

quickly to $10 \times 10^9 \text{ L}^{-1}$. She was treated, successively, with *Helicobacter pylori* eradication, danazol, azathioprine, and cyclosporine, but her platelets remained $<10 \times 10^9 \text{ L}^{-1}$, and she required weekly platelet transfusions. In August 2003, we tried rituximab at a dose of 375 mg/m^2 weekly for 4 doses. The patient's platelet count rose to $>100 \times 10^9 \text{ L}^{-1}$ within 5 weeks and remained at this level for 9 weeks thereafter. However, she relapsed 13 weeks after the rituximab treatment, and was again transfusion-dependent, despite treatment with high-dose dexamethasone (40 mg for 4 days), prednisolone (1 mg/kg), vincristine, danazol, and azathioprine. She received a second rituximab therapy in December 2003, which resulted in a partial platelet response (platelets $30\text{--}50 \times 10^9 \text{ L}^{-1}$) that eliminated the need for platelet transfusion. A monthly pulse cyclophosphamide (300–500 mg) was added, and the patient's platelet count remained $>50 \times 10^9 \text{ L}^{-1}$ for the next 10 months. In November 2004, the patient developed gastrointestinal bleeding and her platelet count was $<10 \times 10^9 \text{ L}^{-1}$, requiring IVIG treatment and platelet transfusion. A third rituximab treatment was given in March 2005, but there was no platelet response. Since then, she has received biweekly IVIG (a single dose of 0.4 g/kg) to maintain a safe platelet count.

Immunologic analyses

Platelet-associated IgG anti-GPIIb/IIIa antibodies and circulating IgG anti-GPIIb/IIIa antibody-producing cells were measured using, respectively, an enzyme-linked immunosorbent assay (ELISA) [10] and an enzyme-linked immunospot (ELISPOT) assay [11]. Human antichimeric antibodies (HACA) were measured using ELISA [12]. Flow cytometric analysis was performed using the following mouse monoclonal antibodies, which were conjugated to fluorescein isothiocyanate, phycoerythrin, or allophycocyanin: anti-CD3, anti-CD138 (Beckman Coulter, Fullerton, CA), anti-CD19 (Sigma, St. Louis, MO); anti-CD20 (BD Biosciences, San Diego, CA), and anti-CD34 (Miltenyi Biotech, Bergisch Gladbach, Germany). Negative controls were cells incubated with an isotype-matched mouse monoclonal antibody to an irrelevant antigen. The cells were analyzed on an FACS[®] Calibur flow cytometer (BD Biosciences) using CellQuest software. In some experiments, peripheral blood mononuclear cells (PBMC) depleted of CD19⁺, CD20⁺, CD138⁺, or CD34⁺ cells were used for the ELISPOT assay. The cell depletion was performed using magnetic bead-coupled anti-CD19, anti-CD20, or anti-CD34 monoclonal antibodies (Miltenyi Biotech) or anti-CD138 monoclo-

nal antibody (Beckman Coulter) in combination with magnetic bead-coupled anti-mouse IgG antibodies (Miltenyi Biotech), followed by separation on a MACS column. Flow cytometric analysis revealed that cells positive for corresponding markers were nearly absent in the treated fractions ($<0.5\%$ for CD19⁺ and CD20⁺ cells, and $<0.01\%$ for CD138⁺ and CD34⁺ cells).

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Excessive exposure to anionic surfaces maintains autoantibody response to β_2 -glycoprotein I in patients with antiphospholipid syndrome

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Antiphospholipid syndrome (APS) is an autoimmune prothrombotic disorder associated with autoantibodies to phospholipid (PL)-binding proteins, such as β_2 -glycoprotein I (β_2 GPI). We have recently reported that binding of β_2 GPI to anionic PL facilitates processing and presentation of the cryptic β_2 GPI epitope that activates pathogenic autoreactive T cells. To clarify mechanisms that induce sustained presentation of the dominant antigenic β_2 GPI determinant in patients with APS, T-cell proliferation induced by β_2 GPI-

treated phosphatidylserine liposome (β_2 GPI/PS) was evaluated in bulk peripheral blood mononuclear cell cultures. T cells from patients with APS responded to β_2 GPI/PS in the presence of immunoglobulin G (IgG) anti- β_2 GPI antibodies derived from APS plasma, and this response was completely inhibited either by the depletion of monocytes or by the addition of anti-Fc γ RI antibody. These findings indicate that efficient presentation of the cryptic determinants can be achieved by monocytes undergoing

Fc γ RI-mediated uptake of β_2 GPI-bound anionic surfaces in the presence of IgG anti- β_2 GPI antibodies. Finally, β_2 GPI-bound oxidized LDL or activated platelets also induced the specific T-cell response. Continuous exposure to these anionic surfaces may play a critical role in maintaining the pathogenic anti- β_2 GPI antibody response in patients with APS. (Blood. 2007;110:4312-4318)

