Recovery of absolute coronary blood flow and microvascular resistance after chronic total occlusion percutaneous coronary intervention

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Recovery of Absolute Coronary Blood Flow and Microvascular Resistance After Chronic Total Occlusion Percutaneous Coronary Intervention: An Exploratory Study

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BACKGROUND: This study aimed to investigate longitudinal physiological changes in the recanalized coronary chronic total occlusion (CTO) vessel and its dependent myocardium after successful percutaneous coronary intervention (PCI).

METHODS AND RESULTS: In this pilot study, 25 patients scheduled for elective CTO PCI with viable myocardium and angiographically visible collaterals were included. Absolute coronary blood flow and absolute microvascular resistance were measured invasively using continuous thermodilution. Measurements were performed immediately after successful CTO PCI and at short-term follow-up. In a subgroup of patients, physiological measurements were performed at the predominant donor vessel before CTO PCI, immediately afterwards, and at follow-up. Absolute coronary blood flow in the recanalized CTO artery increased from 148±53 mL/min immediately after PCI to 221±77 mL/min at follow-up (P<0.001). In agreement, absolute resistance in the myocardial territory perfused by the CTO artery, decreased from 545±255 Wood units immediately after the procedure to 387±128 Wood units at follow-up (P=0.014). There were no significant changes in the absolute coronary blood flow and resistance in the predominant donor between baseline and follow-up. Positive remodeling of the distal CTO vessel with an increase in lumen diameter was observed.

CONCLUSIONS: After successful CTO PCI, blood flow in the recanalized artery and microvascular function of the dependent myocardium are not immediately normal but recover over time.

Key Words: chronic total coronary occlusion ■ coronary flow ■ coronary microvascular function ■ coronary microvascular resistance ■ coronary revascularization

Successful percutaneous coronary intervention (PCI) of chronic total occlusions (CTOs) relieves ischemia, improving patients’ symptoms, functional status, and quality of life. It has been recognized that after successful recanalization of a CTO, there is no immediate complete recovery of the distal artery and the dependent myocardial territory, but its function might improve over time. In most CTOs with viable distal myocardium, collateral blood supply is present and sufficient to maintain resting metabolism (hibernation) or even contraction at rest. However, evident ischemia occurs with increasing demand during stress/exercise. Due to reduced blood flow and perfusion pressure, negative remodeling of the distal part of the occluded vessel occurs, while microvascular changes become present in the distal myocardium, with an increase in microvascular resistance in the CTO territory. Both phenomena
Keulards et al RECOVER CTO contribute to incomplete, delayed, and suboptimal recovery of physiology even after technically successful PCI.

A recent study showed that fractional flow reserve (FFR) of both the coronary artery and the myocardium of a recanalized CTO, increased over time, paralleled by regression of collaterals. These observations strongly suggest that recovery of both the distal coronary artery and the dependent myocardium often occurs because an increase in FFR (without additional intervention in the epicardial artery) implicates a decrease of distal epicardial resistance. Although this study provides indirect evidence of recovery of microvascular function over time along with the positive remodeling of the distal CTO artery, there were no direct measurements of microvascular function or coronary blood flow.

Recently, a novel methodology has become available that allows accurate and quantitative invasive measurements of absolute coronary flow and microvascular resistance during coronary catheterization. The aim of the present study was to perform such serial measurements of absolute coronary flow and microvascular resistance in the recanalized CTO and major donor vessels after successful PCI. Based upon the observations of the present study in combination with the previous work by Karamasis et al., we aimed to propose a complete model explaining the physiologic changes after successful CTO PCI.

## METHODS

### Study Design and Population

This was a prospective observational exploratory study on coronary physiology after CTO percutaneous revascularization. Patients with a clinical indication for elective CTO PCI were recruited in the Catharina Hospital (Eindhoven, The Netherlands) and the Essex Cardiothoracic Centre (Basildon, UK). All patients had stable angina and viability of the myocardial territory supplied by the CTO vessel as evidenced by perfusion cardiac magnetic resonance imaging or preserved wall motion on transthoracic echocardiography at rest. As cardiac magnetic resonance criteria for ischemia and viability, >10% ischemic burden in the CTO territory was the cutoff that we used to justify CTO PCI. Delayed myocardial enhancement was the cardiac magnetic resonance sequence used for viability and >75% with delayed enhancement in the myocardial segment under interrogation was the cutoff for a territory to be considered not viable. In all cases, there were spontaneously visible collateral vessels supplied by a contralateral donor artery. All patients provided written informed consent, and the study was approved by the local ethics committee in Eindhoven (The Netherlands) and Basildon (United Kingdom). The data that support the findings of this study are available from the corresponding author upon reasonable request.

