Incidence and predictors of unplanned non-target lesion revascularisation up to three years after drug-eluting stent implantation: insights from a pooled analysis of the RESOLUTE Global Clinical Trial Program

Mohamed Abdel-Wahab1*, MD; Franz-Joseph Neumann2, MD; Patrick Serruys3, MD, PhD; Sigmund Silber4, MD, PhD; Martin Leon5, MD; Laura Mauri6, MD, MSc; Alan Yeung7, MD; Jorge A. Belardi8, MD; Petr Widimsky9, MD, DrSc; Ian Meredith10, MBBS, PhD; Shigeru Saito11, MD; Gert Richardt1, MD

1. Heart Center, Segeberger Kliniken, Bad Segeberg, Germany; 2. Heart Center Bad Krozingen, Bad Krozingen, Germany; 3. Thoraxcenter-Erasmus University, Rotterdam, The Netherlands; 4. Heart Center at the Isar, Munich, Germany; 5. Columbia University Medical Center and New York-Presbyterian Hospital, New York, NY, USA; 6. Brigham and Women’s Hospital and Harvard Medical School, Boston, MA, USA; 7. Stanford University School of Medicine, Stanford, CA, USA; 8. Instituto Cardiovascular de Buenos Aires, Buenos Aires, Argentina; 9. Third Faculty of Medicine, Charles University, Prague, Czech Republic; 10. Monash University, Melbourne, Australia; 11. Shonan Kamakura General Hospital, Kanagawa, Japan

GUEST EDITOR: Rafael Beyar, MD, DSc, MPH; Director, Rambam Health Care Campus, Women’s Division/Dr Phillip and Sara Gotlieb Chair, Department of Medicine and Biomedical Engineering, Technion, Israel

Abstract

Aims: To compare the incidence and predictors of target lesion revascularisation (TLR) and non-TLR after percutaneous coronary intervention with drug-eluting stents (DES).

Methods and results: We pooled patient-level data on 6,137 patients (Resolute zotarolimus-eluting stent: 5,016, XIENCE everolimus-eluting stent: 1,121) in the RESOLUTE Global Program. At three years, clinically driven TLR, unplanned non-TLR, and no revascularisation occurred in 186, 618, and 5,333 patients, respectively. On multivariate analysis, predictors of both TLR and non-TLR were pre-procedure diameter stenosis (%) (odds ratio [OR] 1.01, 95% confidence interval [CI] [1.01-1.02], and OR 0.99 [0.99-1.00]), diabetes (OR 1.46 [1.07-1.99], and OR 1.37 [1.15-1.64]), and prior PCI (OR 1.42 [1.01-2.00], and OR 1.41 [1.18-1.68]). Baseline characteristics associated with TLR only were prior coronary artery bypass graft surgery (OR 2.85 [1.91-4.27]), in-stent restenosis (OR 2.35 [1.43-3.83]), age (OR 0.98 per year [0.97-1.00]), hypertension (OR 1.64 [1.10-2.44]), and pre-procedure reference vessel diameter (OR 0.74 per mm [0.55-0.99]). Baseline characteristics associated with non-TLR only were lesion location (left anterior descending vs. all others) (OR 0.70 [0.59-0.83]), and hyperlipidaemia (OR 1.42 [1.15-1.75]).

Conclusions: The cumulative incidence of non-TLR at three years in patients treated with current-generation DES was almost three times higher than TLR.

KEYWORDS
• coronary artery disease
• drug-eluting stent
• risk factors
• zotarolimus-eluting stent

* Corresponding author: Herzzentrum, Segeberger Kliniken GmbH, Am Kurpark 1, 23795 Bad Segeberg, Germany. E-mail: mohamed.abdel-wahab@segebergerkliniken.de
Introduction

First-generation drug-eluting stents (DES) substantially decreased angiographic and clinical restenosis after percutaneous coronary intervention (PCI)\(^1\). Newer DES, such as Resolute\textsuperscript{TM} zotarolimus-eluting stents (R-ZES; Medtronic, Minneapolis, MN, USA) and XIENCE V\textsuperscript{®} everolimus-eluting stents (EES) (Abbott Vascular, Santa Clara, CA, USA), further improved the long-term safety and efficacy of DES with reductions of stent thrombosis and target lesion revascularization compared to first-generation DES\(^2\). The highly effective DES-based local therapy of coronary lesions is currently applied in a wide spectrum of patients, including those with advanced coronary artery disease and complex anatomy\(^5,7,8\). Consequently, anticipation and prevention of disease progression in non-intervened non-target lesions is gaining increasing attention.

