**HEMATOPOIESIS**

**Influence of Monoclonal Antiplatelet Glycoprotein Antibodies on In Vitro Human Megakaryocyte Colony Formation and Proplatelet Formation**

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The influence of antiplatelet glycoprotein (GP) antibodies on megakaryocytopenesis in patients with idiopathic or immune thrombocytopenic purpura (ITP) has been well studied. However, the influence of GP antibodies on proplatelet formation is poorly understood. Here we investigated whether in vitro human megakaryocyte colony formation and proplatelet formation are affected by various monoclonal antiplatelet GP antibodies (MoAb). The megakaryocyte colony formation inhibition assay was performed by methylcellulose culture with modifications, using peripheral blood nonadherent mononuclear cells. The proplatelet formation inhibition assay was performed by megakaryocytes derived from CD34+ cells, stimulated with thrombopoietin + stem cell factor, which were then incubated with antiplatelet GP antibodies on proplatelet formation, however, is poorly understood.

In this study, we investigated whether in vitro megakaryocyte colony formation and proplatelet formation are affected by monoclonal antibodies (MoAb) raised against various platelet antigens, using human megakaryocytes, derived from peripheral blood progenitors.

**MATERIALS AND METHODS**

**Human Subjects**

Peripheral blood was obtained from healthy adult volunteers with their informed consent. Serum and plasma were stored at −80°C until use.

**MoAbs for Inhibition Assays**

Mouse IgG, MoAbs, specific for GP-IIIb (CD41; clone 5B12), GP-IIIa (CD61; clone Y2/51), and a negative control (clone DAK-GO1), were purchased from Dako (Glostrup, Denmark). An anti-GP-IIIb MoAb (CD42b; IgG1, clone HIP1) was purchased from PharMingen (San Diego, CA), and an anti-integrin αIbβ3 MoAb (CD51/CD61; IgG1, clone 23C6) was purchased from Southern Biotechnology Associates Inc (Birmingham, AL).

These antibodies contained sodium azide, which we removed by dialysis as described below. After dialysis, the influence of the remaining sodium azide was tested in the megakaryocyte colony formation inhibition assay cell cultures.

**Dialysis of antibodies for inhibition assays.** Antibodies were dialedyzed against Iscove’s modified Dulbecco’s medium (IMDM; Gibco BRL, Life Technologies, Inc, Rockville, MD) using a dialysis cassette (#6450, Slide-A-Lyzer; Pierce Chemical Co, Rockford, IL) for 20 hours. Anti-GP-IIIb MoAb (CD42b; HIP1) slightly inhibited megakaryocyte colony formation (P<.05), and strongly inhibited proplatelet formation (after 24 hours incubation, P<.0002; after 48 hours incubation, P<.00007). Anti-GP-III MoAb (CD41; 5B12) inhibited only proplatelet formation (only after 24 hours incubation, P<.05). Anti-integrin αIbβ3 MoAb (CD51/CD61; 23C6) only slightly inhibited colony size (P<.05). However, anti-GP-IIIa MoAb (CD61; Y2/51) did not inhibit either colony formation or proplatelet formation. These results suggest that antiplatelet GP MoAbs have differing effects on in vitro megakaryocyte colony formation and proplatelet formation.

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hours at 4°C. Dialyzed antibodies were sterilized using an ultracleaning filter (Milllex-GV130S; Millipore Co, Bedford, MA) and were quantified by a sandwich enzyme-linked immunosorbent assay (ELISA). 

**ELISA**

The ELISA plates (EIA Flat Plate I; Sanko Junyaku, Tokyo, Japan) were coated with 1 µg/mL goat affinity-purified F(ab')2 antirat IgG (ICN Pharmaceuticals Inc, Costa Mesa, CA) overnight at 4°C. The plates were then washed with phosphate-buffered saline containing 0.05% Tween 20, pH 7.2 (PBS-Tween). Samples were diluted 1:100, 1:500, and 1:250 with 1% bovine serum albumin (BSA) in PBS-Tween, and the plates were incubated for 1 hour at room temperature. Mouse antihuman von Willebrand factor MoAb (IgG1, clone F8/86, Dako) at known Ig concentrations was used as the standard. After washing, 0.25 µg/mL alkaline phosphatase-conjugated sheep affinity-purified F(ab')2 antirat IgG (ICN Pharmaceuticals Inc) was added to the wells, and the wells were incubated for 1 hour at room temperature. Then, p-nitro phenyl phosphate alkaline phosphatase substrate solution (Sigma, St Louis, MO) was added to each well. After 30 minutes, the reaction was stopped by the addition of 50 µL 4 M NaOH, and the absorbance was measured at 410 nm using an ELISA plate reader (SLeia Reader; Sanko Junyaku).

