Ischaemic heart disease persists as the leading global cause of death and lost life years in adults. Angina is a common clinical presentation of ischaemic heart disease related to a supply-demand mismatch of myocardial blood flow, typically provoked by exertion or stress. Invasive coronary angiography is the reference test for angina and identifies obstructive coronary artery disease (CAD) as a cause for symptoms. In Europe and the US, approximately 4 million elective coronary angiograms are performed each year. However, up to half of all angina patients undergoing elective coronary angiography with symptoms and/or signs of ischaemia have no obstructive epicardial coronary artery disease (INOCA). This large, heterogeneous group includes patients with microvascular angina (MVA), vasospastic angina (VSA) or both conditions together. The burden of these conditions on physical and mental wellbeing can be profound; they are associated with morbidity and a reduction in quality of life. Patients with these conditions commonly attend primary and secondary care, driving up health resource utilisation.

We propose that optimal clinical management starts with the correct diagnosis; hence we begin by summarising the rationale and protocol for invasive tests of coronary function in INOCA patients. We discuss drivers of myocardial ischaemia and reappraise existing consensus guideline-based management in light of the CORonary MICrovascular Angina (CorMicA) study, the first randomised controlled trial of invasive coronary function testing linked to stratified medical therapy in angina. This review aims to educate and empower the invasive cardiologist to perform vasoreactivity testing and to provide them with an understanding of the positive impact of personalised medicine for individual angina patients. We conclude pointing to future directions in care and the benefits of improved diagnosis linked to translational clinical research to develop targeted disease-modifying therapy.

**Background and Aetiology of Angina Without Obstructive Coronary Disease**

INOCA is a recently proposed "umbrella" term conveying the importance of stable coronary syndromes beyond obstructive CAD (Figure 7). INOCA aligns with the sibling term MINOCA, which stands for myocardial infarction with no obstructive CAD. MINOCA is a similarly diverse syndrome with distinct underlying causes. Depending on the patient population studied and the techniques used, between one-third and two-thirds of angina patients with a
negative angiogram have an underlying disorder of coronary vascular function.10 Importantly, the two most common causes of INOCA (MVA and VSA) are not excluded by a negative non-invasive CT coronary angiogram or invasive coronary angiogram.11 For affected patients, symptom burden, morbidity and health resource utilisation can be considerable.12,13 As cardiologists, we often adopt a ‘stenosis-centric’ approach. However, as clinicians we must appreciate the complexity and individual contributors to ischaemia in patients without obstructive epicardial disease (Figure 1). Systemic factors, including heart rate, blood pressure (and their product) and myocardial supply:demand ratio (Buckberg index), are important.12,13 Coronary factors are well recognised, but certain nuances are overlooked. For example, Gould and Johnson recently used their quantitative myocardial perfusion database of over 5,900 patients to show that occult coronary diffuse obstructive coronary disease or flush ostial stenosis may be overlooked on angiography and mislabelled as microvascular angina with suboptimal treatment.14 Other coronary factors that can cause ischaemia and propensity to acute coronary syndromes include structural microvascular dysfunction, endothelial impairment, myocardial bridging and/or epicardial vasospasm.15,16

The final group of factors that can drive INOCA is cardiac, including left ventricular hypertrophy or restrictive cardiomyopathy where subendocardial ischaemia results from challenges with arteriolar vessels penetrating deeper into the myocardial tissue with shorter diastole and enhanced systolic myocardial vessel constriction.17 Heart failure (with reduced or preserved ejection fraction) can lead to elevated left-ventricular end diastolic pressures that reduce the physiological myocardial perfusion gradient.

Valvular heart disease, e.g. aortic stenosis or left ventricular outflow tract obstruction, is a well-recognised cause of INOCA, although controversy exists over whether symptoms in mechanical outflow tract obstruction (aortic stenosis) relate to microvascular dysfunction, supply:demand factors or both.18 Most experts support haemodynamic factors as the main cause of ischaemia here, especially since symptoms and coronary flow reserve improve immediately after valve replacement.19

**Non-invasive Functional Testing**

Non-invasive tests provide indirect assessments of myocardial resistance by assessing perfusion during exercise or pharmacological stress, typically using systemic adenosine. Nevertheless, perfusion assessment lacks the sensitivity to diagnose the relative contributions of epicardial and microvascular disease to myocardial blood flow reduction. In addition, some patients with a propensity to vasospastic chest pain syndromes may have normal findings from pharmacological and exercise stress testing. This review focuses on the invasive diagnosis and related management of angina subjects without obstructive disease; the non-invasive workup is covered elsewhere.20,21

**Diagnosis and Rationale for Invasive Testing**

In the cardiac catheterisation laboratory, coronary vascular function may be assessed ad hoc during the patient’s index coronary angiogram. This often involves an interventional diagnostic procedure (IDP) where a guidewire-based assessment of coronary blood flow is performed at rest and during interrogation with pharmacological probes, typically adenosine and acetylcholine.

