Efficacy of Triple Anti-Platelet Therapy Including Cilostazol in Acute Myocardial Infarction Patients Undergoing Drug-Eluting Stent Implantation

Keun-Ho Park, MD, Myung Ho Jeong, MD, Min Goo Lee, MD, Jum Suk Ko, MD, Shin Eun Lee, MD, Won Yu Kang, MD, Soo Hyun Kim, MD, Doo Sun Sim, MD, Nam Sik Yoon, MD, Hyun Ju Youn, MD, Young Joon Hong, MD, Hyung Wook Park, MD, Ju Han Kim, MD, Won Yu Kang, MD, Soo Hyun Kim, MD, Doo Sun Sim, MD, Nam Sik Yoon, MD, Hyun Ju Youn, MD, Young Joon Hong, MD, Hyung Wook Park, MD, Ju Han Kim, MD, Young Joon Hong, MD, Hyung Wook Park, MD, Ju Han Kim, MD

The Heart Research Center of Chonnam National University Hospital, Cardiovascular Research Institute of Chonnam National University, Gwangju, Korea

ABSTRACT

Background and Objectives: Triple anti-platelet therapy is known to prevent restenosis after drug-eluting stent (DES) implantation. However, there is little available data concerning the efficacy of triple anti-platelet therapy for acute myocardial infarction (AMI). Subjects and Methods: We analyzed 528 consecutive patients with AMI undergoing DES implantation between Nov 2005 and Apr 2008. We compared clinical outcomes in the triple anti-platelet therapy group (group I, n=413: cilostazol combined with aspirin and clopidogrel for at least one month) and dual anti-platelet therapy group (group II, n=115: aspirin and clopidogrel). Results: There were no significant differences in baseline characteristics. However, ST elevation myocardial infarction (STEMI) and use of TAXUS® stents were more common (70.9% vs. 55.7%, p=0.002; 83.5% vs. 73.0%, p=0.011) in Group I. Group I had lower incidences of cardiac death, 6-month target lesion revascularization (TLR), and major adverse cardiac and cerebrovascular events (MACCE) compared to Group II (1.7% vs. 5.7%, p=0.002; 5.7% vs. 11.5%, 0.035; 7.9% vs. 16.0%, p=0.011). On subgroup analysis, the incidence of 6-month TLR was lower among patients with American College of Cardiology/American Heart Association (ACC/AHA) B2 or C lesions and non-STEMI (6.0% vs. 14.9%, p=0.012; 4.3% vs. 19.1%, p=0.002) in Group I as compared to those in Group II. The rates of bleeding complications were no different between the two groups. On multivariate analysis, Killip III or IV and triple anti-platelet therapy were independent predictors of 6-month MACCE (hazard ratio (HR)=3.382; 95% confidence interval (CI)=1.384-8.262, HR=0.436; 95% CI=0.203-0.933). Conclusion: Triple anti-platelet therapy is safe and efficacious, and it prevents TLR in patients with AMI, especially those with complex lesions and non-STEMIs. (Korean Circ J 2009;39:190-197)

KEY WORDS: Platelets; Drug-eluting stents; Myocardial infarction.

Introduction

Percutaneous coronary intervention (PCI) with drug-eluting stents (DES) is well known to reduce the re-stenosis rate to a significant extent compared to bare-metal stents (BMS) in patients with acute myocardial infarction (AMI). It has been reported, however, that DESs increase the incidence of stent thrombosis as time passes. The treatment guidelines from the American College of Cardiology/American Heart Association (ACC/AHA) recommend that double anti-platelet therapy with aspirin and clopidogrel be prescribed for at least 12 months in patients who have undergone DES implantation. Cilostazol selectively inhibits the action of phosphodiesterase type 3, and it thereby has various functions, such as anti-platelet, vasodilatory, anti-atherosclerotic, and anti-proliferative actions. Cilostazol is also known to suppress neointimal hyperplasia following stent implantation. Recent studies have shown that triple anti-platelet therapy including cilostazol reduces the occurrence of stent thrombosis compared to anti-platelet therapy based on aspirin and clopidogrel. Triple anti-platelet...
therapy including cilostazol has also been shown to significantly reduce restenosis and target lesion revascularization, particularly in patients with diabetes mellitus. To date, however, no studies have reported the effect of triple anti-platelet therapy in patients with AMI, a high-risk group for stent thrombosis and re-stenosis.

The aim of this study was to compare the effectiveness of triple anti-platelet therapy with cilostazol with that of standard dual anti-platelet therapy in patients with AMI who have undergone DES implantation.

**Subjects and Methods**

**Subjects**

We conducted a single-center, retrospective study of 528 AMI patients who underwent successful TAXUS® or Cypher® stent implantation at the Heart Center of Chonnam National University Hospital during a 30-month period (November 2005 to April 2008). The mean age of these patients was 61.6 ± 11.72 years, and 395 of them were men.

Patients who received cilostazol in addition to standard dual anti-platelet therapy following successful stent implantation (aspirin + clopidogrel + cilostazol) were assigned to Group I (n=413), and those who received the standard dual anti-platelet therapy (aspirin + clopidogrel) were assigned to Group II (n=115).

