

Advances in the Post-coronary Artery Bypass Graft Management of Occlusive Coronary Artery Disease

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Abstract

Revascularisation of chronic total occlusion (CTO) represents one of the most challenging aspects of percutaneous coronary intervention, but advances in equipment and an understanding of CTO revascularisation techniques have resulted in considerable improvements in success rates. In patients with prior coronary artery bypass grafting (CABG) surgery, additional challenges are encountered. This article specifically explores these challenges, as well as antegrade methods of CTO crossing. Techniques, equipment that can be used and reference texts are highlighted with the aim of providing potential CTO operators adequate information to tackle additional complexities likely to be encountered in this cohort of patients. This review forms part of a wider series where additional aspects of patients with prior CABG should be factored into decisions and methods of revascularisation.

Keywords

Percutaneous coronary intervention, chronic total occlusion, coronary artery bypass grafting, saphenous vein graft, antegrade wire escalation, antegrade dissection re-entry, hybrid algorithm

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Coronary artery bypass grafting (CABG) surgery is a common method of coronary revascularisation and remains the standard of care in patients with multivessel coronary artery disease (CAD), as well as in those with diabetes.¹ Since 2004, CABG numbers have been in decline in the UK, whereas the number of percutaneous coronary interventions (PCI) has shown a consistent increase year-on-year until a recent plateau.² Post-CABG patients now represent a significant proportion of patients who subsequently require PCI and represent a challenging cohort in terms of clinical frailty and anatomical complexity.³ In this review we discuss these challenges and specifically consider the subset of post-CABG patients presenting with chronic total occlusions (CTO) in their native coronary arteries.

Challenges in Post-CABG Patients Graft Failure

Although surgical success rates remain high, venous bypass graft patency rates remain the 'Achilles' heel' of the long-term prognosis of surgical revascularisation. Whereas arterial revascularisation has been demonstrated as a superior, lasting method,⁴⁻⁶ venous bypass grafts do not withstand this test of time. Left internal mammary artery (LIMA) grafts can remain patent in 88–100% of patients at 15 years,^{7,8} a finding echoed by right internal mammary artery (RIMA) use, which retains excellent graft patency up to 10 years, with patency rates quoted at 81% or equivalent to the LIMA for identical coronary territories.^{8,9} However, saphenous vein grafts (SVG) have relatively poor patency rates.^{10,11} In their meta-analysis,

Athanasios et al. compared SVG patency with radial artery graft patency.¹² Of the seven studies examining patency rates after a median 5-year follow-up, four recorded SVG patency rates between 65% and 72%, whereas the others recorded higher rates (72–91%).¹² In a study of 1,074 patients, 10-year SVG patency rates were 61% when compared to LIMA grafts, where the patency rate was recorded at 85%.¹³

More contemporary data are available from the COMPASS-CABG substudy and POPular CABG trials, both of which used CT coronary angiography (CTCA) to assess SVG patency 1 year after surgery.^{14,15} In the COMPASS-CABG substudy, patients were treated with rivaroxaban, with or without aspirin or aspirin alone, with an overall 9.6% occlusion rate of all SVG studied.¹⁴ In the POPular CABG trials, graft occlusion occurred in 9.9% of all grafts, with no significant improvement despite the addition of ticagrelor antiplatelet therapy.¹⁵ CTCA has allowed non-invasive assessment of graft patency, and its wider use may uncover further aspects of post-surgical coronary anatomy and graft viability not previously appreciated.^{16,17}

In particular, attention should be paid to the increased likelihood that patients with existing CTOs and multivessel disease possess higher anatomical and clinical risk scores and are thus more likely to be referred for CABG in the first instance.¹⁸⁻²⁰ Yet, postoperative angiographic assessment in patients who underwent both on- and off-pump CABG for CTOs has revealed that grafts placed on non-left anterior descending artery (LAD) collateralised CTOs suffer from extremely poor patency rates

at 1-year follow-up, as low as 22–24%, which is an unacceptably low graft viability rate that should call into question the rationale for CABG in the presence of a non-LAD CTO.²¹

Revascularisation Complexity

Christopoulos et al. describe the post-CABG population as older, more likely to suffer from diabetes and have suffered from previous MI.²² CTOs are more prevalent in this subgroup of patients than in those without prior CABG, with registry data demonstrating the presence of a CTO in 54% of evaluated post-CABG patients.²³ Patients with a CTO and symptoms relating to ischaemia with myocardial viability do benefit from revascularisation versus optimal medical therapy alone, with improvements in both symptom burden and quality of life.²⁴ However, they are less likely to receive revascularisation therapy, likely due, in part, to the perceived complexity of the procedure.²³ In the Canadian Multicenter CTO Registry published by Fefer et al., of 1,697 patients identified with a CTO (and no prior CABG), medical therapy was opted for in 44% of patients, with 26% undergoing CABG (89% had a bypass graft on the CTO vessel) and 30% undergoing PCI. CTO PCI was attempted in only 31% of these patients and CTO success was achieved in only 24% of all patients undergoing PCI.²³ This registry (2008–2009) suggested the presence of CTOs to be approximately 18% of all patients with CAD, and yet just under half these patients received medical therapy alone, one-quarter received surgical revascularisation and the remainder underwent PCI.²³ This ‘interventional paradox’ will see some patients denied revascularisation for symptoms due to anatomical complexity and the perceived complexity of PCI. Furthermore, post-CABG patients will represent additional challenges when re-presenting with angina pectoris: they are likely to be older, have more comorbidities and have more complex coronary lesion characteristics, and for many, repeat CABG is not feasible due to excessive surgical risk compared with CABG-naïve patients.^{3,25–30}

Saphenous Vein Graft Intervention

Therefore, in post-CABG patients, PCI remains the only strategy for repeat revascularisation, yet the presence of a previous bypass graft creates additional challenges to conventional PCI. While medical therapy can be a good first option for the treatment of angina, PCI for moderate SVG stenoses when compared to optimal medical therapy (OMT) can be effective, with lower rates of major adverse cardiovascular events (MACE) at 1-year follow-up in the VELETI I trial.^{31–33} Although the VELETI I trial was a hypothesis-generating, small, randomised pilot trial, it put forward the concept of ‘plaque sealing’ of moderate, non-significant atheromata in SVGs, which are thought to undergo accelerated atherosclerotic disease progression compared with native vessels.³³ The subsequent larger randomised controlled VELETI II trial did not demonstrate any reduction in clinical endpoints in SVG PCI with drug-eluting stents (DES) at the 3-year follow-up compared with OMT in these so-called ‘intermediate’ lesions, although the pooled analysis of both VELETI trials may yet support the controversial concept of plaque sealing.^{34,35}

Percutaneous treatment of SVGs accounts for between 5% and 10% of all PCIs.^{36–42} Although, unsurprisingly, the vast majority of SVG PCIs are performed within the body of the graft, approximately one-fifth of graft lesions occur at the aorto-ostial anastomosis and one-sixth occur at the distal anastomosis.⁴³ Acute thrombotic SVG occlusion must be managed in the same manner as native coronary occlusion and, although procedural success tends to be high, mortality, recurrent acute coronary syndrome (ACS) and the need for revascularisation within the short to medium term remains significant.^{42,44} Preference is given to revascularising the native coronary artery over SVG by existing guidelines on myocardial

revascularisation.¹ The paucity of data for this recommendation has led to development of the PROCTOR trial, a multicentre, multinational European randomised control trial, which will randomise patients to native vessel or SVG PCI, with results expected in 2027.⁴⁵

The physiology of SVG failure is not fully understood, but it is thought these grafts are poorly adapted to arterial flow and the pathobiology of SVG degeneration results in a friable vessel with atheromatous debris to contend with.^{46,47} Additional challenges include the potential for embolisation of this debris into distal epicardial and coronary microcirculatory vasculature, resulting in the plugging of capillaries, increasing the prospect of the no-reflow phenomenon and associated risk of MI and subsequent in-hospital mortality.^{48–50} The routine use of distal embolic protection devices (DPD) has shown potential to significantly reduce periprocedural MI rates, but no significant reduction in in-hospital mortality could be demonstrated.^{51–54} However, these devices are cumbersome to deploy and, as such, their use has been historically limited.^{55,56} Furthermore, several observational studies and large registry data have shown conflicting results.^{57,58} Thus, the strength of recommendation for the use of DPD for SVG PCI was downgraded in the most recent update of the European Society of Cardiology (ESC)/European Association for Cardio-Thoracic Surgery (EACTS) guidelines on myocardial revascularisation to a Class IIa, level of evidence B, recommendation.¹ Female sex, lesion length, extensive degenerative change and high plaque volume in diseased SVGs predict 30-day MACE.^{59,60}

