Induction of Diabetes Abolishes the Antithrombotic Effect of Clopidogrel in Apolipoprotein E–Deficient Mice

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TH Open 2017;1:e92–e100.

Abstract

Patients with acute coronary syndrome with diabetes mellitus (DM) exhibit an impaired platelet inhibitory response to clopidogrel which is only partially understood. DM was induced by the administration of streptozotocin (STZ) to 9-week-old mice. The antithrombotic effects of clopidogrel (10 mg/kg/d, orally × 5 days) were determined using a FeCl3-induced thrombosis model employing wild-type (WT), apolipoprotein E (apoE)-deficient, and diabetic apoE-deficient mice at 21 weeks. Antiplatelet effects were determined using flow cytometry. The antithrombotic effects of clopidogrel were similar in WT and apoE-deficient mice but were attenuated in diabetic apoE-deficient mice with the percent inhibition of thrombus area (µm²) by clopidogrel being 85.5% (WT mice), 75.0% (apoE-deficient mice), and 1.9% (diabetic apoE-deficient mice). The time to first occlusion and lumen stenosis also reflected a significant loss of the antithrombotic effects of clopidogrel in diabetic apoE-deficient mice. Ex vivo platelet activation, which was assessed using ADP-induced expression of activated glycoprotein IIb/IIIa, was completely inhibited by clopidogrel in these three groups of mice. In contrast, the effect of clopidogrel on the ex vivo expression of platelet P-selectin induced by protease-activated receptor 4–activating peptide was diminished in diabetic apoE-deficient mice compared with that in WT and apoE-deficient mice. These data suggest that diabetic apoE-deficient mice may serve as a useful model to better understand the impaired responses to clopidogrel in patients with DM, which may partially reflect a reduction of the effect of clopidogrel on thrombin-induced platelet activation.

Keywords

► P2Y12
► clopidogrel
► diabetes mellitus
► platelets
► thrombosis
► apolipoprotein E–deficient mouse
► thrombin

Introduction

Dual-antiplatelet therapy using aspirin and an antagonist of the P2Y12 ADP receptor is the first-line treatment for patients with acute coronary syndrome as well as for those undergoing percutaneous coronary intervention.1,2 Despite the development of P2Y12 antagonists with more predictable effects, such as prasugrel3–5 and ticagrelor,6,7 clopidogrel remains a widely used P2Y12 antagonist.8,9 However, the antiplatelet effect of clopidogrel is highly variable among individuals10–13 and patients with higher platelet reactivity on clopidogrel treatment are at higher risk of thrombotic events after percutaneous coronary intervention.14–16 Accordingly, delineating the mechanisms underlying the variable response to clopidogrel is clinically relevant and may provide insights into the reasons for treatment failure and inform selection of the three available oral P2Y12 receptor antagonists administered to patients with acute coronary syndrome.17

Clopidogrel is a prodrug requiring hepatic conversion to an active metabolite (AM) that mediates the antiplatelet effect. Numerous pharmacokinetic (PK) and pharmacodynamic studies


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have shown that CYP2C19, a CYP450 isozyme, is important for generating clopidogrel’s AM; moreover, common genetic polymorphisms that reduce the catalytic activity of CYP2C19 decrease AM production.\textsuperscript{18,19} However, the cause of the variability of the clopidogrel response is multifactorial,\textsuperscript{20–22} and diabetes mellitus (DM) is a major risk factor for a diminished clopidogrel response, independent of the CYP2C19 genotype.\textsuperscript{20,21} Overall, patients with DM have an increased prevalence of low platelet inhibition following clopidogrel treatment and a two- to four-fold increased risk of cardiovascular events compared with patients without DM.\textsuperscript{23–27}

While the cause of these characteristics has been attributed to systemic inflammation and insulin resistance,\textsuperscript{28} impaired inhibition of P2Y\textsubscript{12}-mediated platelet inhibition by clopidogrel is largely attributed to attenuation of clopidogrel’s PK profile.\textsuperscript{29} Animal models for studying the impaired clopidogrel response have been employed; however, to the best of our knowledge, there are no reports of a well-established nonclinical model to study the diabetes-associated reduction of response to clopidogrel.

