Changes in membrane glycoproteins of circulating platelets after coronary stent implantation

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Abstract
Objectives—To evaluate platelet function in patients with coronary stents.
Design—A non-randomised control trial in 30 patients who had immediate implantation of Palmaz-Schatz coronary stents because of a suboptimal angioplasty result. All patients received a standardised anticoagulation regimen including intravenous heparin (activated partial thromboplastin time [APTT] 80 to 120 s), oral vitamin K antagonist (target international normalised ratio [INR] of 3-5), and 100 mg aspirin twice daily. Platelet surface expression of glycoprotein IIb-IIIa, activated fibrinogen receptor, and P-selectin as well as binding of von Willebrand factor and fibrinogen were determined by flow cytometry in peripheral venous blood samples collected before the intervention and then daily for 4 days after it. The results were compared with those in 30 patients undergoing elective coronary balloon angioplasty.
Setting—University hospital.
Results—After coronary stenting surface expression of the activated fibrinogen receptor significantly increased, peaking at day 2 (P < 0.001). Similar results were found for von Willebrand factor binding and P-selectin surface expression, with a maximum at day 2 to 4 after stenting (von Willebrand factor, P < 0.001; P-selectin, P < 0.001). The changes in platelet membrane glycoproteins coincided with a significant drop in peripheral platelet count after stent placement (P < 0.01). No significant change in fibrinogen receptor activity, von Willebrand factor binding, P-selectin surface expression, or platelet count was seen in the control group.
Conclusions—The present study shows that current anticoagulation treatment is inefficient in suppressing platelet activation in patients with coronary stents and, therefore, might not be the best treatment for reducing the incidence of subacute stent thrombosis.

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Keywords: angioplasty, coronary stent implantation, platelets, membrane glycoproteins

Coronary stenting is an effective method to treat abrupt vessel closure after coronary balloon angioplasty (PTCA). Subacute stent thrombosis, however, is a major complication after stent placement and occurs in about 3% to 21%. Thus strict antithrombotic regimens are used in stent patients to prevent thrombotic events, at the cost of increased bleeding complications. Platelet adhesion and aggregation play a fundamental role in acute coronary thrombus formation; however, platelet function in coronary stent patients has not been evaluated.

Platelet adhesion and aggregation depend on the exposure of adhesive glycoproteins on the platelet surface and these mechanisms are the target of newer antiplatelet drugs.

Vascular injury—as it occurs after dilatation and stenting of atherosclerotic lesions—exposes circulating blood to potentially thrombogenic surfaces on the subendothelium. Platelets can then adhere to the injured vessel wall through interaction with immobilised von Willebrand factor (vWF). After they adhere, platelets become activated and induce fibrinogen (fg) binding sites on the glycoprotein complex IIb-IIIa. Plasma fibrinogen can then bind to the activated platelet surface and initiate aggregation and thrombus formation through interplatelet bridging. Concomitantly, platelets degranulate and redistribute granule-stored glycoprotein P-selectin (GMP-140, PADGEM) to their surface; this mechanism supports haemostatic consolidation of the thrombotic plug.

The present study focuses on changes in platelet membrane glycoproteins after coronary stenting. Immunological detection of platelet surface glycoproteins allows a spectrum of specific activation-dependent modifications in the glycoprotein composition of the platelet membrane to be detected with a high degree of sensitivity. Several studies of extracorporeal blood bypass, coronary angioplasty, or multiple organ failure, have shown this method to be a useful tool for assessing prethrombotic states in clinical settings. We studied the hypothesis that platelet function is altered after coronary stent placement by comparing the results with those in a group of patients undergoing elective coronary angioplasty (PTCA).

Patients and methods
STUDY POPULATION
All patients studied had a high-grade coronary artery stenosis (> 80%) and a positive exercise test. Platelet function was investigated in 30 patients who had had a Palmaz-Schatz stent...
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Implanted immediately after angioplasty. The results were compared with 30 patients who had previous angioplasty without angiographically detectable dissection. Written informed consent to coronary stenting and for the repeated collection of blood was obtained from all patients before the intervention. The study was performed after the protocol was approved by the ethics committee of our university.

The basic characteristics and a summary of laboratory variables are shown in tables 1 and 2. Patients were excluded if they had a recent myocardial infarction, unstable angina, or any illness known to influence platelet function including chronic inflammatory disease, active bleeding, severe anaemia, or chronic renal failure. All patients were on long term aspirin treatment before entering the study.