Periprocedural MI (as defined by a rise in creatine kinase (CK)-MB between $\times 1$ and $\times 5$ the upper limit of normal) was a stronger predictor of adverse outcome than similar cardiac enzyme values following SVG PCI and a powerful predictor of late cardiovascular mortality in an albeit dated study, with the overall rate of periprocedural MI reported as 15%.⁶¹ Periprocedural increases in CK-MB following SVG PCI are unsurprisingly significantly greater when no-reflow occurs (43% versus 4%, $p < 0.001$) with probable thrombus, ACS presentation, graft degeneration and graft ulceration independent predictors of no-reflow.⁴⁸ More contemporary studies of SVG PCI tend to use DPD, such as the post hoc analysis of the DIVA trial comparing direct stenting against stent deployment with or without balloon inflation (either prior to and/or after stent implantation).⁶² Patients were recruited to the DIVA trial between 2012 and 2015 and DPD use was $>70\%$ in both groups. Rates of periprocedural MI were low, at 4% of total lesions treated and 5% of patients treated.⁶² The use of DES in SVGs is now supported by a number of trials, all demonstrating poor longevity following treatment with plain old balloon angioplasty and covered stents.^{63–66} DES is advocated for SVG PCI due to lower rates of repeat revascularisation compared with the use of bare metal stents, although clinical outcome data remain conflicted, with only a limited number of randomised trials available.^{1,52,53,67–69} In the absence of randomised control data comparing SVG and native vessel PCI, registry data suggest SVG PCI remains inferior to native vessel PCI, with higher MACE rates, principally driven by MI and revascularisation rates, at 1 year.⁷⁰ A history of previous bypass graft surgery is associated with a higher risk of restenosis, and SVG as the PCI target is independently associated with an increased risk of very-late stent thrombosis.^{71,72}

Chronic Total Occlusion Revascularisation in Post-CABG Patients

Although there remains a paucity of data from randomised control trials supporting CTO revascularisation, symptom- and, where relevant, myocardial viability-driven revascularisation has been established by the EuroCTO Trial.²⁴ This approach is supported by the latest guidelines.¹

The DECISION-CTO trial did not report an improvement in quality of life outcomes, although this trial fell short in recruitment and thus was stopped early.⁷³ Although the trialists have been congratulated for the large number of patients randomised, several limitations have been identified, including a lack of baseline symptoms, cross-over to the CTO PCI group (from the non-CTO PCI group) and the non-inferiority and pre-PCI randomisation design, in addition to the underpowered study.⁷⁴

CTOs in the presence of bypass grafts are often longer in length with a higher calcific burden and diffuse atherosclerotic disease.^{75,76} These CTOs are themselves complex, as graded by the frequently adopted Multicenter CTO Registry of Japan (J-CTO) score, with higher J-CTO scores than in non-CABG patients, and suffer from greater anatomic distortion, with three-dimensional tenting effects exerted on the native vessel at the distal graft anastomosis.^{22,28,77–81} It is unclear whether this is the result of a pre-existing heavy burden of disease that necessitated CABG revascularisation in the first instance, accelerated atherosclerosis or the presence of the distal graft anastomosis resulting in disease progression due to competitive flow.^{82–86} In addition to these anatomical and pathophysiological factors, patient characteristics must also be considered. Evolution of knowledge, techniques and, perhaps most importantly, equipment has facilitated higher rates of success in CTO revascularisation. Among these, the introduction of microcatheters has dramatically altered the ability of operators to safely and successfully cross CTOs and these should be used in all CTO PCI cases regardless of complexity. Microcatheters are further elaborated on below.

Revascularisation by redo CABG in patients with prior CABG is not without jeopardy, with a two- to fourfold increased risk compared with first-time CABG.^{87,88} However, mortality was comparable between PCI and redo-CABG for these patients at 3 years, with higher rates of revascularisation in PCI patients.^{39,89,90} Post-CABG patients can suffer cardiac tamponade at the same frequency as non-CABG patients.⁹¹ In addition, prior CABG is associated with reduced event-free survival, with higher rates of cardiac death and MACE demonstrated by univariable analysis and higher rates of MACE demonstrated by multivariable analysis, driven largely by target vessel revascularisation.²⁸ ‘Dry tamponade’ has been recognised as a significant complication of coronary perforation in post-CABG patients, caused by the extravasation of blood within the myocardial wall or adjacent structures within a pericardium with more adhesions.^{92,93} In-hospital complications are also more frequent in prior CABG patients undergoing CTO PCI than in non-CABG patients, as reported in a multicentre registry of 2,058 patients (prior CABG n=401; non-CABG n=1,657), with higher rates of major complications (3.7% versus 1.5%), any perforation (12% versus 5.2%), periprocedural MI (2.0% versus 0.5%) and procedure-related deaths (0.8% versus 0.1%).²⁷

In a smaller study of 470 patients, contrast-induced nephropathy was more common in prior CABG patients (4.6% versus 1.0%).²⁸ Risk scores developed to predict CTO PCI success, such as the RECHARGE-Score and Clinical and Lesion-related score (CL Score), attribute higher scores to post-CABG patients, reflecting these adverse events.^{94,95} Of note, previous CABG will preclude rapid and safe sternotomy if a complication arises following or during PCI, and this may have contributed to some of the morbidity seen in these scoring systems.⁹⁶ Having a sufficiently experienced team to manage complications in post-cardiotomy patients in high-volume PCI centres is essential. The recognition of longer procedures and older and potentially frailer patients with reduced renal function should be considered when evaluating the benefits of potential percutaneous CTO revascularisation, and the ways in which this can be

mitigated are further elaborated on below. Conversely, the presence of patent grafts, in addition to providing potential retrograde conduits, can reduce ischaemia in the distal target territory and, in the case of a patent LIMA, reduce the consequences of anterior wall ischaemia from inadvertent left main coronary artery dissection.

Factoring in prior CABG, the presence of a non-proximal lesion position, proximal tortuosity (moderate/severe) and distal cap ambiguity, described as the ‘J-CTO+ model’ improved the power of the J-CTO score in predicting successful CTO crossing.⁷⁸ These factors provide additional challenges over and above what may be encountered in native vessel CTO PCI in the absence of graft anastomoses. Understanding these potential challenges up front allows the operator to select appropriate techniques and tools to approach CTO cases where SVGs are involved.

Since early pioneers such as Kaltenbach and Reifart in Frankfurt and Rutherford in Kansas City described their experiences in CTO treatment, significant advances in the understanding of pathology, technology and the formulation of accepted standards and techniques have been made, resulting in significantly improved long-term treatment success rates.^{94,97,98} Original descriptions of CTO PCI were fraught, with difficult, long procedures and prohibitively high reocclusion rates.^{99,100} Early concepts led to the subsequent development of contemporary tools now in use. The formation of ‘CTO Clubs’, such as the Japanese CTO Club in 1991 and the European equivalent in 2006, improved the sharing and dissemination of knowledge and the development of techniques to improve success rates and reduce periprocedural morbidity. The development of registries such as PROGRESS-CTO and RECHARGE, randomised trials and regional consensus documents have provided a basis for understanding accepted techniques and monitoring contemporary practice, including complication and morbidity rates.^{24,73,94,101,102} Among these developments, the hybrid algorithm is the currently accepted consensus strategy being used by high-volume, experienced leaders in the field.¹⁰³ This demands the ability to adopt both antegrade and retrograde approaches to CTO crossing to ensure the optimal use of available techniques with contemporary equipment, with further results from adopting this approach still being reported.

The RECHARGE Registry is thus far the largest of its kind, with over 1,200 patients recruited from European centres to demonstrate both high procedural success rates and low adverse event rates when the hybrid algorithm has been used by experienced operators in high-volume centres.⁹⁷ It is therefore important to recognise the potential benefits gained through the ability to adopt different strategies to recanalise a CTO and, most pertinently, recognise when to opt for a specific strategy. Retrograde techniques are most often necessary in post-CABG patients due to the complexities described above, usually in conjunction with antegrade techniques that then form the hybrid algorithm approach to these patients.²² Retrograde crossing is more common in post-CABG patients where the SVG can often be used as the collateral channel.²²

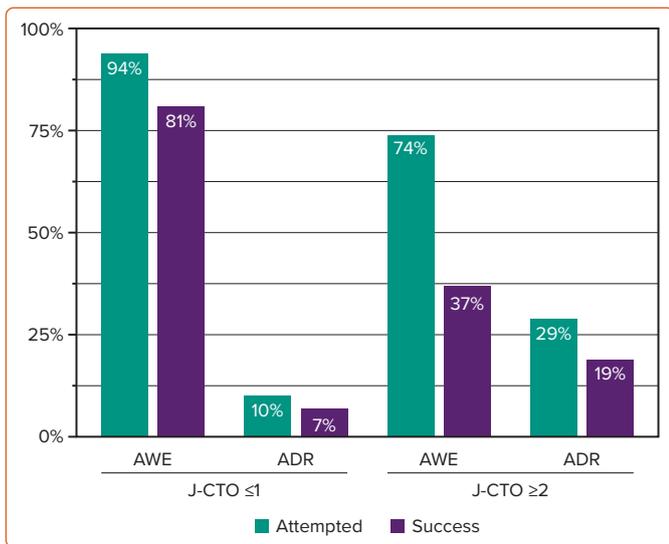
Whether the native vessel CTO or, indeed, the SVG should be treated in post-CABG patients with graft degeneration remains unclear. Current ESC guidelines recommend PCI as the preferred method of revascularisation in patients with a large burden of ischaemia or severe symptoms due to disease progression or late graft failure.¹ However, evidence to support this position is sparse, with limited data implicating prior CABG with poor outcomes, as discussed above, with patient clinical characteristics rather than revascularisation method predominantly determining outcome and anatomical considerations dictating the method of revascularisation.^{29,71,72}

Table 1: Comparison of Chronic Total Occlusion CABG-Naïve and Post-CABG patients

	Prior CABG (n=176)	No Prior CABG (n=320)	p-value
Target vessel (%)			0.07
Right	63	60	
Left anterior descending	13	24	
Left circumflex	18	10	
Other	6	6	
Moderate/severe calcification (%)	74	47	<0.001
Moderate/severe tortuosity (%)	42	26	<0.001
Lesion length (mm); median (IQR)	39 (28–67)	30 (20–40)	<0.001
Lesion age (months); median (IQR)	44 (6–90)	10 (3–42)	<0.01
Previous CTO attempt (%)	16	18	0.61
J-CTO score; mean (±SD)	3.12 ± 1.03	2.41 ± 1.21	<0.001

CABG = coronary artery bypass grafting; CTO = chronic total occlusion; IQR = interquartile range. Source: Christopoulos et al.²² Adapted with permission from Elsevier.

