FLOWER-MI and the root of the problem with non-culprit revascularisation

Matthew E Li Kam Wa,1,2 Kalpa De Silva,1,2 Carlos Collet,3 Divaka Perera1,2

ABSTRACT
How do we reduce cardiac death and myocardial infarction by percutaneous coronary intervention (PCI) in coronary heart disease? Although the interventional community continues to grapple with this question in stable angina, the benefits of PCI for non-culprit lesions found at ST-elevation myocardial infarction are established. Is it then wishful thinking that an index developed in stable coronary disease, for identifying lesions capable of causing ischaemia will show an incremental benefit over angiographically guided non-culprit PCI? This is the question posed by the recently published FLOW Evaluation to Guide Revascularization in Multi-vessel ST-elevation Myocardial Infarction (FLOWER-MI) trial. We examine the trial design and results; ask if there is any relationship between the baseline physiological significance of a non-culprit lesion and vulnerability to future myocardial infarction; and consider if more sophisticated methods can help guide or defer non-culprit revascularisation.

INTRODUCTION
Randomised trials involving more than 6300 patients over the last decade all agree that revascularisation of non-culprit lesions at ST-elevation myocardial infarction (STEMI) is superior to medical therapy alone. However, it is uncertain if fractional flow reserve (FFR) is beneficial in deciding which non-culprit lesions to revascularise. FLOW Evaluation to Guide Revascularization in Multi-vessel ST-elevation Myocardial Infarction (FLOWER-MI) was an investigator-initiated, multicentre trial conducted over 2 years in France, that randomised 1171 patients following STEMI to FFR or angiography groups at 1 year. Why did FFR not perform as well as hoped? It is worth noting that 7 out of 9 (78%) deaths in the FFR group were cardiac, vs only 3 of 10 (30%) in the angiography group. Non-fatal MI was also numerically higher in the FFR group and despite less PCI being performed, the periprocedural MI (type 4a) rate was three times higher. The 0.3% periprocedural MI rate within the angiography arm, however, is almost certainly underreported, given the challenges in diagnosing periprocedural MI around the time of STEMI, when troponin levels are already elevated and/or still rising. Compliance to troponin collection is not reported, nor the number of troponin samples taken per patient. The overall MI rate within deferred patients within the FFR arm was high at 5.6%, with divergence in the Kaplan–Meier curves at 6 months from the index procedure. However, we also do not know which events occurred in deferred lesions and ultimately attempts to draw conclusions are limited by the low numbers of events, which were only one-third of what the authors anticipated.

To understand this, we should examine the trial design in more detail. At first glance, the randomisation of FLOWER-MI bears some resemblance to the stable coronary disease cohort of FAME (Fractional Flow Reserve Versus Angiography for Multivessel Evaluation). Unfortunately, there was no blinded assessment of FFR within the angiography guided arm and it is uncertain why the pre-PCI FFR result was not available for 154 (16%) of lesions. Why is this important? Any study depends on a difference in exposure between the two arms to show a treatment effect and discordance between FFR and angiographic severity is most pronounced in the intermediate (<70%) range. Considering
the inclusion of severe stenoses in this study and that intermediate disease comprised less than half of all lesions, FFR changed the treatment strategy in only 40% of lesions. This is in keeping with both DANAMI-3-PRIMULTI (Third Danish Study of Optimal Acute Treatment of Patients with STEMI: Primary PCI in Multivessel Disease) and Compare-Acute cohorts (31% and 46%, respectively). At the patient level, this translated to only one-third (198 of 586 patients) not having non-culprit PCI. Relying on this minority to show a statistically significant difference in outcomes was always going to be a challenge. Although this trial was designed prior to the results of COMPLETE (Complete Versus Culprit-Only Revascularization Strategies to Treat Multivessel Disease After Early PCI for STEMI); CV, cardiovascular; CVA, cerebrovascular accident; DANAMI-3-PRIMULTI, Third Danish Study of Optimal Acute Treatment of Patients with STEMI: Primary PCI in Multivessel Disease; FFR, fractional flow reserve; FLOWER-MI, FLOW Evaluation to Guide Revascularization in Multi-vessel ST-elevation Myocardial Infarction; MI, myocardial infarction; NR, not formally reported; OMT, optimal medical therapy; PCI, percutaneous coronary intervention; STEMI, ST-elevation myocardial infarction; TIA, transient ischaemic attack.