Streptozotocin (STZ)-treated apolipoprotein E (apoE)-deficient mice are widely used to model atherosclerosis with higher cardiovascular risks compared with apoE-deficient or STZ-treated diabetic mice.\textsuperscript{30–33} The FeCl\textsubscript{3}-induced thrombosis model is widely used as a standard method by thrombosis researchers.\textsuperscript{34,35} In this article, we used this thrombosis model to study the antithrombotic effects of clopidogrel in STZ-treated apoE-deficient mice (diabetic apoE-deficient mice) compared with its antithrombotic effects in wild-type (WT) and nondiabetic apoE-deficient mice (apoE-deficient mice). Further, we determined the antiplatelet effects of clopidogrel in these three groups of mice.

Materials and Methods

Materials

ADP, protease-activated receptor-4 thrombin receptor activating peptide (PAR4 TRAP), and FeCl\textsubscript{3} were purchased from Sigma-Aldrich (St. Louis, Missouri, United States). FITC-labeled rat anti-mouse GPIIb/IIIa (JON/A), and PE-labeled rat anti-mouse P-selectin (Wug.E9) antibodies were purchased from Emfret Analytics GmbH & Co. KG (Eibelstadt, Germany). Clopidogrel bisulfate (clopidogrel) was purchased from Kemprotec Limited (Cumbria, United Kingdom). Gum arabic and STZ were purchased from Kemprotec Limited (Kemprotec Limited, Tokyo, Japan) and Sigma-Aldrich (St. Louis, Missouri, United States). Clopidogrel bisulfate (clopidogrel) was purchased from Kemprotec Limited (Kemprotec Limited, Tokyo, Japan) and Sigma-Aldrich. Receptor nomenclature follows the recommendations of the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology.\textsuperscript{36}

Animals

WT (C57BL/6J; Charles River Laboratories Japan, Inc., Yokohama, Kanagawa, Japan) and apoE-deficient male mice (B6.129P2-Apoem1Unc/J; The Jackson Laboratory, United States), which were acquired at 6 or 8 weeks of age, were quarantined and acclimated for at least 7 days and were observed daily for clinical signs during and after the quarantine period. The ex vivo antiplatelet study was conducted in compliance with the Act on Welfare and Management of Animals of Japan and the Guidance for Animal Care and Use of Ina Research Inc., in accordance with the protocol (study no. IP15213) approved by the Institutional Animal Care and Use Committee of Ina Research Inc., which is fully accredited by AAALAC International (accredited unit no. 001107). The thrombosis study was conducted as approved by the Institutional Animal Experiment Committee of Nissei Bilis Co., Ltd. (permit no. 1505–08). These models of atherothrombosis in mice have been in use for several years.\textsuperscript{30–35} All studies involving mice were reported in accordance with the ARRIVE guidelines.\textsuperscript{37,38}

Drug Administration

Clopidogrel was suspended in 5% gum arabic solution and administered orally to mice once daily for 5 days. The dose of clopidogrel was determined according to the results of pilot ex vivo platelet aggregation studies using platelet-rich plasma, which indicated that administration of 10 mg/kg/d of clopidogrel for 5 days completely inhibited ADP-induced platelet aggregation in WT mice (data not shown).

Induction of Diabetes Mellitus

DM was induced in mice that did not exhibit detectable abnormalities during the quarantine period. Citrate buffer solution or STZ (55 mg/kg/d, intraperitoneally) was administered for 5 days to 9-week-old apoE-deficient and WT mice.