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**Figure 1: Ischaemia with No Obstructive Coronary Artery Disease: A Coronary Syndrome**

**Systemic factors**
- **Microvascular supply-demand ratio** (SEVR or Buckberg index)
- Pulse rate
- Blood pressure
- Heightened sympathetic activation

**Coronary factors**
- **Microvascular dysfunction**
- **Endothelial impairment**
- **Occult diffuse epicardial CAD**
- **Coronary artery spasm**
- **Myocardial bridging**

**Cardiac factors**
- **Impaired diastolic relaxation**
- **Left ventricular hypertrophy**
- **Diastolic dysfunction**
- **Valvular heart disease**
- **Left ventricular outflow tract obstruction**

The traditional paradigm where angina is ubiquitously associated with obstructive epicardial disease overlooks the importance of other determinants of myocardial ischaemia. These three groups of factors combine to determine the physiological myocardial perfusion gradient. CAD = coronary artery disease; INOCA = ischaemia but no obstructive coronary artery disease; SEVR = subendocardial viability ratio.

The rationale for an IDP is three-fold. First, these patients often present with typical angina for invasive coronary angiography, which offers an opportunity for the cardiologist to provide patients with an accurate diagnosis and explanation for their symptoms. Second, discrimination of MVA, VSA and non-cardiac chest pain permits distinct treatment outlined in consensus practice guidelines. Third, evidence of coronary vascular dysfunction carries prognostic insights for patients and their clinicians. However, in contemporary standard practice, additional invasive tests on patients with unobstructed coronary arteries are very rarely performed.

The IDP consists of two steps: assessment of coronary circulation vasorelaxation using invasive coronary physiology at rest and with hyperaemia; and second, assessment of the propensity of the coronary circulation to excessive vasoconstriction using intra-arterial acetylcholine (microvascular and/or epicardial vasospasm) (Table 1). We typically prefer the left anterior descending coronary artery as the target vessel because it subtends the largest myocardial mass. While regional microvascular dysfunction is well recognised, interrogation of multiple vessels increases the procedural duration such that the benefits of testing may be outweighed by the risks.

**Assessment of Coronary Vasorelaxation and Resistance Using Diagnostic Guidewire**

The purpose of step one is to assess the coronary flow reserve (CFR) and microvascular resistance, typically using the index of microcirculatory resistance (IMR; Figure 2). Flow-limiting epicardial coronary disease may be assessed using fractional flow reserve (FFR), which is the ratio of mean distal coronary pressure to mean aortic pressure at maximal hyperaemia. Abnormal FFR is defined as ≤0.80 or alternatively a non-hyperaemic pressure ratio with different cut-off may be used, e.g. diastolic-only pressure ratio.22,23 The CFR is determined by dividing the hyperaemic coronary blood flow by the resting flow. This is also termed the vasodilator capacity and reflects the ability of the coronary circulation to augment blood flow from rest. CFR is calculated using thermodilution as the resting mean transit time divided by hyperaemic mean transit time; an abnormal CFR is defined as ≤2.24,25 Microcirculatory resistance can be assessed using...
Vasospastic angina

Epicardial spasm

Angina symptoms during ACh bolus (e.g. 100 μg acetylcholine over 20 seconds) AND:

- ST-segment deviation on ECG
- No significant epicardial coronary spasm (<90% diameter reduction)

Microvascular angina

Abnormal microvascular resistance • IMR ≥25

Impaired coronary vasorelaxation

Angina symptoms with ACh infusion AND:

- ST-segment deviation on ECG
- No epicardial coronary constriction during ACh reduction

Vasospastic angina

Epicardial spasm

Angina symptoms during ACh bolus (e.g. 100 μg acetylcholine over 20 seconds) AND:

- ST-segment deviation on ECG
- >90% epicardial coronary constriction during ACh reduction

Non-cardiac

- Exclusion of significant epicardial coronary disease (fractional flow reserve >0.8)

ACh = acetylcholine; CFR = coronary flow reserve; IMR = index of microvascular resistance.