To assess the incidence and risk factors associated with clinical plaque progression requiring non-target lesion revascularisation during long-term follow-up after culprit lesion PCI with modern DES, patient-level data were pooled from five clinical trials of the RESOLUTE Global Clinical Trial Program with harmonised endpoint definitions (RESOLUTE First-in-Man, RESOLUTE All Comers, RESOLUTE International, RESOLUTE US and RESOLUTE Japan)\(^3,6\). Baseline lesion characteristics were determined by an independent core laboratory in all studies except RESOLUTE International (site-reported visual estimation). Follow-up was done in-clinic or by telephone.

For this analysis, patients were grouped as follows based on their outcomes up to three years of follow-up: target lesion revascularisation (TLR), non-TLR, and no revascularisation. Patients requiring both TLR and non-TLR were excluded from the analysis. Revascularisations could be percutaneous or surgical. A non-TLR was defined as an unplanned procedure elsewhere in the target vessel or in other coronary arteries.

The studies contributing to this analysis used harmonised, validated endpoint definitions. Study-specific clinical event committees composed of interventional cardiologists not directly involved in the study applied these definitions to all suspected events.

Data safety monitoring boards were composed of non-interventional and interventional cardiologists not directly involved in the study. Each study’s data safety monitoring board reviewed adverse event data and could recommend stopping the study. Clinical event committees and data safety monitoring boards were coordinated by independent academic research organisations.

Statistical analysis

All analyses were conducted on an intention-to-treat basis. Continuous parameters were presented as mean±standard deviation and compared using a t-test or Wilcoxon rank-sum test, as appropriate. Nominal parameters were presented as percentages and compared using Fisher’s exact test. Time to event endpoints were calculated using the Kaplan-Meier method and compared using the log-rank test. Multivariate stepwise regression analysis was conducted to assess independent correlates of TLR and non-TLR at three years of follow-up. A simple logistic regression was performed using the following baseline characteristics: prior coronary artery bypass graft, in-stent restenosis, prior PCI, history of diabetes, hypertension, hyperlipidaemia, pre-procedure diameter stenosis (%), previous myocardial infarction (MI), lesion length (mm), pre-procedure reference vessel diameter (RVD) (mm), bend ≥45\(^\circ\), calcification (moderate/severe vs. none/mild), TIMI flow 3, age (years), male sex, serum creatinine (µmol/L), vessel location (left anterior descending vs. other), lesion class B2/C, unstable angina, and current smoker. Then, a multiple logistic regression was performed with an entry criterion of 0.20 and a stay criterion of 0.10. Analyses were performed using SAS software version 9.1 or later (SAS Institute, Cary, NC, USA). A p-value <0.05 was considered statistically significant.

Results

PATIENT/SUBJECT DISPOSITION AND CHARACTERISTICS

The five RESOLUTE studies contributing to this analysis enrolled 6,282 patients (R-ZES: 5,130, EES: 1,152). At three years, the incidence of clinically driven TLR was 5.3% and unplanned non-TLR was 12.1%. After excluding 145 patients who had both clinically driven TLR and unplanned non-TLR, 6,137 patients (R-ZES:...
5,016, EES: 1,121) were included in the analysis, of whom 186 had clinically driven TLR, 618 had unplanned non-TLR, and 5,333 had no revascularisation at three years. Among the 618 patients with non-TLR, 162 patients (26%) had non-TL TVR and 518 patients (84%) had revascularisation outside the target vessel.

