Monoclonal Antibody against the Platelet Glycoprotein (GP) IIb/IIIa Receptor Prevents Coronary Artery Reocclusion after Reperfusion with Recombinant Tissue-type Plasminogen Activator in Dogs

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Abstract

Localized thrombosis was produced in the left anterior descending coronary artery of open chest dogs by constraining a segment so as to produce >90% stenosis (reducing blood flow to 40±10% of baseline), and placing a thrombus in the segment immediately proximal to the stenosis by inducing endothelial cell injury and instilling a mixture of blood and thrombin.

Intravenous infusion of recombinant tissue-type plasminogen activator (rt-PA) at a rate of 15–30 μg/kg per min for 30 or 60 min in eight dogs induced coronary artery reperfusion within 23±7 min (mean±SD), but reocclusion occurred despite heparin anticoagulation in all but one of these dogs within 7±5 min.

Intravenous injection of 0.8 mg/kg of the F(ab')2 fragment of a monoclonal antibody (7E3) directed against the platelet GPIIb/IIIa receptor, prevented reocclusion in 10/10 dogs during an observation period of 2 h (P < 0.001 vs. rt-PA alone). The antibody abolished ADP-induced platelet aggregation and markedly prolonged the bleeding time. Intravenous aspirin or dipyridamole prevented reocclusion for 1 h or more in only 2/7 and 1/6 dogs, respectively.

We conclude that the monoclonal antibody is very effective in preventing reocclusion after successful thrombolysis of occluded coronary arteries with rt-PA.

Introduction

Reocclusion is a common complication after coronary artery reperfusion with thrombolytic agents, occurring in ~25% of the patients (1–3). The main determinant for reocclusion is the extent of the residual stenosis after thrombolysis (2, 3); thus, with recombinant tissue-type plasminogen activator (rt-PA),1 the majority of patients with a stenosis of >80% develop reocclusion unless treated with a maintenance infusion (2).

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1. Abbreviations used in this paper: GP, glycoprotein; rt-PA, recombinant tissue-type plasminogen activator.

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Fixed atheromatous disease is responsible for much of the residual stenosis after thrombolysis, and reocclusion occurs by the superimposition of thrombi containing both platelets and fibrin (4). Recently, one of us described a murine monoclonal antibody (7E3) that binds to the platelet glycoprotein GPIIb/IIIa receptor and blocks platelet aggregation induced by a variety of physiological agonists (5). Intravenous infusion of F(ab')2 fragments of the antibody at a dose of 0.8 mg/kg in dogs and monkeys induced profound inhibition of platelet aggregation ex vivo and platelet thrombus formation in vivo, without producing severe thrombocytopenia or spontaneous bleeding (6, 7).

In the present study we evaluated the efficacy of F(ab')2 fragments of 7E3 for the prevention of coronary artery reocclusion after thrombolysis with rt-PA in a dog model consisting of a coronary artery thrombus placed adjacent to a fixed high grade (>90%) stenosis in a vessel segment with damaged endothelium.2 In this model, coronary thrombosis can be lysed readily by intravenous rt-PA, but the thrombogenic stimulus is of such magnitude that reocclusion occurs rapidly and consistently when the experimental animals are treated with heparin alone or heparin in combination with either aspirin or dipyridamole.2