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Introduction

Antiphospholipid syndrome (APS) is an autoimmune disorder characterized by arterial and venous thrombosis as well as recurrent intrauterine fetal loss in the presence of antiphospholipid antibodies.¹ β_2 -glycoprotein I (β_2 GPI) is the most common antigenic target recognized by the antiphospholipid antibodies, and anti- β_2 GPI antibodies are shown to be strongly associated with thrombosis and other clinical manifestations of APS.²⁻⁴ β_2 GPI is a plasma protein that binds various anionic substances, including phospholipids (PLs), lipoproteins, and activated platelets and endothelial cells.⁵⁻⁷ Several lines of evidence accumulated from animal models suggest that anti- β_2 GPI antibodies are directly involved in the pathogenic processes of APS.^{8,9}

We have recently identified CD4⁺ T cells responsive to β_2 GPI in patients with APS.¹⁰⁻¹² β_2 GPI-reactive T cells can promote production of pathogenic immunoglobulin G (IgG) anti- β_2 GPI antibodies from autologous B cells in vitro. These T cells respond to bacterially expressed recombinant β_2 GPI fragments and chemically reduced β_2 GPI, but fail to respond to native β_2 GPI,¹⁰ indicating that the epitopes recognized by β_2 GPI-reactive T cells are cryptic determinants that are not generated through processing of native β_2 GPI under normal circumstances. One of the major cryptic determinants recognized by β_2 GPI-reactive T cells is the region spanning amino acids (AAs) 276-290, which contains the major PL-binding site at AA 281-288,^{13,14} in the context of HLA-DRB4*0103 (DR53).¹¹ In our recent study employing β_2 GPI-reactive CD4⁺ T-cell clones generated from patients with APS, dendritic cells or macrophages pulsed with β_2 GPI-bound phosphatidylserine (PS) liposome induced a response of T-cell clones specific for a peptide encoding AA 276-290 (p276-290) in HLA-DR-restricted and antigen-processing-dependent manners. In contrast, those pulsed with β_2 GPI or PS liposome alone failed to induce a response.¹⁵ Together these findings indicate that specialized antigen-presenting cells (APCs) capturing β_2 GPI-coated anionic PLs efficiently present a disease-relevant cryptic T-cell determinant of β_2 GPI as a result of antigen processing.

In patients with APS, anti- β_2 GPI antibody levels are usually stable for many years. However, it remains unclear what mechanisms are responsible for the sustained presentation of the dominant cryptic β_2 GPI determinant that activates β_2 GPI-reactive T cells to subsequently produce pathogenic anti- β_2 GPI antibodies. To elucidate these mechanisms, we examined the cellular and molecular factors required for the sustained activation of β_2 GPI-reactive T cells in patients with APS.

Patients, materials, and methods

Patients and controls

This study examined 5 patients, and all fulfilled the revised Sapporo criteria for APS proposed by the International Workshop.¹⁶ These patients were selected based on the presence of DRB4*0103 (DR53), which is known to present a p276-290 peptide to T cells,¹¹ and positive IgG anti- β_2 GPI antibody. The HLA class II alleles, including DRB1 and DRB4, were determined by restriction fragment length polymorphisms combined with

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locus-specific polymerase chain reaction using peripheral blood granulocyte-derived genomic DNA as a template.¹⁷ IgG anti- β_2 GPI antibody levels were measured with a commercial enzyme-linked immunosorbent assay (ELISA) kit (Yamasa, Choshi, Japan) using immobilized β_2 GPI-cardiolipin complex as an antigen source. A commercial kit based on Russell viper venom test (Gradipore, Sydney, Australia) was used to determine the presence of lupus anticoagulant. At the time of blood examination, all the patients were taking low-dose corticosteroids (< 10 mg/day) and low-dose aspirin. Peripheral blood from healthy volunteers was also used as a control source of plasma. All samples were obtained after the patients and control subjects gave their written informed consent in accordance with the Declaration of Helsinki. The study protocol was approved by Keio University International Review Board.