STUDY DESIGN
Balloon angioplasty was performed identically in both groups by the transfemoral approach. After initial coronary angiography, 15 000 international units of unfractionated heparin (Thrombophob, Nordmark, Germany) and 500 mg of aspirin (Aspisol, Bayer, Germany) were given intravenously just before the catheter system was inserted. None of the patients received any thrombolytic treatment throughout the procedure or thereafter. The decision to implant a stent was made as described recently. Within the first four hours after the intervention we removed the arterial sheath in both study groups, as soon as the APTT fell below 80 s.

All patients were given two 100 mg aspirin tablets on the day of the intervention and then daily thereafter. In patients undergoing balloon angioplasty (PTCA) anticoagulation with intravenous heparin (APTT 60 to 90 seconds) was stopped after 24 hours. Immediately after coronary stenting an oral vitamin K antagonist (Marcumar, Hoffmann-La Roche) was started and intravenous heparin was continued to adjust APTT values of 80–120 s until a target prothrombin time of 3·5 (INR) was achieved.

SPECIMEN COLLECTION
Peripheral venous blood samples were taken with a loose tourniquet through a short venous catheter inserted into a forearm vein just before and then daily in the morning after coronary intervention. A multiple-syringe sampling technique was used and the first 2 ml of blood were discarded. Thereafter, 2-5 ml of blood collected in disodium ethylene-diaminetra-acetic acid (EDTA) was used to measure the platelet and white blood cell count, haemoglobin, and hematocrit with a Coulter Counter (Corning). Four ml of blood was collected in 3·8% citrate to measure coagulation variables and 5 ml was taken for serum chemistry. For flow cytometric analysis 1·6 ml blood was collected into a polypropylene syringe containing 0·4 ml of citrate phosphate dextrose acid (CPDA).

PREPARATION OF SAMPLES FOR FLOW CYTOMETRIC ANALYSIS
Preparation of platelets and immunolabelling of platelets with monoclonal antibodies (mAb) for flow cytometric analysis was performed as described previously. 10-11 In brief, immediately after blood collection CPDA-anticoagulated whole blood was centrifuged at 50 g for 10 min to obtain platelet-rich plasma (PRP). Thereafter, 5 ml of PRP was added to polypropylene tubes (Becton-Dickinson) preloaded with 45 ml of modified HEPES-Tyrode's buffer containing saturating concentrations of fluorescein-isothiocyanate (FITC)- conjugated mAbs. Non-specific membrane immunofluorescence was determined by use of an irrelevant FITC-IgG mAb. Samples were incubated in the dark for 15 min at room temperature without agitation. Immunolabelled samples were fixed by addition of 1 ml 0·5% paraformaldehyde in phosphate buffered saline, pH 7·4, and stored at 4°C until flow cytometry was performed within 24 hours.

The platelet assay we used gives reproducible results without significant artefactual platelet activation and is suitable for platelet analysis in various clinical settings. 10-11

FLOW CYTOMETRY OF PLATELETS
Platelet samples were run on a FACScan cytometer (Becton Dickinson) equipped with a 2W coherent argon-ion laser with an excitation wavelength set at 488 nm. The flow cytometer was calibrated with commercially