Figure 1: Applications and Outcomes Following the Use of the Hybrid Algorithm Stratified by J-CTO Score



Use and success of antegrade wire escalation (AWE) and antegrade dissection re-entry (ADR) in the RECHARGE Registry of Hybrid Algorithm CTO crossing. Source: Maeremens et al.⁹⁷ Adapted with permission from Elsevier.

Retrospective and comparative studies have attempted to address this.^{70,104} However, the PROCTOR trial will be the first randomised trial comparing SVG PCI to native vessel PCI and should help in the decision making for patients with SVG degeneration and stenosis.⁴⁵

Antegrade Techniques

Operators should be able to call on existing, established techniques of antegrade CTO crossing. These will include antegrade wire escalation (AWE) and/or antegrade dissection re-entry (ADR). Some of these techniques are highlighted below, with supporting evidence discussed.

CTOs in the post-CABG cohort exhibit a higher calcific burden, increased tortuosity, longer lesion length and established occlusions for a longer duration, resulting in higher J-CTO scores²² (Table 1).^{22,76} Although AWE can be a successful strategy and is the default in most cases of CTO,

particularly when J-CTO scores are ≤1 (Figure 1), the increased complexity likely to be present in post-CABG patients will often necessitate additional and adjunctive hybrid algorithm techniques.^{78,97} Antegrade techniques are highly useful in many CTO cases with differences in pathology within the CTO body, but must contend with the presence of more complex, calcified distortions of the artery with severe negative remodelling present than in short-duration CTOs in patients without prior CABG.⁷⁶ In the RECHARGE Registry, less complex lesions (J-CTO score ≤1) were successfully crossed using an AWE approach with high success rates (86%), whereas ADR and retrograde techniques were often used as bailout strategies with reasonable success.⁹⁷ However, in more complex lesions (J-CTO score ≥2), AWE was a less successful strategy (50%), requiring ADR and retrograde bailout approaches more frequently.⁹⁷

Planning Revascularisation

Up-front careful analysis of the coronary angiogram is key to understand potential challenges likely to be encountered during the CTO PCI and can improve success rates considerably.¹⁰⁵ The coronary angiogram for the CTO should be acquired without digital magnification in order to visualise the entire course of the vessel, with a long acquisition allowing full visualisation of any antegrade flow either through the CTO or antegrade bridging collaterals into the distal vessel. Large side branches and relevant bifurcations should be noted to help decide which strategy should be used. Where graft anastomoses are present beyond the distal cap of the CTO and where the graft remains patent, antegrade injection along the graft (again without digital magnification) should be used to better visualise the course of the vessel, although this may not be fully appreciated by invasive coronary angiography alone. If dual catheter injections are possible, simultaneous injections first down the patent graft, followed by the native coronary, can provide useful information on occlusion length and potential distal landing zones should an ADR (or, indeed, retrograde) strategy be used. Complex revascularisation attempts, particularly when prior failure has occurred, should be discussed with experienced CTO operators in high-volume centres where familiarity with the hybrid algorithm can be used to establish higher success rates.^{78,97}

Growing evidence supports the use of CTCA as an effective tool for CTO procedural planning in both CABG-naïve and post-CABG patients.^{106–108} However, CT can have limitations here, particularly when high calcium burdens are encountered, making interpretation challenging, which is more likely in post-CABG patients.^{43,109} The CT-RECTOR study assessed the predictive value of successful CTO crossing with prior CTCA and was validated against the J-CTO score, suggesting the CT-RECTOR scoring system provided additive data aiding a successful procedure and optimising procedural time.^{107,108} However, post-CABG patients comprised only 17% and 11% of those included in the studies and, as such, the data should be interpreted with caution in this cohort.

CTOs that have developed in post-CABG patients may have developed multiple native collaterals prior to or since graft degeneration and, as such, visualising the distal vessel may prove challenging. This can be overcome by using retrograde injections from both the diseased graft and contralateral native coronary artery, necessitating the use of an additional vascular access point and a third guide catheter.

Wires

Antegrade techniques use advances in coronary angioplasty wire technology allowing greater options for the operator. These improvements provide the operator with wires with greater torque, steerability and

tactile feedback, as well as improvements in wire tip force, and thus greater potential penetration strength. Wire escalation and success in this manner depends on a good understanding of wire properties. Wires will, in general, be hydrophobic, hydrophilic or polymer jacketed, with the latter providing the greatest lubricity, with the pay-off a reduction in tactile feel. Some wires will combine these features, allowing a balance of both 'slip' through lesions while allowing the tip to grip lesions and still provide some tactile feedback. It is usual to advance a 'workhorse' wire to the lesion, then escalate by exchanging to an appropriate wire, determined by the operator's appreciation of the occlusion characteristics.

Microchannels and loose tissue through the body of the CTO may be accessed via the proximal cap and, as such, a light yet slippery (hydrophilic or polymer-jacketed) wire with high torque response may be selected to successfully traverse the CTO. Histological findings from a sudden coronary death registry have provided insights into CTO lesion morphology in individuals with and without prior CABG.⁷⁶ Histological parameters were used to further subdivide CTOs into those with histological parameters suggestive of a 'short' or 'long' duration and compared with those present in individuals where CABG had been performed at least 2 years prior to autopsy. Although no significant difference was demonstrated between these individuals, a tapered distal cap was more commonly reported in CTOs in prior CABG individuals, whereas an abrupt pattern was noted in the proximal cap, a finding also noted in the 'long'-duration CTOs examined.⁷⁶ Tapered proximal occlusions feature loose fibrous tissue with small microvessel recanalisation and so may be more amenable to wire crossing.¹¹⁰ Therefore, a retrograde approach to cross the cap and access the CTO body may be required in prior CABG patients, with heavier, more penetrative wires necessary to cross abrupt (blunt) caps. Gaia wires (Asahi Intecc) are a dedicated family of CTO wires that improve penetration while retaining tactile feedback due to their design featuring a 'microcone' tip.¹¹¹

Heavier tip force and penetrative wires may be required to engage and cross the proximal and distal caps, which are formed of denser tissue than the body of the CTO. Tip loads vary from workhorse wires, which are typically ≤ 1 g, to gradual increases in tip loads as high as 40 g with the Astatto XS 40 wire (Asahi Intecc), which delivers an equivalent penetration force of 796.2 kg/inches².⁷⁹ Caution must be exercised when traversing the CTO body with highly penetrative wires, particularly in tortuous and ambiguous vessels where tactile feedback is at a minimum. This is more apparent in post-CABG patients, where the distal graft anastomosis can alter vessel anatomy and result in tenting of the distal vessel. AWE demands an appreciation for wire properties so they are selected to tackle anticipated challenges likely to be encountered. Furthermore, AWE demands an understanding of when to escalate, when to de-escalate and subsequently when re-escalate, if and when appropriate. More detailed information regarding the specifics of wire choice when escalating in an antegrade fashion is available in the antegrade CTO book by Spratt et al.⁷⁹

Antegrade Dissection Re-entry

When the CTO plaque cannot be traversed through the proximal cap or through the body of the CTO due to obstructions through an antegrade manner, it is often necessary to switch to an ADR strategy. The higher burden of calcium in post-CABG patients may represent one of these obstructions that cannot, despite the use of high-tip-force wires and adjuncts (elaborated on below), be crossed through in a true lumen-to-lumen fashion.⁷⁶ To perform ADR, the subintimal space must be accessed, and an 'umbrella' shape is often used on a polymer-jacketed wire to drive the wire forward in a knuckle fashion to access the subintimal space with relative safety. Caution should be exercised when the subintimal

space is accessed towards the distal graft anastomosis to ensure the dissection plane does not extend to or beyond this anastomosis, creating haematoma and thereby potentially occluding graft flow into the distal vessel. The plane of dissection created in this manner should be kept to a short distance from this area and re-entry into the distal true lumen should be attempted in a previously identified distal landing zone. This can be facilitated by using the Bridgepoint System (CrossBoss coronary catheter and Stingray LP CTO re-entry system; Boston Scientific), which provides a more controlled manner with which to advance equipment through the subintimal space with a smaller dissection plane created and targeted re-entry into the distal lumen, demonstrating higher rates of success than the less controlled knuckle wire technique.⁹⁷

Dual-injection angiography allows an appropriate distal landing zone to be chosen, ideally proximal to the distal graft anastomosis so as not to compromise graft flow (when patent). ADR may not be the ideal strategy when the re-entry zone from the subintimal space back into the true vessel lumen is within 10 mm before the distal graft anastomosis or important side branches due to the risk of extension of the dissection plane and the resulting occlusion of these branches.¹¹² Whether the subintimal space is accessed intentionally during ADR or inadvertently during attempted AWE, antegrade contrast injections should be avoided in order to minimise hydraulic extension of the subintimal space, resulting in compression of the true lumen and thereby reducing the likelihood of successful re-entry.⁸⁰ Techniques, such as STAR (Sub-intimal TrAcking and Reentry) and LAST (Limited Antegrade Subintimal Tracking), are recognised alternative techniques to traverse the subintimal space and then re-enter into the distal lumen, but are not favoured over the CrossBoss/Stingray system due to a lack of predictable longer-term success.¹¹³⁻¹¹⁵ It is important to recognise the need for at least a 7 Fr system to facilitate the passage and exchange of ADR equipment.