**Influence of sodium azide on cell culture.** We tested the effect of the sodium azide remaining in the dialyzed antibody preparations on the numbers of colonies, other than the megakaryocyte colony. We did this by using the megakaryocyte colony formation inhibition assay cell cultures, as described below. This assay showed that there was no statistically significant difference between the numbers of colonies in the presence or absence of the antibody preparations in the culture medium.

**Megakaryocyte Colony Formation Inhibition Assay**

**Separation of human peripheral blood nonadherent mononuclear cells.** Peripheral blood mononuclear cells were separated by centrifugation on Lymphoprep (density = 1.077 g/mL; Nycomed, Oslo, Norway) at 800g for 20 minutes, then suspended in IMDM with 10% fetal calf serum (GIBCO BRL), and incubated for 1 hour in plastic culture way) at 800 for 20 minutes, then suspended in IMDM with 10% fetal goat serum, and 1 µg/mL antibody for 14 days at 37°C in humidified 5% CO2 in air.

**Chemical Co, Midland, MI), 1% BSA (AlbuMAX I; GIBCO), 10% 5.5 mol/L 2-mercaptoethanol (2-ME; GIBCO), 1% methylcellulose (Dow Chemical Co, Midland, MI), 1% Insulin-Transferrin-Selenium-X (ITS-X; GIBCO), 5.5 × 10−5 mol/L 2-ME, 1% methylcellulose, 1% BSA, and 10% autologous plasma26 at 37°C in humidified 5% CO2 in air. After 9 days, morphologically identified megakaryocyte colonies were collected 900 µL fresh IMDM supplemented with 1% ITS-X, 5.5 × 10−5 mol/L 2-ME, 1% BSA, and 10% autologous plasma, in a tissue culture chamber slide (#4804 Lab-Tek), using a micropipet. The total number of megakaryocyte colonies collected from dishes was 353.2 ± 43.9 colonies (10.3 ± 0.1 megakaryocytes per colony), and the total volume of collected medium was 400 µL in each experiment. Therefore, the remaining concentration of cytokines (TPO and SCF) in this liquid culture medium was always 3 ng/mL.

**Inhibition of proplatelet formation.** The collected megakaryocytes were incubated in the chamber slide for 1 hour at 37°C in humidified 5% CO2 in air. Then, megakaryocytes were aliquoted into the wells of a 96-well tissue culture plate (#430247; Corning Costar, Cambridge, MA) for proplatelet formation. The number of cells evaluated in each well was 360.5 ± 148.7, depending on the individual preparation of cells used. Then, each antibody was added to the wells at a concentration of 10 µg/mL. Megakaryocytes were incubated for 24 or 48 hours at 37°C in humidified 5% CO2 in air. Cells and megakaryocytes with proplatelet formation were counted using an inverted microscope at 100× or 200× magnification. Proplatelets were identified as described previously.26 After the counting was completed, the cells were fixed with buffered formalin-acetone for 30 seconds at 4°C. The cells were exposed to the anti-GP-IIIa MoAb for 30 minutes at room temperature, followed by staining with LSAB kit, and the number of GP-IIIa-positive cells were counted using an inverted microscope.

**Evaluation of inhibition of proplatelet formation.** The percentage of megakaryocytes with proplatelet formation per well (%PFP) was calculated as the number of megakaryocytes with proplatelet formation divided by the number of GP-IIIa-positive cells times 100. The proplatelet formation activity is expressed using the following formula: proplatelet formation activity (%) = (sample %PFP/negative control %PFP) × 100.

**Statistical Analysis**

Statistical significance was determined using the Student’s t-test.