### Chronic Total Occlusion PCI

CTO PCI was performed according to contemporary interventional techniques following the hybrid algorithm. Recanalization strategy was left at the discretion of the CTO operator. A procedure was considered successful if thrombolysis in myocardial infarction flow grade 3 with <30% angiographic residual stenosis in the CTO vessel was achieved. Patients were discharged on statins and on dual antiplatelet therapy (aspirin and clopidogrel) for at least 1 year.

### Measurement of Absolute Blood Flow and Microvascular Resistance

After successful PCI of the CTO, complete physiological assessment in the recanalized vessel and its dependent myocardium was performed by calculating...
absolute coronary blood flow and microvascular resistance using continuous thermodilution. The detailed method was previously published. In brief, after intracoronary administration of nitroglycerin, a guidewire equipped with a pressure and temperature sensor (Pressure Wire X™, Abbott Vascular, St Paul, MN) is equalized at the tip of the guide catheter and positioned in the distal vessel. Thereafter, a multifunctional monorail infusion catheter (RayFlow, Hexacath, Paris, France) is advanced over the wire and placed in the proximal coronary artery. The catheter is connected to an infusion pump and saline infusion at a prespecified flow rate (Qi, usually 20 mL/min) is equalized at the tip of the guide catheter and placed in the proximal coronary artery. The catheter is advanced to place the tip against the vessel wall of the selected guide catheter. The saline infusion itself creates a state of maximum hyperemia within seconds and when a steady hyperemic state is achieved the guidewire is pulled back to the tip of the catheter and the temperature of the infused saline (T) is measured. Absolute coronary flow during steady state maximum hyperemia (Q in mL/min) is calculated by assessment of the changes in distal coronary temperature compared with the infusion temperature of the saline (T) by the equation 

\[ Q = \frac{1.08 \times (T_i - T)}{Qi} \]

The microvascular resistance (R in dyn.s.cm⁻⁵, mm Hg/L per min, or Wood units [WU]) is calculated by dividing the distal coronary pressure (Pd) and the calculated blood flow (Q). Therefore, R is independent of the resistance of the epicardial coronary artery, which can be assessed separately by (Pa-Pd)/Q. Live wireless recording and analysis of coronary pressure traces and temperature and automated calculation of Q and R is performed using a dedicated software system (Coroventis, Uppsala, Sweden).

**Physiological Measurements in the Donor Vessel**

In a subgroup of patients (n=10) with intermediate coronary disease (30%–70% stenosis) in the donor artery, physiological assessment including calculation of absolute flow and resistance as described above was performed in the predominant donor vessel. The measurements were performed at the onset of the PCI procedure (before attempting the CTO vessel) and immediately after successful PCI of the CTO vessel.

**Follow-Up Procedure**

Patients with complete physiological measurements in the recanalized vessel returned for a follow-up procedure within 2 months (7–70 days, mean 46 days) according to institutional practice. During the follow-up procedure, absolute flow and resistance were measured once more in the recanalized CTO vessel following exactly the same method. In cases where measurements were performed in the predominant donor artery during the baseline procedure, these were also repeated. To evaluate recovery of coronary flow and microvasculature function of the CTO artery, we compared the measurements from the index procedure with those at follow-up. Figure 1 illustrates such complete sequence of angiographic and physiologic assessment in 1 patient.

**Quantitative Angiographic Assessment of the Distal CTO Artery**

Collateral circulation to the chronically occluded vessel was graded using collateral connections grading method and the predominant donor vessel was determined. Procedure complexity was graded using the Japanese Multicenter CTO Registry score. After blinded for clinical and procedural data, mean and minimal lumen diameter of the distal coronary artery were assessed by quantitative coronary angiography after the baseline procedure and at follow-up. Quantitative coronary analysis was performed using Philips Intellispace Cardiovascular Software System (Amsterdam, The Netherlands).