Caution must be exercised when using ADR near side branches. Wires, and subsequently microcatheters, will tend to follow the path of least resistance and, as such, can follow subintimal tracks to enter and dissect side branches, particularly hazardous when antegrade contrast injections are prohibited and so these branches cannot be adequately visualised. Targeted re-entry by identifying a suitable distal landing zone for luminal re-entry and utilising the Stingray balloon, for instance, can help avoid dissection extension and side branch compromise. The CrossBoss catheter features a blunt, atraumatic, 1.0 mm tip and can safely traverse the subintimal plane, but it should not be used as an initial strategy to engage the proximal cap, particularly when ambiguous with multiple bridging collaterals, in the presence of extreme vessel tortuosity or when the vessel course is unclear. The use of a knuckled wire, guide catheter support systems (discussed below) and anchor balloons in proximal side branches can enhance support to enable passage of the CrossBoss catheter and delivery of the Stingray balloon. Once in the subintimal space, the CrossBoss will rarely exit due to the low resistance of the surrounding structures, but short segments of intimal tracking can be evident.^{80,97}

The CrossBoss catheter is not steerable and, as such, advancement through the target vessel structure should be regularly monitored with non-contrast fluoroscopy in orthogonal planes during controlled advancement. Retrograde injections can be of use to ensure the CrossBoss moves in synchrony with the architecture of the visualised distal vessel, so-called 'dancing' with the target zone for luminal re-entry. The CrossBoss will track the outer curve of the vessel and, as such, can pass into small side branches, which, if unrecognised

prior to further advancement, can result in vessel exit and coronary perforation, a non-negligible complication contributing to a high burden of morbidity and mortality in this small minority of patients.¹¹⁶ In the event of the CrossBoss entering a side branch, it should be withdrawn and a guidewire used to track beyond the side branch ostium, allowing the CrossBoss to then be delivered beyond this (wire redirect).⁸⁰ In the event that wire crossing beyond the side branch is not possible, a knuckled wire can be used to cross the side branch then redeliver the CrossBoss and attempt advancement once again (knuckle redirect).⁸⁰ Should this also not be successful, a small balloon with a 1:1 ratio to the side branch can be placed in the side branch ostium to deflect passage of a knuckled guidewire into the side branch subintimal space or, alternatively, a dual lumen catheter can be used to allow a second guidewire to cross beyond the side branch, thereby enabling advancement of the CrossBoss also beyond the side branch ostium, where it can then be advanced ahead of the guidewire.⁸⁰

Adjunctive Equipment

Adequate support is key to overcoming the proximal and distal caps. This comes initially from the choice of vascular access. Femoral access affords larger bore access (up to 8 Fr commonly used) with the option of long sheaths to overcome iliac and aortic tortuosity. Biradial access may also allow insertion of a 7 Fr sheath, particularly when using slender sheaths that reduce the outer diameter by 1 Fr. Therefore, appropriate guide catheter selection is imperative, and guide catheters should be selected specifically for graft access where retrograde approaches or distal landing zone visualisation is necessary. Adjunctive support systems, such as guide catheter extensions, provide additional support to aid cap puncture, but are particularly beneficial during ADR.

Advancement of the guide catheter extension into the coronary artery to the point at which endothelial dissection occurred can reduce influx of blood into the subintimal space, thereby reducing haematoma formation and subsequent compression of the true lumen. Re-entry beyond the distal cap is therefore aided using a guide catheter extension, maintaining luminal size for a greater likelihood of success into the true lumen. Post-CABG CTO crossing may require multiple wire changes in AWE or ADR approaches, particularly when using knuckle wires or the CrossBoss catheter; as such, equipment allowing rapid exchange with balloon trapping aids efficiency. The Trapliner (Teleflex) guide extension catheter features a proximal balloon that aids this without the need for additional balloon trapping within the guide catheter system and can be a useful tool in cases such as post-CABG CTO revascularisation.

In longer CTOs, a retrograde approach using both retrograde dissection re-entry (RDR) and ADR techniques may be necessary.^{78,97,98} RDR will involve accessing the subintimal space either distal to or through the distal CTO cap, and the reverse controlled antegrade and retrograde tracking (reverse CART) technique is currently the dominant RDR technique, with high success rates.¹¹⁷ The antegrade aspect to reverse CART involves ADR to facilitate overlapping knuckle wires followed by balloon dilation of the subintimal space to connect the common space between the retrograde and antegrade dissection planes. Following this, retrograde wiring of the guide catheter can be performed should the reverse CART be performed in the proximal portion of the vessel or if more distal a guide catheter extension can be advanced to the point of antegrade dissection (as described above) and facilitate efficient retrograde wiring of the antegrade guide.

Microcatheters have greatly improved the efficiency of wire exchange but also provide additive penetrative forces that can be applied to high-resistance areas within the CTO. Each microcatheter retains specific properties that allow engagement into the proximal cap and can provide support for microcatheter and wire advancement with exchange when necessary. This can reduce friction on the wire through the body of the CTO and allow improved torque transmission.⁷⁹ Microcatheters vary in their construction and so possess specific properties in terms of their size, lubricity, push force and the ability to track the wire and vessel. Microcatheters can be categorised as coil and non-coil based, with braided and non-braided catheters suited to different levels of penetration force and anatomy. Further details and comparisons of selected microcatheters can be found in the antegrade CTO book by Spratt et al. (chapter 7, section 20).⁷⁹

As described above, calcium is a prominent feature in post-CABG patients. As such, it is essential to have calcium modification and imaging tools available and to use them where necessary. Rotational atherectomy ('rotablation') and, more recently, intravascular lithotripsy (IVL; Shockwave Medical, Fremont, California) provide tools to modify calcium, whereas intravascular imaging tools, such as intravascular ultrasound, are critical tools required to understand the calcium burden, pattern and location and the interval effects of calcium modification.¹¹⁸⁻¹²⁰ Other available calcium-modification tools include cutting, scoring and high-pressure balloons. It may be necessary to use these tools in conjunction with each other to allow successful CTO crossing and optimal stent placement in CTO vessels with a high calcium burden.

Deliberate Vein Graft Sacrifice

Following successful revascularisation of CTOs in post-CABG patients, consideration should be given as to whether a patent SVG will provide excessive competitive flow to the distal vessel and therewith reduce long-term patency rates in the reconstructed native vessel. In a retrospective analysis of consecutive post-CABG patients where deliberate SVG sacrifice was performed, mostly by using vascular plugs, Wilson et al. demonstrated this to be a safe and effective method, with high success and low periprocedural complication rates, in these patients.¹²¹ Although more data are still needed to demonstrate whether this technique can improve long-term revascularised CTO vessel patency, consideration should be given to this approach in selected cases.

Conclusion

CTO crossing has improved with available data, advances in technology and techniques, among which the hybrid algorithm has played a crucial role, resulting in high success rates and, importantly, excellent long-term outcomes. Understanding the challenges of CTO revascularisation in post-CABG patients in terms of anatomical and lesion characteristics and clinical patient factors is necessary to prepare operators for selecting appropriate strategies and techniques that it may be necessary to have available in the operator armamentarium for successful CTO crossing and outcomes. Older, frailer patients with multiple comorbidities and more complex, established lesions with increased anatomical variance will need to be appreciated and contended with. Antegrade CTO crossing in these patients is possible, yet it is important to recognise the need to have retrograde options available, particularly because vein grafts can act as excellent conduits to the distal vessel. Experienced operators and high-volume centres will offer these patients a good chance of improvements in symptoms and quality of life, the essence of CTO treatment. □

