinely available 	<ul style="list-style-type: none"> High-quality imaging and expert interpretation routinely available
Likelihood of obstructive CAD	<ul style="list-style-type: none"> Age <65 y 	<ul style="list-style-type: none"> Age ≥65 y
Prior test results	<ul style="list-style-type: none"> Prior functional study inconclusive 	<ul style="list-style-type: none"> Prior CCTA inconclusive
Other compelling indications	<ul style="list-style-type: none"> Anomalous coronary arteries Require evaluation of aorta or pulmonary arteries 	<ul style="list-style-type: none"> Suspect scar (especially if PET or stress CMR available) Suspect coronary microvascular dysfunction (when PET or CMR available)

High-risk CAD means left main stenosis ≥ 50%; anatomically significant 3-vessel disease (≥70% stenosis)

Patients with acute chest pain who have coronary artery stenosis from 40% to 90% on CCTA may benefit from measurement of FFR-CT, especially when the stenosis is proximal or mid-coronary artery



High-risk CAD means left main stenosis $\geq 50\%$; anatomically significant 3-vessel disease ($\geq 70\%$ stenosis)



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Recommendations in 2017 and 2020 versions	Class ^a	LoE ^b	Recommendations in 2023 version	Class ^a	LoE ^b
Recommendations for imaging for patients with suspected NSTEMI-ACS					
In patients with no recurrence of chest pain, normal ECG findings, and normal levels of cardiac troponin (preferably high sensitivity), but still with suspected ACS, a non-invasive stress test (preferably with imaging) for inducible ischaemia or CCTA is recommended before deciding on an invasive approach.	I	B	In patients with suspected ACS, non-elevated (or uncertain) hs-cTn, no ECG changes and no recurrence of pain, incorporating CCTA or a non-invasive stress imaging test as part of the initial workup should be considered.	IIa	A



For intermediate-risk patients with acute chest pain and no known CAD eligible for diagnostic testing after a negative or inconclusive evaluation for ACS, CCTA is useful for exclusion of atherosclerotic plaque and obstructive CAD (1-11).

CTCA

**Rýchle, neinvazívne, bezpečné, prijateľné radiačné zaťaženie -
ED 3-5mSv**

**Vysoká senzitivita > 95% - závažnosť + rozsah a distribúcia AS
postihnutia**

Vysoká NPH 95-99%

Charakter AS plátov – kalcif /nonkalcif + HRP

CT-FFR – funkčné hodnotenie stenóz



**American
Heart
Association.**

**8 randomized controlled
trials and 5 meta-analyses**



European Society
of Cardiology

Éra hs-cTn

CCTA neidentifikuje viac pacientov
vyžadujúcich revaskularizáciu,
neurýchluje prepustenie z ED

CCTA – menší počet následných amb vyš,
nižšie priame náklady, redukcia
opakovaných návštev na ED a
hospitalizácii
stratifikácia rizika – rozsah a charakter AS
plátov

115 Dedic A, Lubbers MM, Schaap J, Lammers J, Lamfers EJ, Rensing BJ, et al. Coronary CT angiography for suspected ACS in the era of high-sensitivity troponins: randomized multicenter study. *J Am Coll Cardiol* 2016;67:16–26. <https://doi.org/10.1016/j.jacc.2015.10.045>
[Google Scholar](#) [Crossref](#) [PubMed](#) [WorldCat](#)

116 Hoffmann U, Truong QA, Schoenfeld DA, Chou ET, Woodard PK, Nagurney JT, et al. Coronary CT angiography versus standard evaluation in acute chest pain. *N Engl J Med* 2012;367:299–308. <https://doi.org/10.1056/NEJMoa1201161>

CCTA u pacientov s podozrením
na NSTEMI-AKS nezlepšila klinické
výsledky po 1 roku

40 – 50 % pacientov s eleváciou
hs-cTn má negat event
neobštručné AS postihnutie
19% redukcia ICA