Methods

Reagents: rt-PA was supplied by Genentech, Inc., South San Francisco, CA. Two preparations were used: one, G11021, is produced in roller bottles and consists predominantly of two-chain rt-PA; the other, G11035, is produced in suspension culture and consists predominantly of single chain rt-PA. The production of the monoclonal antibody 7E3, its purification, and fragmentation into F(ab')2 fragments with pepsin have been described in detail (6, 7). Antibody given to the last five dogs was prepared by a modified technique involving fragmentation with 120 U of pepsin (Cooper Biomedical, Malvern, PA) per mg 7E3 at pH 4.2 (0.15 M NaCl, 0.1 M Na citrate) for 6 h at 37°C. The digestion was stopped by raising the pH to 7.5 with 1 M Tris-HCl, 0.02 M EDTA, pH 8.0, and the F(ab')2 fragments were purified by gel filtration on Superose 12 (Pharmacia Fine Chemicals, Piscataway, NJ) equilibrated with 0.15 M NaCl, and chromatography on Q-Sepharose Fast Flow (Pharmacia) with elution by a linear gradient from 0 to 1.0 M NaCl in 0.05 M Tris·HCl, pH 8.0. The final material was pooled and then both concentrated and diafiltered with 0.15 M NaCl using a YM-10 filter (Amicon Corp., Waltham, MA). Antibody fragments were prepared at concentrations ranging from 0.68 to 3.1 mg/ml in 0.15 M NaCl and frozen until just before use. When analyzed for endotoxin with an


amebocyte lysate clotting assay (Pyrogent; Mallinckrodt, St. Louis, MO) the first antibody preparations were found to contain between 2 and >80 endothelin U/mg protein. The preparations for the last five dogs had lower endothelin values of 0.5–1 endothetin U/mg as judged by a spectrophotometric assay (Whittaker M. A. Bioproducts, Walkersville, MD). A control F(ab')2 fragment of a monoclonal antibody directed against an ovarian carcinoma antigen (OC-125) (Centocor, Malvern, PA) (8) was prepared as described above with only minor modifications.

Animal model. Coronary artery thrombosis and endothelial cell damage was produced in dogs as described before (9). In addition, superimposed high grade stenosis was produced by the application of an external constrictor, and blood flow was continuously monitored with an electromagnetic flow probe. Mongrel dogs weighing ~20–25 kg were anesthetized with a slow intravenous injection of sodium pentobarbital, intubated and placed on an artificial ventilator. A left thoracotomy was performed in the 5th–6th intercostal space, and an arterial catheter was placed in the internal mammary artery for blood pressure monitoring. Procainamide (1.5 g i.m. in two to three sites) was then given, the pericardium opened, and a pericardial cradle prepared. The left anterior descending coronary artery was dissected out from the epicardium, side branches were ligated and a 2.5-cm segment isolated. An electromagnetic flow probe (FM 501; Carolina Medical Electronics, King, NC) was placed on the most proximal portion of the segment and intravenous lidocaine (15-mg bolus followed by a constant infusion at 1 mg/min) was infused. A control left coronary angiogram was performed by injecting ~2 ml of Renograffin 76 by hand through a modified Judkin’s 7 French catheter inserted from a carotid artery. One ml of blood was then removed and kept in a syringe for later use in forming the thrombus, and heparin (5,000 U intravenous bolus) was administered. Additional 1,000 U boluses of heparin were administered at hourly intervals. A permanent 2-mm wide constrictor was placed near the distal end of the segment and adjusted so as to reduce coronary artery blood flow to ~40±10% of control. High resolution postmortem angiograms in selected animals showed that a constriction so placed decreased the luminal diameter by >90%. The 1 cm of coronary artery just proximal to the constriction was then empli bed of blood and isolated between temporary silk snare. Intimal damage was induced by grasping the segment with forceps and then the segment was flushed by releasing the proximal snare and injecting saline retrograde through a cannulated side branch. The segment was then reisolated and 0.1 ml of thrombin (topical thrombin, 1000 U/ml; Parke-Davis Co., Morris Plains, NJ) and 0.3 ml of the stored blood were injected into the isolated segment. After 5 min, first the proximal and then the distal ties were released and the side branch catheter was removed, whereas the permanent constrictor remained in place. Approximately 30 min after injecting the thrombin and blood, a repeat angiogram was performed to confirm the persistence of complete coronary artery occlusion as demonstrated by continuous electromagnetic flow probe monitoring.

