

Chronic Coronary Disease

Diagnostic testing, foundational GDMT, antianginal therapy, revascularization strategy, and antiplatelet therapy for CCD.

COR

I · Recommended

IIa · Reasonable

IIb · May Consider

III · No Benefit / Harm

1 Diagnostic Testing & Risk Stratification

All patients with established CCD — reassess at every visit

I B-NR	New/worsened symptoms despite optimized GDMT : stress PET/SPECT MPI, stress CMR, or stress echo recommended to detect ischemia, estimate MACE risk, and guide therapy. PET preferred over SPECT when available (IIa).
I B-R	Same setting — invasive coronary angiography (ICA) recommended for guiding therapeutic decisions to improve anginal symptoms.
I A	Newly reduced LV systolic function, clinical HF, or both : ICA recommended to assess coronary anatomy and guide revascularization.
III NB/A	ICA not routinely recommended for risk stratification without LV dysfunction, HF, refractory chest pain, or noninvasive testing suggesting significant left main disease. Routine periodic retesting without a change in status is not recommended.
I B-NR	Integrate testing with demographic, social, and clinical variables to classify annual MACE risk as low (<1%) · intermediate (1–3%) · high (>3%) . Testing alone is insufficient to risk-stratify.

2 Foundational GDMT — Start in Every Patient

Optimization of GDMT is recommended to reduce MACE (I, A)

LIPID-LOWERING

I A	High-intensity statin , goal ≥50% LDL-C reduction. If not tolerated, moderate-intensity statin (goal 30–49%).
IIa B-R/A	If very-high-risk and LDL-C ≥70 on max-tolerated statin: add ezetimibe , then PCSK9 mAb . Bempedoic acid/inclisiran reasonable (IIb) if insufficient.
IIb B-R	Persistent fasting TG 150–499 on max statin: icosapent ethyl may be considered.

BLOOD PRESSURE

I B-R	Target <130/<80 mmHg . First-line: ACE-I, ARB, or beta blocker for compelling indications; add CCB/thiazide/MRA as needed.
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ANTIPLATELET (NO OAC INDICATION)

I A	Aspirin 81 mg (75–100 mg) monotherapy. Clopidogrel not additive without recent ACS/PCI indication (III: NB, A).
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SGLT2I / GLP-1 RA

I A	T2DM : SGLT2i or GLP-1 RA with proven CV benefit recommended to reduce MACE. HFrEF (LVEF ≤40%) : SGLT2i recommended regardless of diabetes status.
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LIFESTYLE & FOLLOW-UP

I C-LD	Nutrition, cardiac rehab, weight management, tobacco cessation, immunizations, mental-health screening for all. Follow-up ≥ annually .
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3 Antianginal Therapy Ladder

For angina or an anginal equivalent

I B-R	Step 1 — Beta blocker or CCB or long-acting nitrate (either class as first-line antianginal).
↓ symptoms persist	
I B-R	Step 2 — Add a second antianginal agent from a different class.
↓ symptoms persist	
I B-R	Step 3 — Add ranolazine . Sublingual nitroglycerin recommended for immediate short-term relief (I, B-NR).
↓ refractory, no other options	
IIb B-R	Enhanced external counterpulsation (EECP) may be considered for symptom relief.

2023 update — beta blockers: Not beneficial for MACE reduction without LVEF ≤50%, prior MI (<1 yr), or another primary indication (III: NB). Reassess long-term (>1 yr) use post-MI if no current LVEF ≤50%, angina, arrhythmia, or hypertension (IIb). Avoid adding **ivabradine** with normal LV function — potentially harmful (III: Harm).

4 Revascularization — Goals, Indications & PCI vs. CABG

Symptom relief is the primary driver; survival benefit is anatomy/LV-function specific

I A	Lifestyle-limiting angina despite GDMT with significant, amenable stenosis → revascularization to improve symptoms.	IIa B-NR	Significant left main stenosis where PCI can achieve equivalent revascularization to CABG → PCI reasonable to improve survival.
I B-R	Left main or multivessel disease with severe LV dysfunction (LVEF ≤35%) → CABG + medical therapy over medical therapy alone to improve survival.	I A	Angiographically intermediate stenosis, angina/equivalent, no prior ischemia evaluation → FFR or iFR recommended before PCI.
IIa B-R	Multivessel CAD appropriate for either CABG or PCI → revascularization reasonable to lower risk of spontaneous MI, urgent revascularization, or cardiac death.	I B-NR	Complex 3-vessel disease or unclear optimal strategy → multidisciplinary Heart Team approach recommended.

COR / LOE	CLINICAL SCENARIO	PREFERRED STRATEGY
I · B-R	Left main involvement with high-complexity CAD	CABG preferred over PCI to improve survival
IIa · B-R	Multivessel, complex/diffuse CAD (e.g., SYNTAX >33)	CABG reasonable over PCI to improve survival
I · A	Diabetes + multivessel CAD involving LAD, CABG candidate	CABG (LIMA → LAD) preferred to reduce mortality & repeat revasc.
IIb · B-R	Diabetes + left main, low/intermediate complexity (SYNTAX ≤33)	PCI may be considered as an alternative to CABG
IIa · B-NR	Appropriate for revascularization but poor surgical candidate	PCI reasonable over CABG for symptoms & MACE

5 Antiplatelet Therapy After Revascularization

POST-PCI (NO OAC INDICATION)

I A	DAPT (aspirin + clopidogrel) × 6 months → single antiplatelet therapy (SAPT) to reduce MACE and bleeding.
IIa A	DES with 1–3 month DAPT completed: P2Y12 monotherapy ≥12 months reasonable to reduce bleeding risk.
IIb A	Prior MI + low bleeding risk: extended DAPT up to 3 years may be reasonable to reduce MACE.
IIa B-R	PPI co-therapy with DAPT can be effective to reduce GI bleeding risk.

OAC INDICATED (E.G., CONCOMITANT AF)

I B-R	Elective PCI + OAC required: DAPT for 1–4 weeks → clopidogrel alone to 6 months, in addition to DOAC.
IIb B-R	Low atherothrombotic risk: discontinue aspirin, continue DOAC monotherapy alone ~1 year after PCI to reduce bleeding.

AVOID

III Harm	Vorapaxar or prasugrel with prior stroke/TIA/ICH; chronic NSAIDs — increased bleeding/CV risk.
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