Table 1: Definition and Invasive Diagnostic Criteria for Disorders of Coronary Artery Function

| Disorder                     | Symptoms                                                                 | Clinical measurement                                                                 |
|------------------------------|--------------------------------------------------------------------------|--------------------------------------------------------------------------------------|
| Microvascular angina         | Abnormal microvascular resistance                                       | • IMR ≥25                                                                            |
|                              |                                                                          | • Hyperaemic microvascular resistance ≥2.5 mmHg/cm/s                               |
| Impaired coronary vasorelaxation |                                                                        | • CFR by thermodilution <2.0                                                      |
| Microvascular spasm          | Angina symptoms with ACh infusion AND:                                   | • ST-segment deviation on ECG                                                       |
|                              |                                                                          | • No significant epicardial coronary spasm (<90% diameter reduction)               |
| Vasospastic angina           | Epicardial spasm                                                        | Angina symptoms during ACh bolus (e.g. 100 μg acetylcholine over 20 seconds) AND: |
|                              |                                                                          | • ST-segment deviation on ECG                                                       |
|                              |                                                                          | • >90% epicardial coronary constriction during ACh reduction                       |
| Non-cardiac                  |                                                                          | • Exclusion of significant epicardial coronary disease (fractional flow reserve >0.8) |

without any of the following abnormalities of coronary function: CFR <2.0, IMR ≥25 or positive ACh response.

Assessment for Propensity to Coronary Vasoconstriction: Acetylcholine Provocation

In healthy endothelium, ACh stimulates abluminal release of nitric oxide, mediating vascular smooth muscle relaxation and increased blood flow. At high doses or in patients with endothelial dysfunction, ACh directly stimulates vascular smooth muscle, causing vasoconstriction that can precipitate epicardial vasospasm and/or microvascular vasospasm-induced ischaemia. Typically, infusions of ACh at concentrations approximating 0.182, 1.82, and 18.2 μg/ml (10⁻¹, 10⁻² and 10⁻³ mol/l, respectively) at 1 ml/min for 2–3 minutes are given via a mechanical pump. These doses were historically derived using experiments adopting subselective infusion through an infusion catheter into the left anterior descending artery, assuming a resting flow of 80 ml/min. The effective concentration at tissue level was estimated at 10⁻¹ to 10⁻³ M. The assessment of Doppler response to ACh involves intra coronary infusion catheters in combination with Doppler wire and requires larger guiding catheter sizes (7 Fr) and a 3 Fr infusion catheter into the coronary artery.

Centres in Japan with over four decades of experience with ACh testing adopt a pragmatic and streamlined approach using sequential bolus doses of ACh via the guiding catheter. Doses start from 20 μg, increasing to 50 and 100 up to 200 μg in the left system (or 20, 50 and 80 μg into the right coronary) over 20 seconds followed by up to 3 minutes between doses.

Coronary angiography is performed when either ST segment changes or chest pain (or both) occur, or after 1 minute following the completion of each injection. We routinely use a well-engaged guiding catheter to deliver ACh via a 2-minute infusion using an external mechanical pump without an additional infusion catheter. This approach facilitates smaller guiding catheters and reduces risk, time and procedural cost.
Epicardial coronary artery spasm is defined according to the Coronary Vasomotion Disorders International Study Group criteria whereby chest pain is reproduced with ST segment deviation and ≥90% vasoconstriction to 100 µg of ACh (5.5 ml of 10⁻⁴ M over 20 seconds). Microvascular spasm is defined chest pain and ST segment deviation without significant luminal constriction (<90%) and represents a functional subtype of microvascular angina. Severe epicardial endothelial dysfunction is defined by ≥20% luminal constriction during ACh infusion (up to 10⁻⁴ M); this finding implies a significant reduction in coronary artery blood flow with prognostic implications when compared with patients whose arteries are <20% constricted. Bradycardia with ACh is common and usually self-limiting, although a reduced dose during interrogation of the right coronary artery (maximum 50–80 µg ACh) may reduce occurrence.