117 Gray AJ, Roobottom C, Smith JE, Goodacre S, Oatey K, O'Brien R, et al. Early computed tomography coronary angiography in patients with suspected acute coronary syndrome: randomised controlled trial. *BMJ* 2021;374:n2106. <https://doi.org/10.1136/bmj.n2106>



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Éra hsTc

CCTA neidentifikuje viac pacientov vyžadujúcich revaskularizáciu, neurýchluje prepustenie z ED

CCTA – menší počet následných amb vyš, nižšie priame náklady, redukcia opakovaných návštev na ED a hospitalizácii
stratifikácia rizika – rozsah a charakter AS plátov

115 Dedic A, Lubbers MM, Schaap J, Lammers J, Lamters EJ, Kensing BJ, et al. Coronary CT angiography for suspected ACS in the era of high-sensitivity troponins: randomized multicenter study. *J Am Coll Cardiol* 2016;67:16–26. <https://doi.org/10.1016/j.jacc.2015.10.045>
[Google Scholar](#) [Crossref](#) [PubMed](#) [WorldCat](#)

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CCTA u pacientov s podozrením na NSTEMI-ACS nezlepšila klinické výsledky po 1 roku


40 – 50 % pacientov s eleváciou hs-cTn má negatívny výsledok
neobštručné AS postihnutie
19% redukcia ICA

117 Gray AJ, Roobottom C, Smith JE, Goodacre S, Oatey K, O'Brien R, et al. Early computed tomography coronary angiography in patients with suspected acute coronary syndrome: randomised controlled trial. *BMJ* 2021;374:n2106. <https://doi.org/10.1136/bmj.n2106>

Spokojnosť pacientov v CTCA ramene bola vyššia



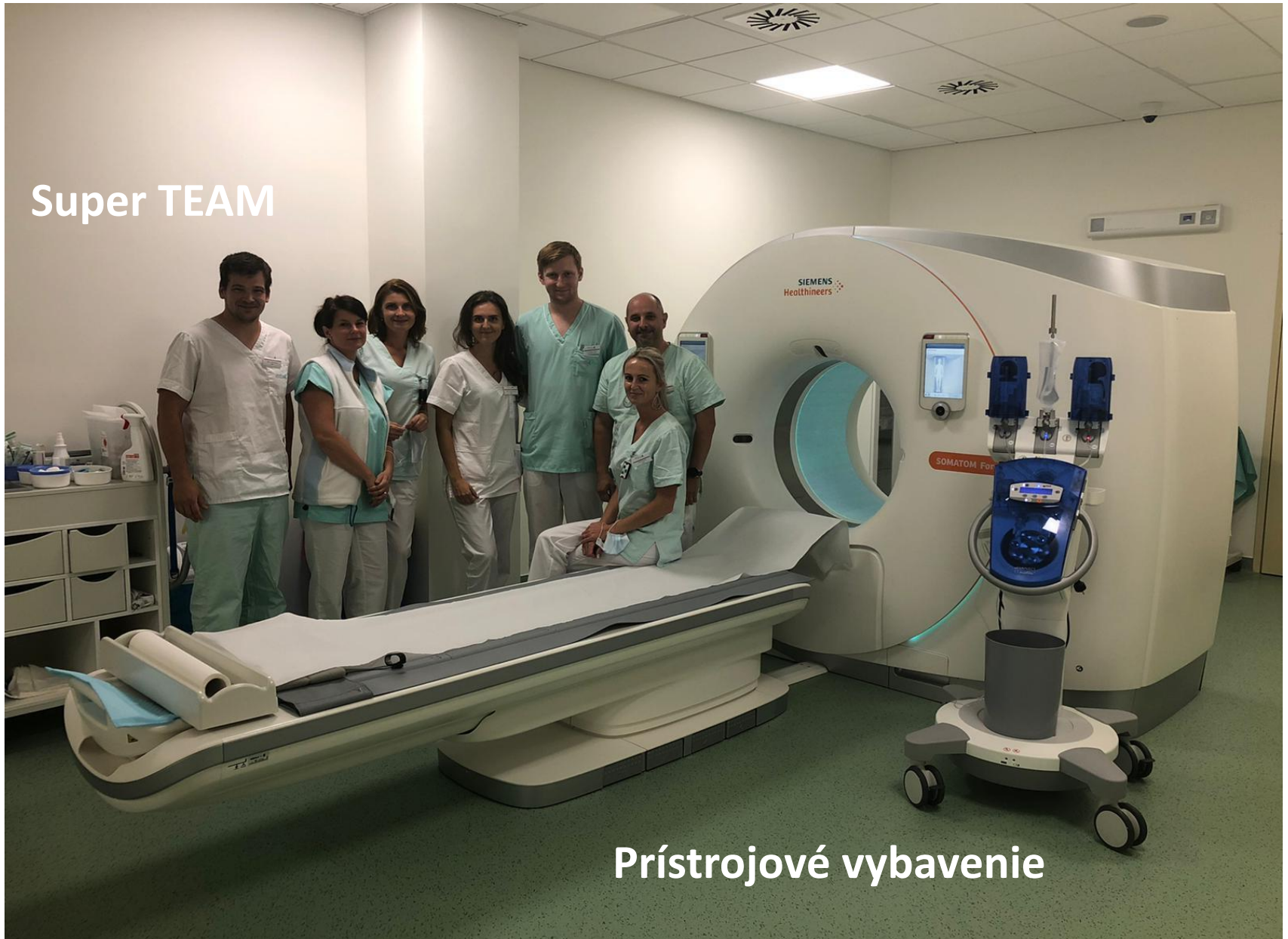
American
Heart
Association.