TIMING AND PREDICTORS OF REPEAT REVASCULARISATION

TLR was numerically more frequent than non-TLR only within the first 20 days after the index procedure (Figure 1A). Thereafter, the Kaplan-Meier curves crossed, and TLR and non-TLR rates further divided over time. The separation was most prominent in the second and third years of follow-up (Figure 1B).

Compared to patients with no revascularisation, those with TLR and with non-TLR were more likely to have diabetes, insulin-dependent diabetes, and a history of cardiovascular procedures and disorders (Table 1). Patients with TLR were also more likely to have Canadian Cardiovascular Society class III angina. Patients with non-TLR had a higher mean BMI, greater number of native vessels with >50% diameter stenosis, and a greater incidence of stable angina and lesser incidence of MI as the reason for revascularisation (Table 1).

Prior in-stent restenosis was significantly more common in TLR patients than in patients with no revascularisation (Table 2). Compared to patients with no revascularisation, those with non-TLR had fewer lesions in the left anterior descending and left main coronary arteries, and fewer lesions with thrombus (Table 2).

A multivariate analysis was conducted to determine predictors of TLR (Figure 2A) and non-TLR (Figure 2B) at three years.

---

**Figure 1.** Cumulative incidence of clinically driven target lesion revascularisation and unplanned non-target lesion revascularisation. **A** Up to 30 days ($p=0.394$) and **B** up to three years ($p<0.001$). The cumulative incidence of clinical events was calculated using the Kaplan-Meier method and compared using the log-rank test. CD-TLR: clinically driven target lesion revascularisation.
Predictors of both TLR and non-TLR were pre-procedure diameter stenosis (%), history of diabetes, and prior PCI. Notably, the degree of pre-procedural diameter stenosis (%) was positively correlated with TLR and negatively correlated with non-TLR. Predictors of TLR only were prior CABG, in-stent restenosis, age (years) (lower age was associated with higher risk for TLR), history of hypertension, and pre-procedure RVD (mm) (lower pre-procedure RVD was associated with higher risk for TLR). Predictors of non-TLR

Table 1. Baseline characteristics of patients with clinically driven target lesion revascularisation (TLR), unplanned non-target lesion revascularisation (Non-TLR), and no revascularisation at three years of follow-up.