### Table 1 Clinical data on study patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Stent group (n = 30)</th>
<th>PTCA group (n = 30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (M/F)</td>
<td>24/6</td>
<td>23/7</td>
</tr>
<tr>
<td>Age (y) (mean (SD), range)</td>
<td>57 (9-6) (39-82)</td>
<td>61 (8-4) (47-80)</td>
</tr>
<tr>
<td>Number of diseased vessels (%)</td>
<td>13 (3-3)</td>
<td>15 (50-0)</td>
</tr>
<tr>
<td>2</td>
<td>10 (33-3)</td>
<td>10 (33-3)</td>
</tr>
<tr>
<td>Single stent (%)</td>
<td>1 (23-4)</td>
<td>5 (16-7)</td>
</tr>
<tr>
<td>Multiple stent (%)</td>
<td>7 (23-3)</td>
<td></td>
</tr>
<tr>
<td>Target vessel (%)</td>
<td>3·7 (0·3) (3·0-4·0)</td>
<td>2·9 (0·4) (2·5-4·0)</td>
</tr>
<tr>
<td>LAD, left anterior descending coronary artery</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RCA, right coronary artery; PTCA, percutaneous transluminal coronary angioplasty</td>
<td></td>
<td></td>
</tr>
<tr>
<td>LV, left ventricle</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Table 2 Laboratory variables of study patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Stent group (n = 30)</th>
<th>PTCA group (n = 30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>WBC (× 10^9/ml)</td>
<td>9·19 (3·00)</td>
<td>8·90 (4·07)</td>
</tr>
<tr>
<td>Haemoglobin (g/dl)</td>
<td>13·42 (0·98)</td>
<td>12·80 (1·31)</td>
</tr>
<tr>
<td>Platelet count (× 10^9/ml)</td>
<td>232·73 (65·13)</td>
<td>227·55 (54·20)</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>0·97 (0·37)</td>
<td>0·94 (0·14)</td>
</tr>
<tr>
<td>Urea (mg/dl)</td>
<td>16·65 (6·33)</td>
<td>11·83 (3·95)</td>
</tr>
<tr>
<td>GPT (U/l)</td>
<td>24·05 (17·09)</td>
<td>15·80 (5·14)</td>
</tr>
<tr>
<td>AP (U/l)</td>
<td>115·50 (56·18)</td>
<td>101·00 (20·52)</td>
</tr>
<tr>
<td>Bilirubin (mg/dl)</td>
<td>0·65 (0·30)</td>
<td>0·60 (0·26)</td>
</tr>
<tr>
<td>CK (U/l)</td>
<td>35·16 (35·29)</td>
<td>22·50 (9·48)</td>
</tr>
</tbody>
</table>

WBC, white blood cell count; GPT, glutamate pyruvate transaminase; AP, alkaline phosphatase; CK, creatine kinase.
Figure 1  Role of membrane glycoproteins in platelet function. Under physiological conditions platelets circulate in a resting state. Activation exposes fibrinogen binding sites on the glycoprotein complex Iib-IIIa that allows plasma fibrinogen to bind to the platelet surface. Fibrinogen bridging between platelets initiates the aggregation process. Thereafter, platelets degranulate and α-granule glycoprotein GMP-140 (P-selectin) is expressed on their surface. von Willebrand factor binds to two distinct glycoprotein complexes, GPIb and GPIb-IIIa, present on the platelet surface and is crucial for platelet adhesion to subendothelium. Use of a panel of mAbs that recognise specific glycoproteins (anti-CD41, anti-LIBS1, anti-CD62P, anti-fg, anti-vWF) identifies the functional state of circulating platelets.

Table 3  Procedural characteristics and angiographic results (mean (SD))

<table>
<thead>
<tr>
<th>Measurement</th>
<th>Stent group</th>
<th>PTCA group</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reference diameter (mm)</td>
<td>3.07 (0.56)</td>
<td>2.73 (0.47)</td>
<td>0.017</td>
</tr>
<tr>
<td>Minimal lumen diameter before (mm)</td>
<td>0.86 (0.41)</td>
<td>0.81 (0.43)</td>
<td>0.603</td>
</tr>
<tr>
<td>Minimal lumen diameter after (mm)</td>
<td>3.16 (0.46)</td>
<td>2.14 (0.55)</td>
<td>0.001</td>
</tr>
<tr>
<td>Diameter stenosis before (%)</td>
<td>70.79 (15.62)</td>
<td>70.85 (13.79)</td>
<td>0.848</td>
</tr>
<tr>
<td>Diameter stenosis after (%)</td>
<td>13.81 (6.23)</td>
<td>25.02 (11.76)</td>
<td>0.001</td>
</tr>
<tr>
<td>Balloon/vessel ratio</td>
<td>1.09 (0.32)</td>
<td>1.09 (0.28)</td>
<td>0.212</td>
</tr>
</tbody>
</table>

*Differences between the two treatment groups.

available standard fluorescent microbeads (Calibrite, Becton-Dickinson). Fluorescence and light scatter data were supplied in lognormal mode and platelets were identified by means of their characteristic size and granularity in the forward versus right angle light scatter plot. For immunofluorescence analysis a narrow gate was set around the platelet population to exclude red cells and cell fragments or background machine noise as described. Samples were analysed at a flow rate of 200 to 300 events per second and a total of 5000 events was acquired and analysed using the FACScan scientific program (Becton-Dickinson). Platelet antigen expression was expressed as relative mean particle fluorescence intensity of total platelet population (GPIIb-IIIa) or as percentage of marker positive platelets (LIBS1, CD62, fg, vWF). The platelet population evaluated was found to be >98% positive for the platelet-specific CD41 antigen.

