Effect of different loading doses of atorvastatin on percutaneous coronary intervention for acute coronary syndromes

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BACKGROUND: Percutaneous coronary intervention (PCI)-induced myocardial damage is associated with late cardiovascular events. Treatment with atorvastatin before PCI can reduce myocardial damage during the peri-PCI period.

OBJECTIVES: To compare the safety and myocardial effects of different atorvastatin loading doses and dosing frequency before PCI in non-ST segment elevation acute coronary syndrome (NSTE-ACS) patients.

METHODS: Eighty NSTE-ACS patients were randomly divided into four groups (20 patients per group). The control group was given 40 mg atorvastatin each night. The three loading dose groups were treated the same as in the control group, but were given 80 mg atorvastatin 12 h before PCI (low-load group) in combination with 40 mg atorvastatin 2 h to 4 h before PCI (mid-load group) or 60 mg atorvastatin 2 h to 4 h before PCI (high-load group). All patients underwent PCI within 48 h to 72 h of admission, and received 40 mg atorvastatin for at least one month after PCI. Changes in myocardial markers and highly sensitive C-reactive protein were analyzed.

RESULTS: No deaths or revascularizations were recorded. The incidences of MACE differed significantly between the four groups (40%, 25%, 10% and 0% for the control, low-load, mid-load and high-load groups, respectively; \( P<0.05 \)). The incidence of MACE and cardiac troponin I level above the normal range, and post-PCI increases in creatine kinase-MB and highly sensitive C-reactive protein were significantly higher in the control group than in the high-load group (all \( P<0.007 \)). The post-PCI alanine aminotransferase levels in all four groups were significantly higher than the pre-PCI levels, but were within normal ranges. No myalgia or myasthenia was observed.

CONCLUSION: The results of the present study show that short-term atorvastatin loading before PCI was well tolerated and had beneficial myocardial effects in patients with NSTE-ACS.

Key Words: Atorvastatin; Interventional therapy; Non-ST segment elevation acute coronary syndrome

Percutaneous coronary intervention (PCI) is now a commonly used revascularization treatment for coronary artery disease. However, although PCI restores blood flow to the heart, it may cause myocardial damage, which is associated with late cardiovascular events (1). Statin drugs, since their introduction in the 1980s, have become the first-choice lipid-lowering therapy and are useful in preventing coronary artery disease. Many studies have found that statins have beneficial pleiotropic cardiovascular effects in addition to their lipid-lowering effects. A number of clinical trials, including the ARMYDA (ARMYDA 1, 2, ARMYDA-Acute Coronary Syndromes (ARMYDA-ACS) and ARMYDA-RECAPTURE (4), have shown that treatment with loading atorvastatin before PCI can significantly reduce PCI-induced myocardial damage. However, no randomized controlled studies have investigated the effects of atorvastatin treatment (including loading dose and frequency) or whether short-term in-hospital atorvastatin treatment is effective for patients with non-ST segment elevation acute coronary syndromes (NSTE-ACS).
**METHODS**

Research subjects and study design

Research subjects were randomly recruited between October 2009 and March 2010 from patients at the Department of Cardiology of the First Affiliated Hospital of China Medical University (Shenyang, China). The study was approved by the ethical committee at The First Affiliated Hospital, and all patients provided informed consent. The patients were included if they presented with NSTE-ACS and were scheduled to undergo PCI within 48 h to 72 h of admission. Patients who met any of the following criteria were excluded: presented with ST segment elevation acute myocardial infarction; presented with high-risk NSTE-ACS requiring emergency PCI; were already on statin therapy or had a history of statin therapy in the previous two months; had a left ventricular ejection fraction below 30%; or had a history of hemorrhage, major surgery, severe renal/hepatic insufficiency, muscular disorders, infectious diseases, autoimmune diseases, a malignant tumour or other contraindications to statin therapy.