**RESULTS**

**Influence of Antiplatelet GP MoAbs on Megakaryocyte Colony Formation**

The influence of antiplatelet GP MoAbs on megakaryocyte colony formation is shown in Fig 1. The number of megakaryocyte colonies and the average colony size were not significantly different between the culture medium control and the negative control. Anti-GP-IIb antibody slightly inhibited colony formation by megakaryocytes (P < .05) compared with the controls, whereas the anti-GP-IIb, anti-integrin α5β1, and anti-GP-IIIa
Fig 1. Influence of antiplatelet GP MoAbs on megakaryocyte colony formation. Peripheral blood nonadherent mononuclear cells (2.5 x 10^5 cells/mL) were cultured in IMDM supplemented with 10 ng/mL TPO, 10 ng/mL SCF, and 1 µg/mL each antiplatelet GP MoAb for 14 days. The number of colonies (A) and colony size (B) were counted using an inverted microscope. The data are the means ± standard error of mean (SEM) from duplicate cultures in two experiments using cells from three donors. *Significantly different compared with the controls (P < .05).

Fig 2. Spontaneous proplatelet formation by megakaryocytes in methylcellulose culture medium. Peripheral blood CD34+ cells were cultured with 10 ng/mL TPO and 10 ng/mL SCF in 35-mm plastic dishes. On day 9 of the culture, megakaryocytes with spontaneous proplatelet formation were observed under an inverted microscope (bar, 50 µm).
antibodies did not inhibit colony formation. Only the anti-integrin αvβ3 antibody slightly reduced the average megakaryocyte colony size (P, 0.05), compared with the controls.

**Influence of Antiplatelet GP MoAbs on Proplatelet Formation**

We first determined the culture time for the preparation of megakaryocytes. Peripheral blood CD34+ cells (4.5 × 10^3 cells/mL) were cultured with 10 ng/mL TPO and 10 ng/mL SCF in 35-mm plastic dishes for 14 days. Megakaryocyte colonies that contained megakaryocytes with proplatelet formation were counted daily on days 1 through 14 of the culture period, using an inverted microscope. Their number peaked between days 9 and 10. Therefore, collection of the megakaryocyte colonies from the culture dishes was determined on the 9th day of culture, before maximal proplatelet production.

We then examined the spontaneous proplatelet formation of the collected megakaryocytes in a 96-well tissue culture plate. Megakaryocyte colonies were derived from peripheral blood CD34+ cells (1.8 ± 0.5 × 10^4 cells/mL) stimulated with 10 ng/mL TPO and 10 ng/mL SCF, and they were collected on the 9th day from culture dishes (343.0 ± 3.0 colonies). Then, collected megakaryocytes were aliquoted into wells of a 96-well tissue culture plate (190.9 ± 11.2 cells per well) and were incubated. The number of megakaryocytes with proplatelet formation was counted after 5, 24, and 48 hours of incubation, respectively, using an inverted microscope. The %PPF calculation is described in detail in the Materials and Methods section. The time 0 corresponds with the 9th day of culture in Fig 3. The purity of the GP-IIIb-positive cells was 98.9% ± 0.3% per well, by indirect immunostaining. The data are the means ± SEM from duplicate incubations of two experiments.

**Inhibition of proplatelet formation by antiplatelet GP MoAbs.**

The influence of antiplatelet GP MoAbs on proplatelet formation is shown in Fig 5. Proplatelet formation activity was not significantly different between the culture medium control and the negative control in any of the incubations. At 24 hours of incubation, anti-GP-Iba or anti-GP-IIIb antibodies inhibited proplatelets in the absence of antibodies (Fig 4). The time 0 corresponds with the 9th day of culture, as shown in Fig 3. The number of proplatelets that developed from megakaryocytes peaked at 24 hours. This peak corresponds with the peak of proplatelet formation shown in Fig 3 (day 10). Thereafter, the total number of proplatelets decreased, fragmenting into smaller platelet-like pieces.
platelet formation by megakaryocytes (P < .0002 and P < .03, respectively), compared with the controls. At 48 hours of incubation, anti-GP-Ibα antibody inhibited platelet formation by megakaryocytes (P < .0007), compared with the controls. During this incubation time, the inhibition of platelet formation by anti-GP-Ibα antibody was not statistically different compared with controls. However, though inhibition by the anti-GP-Ibα antibody decreased a little in comparison with the 24 hours of incubation, this inhibition was still stronger compared with other antibodies. Throughout experimental incubation, formation of morphologically abnormal proplatelets did not occur. The purity of the GP-Ibα-positive cells was 98.6% ± 0.6% per well, as measured by indirect immunostaining (Fig 6).