<table>
<thead>
<tr>
<th></th>
<th>TLR (N=186)</th>
<th>TLR vs. no revascularisation p-value</th>
<th>Non-TLR (N=618)</th>
<th>Non-TLR vs. no revascularisation p-value</th>
<th>No revascularisation (N=5,333)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>63.2±11.8</td>
<td>0.348</td>
<td>64.4±10.8</td>
<td>0.260</td>
<td>63.9±10.9</td>
</tr>
<tr>
<td>Male</td>
<td>78.0% (145/186)</td>
<td>0.437</td>
<td>75.1% (464/618)</td>
<td>0.922</td>
<td>75.3% (4,015/5,333)</td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>28.3±4.8 (186)</td>
<td>0.997</td>
<td>28.3±5.3 (618)</td>
<td>0.008</td>
<td>28.3±5.0</td>
</tr>
<tr>
<td>Prior MI</td>
<td>33.3% (61/183)</td>
<td>0.033</td>
<td>31.0% (190/612)</td>
<td>0.012</td>
<td>26.2% (1,380/5,268)</td>
</tr>
<tr>
<td>Prior PCI</td>
<td>43.0% (80/186)</td>
<td>&lt;0.001</td>
<td>40.0% (247/618)</td>
<td>&lt;0.001</td>
<td>29.1% (1,554/5,333)</td>
</tr>
<tr>
<td>Ejection fraction (%)</td>
<td>59.0±10.7 (65)</td>
<td>0.171</td>
<td>56.5±10.9 (285)</td>
<td>0.294</td>
<td>57.2±10.7 (2,173)</td>
</tr>
<tr>
<td>Prior CABG</td>
<td>18.8% (35/186)</td>
<td>&lt;0.001</td>
<td>11.8% (73/618)</td>
<td>0.001</td>
<td>7.8% (415/5,333)</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>37.6% (70/186)</td>
<td>0.003</td>
<td>35.8% (221/618)</td>
<td>0.001</td>
<td>27.3% (1,458/5,333)</td>
</tr>
<tr>
<td>Insulin-dependent</td>
<td>12.4% (23/186)</td>
<td>0.027</td>
<td>11.2% (69/618)</td>
<td>0.005</td>
<td>7.8% (414/5,333)</td>
</tr>
<tr>
<td>History of hypertension</td>
<td>82.3% (153/186)</td>
<td>0.002</td>
<td>76.9% (475/618)</td>
<td>0.013</td>
<td>72.1% (3,846/5,333)</td>
</tr>
<tr>
<td>History of hyperlipidaemia</td>
<td>78.5% (146/186)</td>
<td>0.007</td>
<td>79.1% (489/618)</td>
<td>&lt;0.001</td>
<td>69.3% (3,698/5,333)</td>
</tr>
<tr>
<td>Current smoker</td>
<td>24.2% (45/186)</td>
<td>&gt;0.99</td>
<td>22.8% (141/618)</td>
<td>0.486</td>
<td>24.1% (1,286/5,333)</td>
</tr>
<tr>
<td>Multivessel disease (&gt;50% stenosis)</td>
<td>54.8% (89/165)</td>
<td>0.142</td>
<td>63.9% (276/432)</td>
<td>&lt;0.001</td>
<td>47.1% (1,553/3,000)</td>
</tr>
<tr>
<td>Reason for revascularisation</td>
<td>Stable angina</td>
<td>44.1% (82/186)</td>
<td>0.076</td>
<td>41.7% (258/618)</td>
<td>0.040</td>
</tr>
<tr>
<td></td>
<td>Unstable angina</td>
<td>26.9% (50/186)</td>
<td>0.435</td>
<td>25.6% (158/618)</td>
<td>0.521</td>
</tr>
<tr>
<td></td>
<td>MI</td>
<td>23.1% (41/186)</td>
<td>0.606</td>
<td>20.6% (127/618)</td>
<td>0.014</td>
</tr>
</tbody>
</table>

Results presented as either mean±SD (n/N) or percent (n/N). CABG: coronary artery bypass graft surgery; MI: myocardial infarction; PCI: percutaneous coronary intervention

Table 2. Lesion characteristics of patients with clinically driven target lesion revascularisation (TLR), unplanned non-target lesion revascularisation (Non-TLR), and no revascularisation at three years of follow-up.