A total of 163 patients met the inclusion criteria, but 49 of these also met one or more exclusion criteria and were excluded, including 10 patients already treated with statins, eight with severe left ventricular dysfunction, 23 with severe hepatic or renal insufficiency, five with a history of hemorrhage or major surgery within the past 6 months, and three high-risk patients who required emergency PCI. Of the remaining 114 patients, four did not provide signed, informed consent and were excluded. After coronary angiography, 30 patients were excluded (17 subsequently underwent surgical bypass and 13 continued medication). Thus, a total of 80 patients who underwent stent implantation were enrolled as subjects in the present study. The patients were randomly divided into four groups (n=20 per group) using a random number table. The groups were a control group, a low-load group, a mid-load group and a high-load group. The control group was given 40 mg atorvastatin each night. The three loading dose groups were treated the same as in the control group, but were given 80 mg atorvastatin 12 h before PCI (low-load group) in combination with 40 mg atorvastatin 2 h to 4 h before PCI (mid-load group) or 60 mg atorvastatin 2 h to 4 h before PCI (high-load group). The operators and clinical staff involved in patient follow-up were blinded to the treatment assigned.

All subjects underwent standard PCI via radial artery puncture. All subjects were given oral acetylsalicylic acid (100 mg/day or 300 mg/day) and clopidogrel (75 mg/day) daily, with an accumulated dose of at least 300 mg before PCI. In some cases, glycoprotein IIb/IIIa receptor antagonists were administered during PCI as deemed necessary by the operators. The PCI operation was considered to be successful if the post-PCI residual stenosis was below 30%. After PCI, the patients were to be treated with low molecular weight heparin for three to five days, clopidogrel for at least 12 months, atorvastatin (40 mg every night) for at least one month, and acetylsalicylic acid for life. In addition, patients without contraindications were given beta-receptor antagonists, and angiotensin-converting enzyme inhibitors, antagonists or receptor blockers.

Venous blood samples were collected from each patient at three time points (before PCI, 8 h after PCI and 24 h after PCI). The creatine kinase-MB (CK-MB) level was assayed using a dry chemical method, cardiac troponin I (cTnI) by chemiluminescence, and highly sensitive C-reactive protein (hs-CRP) by scatter turbidimetry. The normal ranges before PCI, but increased beyond the normal range after PCI, or the pre-PCI cardiac marker levels in a patient were beyond the normal ranges and increased further after PCI.

The post-PCI hs-CRP level was higher than the pre-PCI level.

Statistical analyses

Continuous data are expressed as means ± SDs, and were analyzed by ANOVA. The least significant difference t test was used for multiple comparisons. Categorical data are expressed as n (%), and were analyzed by χ² tests. P<0.05 was considered to be statistically significant. Pairwise multiple comparisons were performed by χ² partitioning with a significance level (α′) of 0.007. All analyses were performed using SPSS version 14.0 (SPSS Inc, USA).

RESULTS

Clinical and procedural features in the four groups are reported in Tables 1 and 2. The four groups had similar baseline characteristics including age, sex, body mass index, history of disease, clinical conditions, left ventricular ejection fraction, in-hospital medication, angiographic features and nature of operation (all P>0.05). All patients were successfully treated by implantation of drug-eluting stents. No deaths or revascularizations were recorded during the follow-up, and the incidence of myocardial infarction was equivalent to the overall rate of MACE. The incidence of MACE within 30 days after PCI was significantly different among the four groups (40%, 25%, 10% and 0% for the control, low-load, mid-load and high-load groups, respectively; P<0.05) and was significantly lower in the high-load group than in the control group (P<0.007). The differences between other pairs of groups were not statistically significant, but the incidence of MACE showed a decreasing trend with increasing loading dose (Table 3).