The image shows two large, scaly, bipedal monsters, resembling Godzilla, engaged in a physical struggle on a grassy hill. The monster on the left is reddish-brown, while the one on the right is dark grey. They are both roaring and pushing against each other. The background is a clear blue sky with some light clouds. A faint circular logo is visible in the bottom right corner of the image.

hs-cTn a koronárna CTA majú komplementárne úlohy a spoločne môžu ponúknuť lepšiu stratégiu triedenia pacientov s akútnou bolesťou na hrudi v porovnaní s akýmkoľvek testom používaným izolovane.

CTCA pri AKS

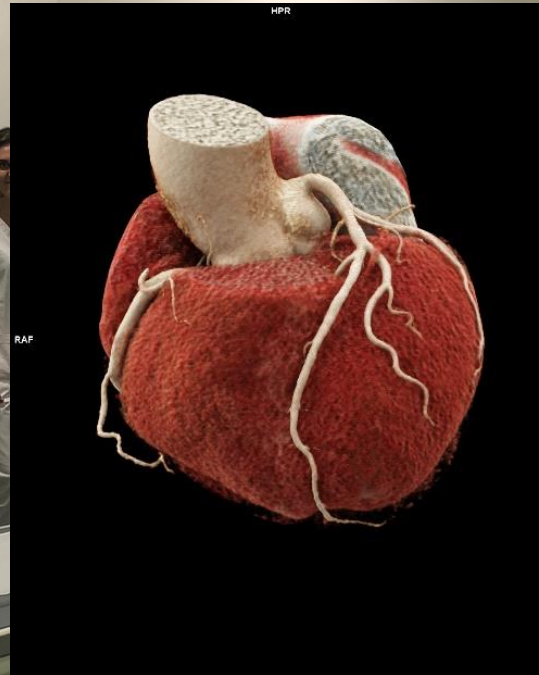
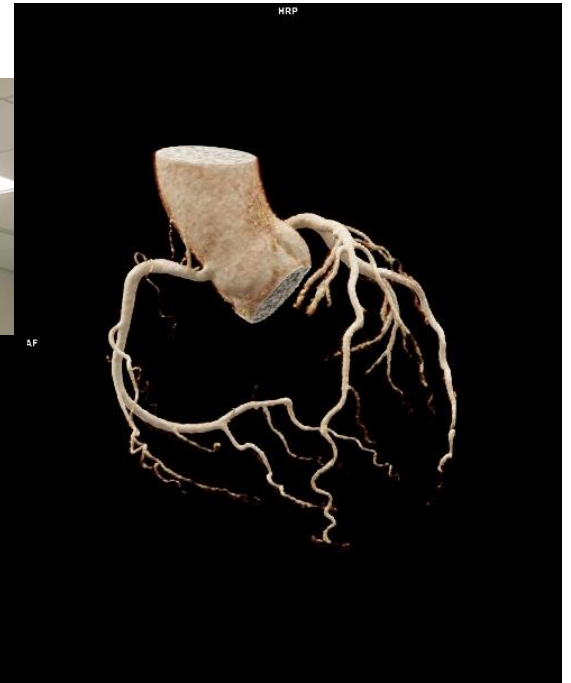
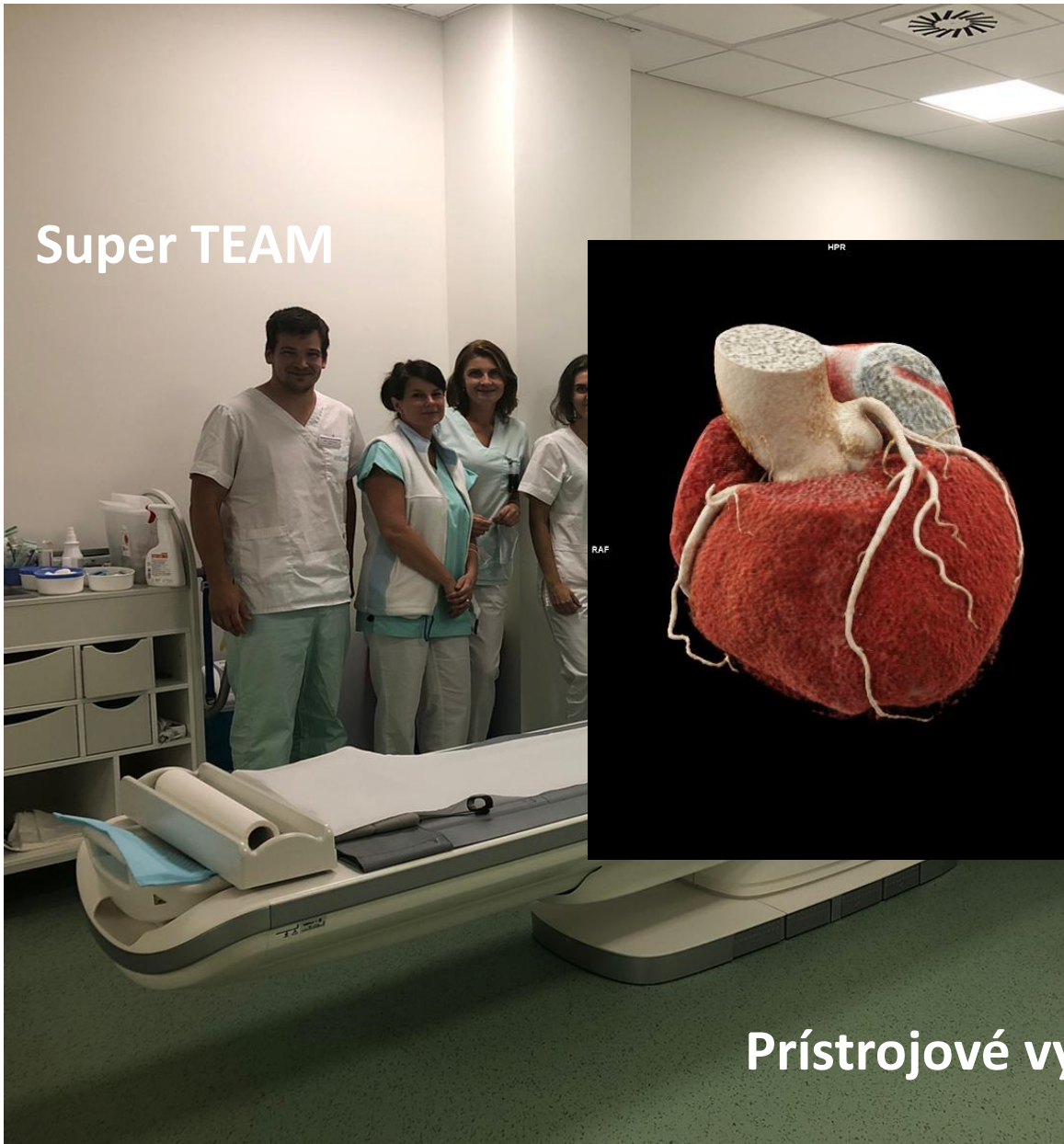
Super TEAM



