

Microvascular Angina

(Angina With No Obstructive Coronary Artery Disease [ANOCA])

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Microvascular angina (previously known as syndrome X) is cardiac microvascular dysfunction or constriction causing angina in patients with normal epicardial coronary arteries on angiography.

(See also [Overview of Coronary Artery Disease](#).)

Patients with microvascular angina have

- Typical [angina](#) that is relieved by rest or nitroglycerin
- Normal coronary angiography (eg, no atherosclerosis, embolism, or inducible arterial spasm)

Some of these patients have ischemia detected during stress testing; others do not. In some patients, the cause of ischemia seems to be reflex intramyocardial coronary constriction and reduced coronary flow reserve. Other patients have microvascular dysfunction within the myocardium: The abnormal vessels do not dilate in response to exercise or other cardiovascular stressors; sensitivity to cardiac pain may also be increased.

This disorder should not be confused with [vasospastic angina](#) due to epicardial coronary spasm.

Prognosis is better than for patients with demonstrable [coronary artery disease](#), although symptoms of ischemia may recur for years. In addition, patients with microvascular angina appear to be at an higher risk for major cardiovascular events than the general population ([1](#)).

The mainstay of treatment is controlling risk factors with lipid-lowering therapy and glycemic control. In many patients, traditional anti-ischemic treatment, including beta-blockers and calcium channel blockers, helps to relieve symptoms ([2](#)).

References

1. [Shimokawa H, Suda A, Takahashi J, et al](#): Clinical characteristics and prognosis of patients with microvascular angina: an international and prospective cohort study by the Coronary Vasomotor Disorders International Study (COVADIS) Group. *Eur Heart J* 42(44):4592–4600, 2021. doi:10.1093/eurheartj/ehab282

2. [Crea F, Camici PG, Bairey Merz CN](#): Coronary microvascular dysfunction: an update. *Eur Heart J* 35:1101–1111, 2014. doi: 10.1093/eurheartj/eh513



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