1. Neuman FJ, Sousa-Uva M, Ahlsson A, et al. 2018 ESC/EACTS guidelines on myocardial revascularization. *Eur Heart J* 2019;40:87–165. <https://doi.org/10.1093/eurheartj/ehy394>; PMID: 25482397.
2. British Heart Foundation. Heart & circulatory disease statistics 2020. <https://www.bhf.org.uk/-/media/files/research/heart-statistics/bhf-statistics-compendium-2020.pdf> (accessed 10 May 2021).
3. Budassi S, Zivelonghi C, Dens J, et al. Impact of prior coronary artery bypass grafting in patients undergoing chronic total occlusion—percutaneous coronary intervention: procedural and clinical outcomes from the Registry of Crossboss and Hybrid procedures in FrAnce, the NetheRlands, BelGium, and UnitEd Kingdom (RECHARGE). *Catheter Cardiovasc Interv* 2021;97:e51–60. <https://doi.org/10.1002/ccd.28954>; PMID: 32369681.
4. Loop FD, Lytle BW, Cosgrove DM, et al. Influence of the internal-mammary-artery graft on 10-year survival and other cardiac events. *N Engl J Med* 1986;314:1–6. <https://doi.org/10.1056/nejm1986101023140101>; PMID: 3484393.
5. Taggart DP, D'Amico R, Altman DG. Effect of arterial revascularisation on survival: a systematic review of studies comparing bilateral and single internal mammary arteries. *Lancet* 2001;358:870–5. [https://doi.org/10.1016/s0140-6736\(01\)06069-x](https://doi.org/10.1016/s0140-6736(01)06069-x); PMID: 11567701.
6. Taggart DP, Benedetto U, Gerry S, et al. Bilateral versus single internal-thoracic-artery grafts at 10 years. *N Engl J Med* 2019;380:437–46. <https://doi.org/10.1056/nejmoa1808783>; PMID: 30699314.
7. Barner HB, Barnett MG. Fifteen- to twenty-one-year angiographic assessment of internal thoracic artery as a bypass conduit. *Ann Thorac Surg* 1994;57:1526–8. [https://doi.org/10.1016/0003-4975\(94\)90114-7](https://doi.org/10.1016/0003-4975(94)90114-7); PMID: 8010797.
8. Tatoulis J, Buxton BF, Fuller JA. Patencies of 2,127 arterial to coronary conduits over 15 years. *Ann Thorac Surg* 2004;77:93–101. [https://doi.org/10.1016/s0003-4975\(03\)01331-6](https://doi.org/10.1016/s0003-4975(03)01331-6); PMID: 14726042.
9. Tatoulis J, Buxton BF, Fuller JA. The right internal thoracic artery: the forgotten conduit – 5,766 patients and 991 angiograms. *Ann Thorac Surg* 2011;92:9–17. <https://doi.org/10.1016/j.athoracsur.2011.03.099>; PMID: 21718825.
10. Taggart DP. Current status of arterial grafts for coronary artery bypass grafting. *Ann Cardiothorac Surg* 2013;2:427–30. <https://doi.org/10.3978/j.issn.2225-319x.2013.07.21>; PMID: 23977618.
11. Tranbaugh RF, Schwann TA, Swistel DG, et al. Coronary artery bypass graft surgery using the radial artery, right internal thoracic artery, or saphenous vein as the second conduit. *Ann Thorac Surg* 2017;104:553–9. <https://doi.org/10.1016/j.athoracsur.2016.11.017>; PMID: 28215422.
12. Athanasiou T, Saso S, Rao C, et al. Radial artery versus saphenous vein conduits for coronary artery bypass surgery: forty years of competition – which conduit offers better patency? A systematic review and meta-analysis. *Eur J Cardiothorac Surg* 2011;40:208–20. <https://doi.org/10.1016/j.ejcts.2010.11.012>; PMID: 21167726.
13. Goldman S, Zalina K, Moritz T, et al. Long-term patency of saphenous vein and left internal mammary artery grafts after coronary artery bypass surgery: results from a Department of Veterans Affairs cooperative study. *J Am Coll Cardiol* 2004;44:2149–56. <https://doi.org/10.1016/j.jacc.2004.08.064>; PMID: 15582312.
14. Lamy A, Eikelboom J, Sheth T, et al. Rivaroxaban, aspirin, or both to prevent early coronary bypass graft occlusion: the COMPASS-CABG study. *J Am Coll Cardiol* 2019;73:121–30. <https://doi.org/10.1016/j.jacc.2018.10.048>; PMID: 30654882.
15. Willemsen LM, Janssen PWA, Peper J, et al. The effect of adding ticagrelor to standard aspirin on saphenous vein graft patency in patients undergoing coronary artery bypass grafting (POPular CABG): a randomized, double-blind, placebo-controlled trial. *Circulation* 2020;142:1799–807. <https://doi.org/10.1161/circulationaha.120.050749>; PMID: 32862716.
16. Chan M, Ridley L, Dunn DJ, et al. A systematic review and meta-analysis of multidetector computed tomography in the assessment of coronary artery bypass grafts. *Int J Cardiol* 2016;221:898–905. <https://doi.org/10.1016/j.ijcard.2016.06.264>; PMID: 27439070.
17. Mushtaq S, Conte E, Pontone G, et al. Interpretability of coronary CT angiography performed with a novel whole-heart coverage high-definition CT scanner in 300 consecutive patients with coronary artery bypass grafts. *J Cardiovasc Comput Tomogr* 2020;14:137–43. <https://doi.org/10.1016/j.jcct.2019.08.004>; PMID: 31405817.
18. Christofferson RD, Lehmann KG, Martin GV, et al. Effect of chronic total coronary occlusion on treatment strategy. *Am J Cardiol* 2005;95:1088–91. <https://doi.org/10.1016/j.amjcard.2004.12.065>; PMID: 15842978.
19. Kappetein AP, Dawkins KD, Mohr FW, et al. Current percutaneous coronary intervention and coronary artery bypass grafting practices for three-vessel and left main coronary artery disease. Insights from the SYNTAX run-in phase. *Eur J Cardiothorac Surg* 2006;29:486–91. <https://doi.org/10.1016/j.ejcts.2006.01.047>; PMID: 16497510.
20. Serruys PW, Kogame N, Katagiri Y, et al. Clinical outcomes of state-of-the-art percutaneous coronary revascularisation in patients with three-vessel disease: two-year follow-up of the SYNTAX II study. *EuroIntervention* 2019;15:e244–52. <https://doi.org/10.4244/eij-d-18-00980>; PMID: 30636684.
21. Widimsky P, Straka Z, Stros P, et al. One-year coronary bypass graft patency. *Circulation* 2004;110:3418–23. <https://doi.org/10.1161/01.cir.0000148139.79580.36>; PMID: 15557371.
22. Christopoulos G, Menon RV, Karpaliotis D, et al. Application of the 'hybrid approach' to chronic total occlusions in patients with previous coronary artery bypass graft surgery (from a contemporary multicenter US registry). *Am J Cardiol* 2014;113:1990–4. <https://doi.org/10.1016/j.amjcard.2014.03.039>; PMID: 24793678.
23. Fefer P, Knudtson ML, Cheema AN, et al. Current perspectives on coronary chronic total occlusions: the Canadian multicenter chronic total occlusions registry. *J Am Coll Cardiol* 2012;59:991–7. <https://doi.org/10.1016/j.jacc.2011.12.007>; PMID: 22402707.
24. Werner GS, Martin-Yuste V, Hildick-Smith D, et al. A randomized multicentre trial to compare revascularization with optimal medical therapy for the treatment of chronic total coronary occlusions. *Eur Heart J* 2018;39:2484–93. <https://doi.org/10.1093/eurheartj/ehy220>; PMID: 29722796.
25. Pereg D, Fefer P, Samuel M, et al. Long-term follow-up of coronary artery bypass patients with preoperative and new postoperative native coronary artery chronic total occlusion. *Can J Cardiol* 2016;32:1326–31. <https://doi.org/10.1016/j.cjca.2016.01.015>; PMID: 27118056.
26. Brilakis ES, O'Donnell CI, Penny W, et al. Percutaneous coronary intervention in native coronary arteries versus bypass grafts in patients with prior coronary artery bypass graft surgery: insights from the Veterans Affairs Clinical Assessment, Reporting, and Tracking Program. *JACC Cardiovasc Interv* 2016;9:884–93. <https://doi.org/10.1016/j.jcin.2016.01.034>; PMID: 27085582.
27. Azzalini L, Ojeda S, Karatasakis A, et al. Long-term outcomes of percutaneous coronary intervention for chronic total occlusion in patients who have undergone coronary artery bypass grafting vs those who have not. *Can J Cardiol* 2018;34:310–8. <https://doi.org/10.1016/j.cjca.2017.12.016>; PMID: 29395703.
28. Dautov R, Nguyen CM, Altisent O, et al. Recanalization of chronic total occlusions in patients with previous coronary bypass surgery and consideration of retrograde access via saphenous vein grafts. *Circ Cardiovasc Interv* 2018;9:e003515. <https://doi.org/10.1161/circinterventions.115.003515>; PMID: 27418611.
29. Brener SJ, Lytle BW, Casserly IP, et al. Predictors of revascularization method and long-term outcome of percutaneous coronary intervention or repeat coronary bypass surgery in patients with multivessel coronary disease and previous coronary bypass surgery. *Eur Heart J* 2006;27:413–8. <https://doi.org/10.1093/eurheartj/ehi646>; PMID: 16272211.
30. Maltais S, Widmer RJ, Bell MR, et al. Reoperation for coronary artery bypass grafting surgery: outcomes and considerations for expanding interventional procedures. *Ann Thorac Surg* 2017;103:1886–92. <https://doi.org/10.1016/j.athoracsur.2016.09.097>; PMID: 28012643.
31. Boden WE, O'Rourke RA, Teo KK, et al. Optimal medical therapy with or without PCI for stable coronary disease. *N Engl J Med* 2007;356:1503–16. <https://doi.org/10.1056/nejmoa070829>; PMID: 17387127.
32. Al-Lamee R, Thompson D, Dehbi HM, et al. Percutaneous coronary intervention in stable angina (ORBITA): a double-blind, randomised controlled trial. *Lancet*. 2018;391:31–40. [https://doi.org/10.1016/s0140-6736\(17\)32714-9](https://doi.org/10.1016/s0140-6736(17)32714-9); PMID: 29103656.
33. Rodés-Cabau J, Bertrand OF, Larose E, et al. Comparison of plaque sealing with paclitaxel-eluting stents versus medical therapy for the treatment of moderate nonsignificant saphenous vein graft lesions. *Circulation* 2009;120:1978–86. <https://doi.org/10.1161/circulationaha.109.874057>; PMID: 19884468.
34. Rodés-Cabau J, Jolly SS, Cairns J, et al. Sealing intermediate nonobstructive coronary saphenous vein graft lesions with drug-eluting stents as a new approach to reducing cardiac events. *Circ Cardiovasc Interv* 2018;9:e004336. <https://doi.org/10.1161/circinterventions.116.004336>; PMID: 27815344.
35. Maes F, Jolly SS, Cairns J, et al. Plaque sealing with drug-eluting stents versus medical therapy for treating intermediate non-obstructive saphenous vein graft lesions: a pooled analysis of the VELET I and VELET II trials. *J Invasive Cardiol* 2019;31:e308–15. [https://doi.org/10.1016/s0735-1097\(10\)61810-4](https://doi.org/10.1016/s0735-1097(10)61810-4); PMID: 31671060.
36. Foster ED, Fisher LD, Kaiser GC, et al. Comparison of operative mortality and morbidity for initial and repeat coronary artery bypass grafting: the Coronary Artery Surgery Study (CASS) registry experience. *Ann Thorac Surg* 1984;38:563–70. [https://doi.org/10.1016/s0003-4975\(10\)62312-0](https://doi.org/10.1016/s0003-4975(10)62312-0); PMID: 6391399.
37. Lytle BW, Loop FD, Cosgrove DM, et al. Fifteen hundred coronary reoperations: results and determinants of early and late survival. *J Thorac Cardiovasc Surg* 1987;93:847–59. [https://doi.org/10.1016/s0022-5223\(19\)37045-x](https://doi.org/10.1016/s0022-5223(19)37045-x); PMID: 3494885.
38. Loop FD. A 20-year experience in coronary artery reoperation. *Eur Heart J* 1989;10(Suppl H):78–84. https://doi.org/10.1093/eurheartj/10.suppl_h.78; PMID: 2627968.
39. Morrison DA, Sethi G, Sacks J, et al. Percutaneous coronary intervention versus repeat bypass surgery for patients with medically refractory myocardial ischemia: AWESOME randomized trial and registry experience with post-CABG patients. *J Am Coll Cardiol* 2002;40:1951–4. [https://doi.org/10.1016/s0735-1097\(02\)02560-3](https://doi.org/10.1016/s0735-1097(02)02560-3); PMID: 12475454.
40. Brodie BR, Wilson H, Stuckey T, et al. Outcomes with drug-eluting versus bare-metal stents in saphenous vein graft intervention: results from the STENT (Strategic Transcatheter Evaluation of New Therapies) Group. *JACC Cardiovasc Interv* 2009;2:1105–12. <https://doi.org/10.1016/j.jcin.2009.08.020>; PMID: 19926052.
41. Brilakis ES, Wang TY, Rao SV, et al. Frequency and predictors of drug-eluting stent use in saphenous vein bypass graft percutaneous coronary interventions: a report from the American College of Cardiology National Cardiovascular Data CathPCI Registry. *JACC Cardiovasc Interv* 2010;3:1068–73. <https://doi.org/10.1016/j.jcin.2010.07.009>; PMID: 20965466.
42. Lee MS, Park SJ, Kandzari DE, et al. Saphenous vein graft intervention. *JACC Cardiovasc Interv* 2011;4:831–43. <https://doi.org/10.1016/j.jcin.2011.05.014>; PMID: 21851895.
43. Brilakis ES, Rao SV, Banerjee S, et al. Percutaneous coronary intervention in native arteries versus bypass grafts in prior coronary artery bypass grafting patients: a report from the National Cardiovascular Data Registry. *JACC Cardiovasc Interv* 2011;4:844–50. <https://doi.org/10.1016/j.jcin.2011.03.018>; PMID: 21851896.
44. Abdel-Karim ARR, Banerjee S, Brilakis ES. Percutaneous intervention of acutely occluded saphenous vein grafts: contemporary techniques and outcomes. *J Invasive Cardiol* 2010;22:253–7. [https://doi.org/10.1016/s0735-1097\(10\)61765-2](https://doi.org/10.1016/s0735-1097(10)61765-2); PMID: 20516502.
45. PeRcutaneous cOronary Intervention of Native Coronary Artery Versus Venous Bypass Graft in Patients With Prior CABG (PROCTOR). 2019. <https://clinicaltrials.gov/ct2/show/NCT03805048> (accessed 28 February 2020).
46. Nordgaard H, Vitale N, Haaverstad R. Transit-time blood flow measurements in sequential saphenous coronary artery bypass grafts. *Ann Thorac Surg* 2009;87:1409–15. <https://doi.org/10.1016/j.athoracsur.2009.02.018>; PMID: 19379875.
47. O'Connor GT, Malenka DJ, Quinton H, et al. Multivariate prediction of in-hospital mortality after percutaneous coronary interventions in 1994–6. *J Am Coll Cardiol* 1999;34:681–91. [https://doi.org/10.1016/s0735-1097\(99\)00267-3](https://doi.org/10.1016/s0735-1097(99)00267-3); PMID: 10483948.
48. Sdringola S, Assali AR, Ghani M, et al. Risk assessment of slow or no-reflow phenomenon in aortocoronary vein graft percutaneous intervention. *Catheter Cardiovasc Interv* 2001;54:318–24. <https://doi.org/10.1002/ccd.1290>; PMID: 11747155.
49. Abbo KM, Dooris M, Glazier S, et al. Features and outcome of no-reflow after percutaneous coronary intervention. *Am J Cardiol* 1995;75:778–82. [https://doi.org/10.1016/s0002-9149\(99\)80410-x](https://doi.org/10.1016/s0002-9149(99)80410-x); PMID: 17717278.
50. Carter LI, Golzar JA, Cavendish JJ, et al. Embolic protection of saphenous vein graft percutaneous interventions. *J Interv Cardiol* 2007;20:351–8. <https://doi.org/10.1111/j.1540-8183.2007.00284.x>; PMID: 17880331.
51. Baim DS, Wahr D, George B, et al. Randomized trial of a distal embolic protection device during percutaneous intervention of saphenous vein aorto-coronary bypass grafts. *Circulation* 2002;105:1285–90. <https://doi.org/10.1161/01.cir.0000012783.63093.0c>; PMID: 11901037.
52. Vermeersch P, Agostoni P, Verheyde S, et al. Randomized double-blind comparison of sirolimus-eluting stent versus bare-metal stent implantation in diseased saphenous vein grafts: six-month angiographic, intravascular ultrasound, and clinical follow-up of the RRISC trial. *J Am Coll Cardiol* 2006;48:2423–31. <https://doi.org/10.1016/j.jacc.2006.09.021>; PMID: 17174178.
53. Brilakis ES, Lichtenwalter C, Lemos JA de, et al. A randomized controlled trial of a paclitaxel-eluting stent versus a similar bare-metal stent in saphenous vein graft