FeCl\textsubscript{3}-Induced Thrombosis

Thrombus formation was induced in 21-week-old mice (12 weeks after STZ administration). Mice were anesthetized using isoflurane, and the right carotid artery was exposed. A soft-cuff blood flow probe (MC0.5PSB, Transonic Systems Inc., New York, United States) was placed on the right carotid artery, and blood flow was continuously monitored using a pulsed Doppler flow meter (TS420, Transonic Systems Inc.). Subsequently, 2 hours after the last dose of clopidogrel, filter paper (qualitative filter paper no. 2; Advantec Toyo Kaisha, Ltd., Tokyo, Japan) trimmed to 1 × 2 mm and saturated with a 10% FeCl\textsubscript{3} solution was placed on the arterial surface for 3 minutes. The filter paper was removed, and the time in seconds to first occlusion (TTO), defined as the time to arrest blood flow for >1 minute, was measured. Blood flow was monitored for 60 minutes after placement of the filter paper. If blood flow was not arrested for >1 minute during the 60-minute observation, TTO was defined as 60 minutes, and monitoring was terminated.

Blood Biochemistry

After completing the measurement of blood flow, a blood sample (~500 μL) was collected from the abdominal vein, transferred to a micro-blood collection tube, and serum was collected after centrifugation (1,609 × g, for 10 minutes at room temperature) using a Model 6000 centrifuge (KUBOTA Manufacturing Corporation, Osaka, Japan). All measurements were performed in duplicate. Insulin was measured using a Mouse Insulin ELISA Kit (TMB)/(AKRIN-011T; Shibayagi Co., Ltd., Gunma, Japan) according to the manufacturer’s instructions.
Glucose, total cholesterol, triglycerides, and free fatty acids were measured using the Glucose C2-Test Wako (mutarotase-GOD method), the Cholesterol E-Test Wako (cholesterol oxidase DAOS method), the triglyceride E-Test Wako (glycerol-3-phosphate oxidase DAOS method), and the NEFA C-Test Wako (acyl-CoA synthetase and acyl-CoA oxidase method; WAKO Pure Chemical Industries, Ltd., Osaka, Japan), respectively. All measurements were performed according to the instructions included in each kit.

Pathological Evaluation
After blood collection, mice were euthanized by exsanguination under isoﬂurane anesthesia with perfusion of saline from the left ventricle to incisions in the abdominal artery and vein. Perfusion ﬁxation was performed using 10% neutral-buffered formalin for 5 to 10 minutes, and the FeCl3-injured regions of the right carotid arteries and the approximately identical regions of the left carotid arteries were collected and ﬁxed in 10% neutral-buffered formalin. Specimens were isolated from three areas of the carotid arteries as follows: the approximate midsection of the FeCl3-injured region, 100-μm distal from the midsection, and 200-μm distal from the midsection. Parafﬁn sections were prepared, and each region was stained using hematoxylin–eosin (HE) and an Elastica van Gieson (EVG). For EVG-stained specimens, the luminal area (A) and the thrombus area (T) were measured using image analysis software (Win ROOF 2013; Mitani Corporation, Fukui, Japan), and lumen stenosis (100 × T/A) was calculated. Mean values of T and lumen stenosis in three positions were calculated.

Detection of Activated Glycoprotein IIb/IIIa and P-Selectin on Platelets
Citrated blood samples were collected from the posterior vena cava of mice 2 hours after the ﬁnal dose of clopidogrel. Blood samples were added to tubes containing an FITC-labeled anti-GPIX antibody, a PE-labeled anti-GPIIb/IIIa antibody or an anti-P-selectin antibody, an agonist, and saline. ADP or PAR4 TRAP was used as agonists. Since preliminary studies showed little signiﬁcant expression of P-selectin in response to ADP stimulation, we utilized PAR4 TRAP to induce platelet P-selectin expression in mice. The tubes were then gently stirred and allowed to stand for 15 minutes at room temperature. FACS lysing solution (Becton, Dickinson and Company, New Jersey, United States) was added, and blood cells were ﬁxed for 30 minutes under refrigeration. A FACS Canto II (Nippon Becton, Dickinson and Company, Tokyo, Japan) was used for the analysis. FITC-positive cells were deﬁned as platelets, and the intensities of PE ﬂuorescence speciﬁc to 10,000 platelets were measured. The geometric mean calculated from the histograms of PE ﬂuorescence intensities was deﬁned as the mean ﬂuorescence intensity (MFI).