**Percutaneous coronary intervention and medical treatment**

In patients with AMI, emergency or early invasive treatments were determined based on patient status, according to the clinical decision of the operator. In interventional treatment, the femoral artery was punctured and a 6 or 7-Fr sheath was inserted. Following catheter insertion, the procedure was performed using a guide wire. DESs were implanted in cases in which coronary artery stenoses were present following balloon angioplasty. The type of DES was decided by the operator. Successful PCI was defined as a target vessel at the treatment site with antegrade thrombolysis in myocardial infarction-3 (TIMI-3) flow and angiographic residual stenosis less than 50% following stent implantation. Anti-platelet agents were administered to all patients prior to intervention: aspirin 300 mg and clopidogrel 300-600 mg. Standard post-intervention treatment was aspirin 100 mg for life and clopidogrel 75 mg for at least one year. In accordance with the subjective decision of each operator, cilostazol was given to patients at a daily dose of 200 mg for at least one month.

Other medical treatments, including angiotensin-converting enzyme inhibitors, beta-blockers, calcium channel blockers, and lipid-lowering agents, were also used in a non-restrictive manner based on the standard treatment regimen for patients with AMI.

**Clinical outcomes**

Patients were told to visit the outpatient clinic one month after discharge and every two to three months thereafter on a regular basis. In “lost to follow-up” cases, specialized personnel monitored the clinical course by means of a telephone call. We measured the incidences of target lesion revascularization (TLR), cardiac death, non-fatal myocardial infarction, coronary artery bypass graft, major adverse cardiac and cerebrovascular accident (MACCE), bleeding, and stent thrombosis in all patients within the 6-month period after the procedure. TLR was defined as any percutaneous revascularization performed on the treated segment. Myocardial infarction was defined as a rise or fall in troponins (with at least one value above the 99th percentile of the upper reference limit) together with evidence of myocardial ischemia with at least one of the following: symptoms of ischemia, ECG change indicative of new ischemia, and imaging evidence of new loss of viable myocardium or new regional wall motion abnormality. Stroke was defined as a brain lesion on MRI or CT within 72 hours after the onset of neurological deficits. Stent thrombosis was defined as definite or probable stent thrombosis according to the criteria of the Academic Research Consortium (ARC). Any death for which the cause was not determined to have a definitive non-cardiac origin was classified as a cardiac death. MACCE was defined as cardiac death, non-fatal myocardial infarction, stroke, stent thrombosis, coronary bypass surgery, or TLR. Bleeding rates were also defined based on the TIMI bleeding classification. TIMI major bleeding was defined as intracranial hemorrhage or bleeding, with a hemoglobin decrease of ≥ 5 g/dL or a hematocrit decrease of ≥ 15%.

If a bleeding site was found, then TIMI minor bleeding was defined as a hemoglobin decrease of > 3 g/dL or a hematocrit decrease of > 10%; if no site was found, then it was defined as a hemoglobin decrease of > 4 g/dL or a hematocrit decrease of > 12%. TIMI minimal bleeding was defined as any clinically overt sign of hemorrhage associated with a hemoglobin decrease of < 3 g/dL or a hematocrit decrease of 9%. Stent thrombosis was defined as definite or probable stent thrombosis according to the criteria of the ARC.

**Statistical analysis**

Continuous variables were expressed as means ± standard deviation (SD), and categorical variables were expressed as frequencies. An analysis of continuous variables was performed using Student’s t-test, and an analysis of categorical variables was performed using the chi-square test or Fisher’s exact test. Cox regression analysis was used to identify the factors affecting the incidence of MACCE following the implantation of DES in patients with AMI. All statistical analyses were performed using Statistical Package for Social Science (SPSS, SPSS Inc,
Chicago, IL, USA) for Windows, version 15.0. P < 0.05 were considered statistically significant.

**Results**

**Clinical characteristics**

In Group I, the mean cilostazol administration period was 107.6 ± 59.0 days, and 211 patients (51%) were prescribed the drug for at least 90 days.

The number of patients diagnosed with ST elevation myocardial infarction (STEMI) at the time of admission was 293 (70.9%) in Group I and 64 (55.7%) in Group II (p = 0.002). However, there were no significant differences in mean age, sex, hypertension, diabetes mellitus, smoking history, hyperlipidemia, family history of cardiovascular disease, or past history of angina pectoris between the two groups. There were also no significant differences in Killip class III or IV ratio, glomerular filtration rate (GFR), or left ventricular ejection fraction (Table 1).

**Coronary angiographic findings and percutaneous coronary intervention**

In our coronary angiography series, infarction-related disease was found in the left anterior descending coronary artery in 275 cases (52.1%), left circumflex artery in 80 cases (15.1%), right coronary artery in 164 cases (31.1%), and left main coronary artery in 9 cases (1.7%). There were no significant differences in these parameters between the two groups (Table 2). Single-vessel disease was seen in 62 (53.9%) patients in Group II and in 193 (46.7%) patients in Group I (p = NS). Triple-vessel disease or left main disease was seen in 104 (25.2%) patients in Group I and in 21 (18.3%) patients in Group II (p = NS). The number of cases in which the degree of blood flow was TIMI flow 0 was 185 (44.8%) in Group I and 42 (36.5%) in Group II. However, this difference was not statistically significant. TAXUS® was the predominant DES used in Group I (83.5% vs. 73.0%, p = 0.011), and Cypher® was the predominant DES used in Group II (16.5% vs. 27.0%, p = 0.011). There was no significant difference in the proportion of cases in which the stent was successfully placed between the two groups (99.8% vs. 98.9%, p = 0.258).