Antigen preparations

Human β_2 GPI was purified from normal pooled plasma,¹⁸ and reduced β_2 GPI was prepared by incubating β_2 GPI with dithiothreitol as previously described.¹⁰ We generated a panel of recombinant maltose-binding protein (MalBP) fusion proteins expressing full-length β_2 GPI (GP-F), domains I and II (GP1), domains III and IV (GP2), and domains IV and V (GP3).¹⁰ MalBP alone was prepared as a control antigen. Two 15-mer peptides, p276-290 and a peptide encoding AA 306–320 of human β_2 GPI (p306-320), were synthesized using a solid-phase multiple synthesizer (Advanced ChemTech, Louisville, KY).¹¹

Liposome containing bovine brain-derived PS (Sigma, St Louis, MO), with a composition of dioleoylphosphatidylcholine (Avanti Polar Lipids, Alabaster, AL) at a molar ratio of 3:7, was prepared and adjusted to a final concentration of 1 μ mol/mL.^{19,20} Low density lipoprotein (LDL) was isolated from freshly prepared normal human plasma by ultracentrifugation, and oxidized LDL (oxLDL) was prepared by incubating LDL with 5 μ M CuSO₄ for 8 hours at 37°C.²⁰ LDL and oxLDL were adjusted to 100 μ g/mL of apoB equivalent. Human platelets were separated from platelet-rich plasma using a modified gel filtration method²¹ to minimize their activation during an isolation procedure. Resting platelets were then activated by incubation with bovine thrombin (1 U/mL; Mochida, Tokyo, Japan) for 15 minutes. All preparations were incubated with or without native β_2 GPI (100 μ g/mL) for 30 minutes at room temperature immediately prior to use in the cultures.

Cell preparations

Peripheral blood mononuclear cells (PBMCs) were isolated from heparinized venous blood by Lymphoprep (Fresenius Kabi Norge AS, Oslo, Norway) density-gradient centrifugation. In some experiments, PBMCs were depleted of CD14⁺ monocytes or CD19⁺ B cells by incubation with anti-CD14 or anti-CD19 monoclonal antibody (mAb)-coupled magnetic beads (Miltenyi Biotecch, Bergisch Gladbach, Germany), respectively, followed by magnetic cell sorting column separation according to the manufacturer's protocol.

Preparation and depletion of IgG from plasma

The IgG fraction was purified or depleted from plasma samples using HiTrap protein G (Amersham Biosciences, Uppsala, Sweden) as described previously.²² Purity of IgG fractions was confirmed to be more than 95% by sodium dodecyl sulfate-polyacrylamide gel electrophoresis, followed by densitometry on Coomassie blue-stained gels. In some experiments, purified IgG was treated with pepsin to prepare F(ab')₂ using a Fab2 preparation kit (Pierce Biotechnology, Rockford, IL). We also prepared IgG fractions depleted of antibodies specific to β_2 GPI. Briefly, purified IgG samples were treated 3 times with cardiolipin-coated 96-well immunoplates (Nunc F96Maxisorp, Roskilde, Denmark), which were preincubated with β_2 GPI or phosphate-buffer saline for 30 minutes. The supernatants were then collected as anti- β_2 GPI antibody-depleted or mock-treated IgG. Removal of anti- β_2 GPI antibody was confirmed by complete loss of antibody reactivity on the anti- β_2 GPI antibody ELISA.

Assays for antigen-specific T-cell response

Antigen-specific T-cell proliferation in the primary cultures was assayed as described previously¹⁰ with some modifications. Briefly, PBMCs (10⁵/well) were cultured with or without antigen in 96-well flat-bottomed culture plates for 7 days. RPMI 1640 supplemented with either 10% fetal bovine serum (FBS; JRH Bioscience, Lenexa, KS) or 8% platelet-poor plasma, which was derived from patients with APS and healthy donors, was used as medium. Prior to use, FBS and plasma samples were heat-inactivated and depleted of β_2 GPI by passing the samples through a HiTrap Heparin column (Amersham Biosciences) twice, to eliminate the potential influence of intrinsic β_2 GPI on the generation of the antigenic peptides. ³H-thymidine (0.5 μ Ci [0.0185 MBq]/well) was added to the cultures during the final 16 hours. The cells were harvested, and ³H-thymidine incorporation was measured in a Top-Count microplate scintillation counter (Packard, Meriden, CT). Native β_2 GPI, reduced β_2 GPI, GP-F, GP1, GP2, GP3, and MalBP were used as antigens at a concentration of 10 μ g/mL. In addition, PS liposome (0.1 μ mol/mL), LDL, oxLDL (10 μ g/mL apoB equivalent), resting platelets, or activated platelets (10⁶/well) were added to the cultures, with or without preincubation with β_2 GPI. To exclude nonspecific unresponsiveness of T cells, all experiments included a culture with phytohemagglutinin at a final concentration of 1 μ g/mL. In some experiments, purified IgG, F(ab')₂, or anti- β_2 GPI antibody-depleted or mock-treated IgG was added at the initiation of the culture. Anti-Fc γ RI (clone 10.1; R&D Systems, Minneapolis, MN), anti-HLA-DR (clone L243; Leinco Technologies, Baldwin, MO), or isotype-matched control mAb was also added to the culture at a final concentration of 2.5 μ g/mL. All experiments were carried out in duplicate or triplicate, and the values are the mean counts per minute (cpm) plus or minus the standard deviation of multiple determinations. In some instances, a T-cell response specific to β_2 GPI-treated PS liposome (β_2 GPI/PS) was expressed as the ratio of cpm in the culture with β_2 GPI/PS to cpm in the culture with PS liposome alone.