Data and Statistical Analysis
The data and statistical analysis comply with the recommendations on experimental design and analysis in pharmacology.29 The data for biochemical parameters, platelet activation markers, TTO, thrombus area, and lumen stenosis are presented as the mean ± standard error (SE). Student’s t-test was conducted for each comparison. A two-way analysis of variance (ANOVA) was performed to evaluate the data for treatment (vehicle and clopidogrel) and genotype (WT mice and apoE-deﬁcient mice) as well as treatment (vehicle and clopidogrel) and diabetes (nondiabetic apoE-deﬁcient mice and diabetic apoE-deﬁcient mice). A statistically signiﬁcant difference was deﬁned as p < 0.05. Statistical analyses were performed using SAS 9.1 or 9.3 for Windows (SAS Institute Inc., North Carolina, United States) with EXSUS Version 8.0 (Arm Systex Co., Ltd., Osaka, Japan) or the INATOX-DP (SAS) system (Ina Research Inc., Nagano, Japan).

Results
Blood Biochemistry
The blood levels of insulin, glucose, total cholesterol, triglyceride, and free fatty acids in WT mice, apoE-deﬁcient mice, and diabetic apoE-deﬁcient mice are shown in -Table 1. The mean insulin levels were as follows: vehicle-treated group, 1.1 ± 0.1 ng/mL; WT mice, 0.4 ± 0.1 ng/mL; apoE-deﬁcient mice and diabetic apoE-deﬁcient mice, 0.0 ± 0.0 ng/mL. Insulin levels of diabetic apoE-deﬁcient mice were lower compared with those of WT mice (p < 0.001) and nondiabetic apoE-deﬁcient mice (p < 0.05). The mean glucose level of the vehicle-treated

Table 1  Effects of clopidogrel on blood physiological parameters in wild-type, apoE-deﬁcient, and diabetic apoE-deﬁcient mice

<table>
<thead>
<tr>
<th>Mice</th>
<th>Compound</th>
<th>n</th>
<th>Insulin (ng/mL)</th>
<th>Glucose (mg/dL)</th>
<th>Total cholesterol (mg/dL)</th>
<th>Triglyceride (mg/dL)</th>
<th>Free fatty acid (mEq/L)</th>
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<td></td>
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<tr>
<td>WT</td>
<td>Vehicle</td>
<td>10</td>
<td>1.1 ± 0.1</td>
<td>329 ± 32</td>
<td>102.8 ± 3.1</td>
<td>77.2 ± 11.4</td>
<td>1.03 ± 0.05</td>
</tr>
<tr>
<td></td>
<td>Clopidogrel</td>
<td>10</td>
<td>1.3 ± 0.2</td>
<td>297 ± 15</td>
<td>99.2 ± 2.4</td>
<td>73.3 ± 9.0</td>
<td>0.96 ± 0.03</td>
</tr>
<tr>
<td>ApoE-deficient</td>
<td>Vehicle</td>
<td>10</td>
<td>0.4 ± 0.1a</td>
<td>261 ± 25</td>
<td>667.6 ± 37.8b</td>
<td>144.1 ± 5.8b</td>
<td>1.58 ± 0.12b</td>
</tr>
<tr>
<td></td>
<td>Clopidogrel</td>
<td>10</td>
<td>0.7 ± 0.1a</td>
<td>279 ± 32</td>
<td>698.6 ± 28.0b</td>
<td>143.9 ± 9.9b</td>
<td>1.49 ± 0.05b</td>
</tr>
<tr>
<td>Diabetic apoE-deficient</td>
<td>Vehicle</td>
<td>10</td>
<td>0.0 ± 0.0b</td>
<td>548 ± 15b</td>
<td>1,671.1 ± 149.7b</td>
<td>192.5 ± 18.6b</td>
<td>3.54 ± 0.54b</td>
</tr>
<tr>
<td></td>
<td>Clopidogrel</td>
<td>10</td>
<td>0.0 ± 0.0b</td>
<td>557 ± 20b</td>
<td>1,714.4 ± 156.2b</td>
<td>190.6 ± 20.2b</td>
<td>3.39 ± 0.56b</td>
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</tbody>
</table>