The rt-PA infusion was then started in the animals not receiving antiplatelet agents. The other animals were given the F(ab')2 fragment of 7E3 (0.7–0.8 mg/kg in 10 dogs), acetylsalicylic acid (35 mg/kg in three dogs), dipryridamole (0.6 mg/kg in six dogs), or fragments of the control monoclonal antibody OC-125 (0.7–0.8 mg/kg in two dogs) by slow intravenous injection. Approximately 10 min later, the last animals received an infusion of rt-PA at 15 μg/kg per min for the predominantly two chain form (G11021), or at 30 μg/kg per min for the predominantly single chain form (G11035) for 30 min. In animals that did not achieve at least partial coronary artery reperfusion near the end of the 30 min infusion, rt-PA infusion was continued for another 30 min. In addition, in four dogs, acetylsalicylic acid (35 mg/kg) was injected intravenously at the time of reperfusion. The blood flow in the vessel was monitored continuously and when flow appeared to be restored, another angiogram was immediately performed. The reperfusion time was taken as the number of minutes from the beginning of the rt-PA infusion until reperfusion was documented by the flow meter and confirmed by the repeat angiogram showing complete antegrade filling of the artery with rapid clearance of the dye (in less than four cardiac cycles). After reperfusion was obtained, blood flow was monitored for evidence of reocclusion, with the final confirmation again obtained by angiography using the same criteria as were used for establishing reperfusion. The reocclusion time was taken as the interval between documented reperfusion and reocclusion.

Blood analyses. Bleeding times were performed before and 30 min after injections of the F(ab')2 fragment of 7E3 (eight dogs), control antibody OC-125 (two dogs) or aspirin (three dogs), with a spring-loaded blade device (Simplate; General Diagnostic Corp., Morris Plains, NJ, or Surgicutt; International Technidyne Corp, Edison, NJ), applied to a shaved foreleg. Venous blood samples for determination of the levels of fibrinogen, activated partial thromboplastin time, ADP-induced platelet aggregation and 125I-7E3 binding were collected into 0.01 M citrate containing 150 KIU/ml aprotinin (Sigma Chemical Co., St. Louis, MO). Platelet counts were performed with an automated particle counter (Coulter, Hialeah, FL) on blood drawn into EDTA. Platelet-rich plasma was prepared as previously described for the aggregation and 125I-7E3 binding studies (6). The number of F(ab')2 molecules bound per platelet in vivo was estimated from the ex vivo binding of 125I-7E3 to platelets removed from the dog before and after giving the F(ab')2 fragments (6). Plasma for the other studies was obtained from blood samples kept on ice until the end of the experiment, then centrifuged at 3,000 g at 22°C for 10 min and stored at -20°C. The assays were performed as previously described (9).

Pathologic examination. At the end of the experiment, the dogs were killed by an overdose of pentobarbital. Thrombosed stenotic and poststenotic segments of the left anterior descending coronary artery of three dogs given rt-PA alone were removed intact and fixed overnight in 5% formaldehyde. The segments were sectioned at 2-mm intervals, stained with hematoxylin and eosin and evaluated microscopically. Five dogs receiving antiplatelet antibody were subjected to perfusion fixation and scanning electron microscopy of the left anterior descending coronary artery. The artery segment was prepared as previously described (10).

Results

Dogs treated with rt-PA alone. 10 animals were studied. Two of these were excluded from analysis; one failed to achieve reperfusion despite 60 min of rt-PA and the second died immediately after reperfusion due to ventricular fibrillation. In the remaining eight dogs (Table I), the permanent constriction reduced the blood flow to 38±10% of the baseline value before thrombus formation. The time to reperfusion after the start of the rt-PA infusion was 23±7 min (mean±SD). No significant differences were observed between the groups receiving the two different forms of rt-PA at the different infusion rates. This finding is consistent with the observation that the specific thrombolytic activity of the predominantly two-chain preparation of rt-PA (G11021) is somewhat greater than that of the predominantly one-chain form (G11035) (11). After reperfusion, seven of the eight dogs rethrombosed rapidly with a mean time to reocclusion of 7±5 min. In some of these animals, followed for a period of time after reocclusion, cyclic reperfusion and reocclusion occurred until either the experiment was terminated or persistent reocclusion occurred.