**CorMicA and Clinical Evidence**

Despite a wealth of clinical evidence from observational studies, until recently there has not been a single randomised controlled trial of coronary function testing. In the absence of randomised trials demonstrating patient benefits, observational evidence has rarely been applied in practice. CorMicA now provides proof-of-concept clinical evidence to support the case for patient benefits when management is guided by invasive tests of coronary artery function (IPD). Ad hoc adoption of coronary function testing for patients with INOCA is currently restricted to a few interested academic centres. In part this relates to lack of evidence that an IPD has clinical utility or improves patient well-being.

Supported by the British Heart Foundation and the patients who kindly agreed to take part, we delivered the CorMicA trial to specifically address this gap in evidence. We hypothesised that stratified medicine, including an IPD with linked medical therapy, would be routinely feasible and lead to improvements in angina and quality of life in patients with no obstructive CAD.

A total of 391 patients with definite or probable angina, as determined on the Rose angina questionnaire, were enrolled over a 12-month period from November 2016 at two large tertiary referral centres serving around half the population of Scotland (approximately 2.5 million people). Coronary angiography revealed no obstructive CAD in 185 (47%) of the patients who completed the Rose questionnaire and 151 individuals were immediately randomised to one of two arms: the intervention group (stratified medical therapy, IPD disclosed) or the control group (standard care, IPD sham procedure, results not disclosed). The mean age of subjects was 62 years and 74% were female.

The diagnostic intervention included a guidewire-based assessment of a major coronary artery, usually the left anterior descending coronary artery, followed by pharmacological coronary reactivity testing (Figure 2). This diagnostic assessment aligned with contemporary guidelines. The IPD involved measurement of CFR (abnormal <2.0), microcirculatory resistance (IMR; abnormal ≥25) and FFR (abnormal ≤0.80). Vasoreactivity testing was then performed by infusing incremental concentrations of ACh followed by a bolus of ACh of up to 100 µg to assess for epicardial or microvascular vasoconstriction. The diagnosis of a clinical endotype (MVA, VSA, both or none) was linked to distinct guideline-based management stratified by diagnosis. The primary endpoint was the mean difference in angina severity at 6 months assessed by the Seattle Angina Questionnaire summary score.

In an all-comers study design, the IPD revealed isolated MVA in 78 subjects (52%), isolated VSA in 25 subjects (17%), mixed angina (both) in 31 subjects (20%) and non-cardiac chest pain in 17 subjects (11%) (Figure 3). The intervention was associated with a mean improvement of 11.7 units in Seattle Angina Questionnaire summary score at 6 months (95% CI [5.0–18.4]; p=0.001). In addition, the intervention led to improvements in the mean quality of life score (EQ-SD index 0.10 units; 95% CI [0.01–0.18]; p=0.024) and visual analogue score (14.5 units; 95% CI [7.8–21.3]; p=0.001). Notably, after the disclosure of coronary function testing, over half of the treating clinicians changed their diagnosis. There were no differences in major adverse cardiac events after 6 months of follow-up (2.6% controls versus 2.6% intervention; p=1.00). Thus, we showed that in patients undergoing invasive coronary angiography, obstructive coronary disease is excluded in half of all patients; and within this large group of patients, the majority have a readily identifiable disorder of coronary vasomotion. Specifically, the IPD with linked medical therapy was routinely feasible and safe, resulting in improvements in angina and quality of life at 6 months in this group of patients. CorMicA was undertaken in a real-world setting and the results appear to be transferable to clinical practice. Future trials are anticipated to determine the wider external validity of this approach.

**Stratified Medicine in Angina**

We start by considering the patient in the context of non-coronary contributors to INOCA (Figure 1). Non-pharmacological therapies encompassing lifestyle modification, risk factor control, evidence-based pharmacological therapy and patient education are also essential for stratifying treatment. Lifestyle recommendations are covered in detail in the recent European Society of Cardiology guidelines. We will focus on the two most common diagnostic groups to guide distinct medical treatments.

**Microvascular Angina**

The diagnosis of MVA may be suspected in angina patients without obstructive CAD who have evidence of microvascular dysfunction. In the IPD above, microvascular dysfunction consists of abnormal CFR (≤2.0), abnormal IMR (≥25) and/or microvascular spasm during ACh provocation. Clearly, this is a heterogeneous entity akin to the syndrome of heart failure with preserved ejection fraction (HFpEF) with diverse aetiology.