**DISCUSSION**

We believe that studying the effects of antiplatelet GP MoAbs on cultured megakaryocytes will provide important information about both megakaryocytopoiesis and the role of antiplatelet autoantibodies in ITP. The present study showed that (1) anti-GP-Ibα MoAb (HIP1) slightly inhibited megakaryocyte colony formation and strongly inhibited proplatelet formation, (2) anti-GP-Ibα MoAb (5B12) inhibited only proplatelet formation (only after 24 hours of incubation), (3) anti-integrin αvβ3 MoAb (23C6) slightly inhibited only colony size, and (4) anti-GP-IIIα MoAb (Y2/51) did not inhibit either colony formation or proplatelet formation.

These results raise two important points. First, our megakaryocyte colony formation inhibition assay data imply that the sites recognized by the anti-GP-Ibα and anti-integrin αvβ3 antibodies play a significant role in megakaryocytopoiesis. The anti-GP-Ibα antibody seemed to affect immature cells, because it reduced the number of colonies, whereas the anti-integrin αvβ3 antibody seemed to affect cell proliferation, because the average colony size was reduced. The GP-Ibα-IIIα complex is expressed on the megakaryocytes’ membrane before GP-III in the early stage of maturation.27 Nevertheless, the anti-GP-Ibα and anti-GP-IIIα antibodies did not affect megakaryocytopoiesis. These results suggest that inhibition of megakaryocytopoiesis varies according to the specific anti-GP antibodies used. Our findings may explain conflicting reports8-11 about the effects of antiplatelet GP antibodies on in vitro megakaryocyte colony formation in patients with ITP.

Second, we have shown that antiplatelet GP antibodies affect not only megakaryocyte maturation,2 as reported previously, but also proplatelet formation. Furthermore, the same antiplatelet GP MoAb affected megakaryocyte colony formation and proplatelet formation. However, the anti-integrin αvβ3 and anti-GP-IIIα antibodies did not inhibit proplatelet formation. These results suggest that inhibition of proplatelet formation varies according to the specific anti-GP antibodies used. The anti-GP-Ibα and anti-GP-Ibβ MoAbs had a more dramatic effect on proplatelet formation than the other antibodies. In particular, the former MoAb, anti-GP-Ibα, strongly and persistently inhibited proplatelet formation. In contrast, inhibition by the latter MoAb, anti-GP-Ibβ, was not persistent. We think that these influences do not delay the kinetics, but the inhibition of proplatelet formation, because proplatelet formation activity did not increase more than the controls after 48 hours of incubation. If this inhibition was to occur in vivo in patients with ITP, platelet production would be inhibited, and these patients might develop severe thrombocytopenia.

In patients with ITP, platelets are frequently bound to endothelial system.1 Assays that measure antiplatelet antibodies have been developed, and patients with ITP usually have increased levels of PAIgG. In most cases, there is a significant correlation between the PAIgG level and the platelet count9; however, this correlation does not always hold true29-31: the PAIgG level is low in patients with severe thrombocytopenia and is high in patients with slight thrombocytopenia. To account for this finding, there are three points to consider in relation to the results of our study. First, when antibodies such as HIP1,
that slightly inhibit colony formation and strongly inhibit proplatelet formation, are present in the patients’ plasma, severe thrombocytopenia may develop. Second, when antibodies such as 5B12, that only inhibit proplatelet formation, or antibodies such as 23C6, that only slightly inhibit colony size, are present in the patients’ plasma, moderate thrombocytopenia may develop. Finally, when antibodies such as Y2/51, that do not inhibit colony formation or proplatelet formation, are present in the patients’ plasma, slight thrombocytopenia may develop. However, these hypotheses are somewhat inconsistent with previous investigators’ reports, because the proliferation of megakaryocytes is not depressed by antiplatelet antibodies in patients with ITP. In fact, the number of megakaryocytes in the bone marrow of ITP patients does not decrease, and the number of megakaryocyte colony-forming cells are increased by the injection of platelet antiserum, in vivo. Therefore, in vivo studies indicate that the inhibition of colony formation is unlikely to be the leading cause of thrombocytopenia.