**Statistical Analysis**

Data were checked for normality of distribution using Kolmogorov–Smirnov testing. Categorical variables are presented as counts and percentages. Continuous variables are presented as mean±SD or median and interquartile range as appropriate. Comparisons of the relative change in absolute blood flow or resistance were made using a paired t-test. Spearman correlation coefficients were used to assess the relationship between 2 variables as appropriate. ANOVA repeated measures test was performed to assess the difference between 3 following measurements in the donor artery. A 2-tailed level of significance of 0.05 was used. All preceding analyses were performed using SPSS 25 software (SPSS Inc, Chicago, IL).

**RESULTS**

Twenty-five patients with successful CTO PCI and complete physiology measurements in the recanalized vessel at baseline and follow-up were included. Baseline and procedural characteristics are listed in Table 1. The mean age was 66±8.7 years, and 19 of 25 patients were male. The most common CTO vessel was the right coronary artery and the mean Japanese–Chronic Total Occlusion score was 1.64±1.25. Antegrade wire escalation was the
The most commonly successful recanalization technique (56%), followed by retrograde dissection/reentry (20%) (Table 2). The follow-up procedure was performed within 2 months (7–70 days, mean 46 days). A subgroup of 10 patients had physiology measurements of the predominant donor vessel at baseline and follow-up.

Figure 1. Typical case example.
Complete assessment of 1 patient at baseline and at follow-up. A and D, The CTO artery (right coronary artery) and the donor artery (left anterior descending) before PCI. The absolute flow in the donor vessel is 256 mL/min, with a resistance of 231 WU. B, E, and G, Flow and resistance measured directly after PCI. C, F and H, At follow-up the CTO vessel showed an increase in flow from 161 mL/min to 300 mL/min, and resistance of the dependent myocardium decreased from 370 to 243 WU. Flow in the donor artery decreased further. CTO indicates chronic total occlusion; FFR, fractional flow reserve; PCI, percutaneous coronary intervention; Q, absolute coronary flow during steady state maximum hyperemia; R, microvascular resistance; and WU, Wood units. Reprinted from Keulards et al17 with permission. Copyright ©2018, Europa Group.

No major side effects of the absolute flow/resistance measurement method were noted. Two patients showed rapidly transient atrioventricular block with a saline infusion rate of 20 mL/min in the right coronary artery. The atrioventricular block resolved immediately after the infusion was stopped. After adjustment of
the infusion pump to a lower infusion rate (ie, less decrease of distal temperature), the measurements were completed without difficulty.

### Absolute Coronary Blood Flow and Resistance Measurement

There was a significant increase in absolute blood flow in the CTO vessel over time in the overall cohort (Figure 2A). Flow in the recanalized CTO artery increased by 49% from 148±53 mL/min immediately after opening to 221±77 mL/min at follow-up ($P<0.001$). Absolute flow increased in 23 of the 25 cases. In agreement, absolute resistance in the myocardial territory perfused by the CTO artery, decreased by 29% from 545±255 WU immediately after the procedure to 387±128 WU at follow up ($P=0.014$) (Figure 2B). There were no significant changes in the absolute coronary blood flow and resistance in the predominant donor between baseline and follow-up. Absolute blood flow changed from 162 mL/min before PCI to 175 mL/min immediately after PCI and to 189 mL/min at follow-up ($P=0.40$). Microvascular resistance was 473 WU before PCI, 542 WU immediately after PCI and 474 WU at follow-up ($P=0.44$; Figure 3). In these 10 patients in whom donor vessel measurements were available, a net gain in summed absolute blood flow to the CTO and donor vessel myocardial territory of 118 mL/min was observed.

### Assessment of Vessel Diameter Changes

All patients had complete follow-up quantitative coronary analysis measurements. The distal CTO vessel lumen dimensions changed significantly from immediately after PCI to follow-up. The minimal lumen diameter increased from 1.17±0.39 mm to 1.42±0.45

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**Table 1. Baseline Characteristics**

| Number of Patients (n=25) | N (%) or Mean±SD |
|--------------------------|------------------|
| Male sex, N (%)          | 19 (76)          |
| Age, mean (SD)           | 65.9±8.7         |
| Medical history          |                  |
| Hypertension, n (%)      | 20 (80)          |
| Current smoking, n (%)   | 4 (16)           |
| Diabetes mellitus, n (%) | 7 (28)           |
| Dyslipidemia, n (%)      | 19 (76)          |
| Previous myocardial infarction, n (%) | 12 (48) |

**Clinical details**

| CCS angina class | I     | 7 (28) |
|                  | II    | 8 (32) |
|                  | III   | 8 (32) |
|                  | VI    | 2 (8)  |

| Left ventricular ejection fraction | <35% | 0 (0) |
|                                   | 35%-50% | 10 (40) |
|                                   | >50%  | 15 (60) |