Dogs treated with rt-PA and 7E3 F(ab')2 fragments. 13 dogs received F(ab')2 fragments of 7E3 at a dose of 0.7–0.8 mg/kg in combination with either 15 μg/kg per min of two-chain rt-PA or 30 μg/kg per min of one chain rt-PA. Three dogs were excluded from analysis; two achieved spontaneous reperfusion of the coronary artery during the 10-min interval between the injection of the antibody and the scheduled start
of rt-PA; the third failed to achieve reperfusion by angio-
graphic criteria despite 60 min of rt-PA.

The left anterior descending coronary artery blood flow in
the remaining 10 dogs (Table I B) was decreased to an average
of 37±11% of the control value by the permanent con-
strictor. Reperfusion occurred after 14±10 min in this group.
None of these animals had coronary reocclusion during an
observation period of ~ 2 h.

Five of the six dogs treated with antibody prepared as pre-
viously described (6) sustained a transient decrease in blood
pressure that responded to saline infusion. In contrast, none of
the five dogs treated with the antibody prepared by the method
described in this paper became hypotensive, suggesting that
the original responses were due to a contaminating vasoactive
agent, perhaps endotoxin.

Two additional dogs were injected with 0.8 mg/kg of the
F(ab')2 fragment of a control monoclonal antibody, OC-125,
and then infused with 30 μg/kg per min of single-chain rt-PA
for 60 min. Reperfusion occurred after 36 and 60 min and then
reocclusion was documented after another 29 and 1 min,
respectively.

Dogs treated with rt-PA and aspirin or rt-PA and dipyrida-
mole. Four dogs were infused with 15 μg/kg per min of two-
chain rt-PA (G11021) for 30 or 60 min and given intravenous
aspirin (35 mg/kg) at the time of reperfusion (Table II). Two of
the dogs suffered reocclusion rapidly (8 and 15 min), one
reoccluded at 116 min, and one remained open for > 120 min.
Three additional dogs were given aspirin 10 min before the
start of the infusion of 30 μg/kg per min of single chain rt-PA
(G11035). All three dogs reoccluded, within 3, 10, and 6 min,
respectively.

Six dogs were treated with dipyridamole and two chain
rt-PA (Table II). The coronary arteries of five of these dogs
reocluded within 11 min, whereas one remained open for 34
min and one remained patent throughout the experiment
(> 55 min).

Analyses of hemostasis and platelet function. Platelet ag-
gregation studies performed on blood obtained before and 30
min after antibody infusion showed essentially complete abo-
lition of aggregation in response to ADP (9 μM), but the
shape-change response remained intact. In all of three dogs
tested after aspirin injection and reperfusion with rt-PA
G11035, reocclusion occurred rapidly despite abolition of the
second wave of ADP-induced platelet aggregation in blood
taken 30 min after the aspirin injection.

Platelet counts were obtained before and at the end of the
experiment in seven animals treated with the 7E3-F(ab')2.
There was a mean reduction in platelet count of 18% (range
3–33%) similar to that observed in our previous study (6).

Blending times were obtained in five dogs before and 20
min after 7E3-F(ab')2 infusion. The values before the antibody
averaged 3.6±2.6 min (range 1.5–5 min). After antibody infu-
sion, one blending time was prolonged to 15 min, whereas the
other four were increased to > 30 min. In contrast, the blend-
ing time after treatment with the control monoclonal antibody
in two dogs did not increase significantly (from 2 and 1.5 min
preinfusion to 2.5 and 5.5 min postinfusion). In the three dogs
with aspirin injection prior to the start of the rt-PA infusion,
were surrounded by lial of de-endothelialized surface (Fig. 1). Tears (Fig. 1) revealed vessels including intimal thickening, which indicated that reocclusion occurs in ~25% of reperfused vessels, especially when the residual stenosis is very pronounced (1–3). The canine model used in the present study was designed to simulate the occurrence of an acute, occlusive thrombus in a coronary artery, superimposed on high grade (>90%) stenosis and endothelial injury. A fixed, high-grade stenosis was produced by a permanent constrictor, endothelial damage was induced by external trauma and an occlusive thrombus was formed just proximal to the site of stenosis. Under these circumstances, the dogs respond in a manner that closely approximates the human response, namely, clot lysis and reperfusion induced by rt-PA infusions at thrombolytic doses, consistently followed by rapid reocclusion despite full heparinization (2).