**Table 1. Baseline characteristics of triple (group I) and dual (group II) anti-platelet therapy**

| Variable                        | Group I (n=413) | Group II (n=115) | p     |
|--------------------------------|----------------|-----------------|-------|
| Age (years)                    | 61.8 ± 11.73   | 61.1 ± 11.71    | 0.586 |
| Gender, male (%)               | 306 (74.1)     | 89 (77.4)       | 0.471 |
| Hypertension (%)               | 193 (46.7)     | 45 (39.1)       | 0.147 |
| Diabetes (%)                   | 65 (15.7)      | 13 (11.3)       | 0.236 |
| Current smoker (%)             | 186 (45.0)     | 63 (54.8)       | 0.073 |
| Dyslipidemia (%)               | 17 (4.1)       | 5 (4.3)         | 0.912 |
| Family Hx of CAD (%)           | 20 (4.8)       | 7 (6.1)         | 0.592 |
| Hx of angina (%)               | 181 (43.8)     | 57 (49.6)       | 0.274 |
| Killip class (%)               |                |                 |       |
| I or II                        | 371 (89.8)     | 106 (92.2)      | 0.452 |
| III or IV                      | 42 (10.2)      | 9 (7.8)         | 0.452 |
| Creatinine clearance (mL/min)  | 70.6 ± 30.1    | 75.1 ± 34.1     | 0.175 |
| Ejection fraction (%)          | 55.6 ± 11.8    | 54.9 ± 11.0     | 0.571 |
| Diagnosis (%)                  |                |                 |       |
| STEMI                          | 293 (70.9)     | 64 (55.7)       | 0.002 |
| NSTEMI                         | 120 (29.1)     | 51 (44.3)       | 0.002 |

Hx of CAD: history of coronary artery disease, STEMI: ST elevation myocardial infarction, NSTEMI: non-ST elevation myocardial infarction

**Major adverse cardiac and cerebrovascular events**

Of the total patients, 511 were available for follow-up observation, excluding eight patients (2.0%) in Group I and nine patients (7.8%) in Group II (Table 3). Two hundred twenty-seven patients (55.5%) in Group I and 49 patients (45.0%) in Group II underwent a 6-month follow-up angiography (p = 0.05). During the 6-month follow-up period, non-fatal myocardial infarction developed in only one patient in Group II. ARC definite or probable stent thrombosis developed in ten patients (2.5%) in Group I and in three patients (2.8%) in Group II, but this difference was not statistically significant. Stroke developed in one patient each in Group I and in Group II (p = 0.372). One patient in Group II underwent coronary bypass surgery at two months following the procedure. Cardiac death occurred in seven patients (1.7%) in Group I and in six patients (5.7%) in Group II (p = 0.022). The cardiac death cases were as follows: three cases of stent thrombosis, one case of severe aortic valve stenosis, one case of aggravated heart failure, and two cases of cardiac death suspected on a telephone interview in Group I, and two cases of stent thrombosis and four cases of cardiac death suspected on a telephone interview in Group II. The incidence of TLR was significantly lower in Group I than it was in Group II (5.7% vs. 11.5%, p = 0.035). The incidence of MACCE within the 6-month period was...
also significantly lower in Group I than it was in Group II (7.9% vs. 16.0%, p = 0.011). Two months following discharge, subarachnoid hemorrhage developed in one patient in Group II. However, there were no significant differences in the incidences of TIMI major or minor bleeding between the two groups (Table 4).

**Subgroup analyses**

Subgroup analyses were performed based on such parameters as lesion severity, ST-segment elevation, and the type of DES used. In accordance with the ACC/AHA lesion classification, in the simple lesions (type A or B1), there were no significant differences in the clinical events between the two groups. However, in the complex lesions (type B2 or C), the incidences of TLR and MACCE were significantly lower in Group I than they were in Group II (6.0% vs. 14.9%, p = 0.012; 8.8% vs. 17.1%, p = 0.037) (Table 5).

In patients diagnosed with non-STEMI, the incidences of TLR and MACCE within a recent 6-month period were significantly lower in Group I than they were in Group II (4.3% vs. 19.1%, p = 0.002; 5.2% vs. 20.4%, p = 0.003). However, in patients diagnosed with STEMI, there were no significant differences between the two groups.