Table 1 Comparison of FLOWER-MI with selected trials of non-culprit revascularisation after STEMI

<table>
<thead>
<tr>
<th></th>
<th>FLOWER-MI</th>
<th>COMPLETE</th>
<th>Compare-Acute</th>
<th>DANAMI-3-PRIMULTI</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Dates</strong></td>
<td>Angio</td>
<td>FFR</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dates</td>
<td>2016–2018</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>No of patients</strong></td>
<td>1171</td>
<td>4041</td>
<td>885</td>
<td>627</td>
</tr>
<tr>
<td>Median follow-up (mths)</td>
<td>12</td>
<td>36</td>
<td>12</td>
<td>27</td>
</tr>
<tr>
<td>Design</td>
<td>Angio (&gt;50%) vs FFR</td>
<td>Angio (≥70%)* vs OMT</td>
<td>FFR vs OMT</td>
<td>FFR vs OMT</td>
</tr>
<tr>
<td>FFR ≥0.80</td>
<td>44%</td>
<td>–</td>
<td>46%</td>
<td>31%</td>
</tr>
<tr>
<td>Composite primary outcome (no/%)</td>
<td>Death, non-fatal MI, unplanned admission and urgent revasc</td>
<td>CV death, MI, ischaemia-driven revasc</td>
<td>Death, non-fatal MI, any revasc, CVA/TIA</td>
<td>Death, non-fatal MI, ischaemia-driven revasc</td>
</tr>
<tr>
<td>Event rate with revascularisation (%)†</td>
<td>24 (4.2)</td>
<td>32 (5.5)</td>
<td>179 (8.9)</td>
<td>23 (7.8)</td>
</tr>
</tbody>
</table>

*Only a minority of patients with moderate stenoses (50%–69%) that would have mandated FFR measurement were enrolled in COMPLETE.
†Event rates for COMPLETE shown as % per person-year as reported by the authors, for comparison with the 12-month follow-up of FLOWER-MI.

COMPLETE, Complete versus Culprit-Only Revascularisation Strategies to Treat Multivessel Disease after Early PCI for STEMI; CV, cardiovascular; CVA, cerebrovascular accident; DANAMI-3-PRIMULTI, Third Danish Study of Optimal Acute Treatment of Patients with STEMI: Primary PCI in Multivessel Disease; FFR, fractional flow reserve; FLOWER-MI, FLOW Evaluation to Guide Revascularization in Multi-vessel ST-elevation Myocardial Infarction; MI, myocardial infarction; NR, not formally reported; OMT, optimal medical therapy; PCI, percutaneous coronary intervention; STEMI, ST-elevation myocardial infarction; TIA, transient ischaemic attack.

There are also differences such as the rates of glycoprotein IIb/IIIa and P2Y12 inhibitor use, although this is reflective of optimal therapy at the time of each study. However, a more significant contributor is likely to be the open label design of these trials.

Open label trials and caveats
In the era of primary PCI and modern medical therapy, it is increasingly difficult to show benefits in objective, ‘hard’ endpoints without large, and therefore, expensive trials. Open label trials conflated with ‘soft’ outcomes such as hospitalisation and urgent revascularisation renders them susceptible to bias, but also makes it difficult to distinguish the true symptomatic and prognostic effects of PCI. In this regard, a strength of COMPLETE was powering for a coprimary endpoint with the objective outcomes of cardiovascular death and MI. We also caution against extrapolating results from any STEMI trial to the non-ST-elevation MI population who have more comorbidities, more complex coronary disease and the culprit lesion is frequently ambiguous or determined by using the angiogram alone.