The data represent mean±SD. * Not included in the calculation of the mean value±SD.

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Discussion

Early reperfusion of thrombosed coronary arteries with thrombolytic agents may salvage ischemic areas of myocardium in patients with acute myocardial infarction, provided sustained recanalization is obtained. Available studies, however, indicate that reocclusion occurs in ~25% of reperfused vessels, especially when the residual stenosis is very pronounced (1–3). The canine model used in the present study was designed to simulate the occurrence of an acute, occlusive thrombus in a coronary artery, superimposed on high grade (>90%) stenosis and endothelial injury. A fixed, high-grade stenosis was produced by a permanent constrictor, endothelial damage was induced by external trauma and an occlusive thrombus was formed just proximal to the site of stenosis. Under these circumstances, the dogs respond in a manner that closely approximates the human response, namely, clot lysis and reperfusion induced by rt-PA infusions at thrombolytic doses, consistently followed by rapid reocclusion despite full heparinization (2).

Because the reoccluding thrombi are rich in platelets, and since both heparin and a standard thrombolytic dose of rt-PA failed to prevent rethrombosis, we tested the efficacy in preventing reocclusion of a new antiplatelet agent, a murine monoclonal antibody to the platelet GpIIb/IIIa receptor that blocks the binding of fibrinogen to platelets (5–7). Whereas platelet aggregation induced by the agonists thought to operate in vivo (e.g., ADP, epinephrine, thrombin, collagen, thromboxane A2) is absolutely dependent upon the binding of fibrinogen (and/or perhaps fibronectin and von Willebrand factor) to this receptor (12), the antibody is capable of completely blocking platelet aggregation induced by these agonists. We chose to use the F(ab')2 fragment rather than the intact anti-
Figure 1. Scanning electron micrographs of the left anterior descending coronary artery after treatment with rt-PA and platelet antibody. 

(A) Low magnification micrographs reveal extensive intimal disruption with circumferentially oriented tears in the segment proximal to the stenosis (P). The intimal surface of the stenotic region (S) appears relatively smooth in comparison (× 25). (B) Prestenotic segment showing deposition of a layer of metamorphizing platelets and body so as to avoid the potential removal of antibody-coated platelets via recognition of the Fc portion of the immunoglobulin molecule.

The present study shows that the 7E3 F(ab')

model from the Folts model. The degree of stenosis in our model is much greater (> 90% vs. ~ 70%) as demonstrated by the marked reduction in blood flow and the lack of such a decrease in the Folts model; in fact, in a study using a nonthrombotic variation of the Folts model involving a level of stenosis comparable to that used in the current study, aspirin failed to inhibit thrombus formation (14). One explanation for the inefficacy of aspirin may be its lack of effect on platelet aggregation induced by shear forces (15). In our model a fully occluding thrombin-induced thrombus is formed and then lysed, raising the possibility that residual thrombin or other platelet activating agents (ADP, thromboxane A2, etc.) may be present in the local environment and provide a greater thrombogenic stimulus. Because the dog's coronary artery is fully occluded for 30 min in our model, the degree of cardiac ischemia is much greater and this may result in release of catecholamines; the latter have been shown to reverse the inhibitory effect of aspirin in the Folts model (7).