<table>
<thead>
<tr>
<th></th>
<th>TLR (N=186)</th>
<th>TLR vs. no revascularisation p-value</th>
<th>Non-TLR (N=618)</th>
<th>Non-TLR vs. no revascularisation p-value</th>
<th>No revascularisation (N=5,333)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vessel location</td>
<td>LAD</td>
<td>51.1% (95/186)</td>
<td>0.882</td>
<td>40.5% (250/618)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>LCX</td>
<td>24.7% (46/186)</td>
<td>0.089</td>
<td>28.8% (178/618)</td>
<td>0.333</td>
</tr>
<tr>
<td></td>
<td>RCA</td>
<td>38.7% (70/186)</td>
<td>0.240</td>
<td>38.2% (236/618)</td>
<td>0.068</td>
</tr>
<tr>
<td></td>
<td>LMCA</td>
<td>2.7% (5/186)</td>
<td>0.425</td>
<td>0.8% (5/618)</td>
<td>0.040</td>
</tr>
<tr>
<td>Lesion length (mm)</td>
<td>15.4±12.5 (243)</td>
<td>0.444</td>
<td>15.7±10.5 (711)</td>
<td>0.048</td>
<td>15.0±9.0 (6,534)</td>
</tr>
<tr>
<td>Prior thrombus</td>
<td>6.3% (16/254)</td>
<td>0.802</td>
<td>4.2% (31/740)</td>
<td>0.003</td>
<td>7.0% (481/6,844)</td>
</tr>
<tr>
<td>Moderate/severe calcification</td>
<td>30.3% (79/261)</td>
<td>0.678</td>
<td>32.2% (244/757)</td>
<td>0.078</td>
<td>29.1% (2,032/6,981)</td>
</tr>
<tr>
<td>Prior TIMI flow 3</td>
<td>76.2% (202/265)</td>
<td>0.403</td>
<td>81.3% (621/764)</td>
<td>0.077</td>
<td>78.5% (5,525/7,041)</td>
</tr>
<tr>
<td>Total occlusion</td>
<td>8.3% (22/265)</td>
<td>0.550</td>
<td>7.9% (60/764)</td>
<td>0.611</td>
<td>7.4% (519/7,041)</td>
</tr>
<tr>
<td>Pre-procedure diameter stenosis (%)</td>
<td>75.4%±17.2% (264)</td>
<td>0.058</td>
<td>72.6%±17.1% (760)</td>
<td>0.285</td>
<td>73.3%±17.4% (6,991)</td>
</tr>
<tr>
<td>Prior ISR</td>
<td>18.4% (26/141)</td>
<td>&lt;0.001</td>
<td>8.7% (38/439)</td>
<td>0.139</td>
<td>6.8% (282/4,162)</td>
</tr>
<tr>
<td>Pre-procedure RVD (mm)</td>
<td>2.7±0.5 (243)</td>
<td>0.145</td>
<td>2.7±0.5 (711)</td>
<td>0.083</td>
<td>2.8±0.5 (6,559)</td>
</tr>
<tr>
<td>Rutherford</td>
<td>19.9% (28/141)</td>
<td>0.914</td>
<td>19.8% (87/439)</td>
<td>0.899</td>
<td>19.6% (814/4,162)</td>
</tr>
<tr>
<td>Modified ACC/AHA lesion class B2/C</td>
<td>67.9% (108/265)</td>
<td>0.893</td>
<td>69.2% (529/764)</td>
<td>0.652</td>
<td>68.4% (4,817/7,043)</td>
</tr>
</tbody>
</table>

Results presented as either mean±SD (n/N) or percent (n/N). ISR: in-stent restenosis; LAD: left anterior descending; LCX: left circumflex artery; LMCA: left main coronary artery; RCA: right coronary artery; RVD: reference vessel diameter; TIMI: Thrombolysis In Myocardial Infarction
only were lesion location (non-left anterior descending [LAD]) (LAD was associated with a lower risk of non-TLR), and history of hyperlipidaemia.

Discussion

The principal finding of our study was that non-TLR occurred more frequently than TLR throughout the three-year observation period of our patient cohort, which comprised more than 6,100 predominantly “all-comer” PCI patients treated with current-generation DES. Nearly 11% of the patients had clinical plaque progression requiring non-TLR at three years, which was approximately three times higher than the observed TLR rate. However, non-TLR, particularly those events that occurred shortly after the index procedure, could be overestimated as some cases might have been due to incomplete coverage during revascularisation of a diffuse lesion.

Non-TLR was infrequently described in the balloon angioplasty and bare metal stent eras, because restenosis rates were high, observation periods were short, and less complex patients were included in PCI studies. Our investigation with modern DES unmasks the true burden of unplanned revascularisations in the PCI landscape.

Other studies have also examined the incidence of TLR vs. non-TLR after PCI. PROSPECT (Providing Regional Observations to Study Predictors of Events in the Coronary Tree) evaluated originally treated (culprit) and untreated (non-culprit) lesion-related adverse events among 697 patients with acute coronary syndromes who successfully underwent PCI during a follow-up of 3.4 years. The study reported revascularisation rates in the treated (10.9%) vs. non-treated (10.5%) lesions at three years of follow-up. Additionally, the EVENT (Evaluation of Drug-Eluting Stents and Ischemic Events) registry included 10,144 patients, of whom 1,207 underwent repeat PCI, and found a similar rate of unplanned TLR vs. unplanned non-TLR at one year (4.5% vs. 4.4%, respectively). In an analysis in SIRIUS (Sirolimus-Eluting Stent in De Novo Native Coronary Lesions) with sirolimus-eluting stents (SES) at five years, TLR was 12.5%, TVR-non TL was 11.7%, and non-target vessel revascularisation was 22.3%. Thus, the ratio of TLR to non-TLR (TVR-non TL and non-target vessel revascularisation) was roughly 1:3 at five years.