- lesions: the SOS (Stenting Of Saphenous Vein Grafts) trial. *J Am Coll Cardiol* 2009;53:919–28. <https://doi.org/10.1016/j.jacc.2008.11.029>; PMID: 19281920.
54. Shoab A, Kinnaird T, Curzen N, et al. Outcomes following percutaneous coronary intervention in saphenous vein grafts with and without embolic protection devices. *JACC Cardiovasc Interv* 2019;12:2286–95. <https://doi.org/10.1016/j.jcin.2019.08.037>; PMID: 31753300.
 55. Pucelikova T, Mehran R, Kirtane AJ, et al. Short- and long-term outcomes after stent-assisted percutaneous treatment of saphenous vein grafts in the drug-eluting stent era. *Am J Cardiol* 2008;101:63–8. <https://doi.org/10.1016/j.amjcard.2007.07.048>; PMID: 18157967.
 56. Mehta SK, Frutkin AD, Milford-Beland S, et al. Utilization of distal embolic protection in saphenous vein graft interventions (an analysis of 19,546 patients in the American College of Cardiology–National Cardiovascular Data Registry). *Am J Cardiol* 2007;100:1114–8. <https://doi.org/10.1016/j.amjcard.2007.04.058>; PMID: 17884373.
 57. Brennan JM, Al-Hejjily W, Dai D, et al. Three-year outcomes associated with embolic protection in saphenous vein graft intervention. *Circ Cardiovasc Interv* 2015;8:e001403; <https://doi.org/10.1161/circinterventions.114.001403>; PMID: 25714391.
 58. Paul TK, Bhatheja S, Panchal HB, et al. Outcomes of saphenous vein graft intervention with and without embolic protection device. *Circ Cardiovasc Interv* 2017;10:e005538. <https://doi.org/10.1161/circinterventions.117.005538>; PMID: 29246912.
 59. Ahmed JM, Dangas G, Lansky AJ, et al. Influence of gender on early and one-year clinical outcomes after saphenous vein graft stenting. *Am J Cardiol* 2001;87:401–5. [https://doi.org/10.1016/s0002-9149\(00\)01391-6](https://doi.org/10.1016/s0002-9149(00)01391-6); PMID: 11179522.
 60. Kirtane AJ, Heyman ER, Metzger C, et al. Correlates of adverse events during saphenous vein graft intervention with distal embolic protection: a PRIDE substudy. *JACC Cardiovasc Interv* 2008;1:186–91. <https://doi.org/10.1016/j.jcin.2008.01.002>; PMID: 19463299.
 61. Hong MK, Mehran R, Dangas G, et al. Creatine kinase-MB enzyme elevation following successful saphenous vein graft intervention is associated with late mortality. *Circulation* 1999;100:2400–5. <https://doi.org/10.1161/01.cir.100.24.2400>; PMID: 10595951.
 62. Latif F, Uyeda L, Edson R, et al. Stent-only versus adjunctive balloon angioplasty approach for saphenous vein graft percutaneous coronary intervention: insights from DIVA trial. *Circ Cardiovasc Interv* 2020;13:e008494. <https://doi.org/10.1161/circinterventions.119.008494>; PMID: 32019343.
 63. Savage MP, Douglas JS, Fischman DL, et al. Stent placement compared with balloon angioplasty for obstructed coronary bypass grafts. *N Engl J Med* 1997;337:740–7. <https://doi.org/10.1056/nejm199709113371103>; PMID: 9287229.
 64. Stankovic G, Colombo A, Presbitero P, et al. Randomized evaluation of polytetrafluoroethylene-covered stent in saphenous vein grafts. *Circulation* 2003;108:37–42. <https://doi.org/10.1161/01.cir.0000079106.71097.1c>; PMID: 12821546.
 65. Turco MA, Buchbinder M, Popma JJ, et al. Pivotal, randomized U.S. study of the Symbiot™ covered stent system in patients with saphenous vein graft disease: eight-month angiographic and clinical results from the Symbiot III trial. *Catheter Cardiovasc Interv* 2006;68:379–88. <https://doi.org/10.1002/ccd.20873>; PMID: 16892434.
 66. Stone GW, Goldberg S, O’Shaughnessy C, et al. 5-year follow-up of polytetrafluoroethylene-covered stents compared with bare-metal stents in aortocoronary saphenous vein grafts: the randomized BARRICADE (Barrier Approach to Restenosis; Restrict Intima to Curtail Adverse Events) trial. *JACC Cardiovasc Interv* 2011;4:300–9. <https://doi.org/10.1016/j.jcin.2010.11.013>; PMID: 21435608.
 67. Mehilli J, Pache J, Abdel-Wahab M, et al. Drug-eluting versus bare-metal stents in saphenous vein graft lesions (ISAR-CABG): a randomised controlled superiority trial. *Lancet*. 2011;378:1071–8. [https://doi.org/10.1016/s0140-6736\(11\)61255-5](https://doi.org/10.1016/s0140-6736(11)61255-5); PMID: 21872918.
 68. Fahrni G, Farah A, Engstrom T, et al. Long-term results after drug-eluting versus bare-metal stent implantation in saphenous vein grafts: randomized controlled trial. *J Am Heart Assoc* 2020;9:e017434. <https://doi.org/10.1161/jaha.120.017434>; PMID: 33032485.
 69. Patel NJ, Bavishi C, Atti V, et al. Drug-eluting stents versus bare-metal stents in saphenous vein graft intervention. *Circ Cardiovasc Interv* 2018;11:e007045. <https://doi.org/10.1161/circinterventions.118.007045>; PMID: 30571204.
 70. Abdelrahman A, Debski M, More R, et al. One-year outcomes of percutaneous coronary intervention in native coronary arteries versus saphenous vein grafts in patients with prior coronary artery bypass graft surgery. *Cardiol J* 2020. <https://doi.org/10.5603/cj.a2020.0131>; PMID: 33001421; epub ahead of press.
 71. Cassese S, Byrne RA, Tada T, et al. Incidence and predictors of restenosis after coronary stenting in 10 004 patients with surveillance angiography. *Heart* 2014;100:153–9. <https://doi.org/10.1136/heartjnl-2013-304933>; PMID: 24270744.
 72. Tada T, Byrne RA, Simunovic I, et al. Risk of stent thrombosis among bare-metal stents, first-generation drug-eluting stents, and second-generation drug-eluting stents results from a registry of 18,334 patients. *JACC Cardiovasc Interv* 2013;6:1267–74. <https://doi.org/10.1016/j.jcin.2013.06.015>; PMID: 24355117.
 73. Lee SW, Lee PH, Ahn JM, et al. Randomized trial evaluating percutaneous coronary intervention for the treatment of chronic total occlusion. *Circulation* 2019;139:1674–83. <https://doi.org/10.1161/circulationaha.118.031313>; PMID: 30813758.
 74. Brilakis ES, Mashayekhi K, Burke MN. How DECISION-CTO can help guide the decision to perform chronic total occlusion percutaneous coronary intervention. *Circulation* 2019;139:1684–7. <https://doi.org/10.1161/circulationaha.119.039835>; PMID: 30933615.
 75. Burke AP, Weber DK, Kolodgy FD, et al. Pathophysiology of calcium deposition in coronary arteries. *Herz* 2001;26:239–44. <https://doi.org/10.1007/pl000202026>; PMID: 11479935.
 76. Sakakura K, Nakano M, Otsuka F, et al. Comparison of pathology of chronic total occlusion with and without coronary artery bypass graft. *Eur Heart J* 2014;35:1683–93. <https://doi.org/10.1093/eurheartj/ehz422>; PMID: 24126875.
 77. Morino Y, Abe M, Morimoto T, et al. Predicting successful guidewire crossing through chronic total occlusion of native coronary lesions within 30 minutes: the J-CTO (Multicenter CTO Registry in Japan) score as a difficulty grading and time assessment tool. *JACC Cardiovasc Interv* 2011;4:213–21. <https://doi.org/10.1016/j.jcin.2010.09.024>; PMID: 21349461.
 78. Wilson WM, Walsh SJ, Yan AT, et al. Hybrid approach improves success of chronic total occlusion angioplasty. *Heart* 2016;102:1486–93. <https://doi.org/10.1136/heartjnl-2015-308891>; PMID: 27164918.
 79. Spratt JC, Hanratty CG, Walsh SJ, Wilson SJ. *A Guide to Mastering Antegrade CTO PCI Part 1*. Optima. 2019. <https://books.apple.com/gb/book/a-guide-to-mastering-antegrade-cto-pci-part-1/id1474956817> (accessed 9 August 2021).
 80. Spratt JC, Hanratty CG, Walsh SJ, Wilson SJ. *A Guide to Mastering Antegrade CTO PCI Part 2*. Optima. 2019. <https://books.apple.com/gb/book/a-guide-to-mastering-antegrade-cto-pci-part-2/id1474965405> (accessed 9 August 2021).