Prístrojové vybavenie

CTCA pri AKS

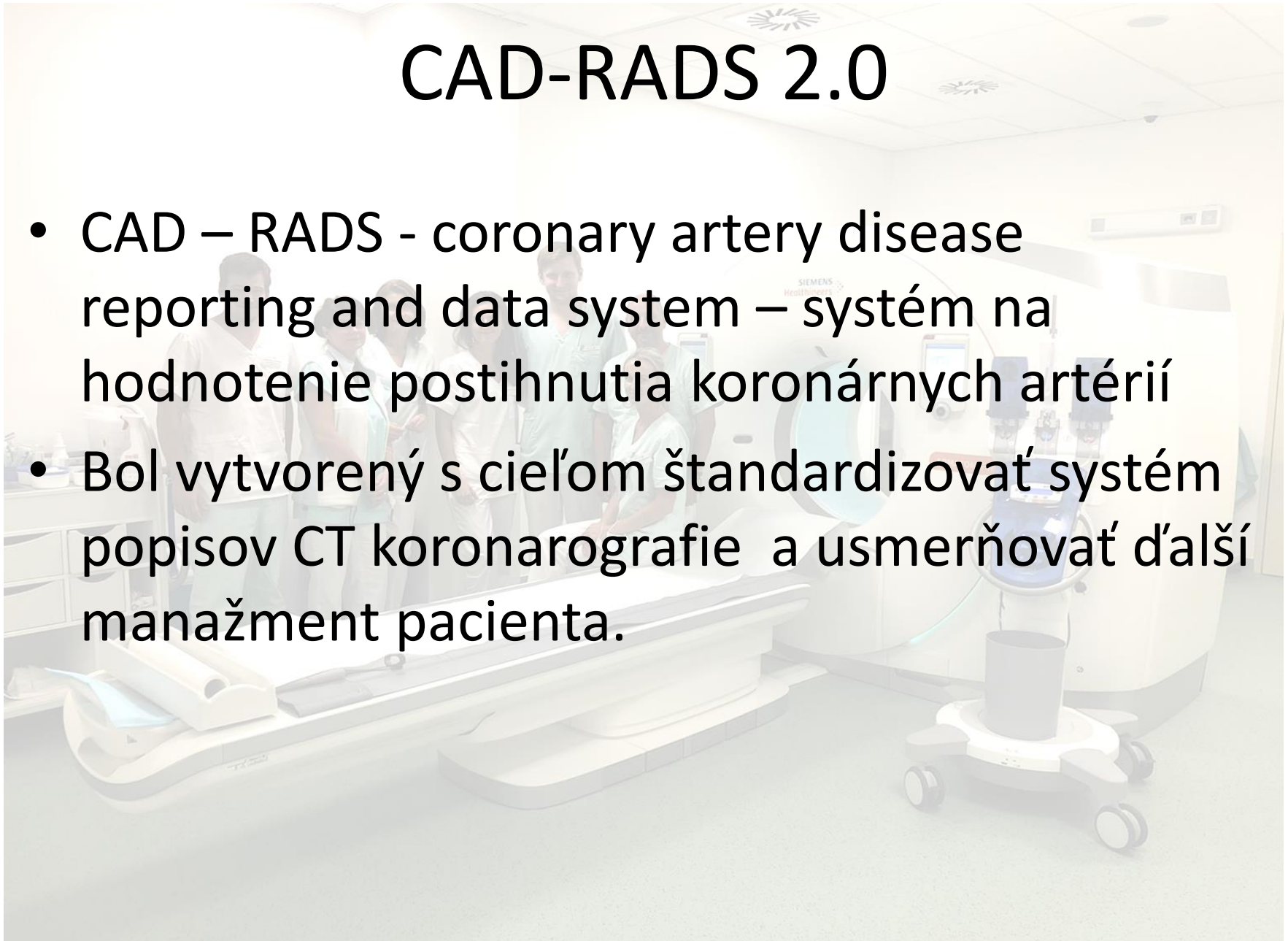
Super TEAM



Prístrojové vybavenie

CAD-RADS 2.0

- CAD – RADS - coronary artery disease reporting and data system – systém na hodnotenie postihnutia koronárnych artérií
- Bol vytvorený s cieľom štandardizovať systém popisov CT koronarografie a usmerňovať ďalší manažment pacienta.



1. Klasifikácia klinicky najrelevantnejšej stenózy (zvyčajne najvyššieho stupňa) u artérii s diametrom >1,5mm

CAD-RADS 0	0%	žiadny plát, stenóza
CAD-RADS 1	1–24%	minimálna stenóza
CAD-RADS 2	25–49%	ľahká stenóza
CAD-RADS 3	50–69%	stredne závažná stenóza
CAD-RADS 4	A - 70–99%	signifikatná stenóza
	B - kmeň LCA ≥50% alebo 3 - cievne postihnutie (≥70%)	
CAD-RADS 5	100%	kompletná oklúzia

2. P - hodnotenie rozsahu atherosklerotického postihnutia koronárnych ciev

P1	ľahké AS postihnutie
P2	stredne závažné AS postihnutie
P3	závažné AS postihnutie
P4	rozsiahle AS postihnutie

3. Modifikátory

- **N** - nediagnostické vyšetrenie - vyšetrenie nie je v plnom rozsahu hodnotiteľné
- **HRP** - rizikový/vulnerabilný plát – aterosklerotický plát asociovaný s vyšším rizikom: a) akútneho kornárneho syndrómu bez ohľadu na stupeň stenózy, b) závažných kardiovaskulárnych príhod u pacientov so stabilnou bolesťou na hrudi, c) léziou podmienenej špecifickej ischemie
- **I** - ischemia – léziou podmienená špecifická ischemia, hodnotená pomocou CT-FFR alebo CT perfúzie. I+ prítomná, I- neprítomná, I+-hraničné hodnoty
- **S** – stent – prítomnosť stentov v koronárnych artériách (stenty vhodné pre CTCA: ≥ 3.0 mm, stenty < 3.0 mm - $> 2,5$ mm môžu byť hodnotiteľné len pri proximálnej a non bifurkačnej lokalizácii)
- **G** – graft/bypass - prítomnosť koronárnych bypassov
- **E** – výnimky – disekcia, aneurizma, pseudoaneurizma, anomálny odstup, vaskulitída, fistula, AV malformácia, externá kompresia