### Table 2. Coronary angiographic findings and procedural characteristics in triple (group I) and dual (group II) anti-platelet therapy group

|                      | Group I (n=413) | Group II (n=115) | p    |
|----------------------|-----------------|------------------|------|
| Infarct-related artery (%) |                 |                  |      |
| LAD                   | 209 (50.6)      | 66 (57.4)        | 0.198|
| LCX                   | 64 (15.5)       | 16 (13.9)        | 0.675|
| RCA                   | 134 (32.4)      | 30 (26.1)        | 0.192|
| LM                    | 6 (1.5)         | 3 (2.6)          | 0.417|
| Involved vessel number (%) |               |                  |      |
| Single vessel         | 193 (46.7)      | 62 (53.9)        | 0.173|
| Two vessel            | 116 (28.1)      | 32 (27.8)        | 0.956|
| Three vessel or LM disease | 104 (25.2)  | 21 (18.3)        | 0.123|
| ACC/AHA classification (%) |               |                  |      |
| A or B1               | 125 (30.3)      | 32 (27.8)        | 0.613|
| B2 or C               | 288 (69.7)      | 83 (72.2)        | 0.613|
| Pre-PCI TIMI flow grade (%) |           |                  |      |
| 0                     | 185 (44.8)      | 42 (36.5)        | 0.113|
| I                     | 23 (5.6)        | 11 (9.6)         | 0.123|
| II                    | 104 (25.2)      | 33 (28.7)        | 0.447|
| III                   | 101 (24.5)      | 29 (25.2)        | 0.867|
| Type of DES used (%)  |                 |                  |      |
| TAXUS®                | 345 (83.5)      | 84 (73.0)        | 0.011|
| Cypher®               | 68 (16.5)       | 31 (27.0)        | 0.011|
| Use of glycoprotein IIb/IIIa inhibitor | 142 (34.4)  | 42 (36.5)        | 0.670|
| Successful PCI (%)    | 401 (99.8)      | 92 (89.8)        | 0.258|

LAD: left anterior descending artery, LCX: left circumflex artery, RCA: right coronary artery, LM: left main, ACC/AHA: American College of Cardiology/American Heart Association, PCI: percutaneous coronary intervention, TIMI: Thrombolysis In Myocardial Infarction, DES: drug-eluting stent, LM: left main artery

### Table 3. Six-month clinical outcomes of triple (group I) and dual (group II) anti-platelet therapy group

|                      | Group I (n=405) | Group II (n=106) | p    |
|----------------------|-----------------|------------------|------|
| Cardiac death (%)    | 7 (1.7)         | 6 (5.7)          | 0.022|
| Non-cardiac death (%)| 0 (0.0)         | 1 (0.9)          | 0.207|
| CVA (%)              | 1 (0.2)         | 1 (0.9)          | 0.372|
| Stent thrombosis (%) | 11 (2.7)        | 3 (2.8)          | 0.999|
| Non-fatal AMI (%)    | 7 (1.7)         | 2 (1.9)          | 0.999|
| Bypass surgery (%)   | 0 (0.0)         | 1 (0.9)          | 0.207|
| TLR (%)              | 23 (5.7)        | 12 (11.3)        | 0.041|
| MACCE (%)            | 32 (7.9)        | 17 (16.0)        | 0.011|

AMI: acute myocardial infarction, CVA: cerebrovascular accident, TLR: target lesion revascularization, MACCE: major adverse cardiac and cerebrovascular events

### Table 4. Bleeding complications after triple (group I) and dual (group II) anti-platelet therapy

|                      | Group I (n=413) | Group II (n=115) | p    |
|----------------------|-----------------|------------------|------|
| Degree of bleeding according to TIMI criteria (%) |                 |                  |      |
| Major                | 0 (0.0)         | 1 (0.9)          | 0.218|
| Minor                | 0 (0.0)         | 0 (0.0)          | 1.000|
| Minimal              | 7 (1.7)         | 1 (0.9)          | 0.999|
| Large-sized hematoma ≥ 5 cm | 16 (3.9) | 8 (7.0)          | 0.160|
| Small-sized hematoma <5 cm | 38 (9.2) | 6 (5.2)          | 0.172|
| Need for blood transfusion | 19 (4.6) | 4 (3.5)          | 0.797|

TIMI: thrombolysis in myocardial infarction
**Table 5. Six-month clinical outcomes on subgroup analysis**

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Patients with ACC/AHA A or B1 lesions (%)    | N=122     | N=30      |       |
| Cardiac death                                | 0 (0.0)   | 3 (10.0)  | 0.007 |
| Non-cardiac death                            | 0 (0.0)   | 0 (0.0)   | 1.000 |
| CVA                                          | 0 (0.0)   | 1 (3.3)   | 0.197 |
| Stent thrombosis                             | 1 (0.8)   | 0 (0.0)   | 0.999 |
| Non-fatal AMI                                 | 1 (0.8)   | 0 (0.0)   | 0.999 |
| Bypass surgery                               | 0 (0.0)   | 0 (0.0)   | 1.000 |
| MACCE                                        | 7 (5.7)   | 4 (13.3)  | 0.229 |
| **Heart death (%)**                          | **194**   | **122**   |       |
| **CVA (%)**                                  | **1 (0.3)**| **1 (1.8)**|       |
| **Non-MACCE (%)**                            | **25 (8.8)**| **13 (17.1)**| 0.037 |