**Angiographic details**

| Occluded target vessel              | Right coronary artery | 20 (80) |
|                                    | Left anterior descending artery | 3 (12) |
|                                    | Left circumflex artery | 2 (8)  |
| Predominant donor vessel            | Right coronary artery | 5 (20)  |
|                                    | Left anterior descending artery | 17 (68) |
|                                    | Left circumflex artery | 3 (12)  |

| Collateral filling                  | Bridge collaterals | 3 (12)  |
|                                    | Retrograde filling | 18 (72) |
|                                    | Both              | 4 (16)  |

| Collateral size                     | CC0               | 2 (8)   |
|                                    | CC1               | 17 (68) |
|                                    | CC2               | 6 (24)  |

| J-CTO score                         | Tapered/blunt     | 5/10 (20/43) |
|                                    | Calcification present | 15 (60) |
|                                    | Bending >45 degrees | 5 (20)   |
|                                    | Length >20 mm      | 11 (44)  |
|                                    | Retry lesion       | 2 (8)    |

**Total J-CTO-score**

| J-CTO 0                  | 7 (28) |
| J-CTO 1                  | 3 (12) |
| J-CTO 2                  | 8 (32) |
| J-CTO 3                  | 7 (28) |

**Total J-CTO score (mean, SD)**

1.64±1.25

Summary values represent number (%) or mean±SD. CC indicates continuous connection; CCS, Canadian Cardiovascular Society; and J-CTO score, Japanese–Chronic Total Occlusion score.

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**Table 2. Procedural Characteristics**

| Number of Patients (n=25) | N (%) or Mean±SD |
|--------------------------|------------------|
| Technique                |                  |
| Antegrade wire escalation| 14 (56)          |
| Retrograde wire escalation| 4 (16)          |
| Antegrade dissection and reentry | 2 (8) |
| Retrograde dissection and reentry | 5 (20) |
| Stent length             | 72.2±26          |

**Other**

| Procedure time, min   | 136.3±78.1 |
| Fluoroscopy time, min | 39.2±26.3  |
| Wire crossing time, min | 53.4    |
| Radiation dose area product, mGy | 200 484±155 497 |

Summary values represent mean±standard deviation. mGy indicates milligray.
The mean lumen diameter increased from 1.73 mm to 1.99 (P<0.05; Table 3). In parallel, epicardial resistance of the CTO artery calculated as (Pa-Pd)/Q, decreased by 17% from 82±55 WU to 68±49 WU (P=0.155; Table 3).

**DISCUSSION**

The present study is the first to perform sequential invasive measurements of absolute hyperemic coronary blood flow and microvascular resistance immediately after successful CTO PCI and at short-term follow-up. The study showed that coronary blood flow and microcirculatory function of the dependent myocardium of a recanalized CTO, are still far from normal immediately after technically successful PCI and significantly improve over time. At the same time, significant luminal increase of the distal CTO vessel was observed. In a subgroup of patients where absolute blood flow and resistance were assessed at the predominant donor vessel, there was no change among measurements before CTO PCI, immediately afterward, and at short-term follow-up.

Novel techniques, advanced equipment, and increasing operators’ experience in contemporary CTO PCI have resulted in high procedural success and low complications rates. However, target lesion failure remains higher compared with non-CTO angioplasty. Better understanding of physiological changes in the recanalized epicardial artery and the dependent myocardium could potentially provide useful insight in optimization of the procedure and improving long-term clinical outcomes.

Our study showed a significant increase in hyperemic coronary blood flow and improvement in microvascular function a few weeks after the procedure: Maximum blood flow increased by 49% and microvascular resistance decreased by 29% at follow-up compared with post-PCI. There was an accompanying increase in distal vessel diameter and a decrease of the epicardial resistance of the CTO artery by 17%. It appears that the increase of coronary flow is explained by 2 factors: decrease in microvascular resistance and distal vessel remodeling reflected by decrease of distal epicardial resistance. The increase in blood flow exceeds the decrease of microvascular resistance as the increase in flow results from both a lower epicardial and lower microvascular resistance.