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**Figure 3: Prevalence and Treatment of Ischaemia with No Obstructive Coronary Artery Disease**

| A | B |
|---|---|
| Vasospastic angina | Consider statin and ACEI |
| MVA | Vasoprotective antagonist |
| VSA | Beta-blocker |
| Mixed (both) | Nitrate |
| Non-cardiac | Ca²⁺ antagonist |
| 1st | Nicorandil |
| 2nd | Ranolazine |
| 3rd | Ranolazine |

A: The overall prevalence of coronary artery vasomotion disorders in the CorMicA study. B: Authors’ interpretation of the evidence for recommended therapy for angina patients without obstructive CAD stratified by diagnosis. This formed the basis of pharmacological treatment for patients in the British Heart Foundation CorMicA study. ACh = angiotensin-converting enzyme inhibitor; MVA = microvascular angina; VSA = vasospastic angina.
Baseline “disease-modifying” therapies that have demonstrated benefit in clinical trials of microvascular angina include angiotensin-converting enzyme inhibitors and statins. We particularly support these baseline therapies in patients with diffuse CAD without epicardial obstruction. European Society of Cardiology guidelines for patients with MVA recommend beta-blockers as first-line and calcium antagonists if the former are not tolerated or efficacious (Table 2). Dihydropyridine calcium blockers, e.g. amiodipine 5–10 mg, may be added to beta-blockers if blood pressure permits. There is accumulating evidence that long-acting nitrates are ineffective or even detrimental in MVA. Lack of efficacy may relate to poor tolerability, steal syndromes through regions of adequately perfused myocardium and/or related to the reduced responsiveness of nitrates within the coronary microcirculation. There is significant clinical overlap between MVA and HFpEF, so inferences about nitrate response may be drawn from the Nitrate’s Effect on Activity, Tolerance in HFpEF (NEAT-HFpEF) study. In this randomised controlled trial, HFpEF patients on isosorbide mononitrate actually did worse with reduced activity levels assessed using an accelerometer.

Ranolazine is a relatively new and well-researched antianginal therapy that may improve myocardial perfusion by decreasing sodium and calcium overload, thereby improving myocyte relaxation and ventricular compliance. In a randomised placebo-controlled clinical trial of ranolazine led by the Women’s Ischemic Syndrome Evaluation investigators, although there were no overall improvements in angina and myocardial perfusion with ranolazine, patients with a reduced CFR (<2.5) benefitted from ranolazine, with significant improvements in myocardial perfusion (p=0.014) and angina frequency (p=0.027). Multiple other drugs that reduce angina may be added, including nicorandil and ibabradine.

**Vasospastic Angina**

VSA is often characterised by rest angina, often with preserved effort tolerance. The poor nitrate response or tolerance seen in MVA is not uncommon. Dual pathology (VSA with underlying microvascular disease) is not uncommon. A positive diagnosis of VSA facilitates treatment using non-dihydropyridine calcium antagonists, e.g. controlled-release diltiazem at up to 500 mg daily, which are usually very effective. High doses of ACE inhibitors (e.g. ramipril 2.5 mg) improve CFR, ↓ workload, may improve small vessel remodelling.