Before PCI, the proportions of patients with levels of myocardial markers above the upper limit of the normal range were similar in the four groups. After PCI, this proportion was higher than the pre-PCI value in the control group (P<0.05) but remained similar to the pre-PCI value in the three groups that received a loading dose of atorvastatin. In addition, the four groups showed statistically significant differences in the proportion of patients with post-PCI cTnI levels above the upper limit of the normal range (80%, 60%, 45% and 35% for the control, low-load, mid-load and high-load groups, respectively; P<0.05). Compared with the control group, the proportion of patients with post-PCI CK-MB or cTnI levels above their respective upper limits was significantly lower in the high-load group (P<0.007). The proportion of patients with post-PCI levels of myocardial markers above the upper limit of the normal range decreased with increasing frequency and dose of atorvastatin treatment (Table 4). The proportion of patients with above-baseline levels of hs-CRP post-PCI was not significantly different among the four groups (75%, 60%, 50% and 25%, respectively; P>0.05), although the proportion was significantly lower in the high-load dose group than in the control group based on χ² partitioning (P<0.007).

Compared with the pre-PCI level, the post-PCI CK-MB level did not significantly increase in the high-load group. Meanwhile, the CK-MB levels in the mid-load and high-load groups were significantly lower than that in the control group. The pre- and post-PCI cTnI levels were similar in the mid-load and high-load groups, and were statistically lower in the high-load group than in the control group. The post-PCI hs-CRP levels in all four groups were significantly higher than their respective pre-PCI levels, and were significantly lower in the high-load groups than in the control and low-load groups. After PCI, the alanine aminotransferase (ALT) levels in all four groups were significantly higher than their respective pre-PCI levels (all P<0.05), but remained within the normal range, except in one patient in the high-load group, and there were no significant differences among the four groups (P>0.05) (Table 5).
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No gastrointestinal reactions (eg, nausea and vomiting), myalgia or myasthenia were observed. One patient in the high-load group had an ALT level beyond the upper limit of the normal range, but by less than 300%. In this patient, atorvastatin treatment was continued and, after treatment with diisopropylamine dichloroacetate, the ALT level decreased to within the normal range.

**DISCUSSION**

PCI-induced myocardial damage does not result in clear clinical symptoms, changes in electrocardiogram readings or effects on cardiac functions; instead, such damage may only appear as an increase in myocardial marker levels. However, such damage is associated with increased rates of MACE (6). Many recent studies have shown that short-term loading of statins has beneficial pleiotropic cardiovascular effects, which are independent of their lipid-lowering effects (7,8) because statins can only lower the lipid level with long-term treatment. The effects of statins include protection of the endothelium (9), dilation of coronary microvessels (10), reduction of thrombogenesis (11), antiplatelet and anti-inflammatory capabilities (12), and significant lowering of post-PCI intercellular adhesion molecule-1 and E-selectin levels (13). Animal studies have also shown that acute statin loading before reperfusion reduced the area of myocardial infarction (14).

Clinical trials, including Myocardial Ischemia Reduction with Aggressive Cholesterol Lowering (MIRACL) (15) and Pravastatin or Atorvastatin Evaluation and Infection Therapy (PROVE-IT) (16), have confirmed that early treatment of ACS patients with high-dose statins can reduce the incidence of cardiovascular events.