 81. Spratt JC. *A Guide to Mastering Retrograde CTO PCI*. Optima. 2015. <https://books.apple.com/fr/book/a-guide-to-mastering-retrograde-cto-pci/id970542167> (accessed 9 August 2021).
 82. Maurer BJ, Oberman A, Holt JH Jr, et al. Changes in grafted and nongrafted coronary arteries following saphenous vein bypass grafting. *Circulation* 1974;50:293–300. <https://doi.org/10.1161/01.cir.50.2.293>; PMID: 4546527.
 83. Hwang MH, Meadows WR, Palac RT, et al. Progression of native coronary artery disease at 10 years: insights from a randomized study of medical versus surgical therapy for angina. *J Am Coll Cardiol* 1990;16:1066–70. [https://doi.org/10.1016/0735-1097\(90\)90533-u](https://doi.org/10.1016/0735-1097(90)90533-u); PMID: 2229749.
 84. Alderman EL, Corley SD, Fisher LD, et al. Five-year angiographic follow-up of factors associated with progression of coronary artery disease in the Coronary Artery Surgery Study (CASS). *J Am Coll Cardiol* 1993;22:1141–54. [https://doi.org/10.1016/0735-1097\(93\)90429-5](https://doi.org/10.1016/0735-1097(93)90429-5); PMID: 8409054.
 85. Chiu J-J, Chien S. Effects of disturbed flow on vascular endothelium: pathophysiological basis and clinical perspectives. *Physiol Rev* 2011;91:327–87. <https://doi.org/10.1152/physrev.00047.2009>; PMID: 21248169.
 86. Galassi AR, Tomasello SD, Crea F, et al. Transient impairment of vasomotion function after successful chronic total occlusion recanalization. *J Am Coll Cardiol* 2012;59:711–8. <https://doi.org/10.1016/j.jacc.2011.10.894>; PMID: 22340262.
 87. Sabik JF, Blackstone EH, Houghtaling PL, et al. Is reoperation still a risk factor in coronary artery bypass surgery? *Ann Thorac Surg* 2005;80:1719–27. <https://doi.org/10.1016/j.athoracsur.2005.04.033>; PMID: 16242445.
 88. Yap C-H, Sposato L, Akowuah E, et al. Contemporary results show repeat coronary artery bypass grafting remains a risk factor for operative mortality. *Ann Thorac Surg* 2009;87:1386–91. <https://doi.org/10.1016/j.athoracsur.2009.02.006>; PMID: 19379870.
 89. Morrison DA, Sethi G, Sacks J, et al. Percutaneous coronary intervention versus coronary artery bypass graft surgery for patients with medically refractory myocardial ischemia and risk factors for adverse outcomes with bypass: a multicenter, randomized trial. *J Am Coll Cardiol* 2001;38:143–9. [https://doi.org/10.1016/s0735-1097\(01\)01366-3](https://doi.org/10.1016/s0735-1097(01)01366-3); PMID: 11451264.
 90. Harskamp RE, Beijik MA, Damman P, et al. Clinical outcome after surgical or percutaneous revascularization in coronary bypass graft failure. *J Cardiovasc Med* 2013;14:438–45. <https://doi.org/10.2459/jcm.0b1013e328356a4fc>; PMID: 22828774.
 91. Kinnaird T, Anderson R, Ossei-Gerning N, et al. Coronary perforation complicating percutaneous coronary intervention in patients with a history of coronary artery bypass surgery. *Circ Cardiovasc Interv* 2017;10:e005581. <https://doi.org/10.1161/circinterventions.117.005581>; PMID: 28916604.
 92. Vetrugno V, Sharma H, Townend JN, Khan SQ. What is the cause of hypotension? A rare complication of percutaneous coronary intervention of a chronic total occlusion: a case report. *Eur Heart J Case Rep* 2019;3:1–5. <https://doi.org/10.1093/ehjcr/ytz184>; PMID: 32123803.
 93. Vecovo GM, Zvelonghi C, Scott B, Agostoni P. Percutaneous coronary intervention for chronic total occlusion. *US Cardiol* 2020;14:e11. <https://doi.org/10.15420/usc.2020.10>.
 94. Maeremans J, Spratt JC, Knaepen P, et al. Towards a contemporary, comprehensive scoring system for determining technical outcomes of hybrid percutaneous chronic total occlusion treatment: The RECHARGE score. *Catheter Cardiovasc Interv* 2018;91:192–202. <https://doi.org/10.1002/ccd.27092>; PMID: 28471074.
 95. Guelker JE, Bansemir L, Ott R, et al. Validity of the J-CTO score and the CL-score for predicting successful CTO recanalization. *Int J Cardiol* 2017;230:228–31. <https://doi.org/10.1016/j.ijcard.2016.12.165>; PMID: 28041697.
 96. Potter BJ, Matteau A, Noisueux N, Mansour S. High stakes: CTO-PCI in the post-CABG patient. *Can J Cardiol* 2018;34:238–40. <https://doi.org/10.1016/j.cjca.2017.12.022>.
 97. Maeremans J, Walsh S, Knaepen P, et al. The hybrid algorithm for treating chronic total occlusions in Europe: the RECHARGE registry. *J Am Coll Cardiol* 2016;68:1958–70. <https://doi.org/10.1016/j.jacc.2016.08.034>; PMID: 27788851.
 98. Wilson WM, Walsh SJ, Bagnall A, et al. One-year outcomes after successful chronic total occlusion percutaneous coronary intervention: the impact of dissection re-entry techniques. *Catheter Cardiovasc Interv* 2017;90:703–12. <https://doi.org/10.1002/ccd.26980>; PMID: 28296045.
 99. Kahn JK, Hartzler GO. Retrograde coronary angioplasty of isolated arterial segments through saphenous vein bypass grafts. *Cathet Cardiovasc Diagn* 1990;20:88–93. <https://doi.org/10.1002/ccd.1810200205>; PMID: 2354520.
 100. Kaltenbach M, Hartmann A, Vallbracht C. Procedural results and patient selection in recanalization of chronic coronary occlusions by low speed rotational angioplasty. *Eur Heart J* 1993;14:826–30. <https://doi.org/10.1093/eurheartj/14.6.826>; PMID: 8325312.
 101. Prospective Global Registry for the Study of Chronic Total Occlusion Intervention (PROGRESS-CTO). 2014. <https://clinicaltrials.gov/ct2/show/NCT02061436> (accessed 27 February 2021).
 102. Galassi AR, Werner GS, Boukhris M, et al. Percutaneous recanalization of chronic total occlusions: 2019 consensus document from the EuroCTO Club. *EuroIntervention* 2019;15:198–208. <https://doi.org/10.4244/eij-d-18-00826>; PMID: 30636678.
 103. Brilakis ES, Grantham JA, Rinfret S, et al. A percutaneous treatment algorithm for crossing coronary chronic total occlusions. *JACC Cardiovasc Interv* 2012;5:367–79. <https://doi.org/10.1016/j.jcin.2012.02.006>; PMID: 22516392.
 104. Li X, Liu Y, Gao J, et al. Comparison of graft vessel versus native vessel strategies for late saphenous vein graft disease after coronary artery bypass grafting. *Zhonghua Xin Xue Guan Bing Za Zhi* 2020;48:367–72 [in Chinese]. <https://doi.org/10.3760/cma.j.cn112148-0190827-00523>; PMID: 32450652.
 105. Lembo NJ, Karpapaliotis D, Kandzari DE. CTO PCI procedural planning. *Interv Cardiol Clin* 2012;1:299–308. <https://doi.org/10.1016/j.iccl.2012.04.002>; PMID: 28582014.
 106. Bluemke DA, Achenbach S, Budoff M, et al. Noninvasive coronary artery imaging. *Circulation* 2008;118:586–606. <https://doi.org/10.1161/circulationaha.108.189695>; PMID: 18586979.
 107. Opolski MP, Achenbach S, Schubbäck A, et al. Coronary computed tomographic prediction rule for time-efficient guidewire crossing through chronic total occlusion insights from the CT-RECTOR Multicenter Registry (Computed Tomography Registry of Chronic Total Occlusion Revascularization). *JACC Cardiovasc Interv* 2015;8:257–67. <https://doi.org/10.1016/j.jcin.2014.07.031>; PMID: 25700748.
 108. Tan Y, Zhou J, Zhang W, et al. Comparison of CT-RECTOR and J-CTO scores to predict chronic total occlusion difficulty for percutaneous coronary intervention. *Int J Cardiol* 2017;235:169–75. <https://doi.org/10.1016/j.ijcard.2017.02.008>; PMID: 28274578.
 109. Malagutti P, Nieman K, Meijboom WB, et al. Use of 64-slice CT in symptomatic patients after coronary bypass surgery: evaluation of grafts and coronary arteries. *Eur Heart J* 2007;28:1879–85. <https://doi.org/10.1093/eurheartj/ehf155>; PMID: 16847009.