**NSTEMI group (%)**

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Cardiac death                                | 7 (2.5)   | 3 (3.9)   | 0.447 |
| Non-cardiac death                            | 0 (0.0)   | 1 (1.3)   | 0.212 |
| CVA                                          | 1 (0.4)   | 1 (0.0)   | 0.297 |
| Stent thrombosis                             | 9 (3.2)   | 3 (3.9)   | 0.724 |
| Non-fatal AMI                                 | 6 (2.1)   | 2 (2.6)   | 0.678 |
| Bypass surgery                               | 0 (0.0)   | 1 (1.3)   | 0.212 |
| MACCE                                        | 17 (6.0)  | 11 (14.9) | 0.022 |
| **Heart death (%)**                          | **25 (5.7)**| **15 (5.3)**|       |
| **CVA (%)**                                  | **1 (0.3)**| **1 (1.5)**|       |
| **Non-MACCE (%)**                            | **25 (8.8)**| **13 (17.1)**| 0.037 |

**STEMI group (%)**

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Cardiac death                                | 6 (2.1)   | 3 (5.3)   | 0.171 |
| Non-cardiac death                            | 0 (0.0)   | 0 (0.0)   | 1.000 |
| CVA                                          | 1 (0.3)   | 1 (1.8)   | 0.307 |
| Stent thrombosis                             | 8 (2.8)   | 1 (1.7)   | 0.999 |
| Non-fatal AMI                                 | 6 (2.1)   | 1 (1.8)   | 0.999 |
| Bypass surgery                               | 0 (0.0)   | 1 (1.8)   | 0.165 |
| MACCE                                        | 18 (6.2)  | 3 (5.3)   | 0.999 |
| **Heart death (%)**                          | **26 (9.0)**| **7 (12.3)**| 0.440 |
| **CVA (%)**                                  | **1 (0.3)**| **1 (1.8)**|       |
| **Non-MACCE (%)**                            | **26 (9.0)**| **7 (12.3)**| 0.440 |

**Use of Cypher® stent (%)**

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Cardiac death                                | 1 (1.5)   | 0 (0.0)   | 0.171 |
| Non-cardiac death                            | 0 (0.0)   | 0 (0.0)   | 1.000 |
| CVA                                          | 0 (0.0)   | 0 (0.0)   | 1.000 |
| Stent thrombosis                             | 3 (4.6)   | 1 (3.4)   | 0.999 |
| Non-fatal AMI                                 | 2 (3.1)   | 1 (3.4)   | 0.999 |
| Bypass surgery                               | 0 (0.0)   | 0 (0.0)   | 1.000 |
| MACCE                                        | 6 (9.2)   | 1 (3.4)   | 0.431 |
| **Heart death (%)**                          | **6 (1.8)**| **6 (1.8)**|       |
| **CVA (%)**                                  | **1 (0.3)**| **1 (1.8)**|       |
| **Non-MACCE (%)**                            | **6 (1.8)**| **6 (1.8)**|       |

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Cardiac death                                | 6 (1.8)   | 6 (7.8)   | 0.004 |
| Non-cardiac death                            | 0 (0.0)   | 1 (1.3)   | 0.185 |
| CVA                                          | 1 (0.3)   | 1 (1.8)   | 0.336 |

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Cardiac death                                | 6 (1.8)   | 6 (7.8)   | 0.004 |
| Non-cardiac death                            | 0 (0.0)   | 1 (1.3)   | 0.185 |
| CVA                                          | 1 (0.3)   | 1 (1.8)   | 0.336 |

**Table 5. Continued**

| Category                                      | Group I   | Group II  | p     |
|-----------------------------------------------|-----------|-----------|-------|
| Stent thrombosis                             | 7 (2.1)   | 2 (2.6)   | 0.674 |
| Non-fatal AMI                                 | 5 (1.5)   | 1 (1.3)   | 0.999 |
| Bypass surgery                               | 0 (0.0)   | 1 (1.3)   | 0.185 |
| TLR (%)                                      | 18 (5.3)  | 12 (15.6) | 0.002 |
| MACCE (%)                                    | 26 (7.6)  | 16 (20.8) | 0.001 |

ACC/AHA: American College of Cardiology/American Heart Association, CVA: cerebrovascular accident, AMI: acute myocardial infarction, TLR: target lesion revascularization, MACCE: major adverse cardiac and cerebrovascular events, NSTEMI: non-ST elevation myocardial infarction, STEMI: ST elevation myocardial infarction groups (Table 5).

In the Cypher® stent group, there were no significant differences in the incidences of clinical events between Groups I and II. However, in the TAXUS® stent group, the incidences of TLR and MACCE within a recent 6-month period were significantly lower in Group I than they were in Group II (5.3% vs. 15.6%, p=0.002; 7.6% vs. 20.8%, p=0.001) (Table 5).

### Independent predictors for major adverse cardiac and cerebrovascular events

Cox proportional regression analysis was performed to identify predictors of MACCE. Variables included age, sex, hypertension, diabetes mellitus, hyperlipidemia, a past history of smoking, a family history of cardiovascular disease, a past history of angina pectoris, left ventricular ejection fraction, creatinine clearance, Killip classification, classification of ACC/AHA lesions, ST-segment elevation, the number of diseased vessels, the type of DES used, and triple anti-platelet therapy. Independent predictors for MACCE in AMI patients undergoing DES implantation included Killip class III or IV (HR=3.382; 95% confidence interval (CI)=1.384-8.262, p=0.007) and triple anti-platelet therapy (HR=0.436; 95% CI=0.203-0.933, p=0.017) (Table 6).