Abbreviations: apoE, apolipoprotein E; WT, wild-type.
Notes: Clopidogrel (10 mg/kg/d) was orally administered once a day for 5 days. There were no statistical differences between each vehicle and clopidogrel groups in WT, apoE-deficient, and STZ-induced diabetic apoE-deficient mice.

*p < 0.01.

*p < 0.001 versus each WT group.
diabetic apoE-deficient mice was 548 ± 15 mg/dL, which was significantly higher compared with those of the vehicle-treated WT mice (329 ± 32 mg/dL, *p* < 0.001) and the vehicle-treated nondiabetic apoE-deficient mice (261 ± 25 mg/dL, *p* < 0.001). The total cholesterol, triglyceride, and free fatty acid levels in the vehicle-treated diabetic apoE-deficient mice were 1671.1 ± 149.7 mg/dL, 192.5 ± 18.6 mg/dL, and 3.54 ± 0.54 mEq/L, respectively, which were significantly higher compared with those of the vehicle-treated WT mice (*p* < 0.001). There were no statistically significant differences between any vehicle and clopidogrel group among the WT, apoE-deficient, and diabetic apoE-deficient mice.

**Effect of Clopidogrel on Thrombus Formation: Blood Flow Analyses**

The patency of the carotid artery following FeCl₃-induced injury in each animal is shown in Fig. 1. In the three vehicle groups, blood flow stopped in most mice soon after the application of FeCl₃, and arterial occlusion persisted thereafter. In contrast, in clopidogrel-treated WT and apoE-deficient mice, blood flow did not stop in all mice. In diabetic apoE-deficient mice, blood flow was disrupted in 6 of 10 mice. Typical tracings of carotid artery blood flow after the arterial injury show that blood flow stopped in diabetic apoE-deficient mice despite clopidogrel pretreatment (supplementary Fig. S1, available online).

We further evaluated TTO as a marker of thrombus formation. The mean TTO values were as follows: vehicle group, 594 ± 37 seconds in WT mice; 530 ± 21 seconds in apoE-deficient mice; and 487 ± 13 seconds in the diabetic apoE-deficient mice (Fig. 2). The mean TTO of diabetic apoE-deficient mice was significantly shorter compared with that of the WT mice (*p* < 0.05). In the clopidogrel-treated groups, the mean TTO values were 3,600 ± 0 seconds

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**Fig. 1** Effects of clopidogrel on the patency of carotid arteries. Clopidogrel (10 mg/kg/d, orally) or vehicle was administered once daily for 5 days to wild-type (WT), apoE-deficient, and diabetic apoE-deficient mice (*n* = 10 per group). Carotid artery occlusion was initiated by application of FeCl₃ 2 hours after the last dose of clopidogrel, arterial blood flow was monitored, and the time in seconds to first occlusion was determined. The results are presented as the mean ± SE (*n* = 10). ***p* < 0.001 versus each vehicle group, †*p* < 0.05 versus the vehicle group of WT mice, #*p* < 0.05 (Student’s t-test).