As for the platelet production in patients with ITP, there are differences of opinion among specialists. Branehog et al and Harker et al reported platelet production (platelet turnover) to be increased in ITP patients. According to Harker et al, the total megakaryocyte mass increases 2 to 8 times from normal, indicating that thrombopoiesis is effective in ITP. In contrast, more recent studies of autologous platelet turnover have shown that platelet production is not increased in all ITP patients. Ballem et al have reported that 30% of ITP patients show decreased platelet production, 43% have production rates within the normal range, and 27% have increased platelet production. Similarly, Tomer et al have shown that the platelet turnover is less than normal in some ITP patients. Stoll et al have shown that the rate of platelet production is not increased in most patients with moderate ITP. Thus, the role of platelet production in the pathogenesis of ITP is controversial. We estimate that if proplatelet formation is inhibited by antiplatelet antibodies, in vivo, then platelet production may be depressed. In particular, the reports by Ballem et al and Tomer et al are not inconsistent with our data on inhibited proplatelet formation and may explain depressed platelet production in patients with ITP. In other words, the cause of thrombocytopenia may be a result of the inhibition of proplatelet formation of megakaryocytes by the antiplatelet antibodies, as well as of the destruction of the platelets in the reticuloendothelial system. It is necessary, however, that more studies be performed using various approaches, because the inhibition of proplatelet formation in patients with ITP is poorly understood.

The results of this study suggest that the degree of thrombo-
cytopenia may be related to the epitope recognized by the patients’ antibodies, because the antibodies we used were MoAbs. Nagasawa et al. reported that the proliferation of CFU-Meg was suppressed by an anti-GP-IIb/-IIIa MoAb (TM83), whereas the effect of an anti-GP-Ib MoAb (TM60) was negligible. Handagama et al. made a heterologous antiplatelet antibody and reported that rat megakaryocyte proplatelet formation is inhibited by the antibody. However, which epitope was recognized by the antibody is unknown, because the antibody used was polyclonal. The effects on guinea pig megakaryocyte proplatelet formation by antiplatelet MoAbs have been studied previously, and an anti-integrin αvβ3 MoAb (LM609), an anti-integrin αv MoAb (LM142), and an anti-GP-IIb/IIIa MoAb (PG2) each inhibited proplatelet formation. We suspect that the differences between the results of our study and those of other studies are a result of the antibodies used, because they recognize different epitopes on the same GP. Unlike previous studies, we used the same MoAb to examine both colony formation and proplatelet formation. Thus, we were able to compare inhibition of colony formation and the inhibition of proplatelet formation.

In conclusion, our in vitro study has shown that both human megakaryocyte colony formation and proplatelet formation are inhibited by an anti-GP-Ibα antibody, suggesting that GP-Ibα plays an important role in megakaryocytopoiesis. Several investigators have reported that thrombocytopoiesis is more severe in patients with anti-GP-Ib autoantibodies, whereas ITP patients with anti-GP-IIb/IIIa autoantibodies do not develop severe thrombocytopoiesis. Moreover, Hasegawa et al. reported that antiplatelet GP-Ib antibody may impair platelet production by megakaryocytes in ITP. These clinical reports strongly support our experimental findings. However, the inhibition of colony formation in vitro did not agree with in vivo studies because the number of megakaryocytes in the bone marrow of ITP patients did not decrease. Also, the megakaryocyte colony-forming cells increased when platelet antiserum was injected. From the above mentioned facts and discussion, the inhibition of proplatelet formation by the antiplatelet antibodies may be at the basis of thrombocytopoiesis in patients with ITP, in which depressed platelet production occurs. Our present findings suggest that autoantibodies against different platelet GPs in patients with ITP may have differing effects on proplatelet formation. This subsequently results in the development of different degrees of thrombocytopoiesis. Further studies are needed to obtain in vitro evidence that proplatelet formation is inhibited by antiplatelet serum, and the antiserum must be analyzed.

ACKNOWLEDGMENT

The authors thank Dr. D. B. Douglas for comments on the manuscript. The authors appreciate technical assistance in ELISA from Mr. H. Miyazawa.

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