TIME COURSE OF REVASCULARISATION

The current analysis was unique from PROSPECT and EVENT in that the Kaplan-Meier curves of TLR and non-TLR in our analysis separated early after the index procedure and non-TLR became more frequent than TLR as early as 30 days after the procedure. Such an early dissociation of curves is unexpected, as one would expect that they would separate later. The curves further divided...
over time, with a slower proportional increase of TLR than non-TLR after one year. Obviously, plaque progression in non-target lesions occurred in a continuous manner, while pathological processes in the target lesions cooled down over time. The risk of TLR was highest in the first year. Afterwards, the event rate for TLR was very low (roughly 0.7% per year). The fact that TLR is most common in the first year after PCI is seen across DES studies20–22. In EVENT, the risk of non-TLR was constant over time, while the hazard ratio for TLR was highest between two to nine months after PCI14. This trend in the first year could be due to trauma to the vessel during stent delivery.

**PREDICTORS OF REVASCULARISATION**

For TLR, features of lesion morphology – such as the presence of restenotic lesions, a smaller reference vessel diameter and larger pre-procedure diameter stenosis – were identified as independent predictors of repeat TLR. Moreover, previous revascularisations with either prior PCI or CABG were independent predictors of TLR. With respect to clinical characteristics, hypertension and diabetes mellitus, in addition to age, continue to represent significant predictors of TLR. Predictors of non-TLR included lesion characteristics (smaller pre-procedure diameter stenosis, and non-LAD lesion location), and clinical characteristics (history of diabetes, hyperlipidaemia, and prior PCI). Diabetes mellitus (along with prior PCI and pre-procedure diameter stenosis) was a predictor of both TLR and non-TLR. Given the prevalence of diabetes mellitus and its clinical significance, analysis on subjects with diabetes mellitus was a pre-specified secondary objective in the RESOLUTE Global Clinical Program15.

Other studies are available on predictors of TLR and non-TLR among patients treated with coronary stent implantation. Similar to this analysis, the EVENT registry found predictors of TLR to include age, prior PCI, and minimum stent diameter (which is similar to our finding of pre-procedure RVD)19. EVENT also found predictors of TLR to include sex, smoking status, prior MI, PCI of the left main or saphenous vein graft, total stent length, and use of a bare metal stent (18% of patients did not receive a DES but only BMS)18. Predictors of non-TLR in EVENT were different from this study and included age, number of diseased coronary arteries, and vein graft PCI14. Similar to our study, in PROSPECT, insulin-dependent diabetes and prior PCI were independent predictors of MACE related to non-culprit lesions, as well as plaque burden, thin-cap fibroatheroma, and a small minimal lumen area, which were not examined in our study17. Our results are consistent with the PROSPECT findings with respect to the impact of diabetes and prior PCI on any revascularisation. The TAXUS pooled programme also looked at predictors of TLR and non-TL TVR in years two to five. Predictors of TLR were (similar to this analysis) younger age, and pre-procedure vessel diameter and (different from this analysis) lesion length24. Predictors of non-TL TVR were diabetes (similar to this analysis), and hypertension treatment (different from this analysis)24. We identified several additional predictors of disease progression beyond the target lesion, such as hyperlipidaemia, target lesion location, diameter stenosis, and a trend for coronary calcification at baseline. In addition to baseline metabolic conditions, angiographic complexity plays a role in the progression of coronary artery disease. Coronary calcification has recently been identified as a strong indicator of adverse outcomes of PCI patients both in acute coronary syndromes and in the elective setting25,26. Interestingly, in our cohort, calcification in the target lesion had a stronger impact on non-TLR than on TLR. Another finding is the importance of the target lesion location. Interventions outside the LAD bear a higher risk of non-TLR. In other words, an untreated LAD is more often prone to revascularisations than the left circumflex and the right coronary artery combined. Two possible hypothetical explanations could be that LAD lesions progress more rapidly than non-LAD lesions, or that LAD lesions are more commonly underestimated and left untreated during the index procedure. Moreover, the LAD has such a large territory to supply that any progression has a clinical impact. The latter assumptions are underscored by fractional flow reserve studies27.