Notably, almost all patients showed the phenomena described above, but in a few patients (generally mildly) increased resistance or decreased absolute flow at follow-up was observed despite normal cardiac magnetic resonance. Of the 5 patients with increased resistance at follow-up, 4 patients had diabetes mellitus. Werner et al also reported less or delayed recovery in such patients, but these numbers are too small to justify
conclusions. Further improvement in flow and resistance may happen over time and will be further investigated in the presently ongoing IMPACT-CTO 2 (Impact of Coronary Chronic Total Occlusion Percutaneous Coronary Intervention on Culprit Vessel Physiology) trial (ClinicalTrials.gov Identifier: NCT03830853).

Previous studies have suggested a sort of “vascular wall hibernation” at the distal coronary segments of a recanalized CTO and failed response to endothelium-dependent and -independent stimuli. In these studies, different hyperemic stimuli were applied immediately after successful CTO PCI and vasomotion was assessed; intracoronary administration of nitroglycerin, adenosine, and acetylcholine or incremental atrial pacing failed to provoke the expected vessel dilatation. The one study that included a follow-up assessment suggested transient impairment of vasomotion function. The increase in hyperemic absolute coronary blood flow demonstrated in our study can be partially explained by the above findings and strongly argues for improved distal coronary artery wall function. As in our study, it has been previously shown that restoration of flow, shear stress, and vasomotor properties lead to increase in distal vessel luminal dimensions mainly via external elastic membrane enlargement and plaque regression. Regarding microvascular function, a previous study by Werner et al suggested that microvascular dysfunction was frequently observed immediately after recanalization of a CTO and recovered over time in most of the cases. In that study, Doppler-derived coronary flow reserve (CFR) was used to quantify microvascular function. However, coronary flow reserve is determined by both the epicardial and microcirculatory function. Our study remains the first to report sequential invasive measurements of absolute microvascular resistance.

In a recently published study by Karamasis et al., comprehensive FFR and collateral FFR measurements were performed immediately after CTO PCI and at short-term follow-up. The study showed an FFR increase overtime accompanied by significant reduction of collateral function as expressed by collateral FFR. Furthermore, increased distal vessel diameter and "positive remodeling" with a significant increase in distal luminal diameter was observed. Our study adds essential information regarding microvascular resistance to these observations, completing the physiological concept.

In the subgroup of patients in whom physiological measurements were performed in the predominant donor vessel, there was no significant change in the hyperemic coronary blood flow immediately after CTO or at follow-up. Two recent studies assessed changes in the donor artery FFR after CTO PCI and reported a small but statistically significant increase. One of them suggested a concomitant reduction in coronary flow assessed by intracoronary Doppler. This change was explained by the decrease in the myocardial territory supplied by the donor artery after successful recanalization of the CTO vessel. In our study, possibly the small number of patients in whom donor measurements were performed did not allow a similar observation. Regarding microvascular resistance, as expected there were no changes in the remote to CTO myocardium.

**LIMITATIONS**

Our study has a number of limitations. First, it involved a small number of patients, and the results should be considered hypothesis generating and cannot be used to generalize the findings to a larger complex
population undergoing CTO-PCI. The magnitude of the changes was variable presumably because of the variability in collateral flow, microvascular and coronary artery wall dysfunction, and the amount of viable tissue perfused by the CTO artery. Nevertheless, the observations were consistent among most of the patients. The vast majority of the recanalized CTO vessels involved the right coronary artery. Although this reflects current CTO PCI practice, a more equal CTO vessel distribution would be more representative for the different vascular beds. We did not measure coronary wedge pressure, and therefore we couldn’t calculate collateral FFR and provide a quantitative assessment of collateral blood flow enhancing further the interpretation of the observed physiological changes.

Also, the follow-up interval was not regular, which should be recognized. However, no further procedures were performed before physiological measurements, and if staged PCI was indicated, it took place after all study measurements so as not to affect these.

Finally, changes in the distal vessel lumen were assessed by quantitative coronary analysis. Intracoronary imaging would be more informative, as it would allow evaluation of vessel remodeling, plaque modification, and stent deployment features. The ongoing IMPACT-CTO 2 study combines sequential advanced physiological measurements, including absolute blood flow and resistance with optical coherence tomography in an attempt to provide further insight in coronary physiology and anatomy after CTO PCI.

CONCLUSIONS

This study demonstrates that after successful PCI of a chronically occluded coronary artery, physiology of the distal coronary artery and microvasculature does not normalize immediately but improves significantly within a few months. This is reflected by an increase of maximum coronary blood flow of 49%, decrease of microvascular resistance of the myocardium by 29%, and positive remodeling of the distal coronary.

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