**TABLE 1**

Main clinical features of patients in the four groups

| Variable                        | Control group (n=20) | Low-load group (n=20) | Mid-load group (n=20) | High-load group (n=20) |
|---------------------------------|----------------------|-----------------------|-----------------------|------------------------|
| Age, years                      | 60.80±7.52          | 61.65±8.24            | 61.90±6.40            | 61.40±6.82             |
| Male sex                        | 12 (60)             | 14 (70)               | 13 (65)               | 11 (55)                |
| Body mass index, kg/m²          | 23.98±2.31          | 24.26±2.27            | 23.96±2.40            | 24.50±2.40             |
| Diabetes mellitus               | 4 (20)              | 5 (25)                | 6 (30)                | 6 (30)                 |
| Hypertension                    | 12 (60)             | 10 (50)               | 11 (55)               | 12 (60)                |
| Smokers                         | 6 (30)              | 7 (35)                | 8 (40)                | 8 (40)                 |
| Previous myocardial infarction  | 1 (5)               | 0 (0)                 | 0 (0)                 | 0 (0)                  |
| Previous coronary intervention  | 2 (10)              | 0 (0)                 | 1 (5)                 | 0 (0)                  |
| Previous bypass surgery         | 0 (0)               | 0 (0)                 | 0 (0)                 | 0 (0)                  |
| LVEF, %                         | 56.00±9.75          | 58.95±7.29            | 55.25±8.41            | 59.85±4.68             |
| LDL-C, mmol/L                   | 2.74±0.93           | 2.83±1.09             | 2.82±0.63             | 2.78±0.63              |
| Triglycerides, mmol/L           | 2.00±1.25           | 1.96±0.93             | 1.82±0.67             | 1.90±0.90              |
| Cholesterol, mmol/L             | 4.94±1.03           | 4.97±1.32             | 4.66±0.98             | 5.07±1.40              |
| HDL-C, mmol/L                   | 1.19±0.47           | 1.35±0.56             | 1.30±0.56             | 1.18±0.35              |
| Creatinine, mg/L                | 78.95±17.17         | 79.2±18.21            | 81.40±13.24           | 72.00±16.99            |

Other medical therapy

- Acetylsalicylic acid: 20 (100)
- Clopidogrel: 20 (100)
- Beta-blockers: 14 (70)
- ACEI/ARB: 15 (75)
- GP IIb/IIIa receptor inhibitors: 2 (10)
- Unstable angina: 17 (85)
- NSTEMI: 3 (15)
- Multivessel coronary disease: 6 (30)

Data presented as n (%) or mean ± SD. ACEI/ARB Angiotensin-converting enzyme inhibitor or angiotensin receptor blocker; GP Platelet glycoprotein; HDL-C High-density lipoprotein cholesterol; LDL-C Low-density lipoprotein cholesterol; LVEF Left ventricular ejection fraction; NSTEMI Non-ST segment elevation myocardial infarction

**TABLE 2**

Procedural features in the four groups

| Variable                          | Control group (n=20) | Low-load group (n=20) | Mid-load group (n=20) | High-load group (n=20) |
|-----------------------------------|----------------------|-----------------------|-----------------------|------------------------|
| Vessels treated                   |                      |                       |                       |                        |
| Left main artery                  | 0 (0)                | 0 (0)                 | 0 (0)                 | 0 (0)                  |
| Left anterior descending artery   | 8 (40)               | 10 (50)               | 8 (40)                | 7 (35)                 |
| Left circumflex artery            | 6 (30)               | 7 (35)                | 7 (35)                | 9 (45)                 |
| Right coronary artery             | 10 (50)              | 7 (35)                | 8 (40)                | 9 (45)                 |
| Multivessel intervention          | 4 (20)               | 4 (20)                | 3 (15)                | 5 (25)                 |
| Number of stents                  | 1.60±0.68            | 1.50±0.69             | 1.45±0.60             | 1.55±0.76              |
| Stent diameter, mm                | 4.89±1.90            | 4.61±1.96             | 4.38±1.71             | 4.50±2.57              |
| Total length, mm                  | 29.50±13.74          | 32.10±13.55           | 31.00±14.28           | 29.25±14.48            |
| Stent deployment pressure, atm    | 21.05±11.05          | 21.00±11.49           | 21.15±11.37           | 22.70±11.07            |
| Duration of stent deployment, s   | 8.00±3.40            | 7.50±3.44             | 7.25±3.02             | 7.75±3.80              |
| Use of postdilatation procedure   | 2 (10)               | 2 (10)                | 3 (15)                | 1 (5)                  |

Data presented as n (%) or mean ± SD
In the ARMYDA-ACS trial (3), patients with NSTE-ACS who underwent PCI within 48 h of admission were treated with atorvastatin (80 mg and 40 mg atorvastatin at 12 h and 2 h before PCI, respectively) or placebo. The results revealed that pre-PCI loading with atorvastatin reduced the incidence of myocardial damage. The subsequent ARMYDA-RECAPTURE trial (4) studied the effect of atorvastatin in patients with stable angina or NSTE-ACS. In that study, patients underwent PCI after treatment with statins for at least 30 days, and were given atorvastatin (80 mg and 40 mg at 12 h and 2 h before PCI, respectively) or placebo. Similarly, that trial found that pre-PCI treatment with atorvastatin significantly reduced myocardial damage.