MONOCLONAL ANTIBODIES
All mAbs used in this study were either commercially obtained as FITC-conjugates or were labelled with the fluorescent dye according to standard methods. Anti-CD41 (Dianova, Hamburg) is raised against the glycoprotein complex IIb-IIIa and detects the receptor in its resting or activated forms. Anti-LIBS1 mAb (generously provided by Dr Mark Ginsberg, Scripps Clinic, La Jolla, CA) recognises a cryptic epitope on GPIIIa that becomes exposed only on the activated and ligand-occupied GPIIb-IIIa complex. Thus anti-LIBS1 binding indicates fibrinogen receptor activity on the platelet surface. Anti-fg mAb recognises the E-fragment of the fibrinogen molecule and was used to detect fg molecules bound to the activated platelet surface. Anti-vWF mAb is directed against human von Willebrand factor and was used to determine von Willebrand factor bound to platelets. Anti-CD62P recognises the α-granule membrane glycoprotein P-selectin (GMP-140, PAGEM) that is exclusively expressed on the surface of activated platelets and was used as marker for α-degranulation. The antigens determined in the present study and the physiological significance of platelet membrane glycoproteins for platelet function are summarised in figure 1.

STATISTICAL ANALYSIS
To test for differences between values obtained before and then daily after the intervention we used non-parametric tests (Friedman test followed by Wilcoxon rank sum test). Data are presented in tables as median (25, 75th quartile) for N observations. In the figures data are presented as line graphs showing medians and quartiles.

Results
PATIENTS
Between January and May 1994, 30 patients undergoing balloon angioplasty and 30 patients that were treated with coronary stent implantation were studied. Both groups of patients had similar basic demographic characteristics and laboratory variables (tables 1 and 2). In all patients angioplasty of the stenosed artery was successful, resulting in a significant reduction in residual stenosis without angiographically detectable dissection (table 3). Minimal lumen diameter before,
Table 4 Medication given before and after intervention

<table>
<thead>
<tr>
<th>Medication</th>
<th>Before Stent group (n = 30)</th>
<th>PTCA group (n = 30)</th>
<th>After Stent group (n = 30)</th>
<th>PTCA group (n = 30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anti-anginal</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>medication:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aspirin</td>
<td>26</td>
<td>27</td>
<td>29</td>
<td>30</td>
</tr>
<tr>
<td>Ticlopidine</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Unfractionated heparin</td>
<td>30 (14 000 to 43 000 IU/day)</td>
<td>30 (19 000 to 38 000 IU/day)</td>
<td>Until target INR 3.5</td>
<td>For 24 hours</td>
</tr>
<tr>
<td>Anti-anginal</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>medication:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>β-receptor blockers</td>
<td>24</td>
<td>25</td>
<td>29</td>
<td>28</td>
</tr>
<tr>
<td>Nitric (long-acting)</td>
<td>21</td>
<td>23</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

None of the differences between the two groups reached statistical significance (P > 0.05).

Figure 2 Fibrinogen receptor function on circulating platelets in patients undergoing stent placement and PTCA. Graphs show surface expression of GPIIb-IIIa complex and of the activated fibrinogen receptor, and show fibrinogen binding on circulating platelets in patients after coronary stent implantation (n = 30) or balloon angioplasty (n = 30). The relative mean intensity of immunofluorescence is shown and was used as measure of mAb binding and glycoprotein surface expression. An asterisk indicates significance compared with starting values before intervention (bound fibrinogen day 1 to 3, P < 0.001; day 4, P < 0.01; activated fibrinogen receptor day 2 and 3, P < 0.01).

Figure 3 von Willebrand factor binding to circulating platelets in patients undergoing stent placement and PTCA. Graphs show binding of von Willebrand factor to circulating platelets in patients after coronary stent implantation (n = 30) or elective balloon angioplasty (n = 30). An asterisk indicates significance compared with starting values before intervention (day 1 to 4, P < 0.01).

