Percutaneous coronary intervention versus bypass grafting in left main coronary artery disease

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Without revascularization, patients with left main coronary artery disease (LMCAD) have a poor prognosis (1). There is a clear survival benefit from revascularization over medical management (1). Percutaneous coronary intervention (PCI) has been traditionally deferred in preference for coronary artery bypass grafting (CABG), due to the anatomic complexity (2). However, with the evolution of drug-eluting stents (DES), there has been a reduction in restenosis rates and mortality, repeat revascularization and major adverse cardiac and cerebrovascular events (MACCE) compared with bare metal stents (3). There has been a renewed interest in expanding the indication for PCI in patients with LMCAD (3). Despite this heightened interest in the comparative outcomes of DES versus CABG for patients with LMCAD, the choice for the optimal revascularization technique remains controversial.

A recent meta-analysis by Li and co-workers indicated that DES was associated with lower peri-procedural risks than CABG, but was inferior to CABG in terms of repeated revascularization in patients with LMCAD at 5 years (4). There was no difference in death, myocardial infarction, cerebrovascular events or revascularization between randomized controlled trials and observational groups (4). It is important to point out that LMCAD includes a wide spectrum of anatomic features and it may be associated with concurrent multi-vessel disease. Previous studies have shown that 70% to 90% of patients with LMCAD will present with multi-vessel coronary artery disease (5,6). A recent meta-analysis by Cao et al. demonstrated that patients with LMCAD and concomitant multi-vessel disease may be underrepresented in the comparative studies (7). In the real world, patients are more likely to undergo CABG, if they are found to have LMCAD in combination with multi-vessel disease (7).

Studies evaluating more than 5-year outcomes of PCI versus CABG in patients with LMCAD are limited. Subgroup analysis of LMCAD patients in the SYNTAX randomized trial recently reported comparable 5-year cardiovascular outcomes for PCI and CABG (8). The randomized control trial is the gold standard for comparing treatments, but generally enrolls selective patients without high-risk profiles. Perhaps more importantly, it should be highlighted that only a minor proportion of patients assessed for eligibility were eventually randomized in the SYNTAX trial. Of the patients who were excluded from randomization, 198 patients who underwent DES and 1,077 patients who underwent CABG were included in a separate nested registry, which found higher incidences of MACCE, mortality, myocardial infarction, and repeat revascularization for patients who were treated by DES at 12 months. The main reason for registry allocation to CABG (70.9%) was the complexity of anatomy, whereas the main reason for PCI allocation was increased comorbidity (70.7%). Likewise, a randomized controlled trial by Boudriot and colleagues compared 100 patients who underwent DES with 101 patients who underwent CABG (9). The authors reported similar combined incidences of cardiac death, myocardial infarction, and repeat revascularization at 12 months (19.0% vs. 13.9%, P=0.19 for non-inferiority). However, of the 229 patients with LMCAD who were considered ineligible for randomization, a significantly lower incidence of
MACCE was reported for CABG compared with DES and conservative therapy (17.8% vs 27.5% vs 43%). Although reasons for exclusion from randomization differ between trials, it should be emphasized that results derived from patients selected for randomization in these tertiary referral centers do not necessarily represent the target population of patients diagnosed with LMCAD, especially those with more complex disease.

The 5-year follow-up results from the SYNTAX randomized trial and the 5-year outcome from CREDO-Kyoto PCI/CABG Registry Cohort-2—confirmed the utility of the SYNTAX score for risk stratification and selection of mode of revascularization procedure, supporting the current updated clinical guidelines for LMCAD (8,10-13). In LMCAD patients with relatively less anatomical complexity, represented by a low or intermediate SYNTAX score, PCI using DES is a reasonable alternative to CABG in real-world clinical practice. However, CABG still remains the preferable treatment for LMCAD patients, especially those with high anatomical complexity (i.e., high SYNTAX score or complex multi-vessel disease).

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

References

4. Li Q, Zhang Z, Yin RX. Drug-eluting stents or coronary artery bypass grafting for unprotected left main coronary artery disease: a meta-analysis of four randomized trials and seventeen observational studies. Trials 2013;14:133.
13. Authors/Task Force members, Windecker S, Kolh P,

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