In China, most NSTE-ACS patients only start statin therapy during an in-hospital stay and undergo PCI after appropriate examinations (generally two to three days after admission). The optimum dose and frequency of atorvastatin treatment in this patient group have not been formally evaluated. In the present study, we compared the myocardial effects of short-term atorvastatin treatment before PCI at four loading doses. Our findings confirm that, in these patients, the incidence of myocardial damage and inflammatory reactions decreased with increasing dose and frequency of atorvastatin administration. Our findings differ from those of the ARMYDA-RECAPTURE trial in that the mid-load dose (80 mg atorvastatin 12 h before PCI plus 40 mg atorvastatin 2 h to 4 h before PCI) was associated with a lower incidence of MACE compared with the control group (although the difference was not statistically significant in our study), and that the high-load dose (80 mg atorvastatin 12 h before PCI plus 60 mg atorvastatin 2 h to 4 h before PCI) seemed to be more effective for this patient group. The differences may be attributed to the length of atorvastatin treatment before PCI and the sample size. Nevertheless, dose-dependent antiplatelet and anti-inflammatory effects of atorvastatin have been demonstrated (17). For ACS patients, the thrombogenesis and inflammatory reactions are closely related to the PCI-related myocardial damages, and our results indicate that a high dose of atorvastatin should be given before PCI.

The incidence of transaminase elevations during statin therapy was reported to be approximately 1% to 2%. These elevations usually occurred within the first three months of treatment in a dose-dependent manner, and could be restored to the baseline level by reducing the statin dose or discontinuing statin treatment (18). The maximum short-term pre-PCI loading dose was 120 mg in the ARMYDA studies, and no clear adverse reactions were observed. Similarly, the study by Ge et al (19) showed that 80 mg (12 h pre-PCI) and 40 mg (2 h pre-PCI) of atorvastatin was well tolerated in Asian patients with NSTE-ACS and reduced the incidence of MACE. However, other studies (20) do not support high doses of statins, particularly rosuvastatin, in Asian populations. The doses currently recommended for statins are effective only for lowering blood lipid levels, and the appropriate dose for myocardial protection during the peri-PCI period remains unclear. Therefore, in the present study, we explored the effects of higher loading doses on preventing myocardial damage. The highest pre-PCI dose was 140 mg in the high-load group (80 mg 12 h before PCI plus 60 mg 2 h to 4 h before PCI).
We also evaluated the benefits and risks. Significant elevations in transaminase levels occurred after PCI in all four groups, but with similar magnitudes, and were within the upper limit of the normal range. No patient experienced myalgia or myasthenia. These observations indicate that high-dose pre-PCI atorvastatin therapy may temporarily increase the workload on the liver and result in a transient elevation of the transaminase level, but this elevation can be reversed by discontinuing the atorvastatin loading, suggesting that the benefits outweigh the potential risks.

In summary, our study revealed that pre-PCI loading with statins may reduce the incidence of PCI-induced myocardial damage, and that these beneficial myocardial effects increased with increasing dose and frequency of statin loading. This loading treatment might be associated with relatively mild adverse reactions and be well tolerated. The present study suggests that different pre-PCI loading protocols may be used clinically depending on the specific conditions of the patient. The limitations of the present study include a relatively small sample size and the discontinuous observation of indicators after PCI. Larger-scale controlled, randomized trials with extended monitoring of PCI indexes are needed to confirm the clinical efficacy of this approach.

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