Figure 4 α-Degranulation of circulating platelets in patients undergoing stent placement and PTCA. Surface expression of P-selectin was used as marker of degranulation of α-granules. Graphs show surface exposure of P-selectin on circulating platelets in patients after coronary stent implantation (n = 30) or elective balloon angioplasty (n = 30). An asterisk indicates significance compared with starting values before intervention (day 1 to 4, P < 0.01).

Figure 5 Peripheral platelet count in patients undergoing stent placement and PTCA. Graphs show time course of peripheral platelet count in patients after coronary stent implantation (n = 30) or elective balloon angioplasty (n = 30). An asterisk indicates significance compared with starting values before intervention (day 2 to 4, P < 0.01).

Diameter stenosis before, and balloon/vessel ratio were not different in the two groups (table 3). In the PTCA group the diameter of the treated vessel was smaller and the degree of residual stenosis after the intervention was significantly higher than in the stent group (table 3). Within the observation period none of the patients had an acute ischaemic event that required repeat angioplasty. Twenty six of 30 patients undergoing stenting and 27 of the PTCA group were on long term aspirin treatment (table 4). Similarly, antianginal treatment before the procedure did not differ significantly between the two study groups (table 4). In the post-interventional observation time (4 days) both groups were receiving similar antithrombotic or β-blocker therapy (table 4). There were major differences between groups in the doses of heparin given (table 4).

PLATELET FUNCTION

Fibrinogen receptor function
Fibrinogen binding to platelets increased significantly above starting values the day after stent placement and remained raised for four
days (fig 2). No significant change in fibrinogen binding was observed in the angioplasty (PTCA) group (fig 2). Similar results were found for anti-LIBS1 binding with increased anti-LIBS1 immunofluorescence signals above baseline values on days 2 and 3 whereas no significant change in LIBS1 surface expression was found in the angioplasty (PTCA) group (fig 2). The increase in fibrinogen binding and LIBS1 surface exposure after stent implantation was not due to altered surface expression of glycoprotein IIb-IIIa complexes because binding of anti-CD41 remained unchanged throughout the observation period in both groups of patients (fig 2).

von Willebrand factor binding
As described above for fibrinogen receptor function, the percentage of platelets positive for von Willebrand factor significantly increased above starting values the day after coronary stenting and remained raised thereafter (fig 3). No change in von Willebrand factor binding to circulating platelets was found in the angioplasty group (fig 3).

α-Degranulation
After coronary stenting degranulation of platelet P-selectin increased above starting values peaking at day 2 to 4 (fig 4). In contrast, no change in P-selectin surface expression was noted in the PTCA group (fig 4).

Peripheral platelet count
The day after coronary stenting the peripheral platelet count dropped below pre-stent values and remained decreased throughout the observation period (fig 5). No significant change in platelet count was noted in patients undergoing PTCA (fig 5).

Discussion
The present data show that coronary stenting with subsequent strict anticoagulation results in enhanced platelet activation. The major findings of this study are: (1) After coronary stent implantation fibrinogen receptor activity, von Willebrand factor binding, and surface expression of P-selectin on circulating platelets is increased for days after the procedure. (2) The peripheral platelet count decreases after coronary stenting. (3) No significant change in peripheral platelet function and platelet count was found in patients undergoing angioplasty without stenting. These findings indicate that significant changes in platelet membrane glycoproteins occur in patients with coronary stent implantation. Platelet activation after coronary stenting coincides with a decrease in peripheral platelet count, suggesting increased sequestration of hyperactive circulating platelets. Enhanced platelet activation might be of pathophysiological importance in development of subacute stent thrombosis.

EFFECTS OF CORONARY STENTING ON FIBRINOGEN RECEPTOR ACTIVITY
We found that fibrinogen receptor activity is significantly increased for days in patients after coronary stenting but not after balloon angioplasty (PTCA). Thus platelets circulate in a hyperreactive state after coronary stenting. Since fibrinogen receptor on platelets has a central role in platelet aggregation and thrombus formation, this may contribute to the risk of subacute stent thrombosis. Although the present study does not provide direct evidence, this assumption is supported by studies showing that patients with increased platelet activation before coronary angioplasty are at increased risk of acute ischaemic events after the procedure. Moreover, antagonism of platelet fibrinogen receptor during angioplasty when the risk of coronary thrombosis is increased was shown to reduce acute coronary events significantly.