Dipyridamole was also of only minimal benefit in preventing reocclusion in this model, despite using the maximal dose (0.6 mg/kg) that can be administered without hemodynamic
compromise. The lack of effect is not surprising given the recent evidence calling into question the efficacy of dipyrindamole as an antithrombotic agent, either alone or in combination with aspirin. In particular, recent results of several large scale clinical trials which have indicated that aspirin alone is equivalent to combinations with dipyrindamole for the prevention of arterial occlusion in patients with established cardiovascular disease (16) and (b) two recent animal studies showed that dipyrindamole alone had no effect and that its combination with aspirin was no more effective than aspirin alone at a dosage > 2 mg/kg in preventing platelet thrombus formation on partially stenosed dog coronary arteries (17, 18).

The two models with moderate and severe coronary artery stenosis may have complementary features rendering them more appropriate for different applications. The model with less severe stenosis (14, 19) may be more adapted to the investigation of pharmacological agents suitable for the prevention of arterial occlusion in patients with atherosclerotic disease, where aspirin has indeed been shown to be effective. The model used in the present study may be more suitable for the investigation of therapeutic approaches for the prevention of acute coronary artery reocclusion in patients with myocardial infarct, where a high grade residual stenosis after successful reperfusion with thrombolytic agents persists. In this condition the anatomical substrate consists of plaque hemorrhage, mural thrombus, endothelial damage, and fixed atherosclerotic narrowing, which in concert constitute an intense thrombogenic stimulus.

We were concerned about the potential hemorrhagic effect of the combination of agents used in this study, since the animals were subjected to extensive surgery with open operative wounds, and the agents employed [rt-PA, heparin, and 7E3-F(ab')2] inhibit all phases of hemostasis. We therefore anticipated that the addition of the monoclonal antiplatelet antibody to the heparin/rt-PA combination might provoke excessive hemorrhage, especially since the antibody produced a dramatic increase in the bleeding time. Several factors, however, may have contributed to the relative absence of excessive hemorrhage during the combined infusion of rt-PA and 7E3 F(ab')2. Firstly, most of the invasive procedures were completed before the agents were administered, thus possibly permitting hemostatic plugs to mature. Moreover, the antibody only blocks the receptor-mediating platelet aggregate formation, leaving intact other platelet receptors for collagen, von Willebrand factor, and perhaps fibronectin, that probably are more important in mediating adhesion of platelets to the subendothelium (13, 14, 20). In support of this hypothesis, similar antibodies have had a more dramatic effect on platelet aggregation than on platelet adhesion in model systems (20–22). In addition, platelets treated with a similar antibody also retained the ability to undergo the release reaction and activate fibrin formation (22). Finally, although prolongation of the bleeding time must be considered a significant risk factor for excessive hemorrhage, patients who have long bleeding times because they are congenitally deficient in the GPIIIb/IIIa receptor (Glanzmann thrombasthenia) usually only bleed excessively with trauma or other provocations (23), and patients with long bleeding times as a result of treatment with the drug ticlopidine, which interferes with the GPIIIb/IIIa receptor by an unknown mechanism, appear to be at low risk for spontaneous hemorrhage (24, 25). There is thus reason to hope that with the antiplatelet antibody it may be possible to completely inhibit platelet aggregation, the process most likely to produce vascular occlusion, without significantly inhibiting platelet adhesion and other platelet functions that contribute to maintaining hemostasis. The potential hemorrhagic risk of short term therapy with the combination of 7E3 F(ab')2, with a rapidly cleared thrombolytic agent such as rt-PA and a rapidly reversible anticoagulant, thus may be justified if the combination is found to be highly effective in treating this life-threatening illness.

In conclusion, the present study indicates that infusion of F(ab')2 fragments of monoclonal antibody 7E3, which is directed against the platelet GPIIIb/IIIa receptor, can efficiently protect against coronary artery reocclusion that follows thrombolysis with rt-PA, despite the presence of high-grade residual stenosis. Whether such treatment can be administered safely to humans or has value as an alternative to maintenance rt-PA infusion remains to be further investigated.

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