### Discussion

This study was conducted in order to evaluate the effect of triple anti-platelet therapy following DES implantation in AMI patients. Our study showed that, although there were no significant differences in non-fatal myocardial infarction, stroke, or stent thrombosis between the group receiving dual anti-platelet therapy and the group receiving dual anti-platelet therapy, the incidences of cardiac death and TLR were significantly lower in the group receiving triple anti-platelet therapy. These findings indicate that triple anti-platelet therapy was effective in lowering the incidence of MACCE. It was particularly effective in lowering the incidence of TLR in patients with complex lesions and those diagnosed with non-STEMI.

Recent studies have shown that DES significantly reduces repeat target-vessel revascularization in patients with...
AMI and also significantly lowers the two-year mortality.\textsuperscript{13} However, varying opinions still exist regarding the use of DES in patients with AMI, as an off-label indication.\textsuperscript{14} The rate of in-stent restenosis is markedly decreased with DES compared to BMS. Some investigators have reported that DESs are relatively free from complications of stent thrombosis,\textsuperscript{10,15} but this opinion is still controversial. To avoid this complication, medical treatment plays a crucial role in addition to the development of novel stents\textsuperscript{16,17} and the development of technology for stent implantation.

In the setting of percutaneous intervention, dual anti-platelet agents including aspirin and clopidogrel can prevent the occurrence of acute or subacute stent thrombosis and have been reported to lower the incidence of major adverse cardiac events.\textsuperscript{18,19} Therefore, at present, the use of dual anti-platelet agents prior to or following the procedure has been accredited as the standard treatment.

Cilostazol selectively inhibits phosphodiesterase type 3, which is released from the platelet, and thereby raises the intracellular concentration of cyclic adenosine monophosphate (cAMP) and calcium. This leads to suppression of platelet aggregation and relaxation of vascular smooth muscle cells, which ultimately have a vasodilatory effect.\textsuperscript{20,21} Expression of the p53 gene is enhanced, and thereby the progression of the cell cycle is inhibited and apoptosis is induced. Hepatocyte growth factor (HGF) secretion is enhanced, and vascular endothelial cell function is improved. Hence, restenosis is improved following angioplasty.\textsuperscript{22}

This anti-proliferative effect plays a crucial role in reducing in-stent restenosis, along with the anti-platelet and vasodilatory effects. Therefore, triple anti-platelet therapy including cilostazol has been reported to significantly reduce in-stent restenosis in patients with diabetes mellitus or long lesions, who are at high risk of restenosis.\textsuperscript{9,10,23}

Myocardial infarction is also associated with coronary artery stenosis due to severe atherosclerosis or coronary occlusion due to atherosclerotic plaque rupture.\textsuperscript{24} Restenosis remains problematic following stent implantation. In this study, we showed that triple anti-platelet therapy including cilostazol reduced the incidence of TLR in patients with myocardial infarction (relative risk reduction: 50%). These findings indicate that cilostazol has an anti-proliferative effect, even in patients with myocardial infarction.

In our study, subgroup analysis showed that triple anti-platelet therapy significantly lowered the incidence of TLR in ACC/AHA type B2 or C lesions compared to ACC/AHA type A or B1 lesions, and in patients diagnosed with non-STEMI compared to patients diagnosed

### Table 6. Independent predictors of 6-month MACCE

| Variables                          | HR     | 95.0% CI  | p       |
|------------------------------------|--------|-----------|---------|
| Killip III or IV on admission      | 3.382  | 1.384–8.262 | 0.007  |
| Triple antiplatelet therapy        | 0.436  | 0.203–0.933 | 0.033  |
| Ejection fraction                  | 0.979  | 0.950–1.008 | 0.154  |
| Family Hx of CAD                   | 1.200  | 0.561–2.565 | 0.163  |
| Single vessel disease              | 0.596  | 0.273–1.299 | 0.193  |
| Diagnosed STEMI                    | 1.200  | 0.561–2.565 | 0.221  |
| ACC/AHA B2 or C lesions            | 1.586  | 0.728–3.455 | 0.245  |
| Current smoker                     | 1.347  | 0.532–3.412 | 0.390  |
| Female gender                      | 1.487  | 0.636–3.478 | 0.360  |
| Dyslipidemia                       | 0.392  | 0.046–3.315 | 0.390  |
| Diabetes                           | 1.347  | 0.532–3.412 | 0.529  |
| Use of TAXUS\textsuperscript{®} stent | 1.294  | 0.505–3.317 | 0.591  |
| Age                                | 0.990  | 0.952–1.029 | 0.599  |
| Hypertension                       | 0.829  | 0.394–1.747 | 0.623  |
| Creatinine clearance               | 0.998  | 0.998–1.013 | 0.827  |
| Three vessel or LM disease         | 0.917  | 0.398–2.114 | 0.840  |
| Hx of angina                       | 1.065  | 0.533–2.129 | 0.859  |