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**Fig. 2** Effect of clopidogrel on time to first occlusion. Clopidogrel (C) at 10 mg/kg/d (orally) or vehicle (V) was administered once daily for 5 days to wild-type (WT), apoE-deficient, and diabetic apoE-deficient mice (*n* = 10 per group). Carotid artery occlusion was initiated by application of FeCl₃ 2 hours after the last dose of clopidogrel, arterial blood flow was monitored, and the time in seconds to first occlusion was determined. The results are presented as the mean ± SE (*n* = 10). ***p* < 0.001 versus each vehicle group, †*p* < 0.05 versus the vehicle group of WT mice, #*p* < 0.05 (Student’s t-test).
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(p < 0.001) in WT mice, 3,600 ± 0 seconds (p < 0.001) in apoE-deficient mice, and 2,421 ± 481 seconds (p < 0.001) in diabetic apoE-deficient mice (►Fig. 2). The mean TTO values of the clopidogrel-treated groups were significantly prolonged in WT mice, apoE-deficient mice, and diabetic apoE-deficient mice compared with those of each vehicle group; however, the mean TTO of clopidogrel-treated diabetic apoE-deficient mice was significantly shorter compared with those of clopidogrel-treated WT and apoE-deficient mice (each p < 0.05).

Effect of Clopidogrel on Thrombus Formation: Pathological Evaluation
HE-stained cross-sections of injured carotid arteries show the presence of large thrombi in all vehicle-treated mice and clopidogrel-treated diabetic apoE-deficient mice (►Fig. 3). There were no significant differences among the mean thrombus areas of the vehicle groups: 147,174 ± 9,936 μm² in WT mice, 149,814 ± 8,606 μm² in apoE-deficient mice, and 121,557 ± 14,916 μm² in diabetic apoE-deficient mice (►Fig. 4). The mean thrombus areas of the clopidogrel groups were significantly reduced in WT mice (21,272 ± 7,787 μm², p < 0.001) and in apoE-deficient mice (37,501 ± 7,034 μm², p < 0.0001) compared with each vehicle group. The mean thrombus area of the clopidogrel-treated diabetic apoE-deficient mice was 119,280 ± 16,835 μm², and a statistically significant effect of clopidogrel was not observed compared with the corresponding vehicle treatment. The mean thrombus area in diabetic apoE-deficient mice was significantly larger compared with those of WT mice (p < 0.001) and nondiabetic apoE-deficient mice (p < 0.001).

Mean lumen stenosis was not significantly different among the vehicle groups as follows: 95.0 ± 2.2% in WT mice, 96.2 ± 1.1% in apoE-deficient mice, and 87.7 ± 6.5% in diabetic apoE-deficient mice (►Fig. 4). Clopidogrel significantly reduced mean lumen stenosis in WT mice (18.1 ± 6.0%, p < 0.001) and in apoE-deficient mice (34.5 ± 6.4%, p < 0.001) compared with each vehicle group. Mean lumen stenosis in clopidogrel-treated diabetic apoE-deficient mice was 78.4 ± 7.0%, which was not significantly different from that of the corresponding vehicle-treated mice. Mean lumen stenosis in diabetic apoE-deficient mice was significantly higher compared with those of WT mice (p < 0.001) and apoE-deficient mice (p < 0.001).

Fig. 3 Hematoxylin–eosin-stained cross-sections of injured carotid arteries. (A) Vehicle-treated wild-type (WT) (animal ID no. 105 in ►Fig. 1); (B) clopidogrel-treated WT (no. 202); (C) vehicle-treated apoE-deficient (no. 308); (D) clopidogrel-treated apoE-deficient (no. 406); (E) vehicle-treated diabetic apoE-deficient (no. 501); and (F) clopidogrel-treated diabetic apoE-deficient mice (no. 607). Scale bar = 100 μm.
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Fig. 4 Effects of clopidogrel on morphometric parameters of injured carotid arteries, thrombus area (A) and lumen stenosis (B). Clopidogrel (C) at 10 mg/kg/d (orally) or vehicle (V) was administered once daily for 5 days to wild-type (WT), apoE-deficient, and diabetic apoE-deficient mice (n = 10 per group) and arterial occlusion was induced 2 hours after the last dose of clopidogrel. After completing blood flow monitoring, mice were euthanized. The FeCl3-injured regions of the carotid arteries were collected, further fixed in 10% neutral-buffered formalin and paraffin sections prepared, and stained. The lumen and thrombotic areas were measured using an image analysis software (WinROOF 2013), and lumen stenosis (100 x thrombus area/lumen area) was calculated. Results are presented as the mean ± SE (n = 10). ***p < 0.001 versus each vehicle group, ###p < 0.001 (Student’s t-test).