CAD-RADS 1	P1 a 2	<p>Ďalšie vyšetrowanie AKS nie je potrebné.</p> <p>Ak Tn + je potrebné prehodnotiť inú príčinu zvýšeného troponínu. Odporučenie pre ambulantné sledovanie + modifikáciu rizikových faktorov a preventívnu farmakoterapiu</p>
	P3 a P4	<p>Ďalšie vyšetrowanie AKS nie je potrebné.</p> <p>Ak Tn + je potrebné prehodnotiť inú príčinu zvýšeného troponínu. Odporučenie pre ambulantné sledovanie + agresívnu modifikáciu rizikových faktorov a preventívnu farmakoterapiu</p>
CAD-RADS 2	P1 a 2	<p>Ak je vysoká klinická suspekcia pre AKS, Tn + alebo prítomná HRP morfológia AS plátu zväžiť hospitalizáciou s kardiologickou konzultáciou.</p> <p>Ak Tn + je potrebné prehodnotiť inú príčinu zvýšeného troponínu. Odporučenie pre ambulantné sledovanie + modifikáciu rizikových faktorov a preventívnu farmakoterapiu</p>
	P3 a P4	<p>Ak je vysoká klinická suspekcia pre AKS, Tn + alebo prítomná HRP morfológia AS plátu zväžiť hospitalizáciou s kardiologickou konzultáciou.</p> <p>Ak Tn + je potrebné prehodnotiť inú príčinu zvýšeného troponínu. Odporučenie pre ambulantné sledovanie + agresívnu modifikáciu rizikových faktorov a preventívnu farmakoterapiu</p>
CAD-RADS 3	P1, P2, P3, P4	<p>Zväžiť hospitalizáciou s kardiologickou konzultáciou.</p> <p>Zväžiť CT-FFR, CTP (CT perfúzia), záťažové testovanie.</p> <p>Preventívny manažment vrátane agresívnej preventívnej farmakoterapie.</p> <p>Iná liečba (vrátane antianginóznej liečby) by sa mala zväžiť podľa aktuálnych usmerení.</p>
	I+	Zväžiť ICA
CAD-RADS 4A	P1, P2, P3, P4	<p>Hospitalizácia s kardiologickou konzultáciou. Zväžiť ICA.</p> <p>Preventívny manažment vrátane agresívnej preventívnej farmakoterapie.</p> <p>Antianginózna liečba a revaskularizácia by sa mala zväžiť podľa aktuálnych usmerení.</p>
CA-RADS 4B	P1, P2, P3, P4	<p>Hospitalizácia s kardiologickou konzultáciou. ICA je doporučená. Preventívny manažment vrátane agresívnej preventívnej farmakoterapie.</p> <p>Antianginózna liečba a revaskularizácia by sa mala zväžiť podľa aktuálnych usmerení.</p>
CAD-RADS 5	P1, P2, P3, P4	<p>Hospitalizácia s kardiologickou konzultáciou.</p> <p>ICA a revaskularizácia pri suspektnej akútnej oklúzii.</p> <p>Preventívny manažment vrátane agresívnej preventívnej farmakoterapie.</p> <p>Antianginózna liečba a revaskularizácia by sa mala zväžiť podľa aktuálnych usmerení.</p>
CAD-RADS N		Zväžiť doplňujúce alebo alternatívne vyšetrenie

a) Funkčná diagnostika : CT-FFR, CTP (CT perfúzia), záťažové testovanie (ETT, záťažové ECHO, SPECT, PET, MRI srdca) alebo invazívne FFR.

b) ICA – invazívna koronárna angiografia môže byť uprednostnená, pri kritickej stenóze (> 90 %), pri plátoch s rizikovou morfológiou alebo I + (prítomnosť špecifickej ischémie pre léziu na CT FFR alebo CTP) alebo ischémia potvrdená inými funkčnými vyšetreniami alebo u kandidátov na revaskularizáciu.

Ďakujem za pozornosť