MACCE: major adverse cardiac and cerebrovascular events, HR: hazard ratio, CI: confidence interval, Hx of CAD: history of coronary artery disease, STEMI: ST elevation myocardial infarction, ACC/AHA: American College of Cardiology/American Heart Association, LM: left main artery.
with STEMI (Table 4). These findings indicate that the anti-proliferative effect of cilostazol is greater in more severe atherosclerotic lesions. This is also demonstrated by the fact that the incidence of TLR was decreased in NSTEMI (where atherosclerosis is more prevalent and there are more AC/AHA type B2 or C lesions) compared to STEMI (where there are more lesions with thrombotic total occlusion). However, this merits further study. The incidence of TLR was significantly lower in the TAXUS® stent group compared to the Cypher® stent group. This finding also merits further study, because the number of enrolled patients who had Cypher® stents was much smaller.

The incidence of stent thrombosis has been reported to increase in cases in which primary stent implantation is performed for AMI.6) Lee et al.10) reported that triple anti-platelet therapy including cilostazol significantly lowered the incidence of stent thrombosis following stent implantation. In our study, a total of 13 patients (2.5%) had definite or probable stent thrombosis. Of these, nine patients were diagnosed with STEMI and underwent DES implantation. There was no significant difference in the incidence of stent thrombosis between the triple anti-platelet therapy group and the dual anti-platelet therapy group (2.5% vs. 2.8%, p=0.738). Considering that the triple anti-platelet therapy group had a greater number of patients with STEMI who had a greater thrombotic burden, we believe triple anti-platelet therapy offers an effective treatment for the prevention of stent thrombosis.

Cilostazol therapy was not associated with any adverse effects, such as TIMI major hemorrhage. This implies that triple anti-platelet therapy following DES implantation is safe.

Our study was conducted in patients to whom cilostazol was administered for more than one month. In approximately 51% of the patients, however, cilostazol was administered for more than three months. A minimal treatment period of one month is not sufficient for determining if cilostazol is an effective drug. Hence, further large-scale studies are warranted to examine the appropriate period of cilostazol treatment.

The greatest limitation of our study is that it was conducted as a single-center, retrospective, non-randomized, comparative study. Furthermore, regular follow-up coronary angiography was performed in only approximately 53% of the patients. TLR were performed based on the operators’ judgment of regular follow-up coronary angiography, rather than on ischemic symptoms. Because of this, the possibility of selection bias cannot be completely ruled out. Intravascular ultrasonography, which is capable of assessing in-stent restenosis and neointimal hyperplasia more accurately, was not performed in all the patients.

The current study is significant, however, in that it included a relatively large number of patients and the incidence of TLR was further decreased. This occurred despite the presence of an "ocular-stenotic reflex" in the triple anti-platelet therapy group secondary to the higher prevalence of follow-up coronary angiography in this group.

In conclusion, triple anti-platelet therapy following DES implantation in AMI patients was safe and effective in reducing the occurrence of TLR and MACCE within six months, as compared to dual anti-platelet therapy. This reduction was particularly significant in patients with complex lesions and those diagnosed with non-STEMIs.

Acknowledgments

This study was supported by a grant of the Korea Healthcare technology R&D Project, Ministry for Health, Welfare and Family Affairs (A084869), Republic of Korea.