Two-Way Analysis of Variance
Two-way ANOVA revealed associations between DM and clopidogrel according to TTO (p < 0.05), thrombus (p < 0.001), and lumen stenosis (p < 0.001), but not with all biochemical parameters. These results confirm that the antithrombotic effects of clopidogrel were attenuated in diabetic apoE-deficient mice. There was no significant association between genotype of mice and clopidogrel for all parameters, indicating that apoE deficiency did not influence the effects of clopidogrel.

Platelet Surface Expression of ADP-Induced Activated GPIIb/IIIa
In the vehicle groups, treatment with ADP (5 and 20 μM) significantly increased the expression of activated GPIIb/IIIa on platelets compared with no agonist (►Fig. 5) in WT, apoE-deficient, and diabetic apoE-deficient mice (p < 0.01). There was no significant difference in levels of activated GPIIb/IIIa among the vehicle-treated WT, apoE-deficient, and diabetic apoE-deficient mice. Clopidogrel treatment resulted in near-complete inhibition of ADP-induced activated GPIIb/IIIa expression compared with each vehicle group (p < 0.01 each).

Effect of Clopidogrel on PAR4-Activating Peptide-Induced Expression of Platelet P-Selectin
In the vehicle groups, treatment with PAR4 TRAP (100 and 300 μM) significantly increased the expression of platelet P-selectin compared with those of the controls (p < 0.001, all groups; ►Fig. 6). Of note, the level of P-selectin expression in diabetic apoE-deficient mice induced by 100 μM PAR4 TRAP was significantly increased compared with those in WT mice and non-diabetic apoE-deficient mice (p < 0.05 each). Clopidogrel treatment significantly inhibited PAR4 TRAP-induced platelet P-selectin expression compared with that of each vehicle group (p < 0.05, diabetic apoE-deficient mice, 300 μM PAR4 TRAP; p < 0.001 for the other groups). However, the degree of the inhibition differed among the groups as follows: the expression of P-selectin induced by PAR4 TRAP in the diabetic apoE-deficient mice was significantly higher compared with that of WT or nondiabetic apoE-deficient mice (p < 0.001, all groups).

Discussion
In this study, blood biochemistry values revealed unmeasurable insulin levels and a significant increase in glucose in diabetic apoE-deficient mice compared with those of WT and

Fig. 5 Effects of clopidogrel on platelet expression of ADP-induced activated GPIIb/IIIa expression. Clopidogrel (10 mg/kg/d) or vehicle was administered orally once daily for 5 days to wild-type (WT), apoE-deficient, and streptozotocin-induced diabetic apoE-deficient mice, and blood collected 2 hours after the last dose. Ex vivo expression of activated GPIIb/IIIa on platelets induced by 5 and 20 μM ADP was assessed using a flow cytometer. In the vehicle groups, ADP stimulation resulted in a significant increase in the expression of activated platelet GPIIb/IIIa compared with no agonist (p < 0.01 for diabetic apoE-deficient mice, p < 0.001 for other groups). The results are presented as mean ± SE (n = 10). ***p < 0.01 versus corresponding vehicle group (Student’s t-test).
nondiabetic apoE-deficient mice, thus indicating induction of DM. Repeated oral doses of clopidogrel markedly inhibited acute thrombus formation following FeCl3-induced injury of carotid arteries in WT and apoE-deficient mice, consistent with previous reports. In contrast, the antithrombotic effect of clopidogrel, determined by measurements of blood flow, significantly decreased in diabetic apoE-deficient mice compared with those of WT and nondiabetic apoE-deficient mice. To the best of our knowledge, the present study is the first to report a mouse model for studying the diabetes-associated reduction of the response to clopidogrel and therefore represents a unique and potentially valuable approach to identify mechanisms underlying DM-associated impairment of antiplatelet and antithrombotic responses to clopidogrel.