Impact on clinical practice
Could less be more?
Given the similar risk of MI and revascularisation in previous FFR and angiographic guided trials versus medical therapy (table 1), FLOWER-MI certainly addresses clinically important questions. Is there a middle ground where the benefits of ‘complete’ (or...
COMPLETE) revascularisation can be gained, without stenting every angiographic lesion? Unfortunately, given the lack of statistical difference in its outcomes, the answers remain unclear. We believe it continues to remain reasonable to offer either an angiographic or FFR guided strategy. Based on expected discordance rates, an FFR strategy versus the (almost entirely) angiographic strategy of COMPLETE for >70% stenoses would be expected to still revascularise 85% of lesions.

If this is the case, should we quibble over a 15% reduction in lesion revascularisation? In our view, it is important to remember that PCI should not be regarded as a benign procedure. Aside from the periprocedural risk, underreported in FLOWER-MI, at a mean age of 62 years these patients would likely live another 20 years, accruing risk of very late stent-related events year-on-year. This may be an acceptable trade off on an individualised basis, but not if stent implantation has no prospect of improving their prognosis in the first place. We believe that FLOWER-MI provides a potentially simple and important message, FFR can safely defer 1/3 of non-culprit lesions noted at the time of STEMI.

Is physiological assessment valid at the time of STEMI?
FLOWER-MI aside, we should consider the validity of physiological coronary assessment in acute coronary syndromes (ACS). We know the measurement of intracoronary pressure as a pragmatic surrogate for flow reduces urgent revascularisation, spontaneous MI and is cost effective in those with stable coronary artery disease planned for PCI. Physiological assessment to improve symptoms has a clear rationale here: if ischaemia is required to cause angina, intervention in the absence of flow limitation exposes the patient to the risk of a procedure, while also being unhelpful for their angina.

Due to changes in microvascular physiology at the time of STEMI, concern exists whether indices such as FFR or iFR, (both developed and validated within the stable coronary artery disease population) accurately reflect functional significance. This is evaluated in detail by a recent review, but importantly, flow as well as hyperaemic and resting pressure indices can be overestimated or underestimated during the index admission. These changes could have significant impact around the dichotomous treatment thresholds established for stable angina. Although studies are small, conflicting, and the long-term clinical implications are not characterised - up to one in five lesions deferred by FFR, and one in three lesions treated by iFR could have a different treatment if reassessed 30 days later.

Why do patients have adverse events after culprit revascularisation?
If physiological indices are prone to inaccuracy after STEMI, why not simply pursue angiographic-guided revascularisation? Serial angiography and pathological studies certainly support that culprit lesions at the time of STEMI are severely stenotic. However, to adopt this approach would be to disregard compelling evidence about the origin of adverse events after STEMI.

FLOWER-MI does not allow us to attribute events to culprit or non-culprit lesions (deferred by FFR or otherwise) but vessel level outcomes were reported in a sub study of COMPARE-Acute, which had the benefit of measuring non-culprit FFR in both arms. In this study, non-culprit, angiographically insignificant vessels (that did not warrant FFR measurement in the first place) were responsible for two thirds of the spontaneous MIs within the follow-up period.

For a systematic multimodality evaluation and natural history of non-culprit lesions, we should turn our attention to the contemporary PROSPECT II study. After ACS (22% STEMI), 898 patients underwent FFR guided revascularisation of all >40% non-culprit lesions, followed by three-vessel intravascular ultrasound (IVUS) and follow-up for 4 years. Although we must be cautious in extrapolating data from a combined ACS cohort, of the major adverse cardiac events, two thirds were attributable to non-culprit lesions. Of the non-culprit lesions which underwent intracoronary imaging characterisation at baseline, the mean diameter stenosis was only 47%. Clearly, a strategy of angiographic-guided revascularisation for >50%–70% stenoses is liable to miss a significant number of potentially clinically relevant lesions.

Conversely, not all severely stenotic lesions at baseline are destined to cause a future adverse event. This broad heterogeneity in coronary plaque vulnerability, haemodynamic stresses and the risk profile of patients with coronary artery disease explains why some lesions lead to MI and others do not. Plaques that are more likely to rupture or erode are termed ‘vulnerable’ and have distinct features that can be identified by dedicated imaging techniques. When combined with patient level factors such as diabetes, lifestyle factors or the pro-inflammatory state of STEMI, these unstable plaques can lead to acute coronary syndromes (ACS).