REFERENCES

1) Lee SR, Jeong MH, Ahn YK, et al. Clinical safety of drug-eluting stents in the Korea acute myocardial infarction registry. Circ J 2008;72:392-8.
2) Pasceri V, Patti G, Speciale G, Pristipino C, Richichi G, Di Sciascio G. Meta-analysis of clinical trials on use of drug-eluting stents for treatment of acute myocardial infarction. Am Heart J 2007;153:749-54.
3) Melkian N, Wijns W. Drug-eluting stents: a critique. Heart 2008; 94:145-52.
4) Antman EM, Hand M, Armstrong PW, et al. 2007 Focused update of the ACC/AHA 2004 guidelines for the management of patients with ST-elevation myocardial infarction: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines: developed in collaboration with the Canadian Cardiovascular Society endorsed by the American Academy of Family Physicians: 2007 writing group to review new evidence and update the ACC/AHA 2004 guidelines for the management of patients with ST-elevation myocardial infarction, writing on behalf of the 2004 writing committee. Circulation 2008;117:296-329.
5) Goto S. Cilostazol: potential mechanism of action for antithrombotic effects accompanied by a low rate of bleeding. Atheroscler Suppl 2005;6:3-11.
6) Douglas JS Jr. Role of adjunct pharmacologic therapy in the era of drug-eluting stents. Atheroscler Suppl 2005;6:47-52.
7) Douglas JS Jr, Holmes DR Jr, Kereakesi DJ, et al. Coronary stent restenosis in patients treated with cilostazol. Circulation 2005;112:2826-32.
8) Lee SW, Park SW, Hong MK, et al. Triple versus dual antiplatelet therapy after coronary stenting: impact on stent thrombosis. J Am Coll Cardiol 2005;46:1833-7.
9) Ahn Y, Jeong MH, Jeong JW, et al. Randomized comparison of cilostazol vs clopidogrel after drug-eluting stenting in diabetic patients: cilostazol for diabetic patients in drug-eluting stent (CIDES) trial. Circ J 2008;72:35-9.
10) Lee SW, Park SW, Kim YH, et al. Drug-eluting stenting followed by cilostazol treatment reduces late restenosis in patients with diabetes mellitus the DECLARE-DIABETES Trial (A Randomized Comparison of Triple Antiplatelet Therapy with Dual Antiplatelet Therapy After Drug-Eluting Stent Implantation in Diabetic Patients). J Am Coll Cardiol 2008;51:1181-7.
11) Mauri L, Hsieh WH, Massaro JM, Ho KK, D’Agostino R, Cutlip DE. Stent thrombosis in randomized clinical trials of drug-eluting stents. N Engl J Med 2007;356:1020-9.
12) Rao AK, Pratt C, Berke A, et al. Thrombolysis in Myocardial Infarction (TIMI) Trial phase I. hemorrhagic manifestations and changes in plasma fibrinogen and the fibrinolytic system in patients treated with recombinant tissue plasminogen activator and streptokinase. J Am Coll Cardiol 1988;11:1-11.
13) Mauri L, Silbaugh TS, Garg P, et al. Drug-eluting or bare-metal stents for acute myocardial infarction. N Engl J Med 2008;359:1330-42.
14) Nakazawa G, Finn AV, Joner M, et al. Delayed arterial healing and increased late stent thrombosis at culprit sites after drug-eluting stent placement for acute myocardial infarction: an autopsy study. Circulation 2008;118:1138-45.
15) Stone GW, Moses JW, Ellis SG, et al. Safety and efficacy of sirolimus- and paclitaxel-eluting coronary stents. N Engl J Med 2007;356:998-1008.
16) Kim W, Jeong MH, Kim KH, et al. The clinical results of a platelet glycoprotein IIb/IIIa receptor blocker (abciximab: ReoPro)-coated stent in acute myocardial infarction. J Am Coll Cardiol 2006;47:933-8.
17) Mitka M. New drug-eluting stents under study. JAMA 2007;297:2064-7.
18) Mehta SR, Yusuf S, Peters RJ, et al. Effects of pretreatment with clopidogrel and aspirin followed by long-term therapy in patients undergoing percutaneous coronary intervention: the PCI-CURE study. Lancet 2001;358:527-33.
19) Yusuf S, Zhao F, Mehta SR, Chrolavicius S, Tognoni G, Fox KK. Effects of clopidogrel in addition to aspirin in patients with acute coronary syndromes without ST-segment elevation. N Engl J Med 2001;345:494-502.
20) Tanaka T, Ishikawa T, Hagiwara M, Onoda K, Itoh H, Hidakah H. Effects of cilostazol, a selective cAMP phosphodiesterase inhibitor on the contraction of vascular smooth muscle. Pharmacology 1988;36:313-20.
21) Chu YH, Park SW, Lee CW, et al. Effects of cilostazol treatment on angiographic restenosis after coronary stent placement. Korean Circ J 2000;30:1494-500.
22) Ikeda Y, Kikuchi M, Murakami H, et al. Comparison of the inhibitory effects of cilostazol, acetylsalicylic acid and ticlopidine on platelet functions ex vivo: randomized, double-blind cross-over study. Arzneimittelforschung 1987;37:563-6.
23) Ahn JC, Song WH, Kwon JA, et al. Effects of cilostazol on platelet activation in coronary stenting patients who already treated with aspirin and clopidogrel. Korean J Intern Med 2004;19:230-6.
24) Kim SM, Kim DI, Cho HJ, et al. Effect of cilostazol on the drug-eluting stent in native coronary arteries. Korean Circ J 2007;37:304-11.
25) Takahashi S, Oida K, Fujiwara R, et al. Effect of cilostazol, a cyclic AMP phosphodiesterase inhibitor, on the proliferation of rat aortic smooth muscle cells in culture. J Cardiovasc Pharmacol 1992;20:900-6.
26) Lee KJ, Yun SW, Kim SW, Kim TH, Kim CI, Byu WS. The effect of cilostazol on proliferation of vascular smooth muscle cells and expression of iNOS and p21. Korean Cir J 2004;34:500-6.
27) Morishita R. A scientific rationale for the CREST trial results: evidence for the mechanism of action of cilostazol in restenosis. Atheroscler Suppl 2005;6:41-6.
28) Lee SW, Park SW, Kim YH, et al. Comparison of triple versus dual antiplatelet therapy after drug-eluting stent implantation (from the DECLARE-Long trial). Am J Cardiol 2007;100:1103-8.
29) Viles-Gonzalez JF, Fuster V, Badimon JJ. Atherothrombosis: a widespread disease with unpredictable and life-threatening consequences. Eur Heart J 2004;25:1197-207.
30) Ong AT, Hoye A, Aoki J, et al. Thirty-day incidence and six-month clinical outcome of thrombotic stent occlusion after bare-metal, sirolimus, or paclitaxel stent implantation. J Am Coll Cardiol 2005;45:947-53.