Additionally, a previous analysis by Taniwaki et al. on predictors of TLR and predictors of all revascularisation was conducted from the RESOLUTE All Comers study15. Similar to our study, predictors of TLR were age, insulin-treated diabetes, in-stent restenosis, and CABG15. Our analysis probably benefited from the larger sample size of 6,137 patients and the clear differentiation between TLR and non-TLR.

**Limitations**

The present analysis has several limitations. It was a post hoc analysis of trials not primarily intended to investigate coronary artery disease progression, and almost 40% of patients came from RESOLUTE International, which used visual estimation of baseline angiographic characteristics and was not 100% monitored. However, both TLR and any revascularisation were pre-specified secondary endpoints of the trial, and events were adjudicated by a clinical events committee which used definitions harmonised across the RESOLUTE Global Clinical Program. Furthermore, compliance to cardiovascular medication, apart from dual antiplatelet therapy, was not available and therefore it was not possible to evaluate the impact of compliance on restenosis and progression of coronary artery disease. In addition, if a patient had a TLR and a non-TLR, that patient was excluded from the study to sharpen differences between both groups of patients. This analytic approach underestimates the actual rate of TLR and non-TLR. Finally, we do not know whether the clinical indication for repeat revascularisation differed between TLR and non-TLR, as these data are not available.

**Conclusions**

The RESOLUTE Global Clinical Program was associated with a low TLR rate at three years. Given this low rate of TLR, the cumulative incidence of non-TLR in patients treated with second-generation DES was almost three times higher than TLR. Predictors of non-TLR were often different from predictors of TLR. Further improvements in PCI outcomes will largely depend on prospectively identifying ischaemia-producing lesions and intensifying preventive therapies.
Impact on daily practice

With contemporary drug-eluting stents, target lesion-related events after percutaneous coronary intervention (PCI) are markedly suppressed. The highly effective DES-based local therapy of coronary lesions is currently applied to a wide spectrum of patients, including those with more advanced coronary artery disease and complex anatomies. Consequently, revascularisation events that are not stent-related are prominent in the follow-up of contemporary PCI populations, and are three times higher than target lesion-related events. Therefore, further improvements in PCI outcomes will largely depend on prospectively identifying ischaemia-producing lesions and intensifying preventive therapies.

Guest Editor

This paper was guest edited by Rafael Beyar, MD, DSc, MPH; Director, Rambam Health Care Campus, Women’s Division/Dr Phillip and Sara Gotlieb Chair, Department of Medicine and Biomedical Engineering, Technion, Israel.

Acknowledgements

We thank Colleen Gilbert, PharmD, CMPP, Tim Peoples, MA, ELS, and Nicole Brilakis, MS, MBA, for editorial assistance, and Minglei Liu, PhD, and Yun Peng, MS, for statistical analysis and oversight (all from Medtronic).

Funding

This analysis and the studies contributing to it were sponsored by Medtronic, Inc.

Conflict of interest statement

M. Abdel-Wahab has received an institutional research grant from Medtronic and speaker honoraria from Boston Scientific. S. Silber has received grants from Medtronic. L. Mauri receives grants from Medtronic, Abbott Vascular, Boston Scientific, Eli Lilly/Daiichi Sankyo, Bristol Myers Squibb/Sanoﬁ-Aventis, and Biotronik. A. Yeung has served as an advisor to Medtronic. J. Belardi is a consultant for Medtronic and for Eli Lilly. P. Widimsky receives speaker honoraria from Medtronic. I. Meredith serves as a consultant to Boston Scientific and Medtronic. S. Saito is a speaker for Medtronic. Abbott Vascular, and Boston Scientific. G. Richardt has received an institutional research grant from Medtronic, and receives honoraria from Abbott Vascular for lectures and advisory board activities. The other authors have no conﬂicts of interest to declare. The guest editor has no conﬂicts of interest to declare.

References