Vulnerable plaque and vulnerable patient assessment
Intracoronary imaging has focused on identifying the pathological features associated with thrombotic occlusions: virtual histology IVUS (VH-IVUS: thin cap fibroatheroma, plaque burden and minimum lumen area), near-infrared spectroscopy (NIRS: lipid rich plaque) and optical coherence tomography (OCT: lipid arc, cap thickness and macrophages) which seem to be more prevalent in angiographically severe lesions. However, only a minority of those possessing a full house of morphological high-risk features progress to an ACS. The addition of shear and axial stress assessment and the quantification of trans-lesional gradients by computational fluid dynamics analysis appears to also provide additional information regarding risk of future rupture. It also may help predict plaque erosion, which comprises an increasing proportion of ACS, particularly in women.
FFR appears to have an association with these known plaque vulnerability features. Could it be that FFR provides a ‘good enough’ measure of vulnerability and lipid burden that avoids the excess intervention with an angiographic guided approach? A pooled analysis of three randomised trials (with half of its patients drawn from the STEMI cohort of DANAMI-3-PRIMULTI), assessed the ability of the residual SYNTAX (Synergy between PCI with TAXUS and Cardiac Surgery) score after FFR guided ‘complete’ revascularisation to predict adverse events. Bearing in mind the limitations of post-hoc analyses and the mixed cohort, leaving angiographically significant lesions that were FFR negative did not appear to be associated with ischaemic outcomes after 2 years. It may be that physiological indices are a crude, but familiar and reproducible surrogate for abnormal plaque stress, but also that a significant component of a patient’s risk may potentially be predicted by total coronary atheroma burden alone. Although it appears the functional characteristics of a lesion may describe some measure of its vulnerability, whether this relationship is associative, or causative, remains unknown.

**FUTURE DIRECTIONS**

Despite decades of advancement, vulnerable plaque identification still has several obstacles to overcome. First, a method of identifying plaques with a sufficient positive predictive value for adverse events is needed. We may find more success with the alternative approach of identifying the truly stable plaque where PCI could be safely deferred, or where novel and expensive therapies could be withheld.

Second, plaque assessment at baseline may be of limited value given that plaques can both deteriorate as well as stabilise over time on medical therapy. Invasive techniques or the use of ionising radiation are not ideal for serial assessment and magnetic resonance angiography may hold some promise in this area.

Third, the identification of vulnerability on a lesion rather than patient level seems valuable only if an effective targeted local intervention is available. With the withdrawal of the absorb bioresorbable vascular scaffold (BVS) from the market, we are left with encouraging signs, but no conclusive evidence to support a strategy of PCI to vulnerable plaques. The PREVENT trial (NCT02316886) continues to recruit, randomising 1600 patients to drug eluting stent insertion versus optimal medical therapy for non-flow limiting disease with high-risk intracoronary imaging features, and OCT-CONTACT (NCT04878133) will randomise 460 patients to an OCT-guided strategy versus complete revascularisation. However, both studies are limited to only lesions with ≥50% angiographic stenosis. Whether contemporary drug eluting stents with ultrathin struts, the next generation of bioresorbable vascular scaffolds, or PCI to vulnerable plaque regardless of angiographic stenosis severity will be beneficial, remains unanswered.

**CONCLUSIONS**

FLOWER-MI is ultimately limited by its low event rate in evaluating an FFR versus an angiography-guided revascularisation strategy for non-culprit lesions after STEMI, and therefore, either a physiology or angiography guided approach still seem reasonable. However, trying to reduce adverse hard outcomes with physiology versus the blunt tool of angiography may only be a partial solution. Risk stratifying individual lesions with new and emerging techniques could allow a truly personalised approach to STEMI care. The pursuit of a clinically viable method to detect, and then intervene on the vulnerable plaque continues.

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