**Table 2:** Pharmacological and Non-pharmacological Treatment Options for Angina

| Diagnosis                        | Investigation   | Pathophysiology                      | Treatment | Effects                                |
|----------------------------------|-----------------|--------------------------------------|-----------|----------------------------------------|
| Microvascular angina ↓ vasorelaxation | ↓ CFR and/or ↑ microvascular resistance | Anatomical remodelling, vascular rarefaction, disturbed coronary regulation | Beta-blockers (e.g. nebivolol 2.5–10mg) | ↓ myocardial oxygen consumption |
|                                  |                 |                                      | ACE inhibitors (e.g. ramipril 2.5 mg) | Improve CFR, ↓ workload, may improve small vessel remodelling |
|                                  |                 |                                      | Ranolazine (e.g. 375 mg twice daily) | Improves microvascular perfusion reserve index in patients with MVA and reduced CFR |
| Microvascular angina ↑ vasoconstriction | Hyper-reactivity to stimuli (e.g. acetylcholine, exercise, stress) | Endothelial dysfunction, inappropriate pre-arteriolar vasoconstriction | Calcium antagonists (e.g. amlodipine 10 mg) | Vascular smooth muscle relaxation, ↓ myocardial oxygen consumption |
|                                  |                 |                                      | ACE inhibitors (e.g. ramipril 2.5 mg) | Improves endothelial vasomotor dysfunction |
|                                  |                 |                                      | Nicorandil (e.g. 5–10mg twice daily) | Potassium-channel activator with coronary microvascular dilatory effect |
|                                  |                 |                                      | Statins (e.g. rosuvastatin 10–20 mg) | Improve coronary endothelial function, pleiotropic effects including reduced vascular inflammation |
|                                  |                 |                                      | Calcium channel blockers (e.g. amiodipine 10 mg or verapamil 240 mg SR) | ↓ spontaneous and inducible coronary spasm via vascular smooth muscle relaxation and ↓ oxygen demand |
|                                  |                 |                                      | Nitrates (e.g. isosorbide mononitrate XL 30 mg) | ↓ spontaneous and inducible coronary spasm via large epicardial vasodilation, ↓ oxygen demand, lack of efficacy in microvascular angina with potential deleterious effect |
| Microvascular angina Abnormal pain processing | ↑ nociception | Dysfunctional cortical pain processing | Tricyclic antidepressants (e.g. imipramine up to 25 mg) | Improved symptom burden potentially through ↓ visceral pain |
|                                  |                 |                                      | Xanthine derivatives (e.g. aminophylline 225 mg twice daily) | Anti-algogenic effect (due to the direct involvement of adenosine in cardiac pain generation) |
| Vasospastic angina                | Propensity to coronary vasospasm | Vascular smooth muscle hyper-reactivity | Calcium channel blockers (β blockers) | ↓ spontaneous and inducible coronary spasm via vascular smooth muscle relaxation and ↓ oxygen demand |
|                                  |                 |                                      | Nitrates (e.g. isosorbide mononitrate XL 30 mg) | ↓ spontaneous and inducible coronary spasm via large epicardial vasodilation, ↓ oxygen demand, lack of efficacy in microvascular angina with potential deleterious effect |
| Adjunctive non-pharmacological interventions | May be useful in all endotypes | Metabolic syndrome, endothelial dysfunction, cardiovascular risk factors, anxiety/depression | Smoking cessation, exercise, cardiac rehabilitation, Mediterranean diet, cognitive behavioural therapy |

ACE = angiotensin converting enzyme; CFR = coronary flow reserve; MVA = microvascular angina; SR = sustained-release preparation.
Beyond Pharmacotherapy

The importance of addressing lifestyle factors cannot be overemphasized, particularly given that half of the patients in the CorMiCA study were clinically obese. Strategies to help address this, including exercise programmes and cardiac rehabilitation, may help facilitate important long-term lifestyle changes. Additionally, a new diagnosis of angina may increase the use of non-pharmacological therapies, including cardiac rehabilitation which may benefit patients with ischaemic heart disease.

In the CorMiCA study, we noted significantly lower illness perception scores at 6 months among the intervention arm, representing a less threatening view of illness. Angina reduction and improved quality of life scores could be in part related to better patient understanding and a less threatening perception of the illness. Longitudinal studies of other cardiovascular diseases have shown that illness perception is an important predictor of longer-term outcomes, including disability and returning to work. The Objective Randomised Blinded investigation with optimal Medical Therapy of Angioplasty in stable angina (ORBITA) trial highlights a placebo effect and we support that the positive diagnosis may be therapeutic in itself. Angina symptoms are often subjective and multifactorial in origin, so patient education and validation of symptoms may facilitate further improvement.

Future Directions

Our hope is for a personalised medicine approach whereby patients with different angina subtypes, defined by the results of coronary function tests, may benefit from targeted therapy. Further research is needed to determine whether this approach may lead to patient benefits. More widespread invasive testing allows identification of diagnostic subgroups for the development of targeted therapies guided by mechanistic studies. We recently identified systemic vascular abnormalities in patients with MVA and VSA, highlighting a potential therapeutic role for endothelin-receptor antagonists targeting the ET₁ receptor. In addition, Rho-kinase inhibitors represent a potential future therapeutic option with anti-effects in patients with excessive vascular smooth muscle constriction. More research is needed in well-defined patient endotypes (subgroups).

Conclusion

Patients with INOCA present a diagnostic and therapeutic challenge to physicians. MVA and/or VSA are the two most common causes of INOCA and may be overlooked using anatomical coronary tests alone. Invasive diagnostic testing permits a positive diagnosis to be made, or excluded, during the patients’ index presentation. Correct diagnosis of the underlying cause of angina permits stratified treatment of the distinct disorders (MVA, VSA or non-cardiac chest pain). CorMiCA has shown this approach to be safe, feasible with demonstrable benefit for patients.
Coronary

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