EFFECTS OF CORONARY STENTING ON VON WILLEBRAND FACTOR BINDING
As well as enhanced fibrinogen receptor activity, we also found increased binding of von Willebrand factor to platelets after stent implantation: no detectable change in von Willebrand factor binding was found in patients undergoing PTCA. Since interaction of von Willebrand factor with platelets is crucial for platelet adhesion the increase in von Willebrand factor binding to platelets implies that platelets circulate in a hyperadhesive state after coronary stenting. Thrombotic plug formation requires von Willebrand factor for two reasons: the glycoprotein is necessary for platelet adhesion to subendothelium and it is also necessary as a carrier for the procoagulant factor VIII molecule, which serves as an essential cofactor in factor Xa generation. We therefore speculate that the increased platelet binding of von Willebrand factor in coronary stent patients might enhance both platelet adhesion to the injured vessel wall and the procoagulant activity of circulating platelets.

EFFECTS OF CORONARY STENTING ON α-DEGRANULATION
P-selectin associated with the platelet membrane was used as marker for platelet degranulation. We found that surface expression of P-selectin on platelets was significantly increased after stenting. Thus coronary stenting is associated with an increased granule secretion and liberation of biologically potent platelet-stored compounds which might trigger biochemical mechanisms involved in restenosis as suggested previously. This includes release of growth factors such as platelet-derived growth factor (PDGF) or tumour growth factor that have been shown to initiate proliferation of smooth muscle cells and fibroblasts and thus have been implicated in the pathophysiology of restenosis after angioplasty.

PATHOPHYSIOLOGICAL CONSIDERATIONS
Several factors that might have an impact on platelet function may account for the differences between the two patient groups: (1) prolonged and intensified antithrombotic therapy,
Changes in membrane glycoproteins of circulating platelets after coronary stent implantation

(2) degree of vascular injury at the puncture site after angioplasty, (3) contact between circulating blood and an artificial surface, and (4) bleeding complications.

Because heparin has been shown to induce platelet activation in vitro and in vivo,70 the enhanced platelet activity described in stent patients might be at least partially due to prolonged administration of high doses of heparin. Comparative studies using different anticoagulation regimens may help to clarify this. Furthermore, in our patients we performed coronary stent implantation because of a suboptimal immediate result after PTCA. Thus enhanced platelet activation after stenting might be a consequence of increased vessel injury at the plaque. We and others showed that the atherosclerotic lesion is a major source of platelet activation during PTCA.79 Although it is the least likely factor, the contact of circulating blood with the artificial surface of stents might trigger activation of the haemostatic system.

STUDY LIMITATIONS

The study of platelet function is always hampered by the possibility of artefactual platelet activation in vitro during blood sampling or further processing. In our study these variables were kept constant in all the individuals investigated. The reproducibility of the platelet assay was confirmed by evaluation of samples obtained from healthy controls on the day of patient evaluation.

The platelet assay described here has been shown to be useful in evaluating platelet function in various diseases.66-68 We did not use fixed samples since we (unpublished observation) and others23 found that the immunoreactivity of platelet membrane glycoproteins was significantly decreased after fixation. Moreover, the fixative paraformaldehyde increases platelet degranulation.24

The present study was designed to evaluate platelet function in patients undergoing coronary stenting who were treated with conventional anticoagulation. At present we are not able to draw definite conclusions about what causes platelets to become activated in patients with coronary stents. Specifically, we cannot rule out that differences in angiographic results (for example, less obvious opening up of a stenosis) may be partly due to the observed differences in platelet activation in both groups of patients.

THERAPEUTIC IMPLICATIONS AND CONCLUSION

Platelets play a central part in acute coronary thrombotic events.12,13 The results that we describe question the effectiveness of currently applied antithrombotic regimens in coronary stent patients. The increase in platelet activation after coronary stenting emphasises the need for potent antplatelet regimens in patients undergoing coronary stenting. Platelet membrane glycoproteins are increasingly becoming the target of novel antplatelet drugs. Fibrinogen receptor antagonism in high risk PTCA procedures has been shown to reduce early ischaemic complications.71 Moreover, recent experimental data suggest that pharmacological interference in the interaction of von Willebrand factor with platelets inhibits thrombotic plug formation in vivo.75 Thus the changes in specific platelet membrane glycoproteins described here may help to develop new antithrombotic strategies in patients with stent implantation.

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