110. Katsuragawa M, Fujiwara H, Miyamae M, et al. Histologic studies in percutaneous transluminal coronary angioplasty for chronic total occlusion: comparison of tapering and abrupt types of occlusion and short and long occluded segments. *J Am Coll Cardiol* 1993;21:604–11. [https://doi.org/10.1016/0735-1097\(93\)90091-e](https://doi.org/10.1016/0735-1097(93)90091-e); PMID: 8436741.
111. Asahi Intecc. <http://www.asahi-intecc.co.jp/en/medical/pci/gaia.html> (accessed 30 April 2021).
112. Sapontis J, Marso SP, Lombardi WL, et al. How to fix common problems encountered in CTO PCI: the expanded hybrid approach. In: Rinfret S, ed. *Percutaneous Intervention for Coronary Chronic Total Occlusion*. Cham: Springer, 2016;141–59. https://doi.org/10.1007/978-3-319-21563-1_11.
113. Colombo A, Mikhail GW, Michev I, et al. Treating chronic total occlusions using subintimal tracking and reentry: the STAR technique. *Catheter Cardiovasc Interv* 2005;64:407–11. <https://doi.org/10.1002/ccd.20307>; PMID: 15789384.
114. Lombardi WL. Retrograde PCI: what will they think of next? *J Invasive Cardiol* 2009;21:543; PMID: 19805844.
115. Valenti R, Vergara R, Migliorini A, et al. Predictors of reocclusion after successful drug-eluting stent-supported percutaneous coronary intervention of chronic total occlusion. *J Am Coll Cardiol* 2013;61:545–50. <https://doi.org/10.1016/j.jacc.2012.10.036>; PMID: 23273395.
116. Azzalini L, Poletti E, Ayoub M, et al. Coronary artery perforation during chronic total occlusion percutaneous coronary intervention: epidemiology, mechanisms, management, and outcomes. *EuroIntervention* 2019;15:e804–11. <https://doi.org/10.4244/eij-d-19-00282>; PMID: 31217142.
117. Matsuno S, Tsuchikane E, Harding SA, et al. Overview and proposed terminology for the reverse controlled antegrade and retrograde tracking (reverse CART) techniques. *EuroIntervention* 2018;14:94–101. <https://doi.org/10.4244/eij-d-17-00867>; PMID: 29360064.
118. Brinkmann C, Eitan A, Schwencke C, et al. Rotational atherectomy in CTO lesions: too risky? Outcome of rotational atherectomy in CTO lesions compared to non-CTO lesions. *EuroIntervention* 2018;14:e1192–8. <https://doi.org/10.4244/eij-d-18-00393>; PMID: 30175961.
119. Yeoh J, Hill J, Spratt JC. Intravascular lithotripsy assisted chronic total occlusion revascularization with reverse controlled antegrade retrograde tracking. *Catheter Cardiovasc Interv* 2019;93:1295–7. <https://doi.org/10.1002/ccd.28165>; PMID: 30838746.
120. del Olmo VV, Rodríguez-Leor O, Redondo A, et al. Intracoronary lithotripsy in a high-risk real-world population. First experience in severely calcified, complex coronary lesions. *REC Interv Cardiol* 2020;2:76–81. <https://doi.org/10.24875/RECICE.M19000083>.
121. Wilson SJ, Hanratty CG, Spence MS, et al. Saphenous vein graft sacrifice following native vessel PCI is safe and associated with favourable longer-term outcomes. *Cardiovasc Revasc Med* 2019;20:1048–52. <https://doi.org/10.1016/j.carrev.2019.01.025>; PMID: 30745059.