Pathological evaluation, which measured thrombus area and stenosis, further confirmed a significant reduction of the antithrombotic effect of clopidogrel; indeed the extent of the loss of protection against thrombus formation and stenosis was greater than that of blood flow measurements using TTO. The TTO is a commonly used index for the dynamic assessment of thrombus formation in several models of thrombosis. However, our results showed that at least in this model, direct pathological evaluation may be more sensitive for detecting impaired antithrombotic responses.

Impaired inhibition of P2Y12 by clopidogrel in DM is largely attributable to attenuation of the PK profile of clopidogrel. Thus, when we determined the antiplatelet effects of clopidogrel using the present model, we found that ADP-induced platelet activation was similarly inhibited in each group of mice, suggesting that in this model, production of clopidogrel AM was similar among the mice of each genotype/phenotype. Therefore, a mechanism other than those that alter the PK profile may contribute to the impaired response by clopidogrel in STZ-treated diabetic apoE-deficient mice.

Platelet activation is enhanced both in patients with DM and in animal models of diabetes. Although the cause of increased platelet activation is multifactorial, enhancement of ADP-induced platelet activation may contribute to the higher rates of ischemic complications in patients with DM. In this study, levels of ADP-induced platelet GPIIb/IIIa activation among the three genotypes of mice treated with vehicle were not significantly different. Therefore, diabetes-associated enhancement of ADP-induced platelet activation likely does not underlie the impaired antithrombotic effect in diabetic apoE-deficient mice.

In contrast, upon examination of the effects of clopidogrel on PAR4 TRAP-induced activation of platelets, we found an interesting reduction in the antiplatelet effect in diabetic apoE-deficient mice compared with those in WT and nondiabetic apoE-deficient mice. These data suggest that any changes in the effect of clopidogrel on thrombin-induced platelet activation may, in part, mediate the impaired antithrombotic effect of clopidogrel in diabetic apoE-deficient mice. In this study, significant enhancement of PAR4 TRAP-induced platelet activation was observed in diabetic platelets stimulated with PAR4 TRAP (100 μM), which is consistent with previous clinical findings that significant enhancement of PAR4 TRAP-induced platelet aggregation occurs in patients with DM. Moreover, the effects of thienopyridines on thrombin-induced platelet activation are dependent on thrombin concentrations because increased activation of the thrombin receptor of platelets may reduce the antithrombotic effects of thienopyridines. These findings suggest that greater DM-associated enhancement of enhanced platelet activation via thrombin receptors in diabetic mice may reduce the antithrombotic effects of clopidogrel, Prasugrel, and ticagrelor, which exert more potent pharmacodynamic inhibition than previous P2Y12 antagonists, are associated with increased reduction of cardiovascular risk in patients with DM compared with clopidogrel. These findings may be partially explained by the more potent inhibition of thrombin-induced platelet activation by prasugrel and ticagrelor.

These data suggest that apoE-deficient mice with STZ-treated diabetes will serve as a useful model to conduct further studies of the impaired responses to clopidogrel found in patients with DM, which may partially reflect a reduction of the effect of clopidogrel on thrombin-induced platelet activation.

Conflict of Interest
Joseph A. Jakubowski (retired) is a minor shareholder of Eli Lilly and Company. Other authors are employees of Daiichi Sankyo Co., Ltd.

Acknowledgments
We thank Nissei Bilis Co., Ltd. and Ina Research Inc. for their expert technical contributions.
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