Secondary stimulation of peripheral blood T cells was also performed as described.¹⁰ PBMCs were primed with β_2 GPI/PS in medium supplemented with 8% autologous plasma for 10 days. Viable cells were then cultured for an additional 3 days in the presence of 50 U/mL recombinant interleukin-2 (Biogen Idec, San Diego, CA) and irradiated (3000 rad) autologous monocyte-derived dendritic cells in medium supplemented with 10% FBS in the absence or presence of β_2 GPI, reduced β_2 GPI, GP-F, GP1, GP2, GP3, MalBP (10 μ g/mL), p276-290, or p306-320 (5 μ g/mL). Frequencies of β_2 GPI-reactive T cells in peripheral blood T cells were estimated by limiting dilution analysis using GP-F as an antigen.²³ The recognition of p276-290 by peripheral blood T cells was determined based on the specific response to p276-290 by at least 2 T-cell clones established by repeated stimulation of peripheral blood T cells with GP-F.¹¹

Results

Clinical and immunologic characteristics of patients with APS

As shown in Table 1, all patients with APS had thrombosis and/or loss of pregnancy, and were positive for lupus anticoagulant. IgG anti- β_2 GPI antibody titer was high in all but one patient (APS1). Frequencies of β_2 GPI-reactive T cells were variable among patients, and ranged from 2.9 to 12.4 per 10⁴ peripheral blood T cells. In addition, T-cell recognition of p276-290 was detected in all 3 patients examined.

T-cell response induced by β_2 GPI/PS in PBMC cultures

We first examined the responses of peripheral blood T cells to β_2 GPI/PS using regular medium supplemented with FBS (Figure 1A). T cells from all 5 patients responded to GP-F, but failed to proliferate in the presence of β_2 GPI/PS. Interestingly, a T-cell response to β_2 GPI/PS, as well as to GP-F, was detected when a patient's autologous plasma was used instead of FBS to supplement the culture medium. This response was blocked by anti-HLA-DR

Table 1. Clinical and immunologic characteristics of patients with APS analyzed in this study

Patient no.	Age/sex	Thrombosis	Loss of pregnancy	IgG anti- β_2 GPI antibodies (U/mL)†	HLA class II alleles: DRB1	Frequency of β_2 GPI-reactive T cells in circulation/10 ⁴ T cells	Recognition of p276–290 by peripheral blood T cells
APS1	51/F	None	+	16	*1502/*0405	4.5	NT
APS2	43/F	DVT, stroke	+	>120	*0405/*1202	2.9	NT
APS3	46/F	DVT, PE, retinal artery thrombosis	+	>120	*1502/*0901	6.8	+
APS4	47/F	Stroke	+	>120	*1501/*0403	8.1	+
APS9	46/F	DVT, PE, stroke, amaurosis fugax	NA	>120	*0901	12.4	+

All patients were lupus anticoagulant positive; all DRB4 alleles were *0103.

DVT indicates deep venous thrombosis of lower extremity; PE, pulmonary embolism; NA, not applicable; and NT, not tested.

†Normal range less than 3.5 U/mL.

mAb, but not by control mAb (data not shown). However, a β_2 GPI/PS-induced response was not detected in the culture with allogenic plasma from a healthy individual. This finding was reproducible in a total of 7 PBMC samples obtained from 5 patients with APS.

Next, PBMCs from a patient with APS were cultured with β_2 GPI/PS or PS liposome alone in medium supplemented with 2 different lots of FBS, plasma samples from 4 patients with APS, or samples from 3 healthy donors (Figure 1B). The β_2 GPI/PS-specific response was exclusively detected in cultures with autologous and allogenic plasmas derived from patients with APS, although the degree of response was variable among APS plasmas. Analogous findings were obtained with PBMCs from 3 additional patients with APS. In all cases, the lowest response was detected in the culture supplemented with APS1 plasma, which contained low-titer anti- β_2 GPI antibodies.

We next sought to confirm whether T-cell responses induced by β_2 GPI/PS in cultures with APS plasma were specific to β_2 GPI. Peripheral blood T cells primed with β_2 GPI/PS in medium supplemented with autologous plasma were further examined for their reactivity to various β_2 GPI preparations in the secondary culture with FBS (Figure 2). β_2 GPI/PS-primed T cells from all 5 patients specifically responded to reduced β_2 GPI, GP-F, and GP3, indicating a specific recognition of β_2 GPI-derived peptides. More important, the cryptic p276–290 was efficiently presented by APCs in culture with β_2 GPI/PS and APS plasma. T-cell recognition of GP1 was detected in APS2, APS4, and APS9 samples, whereas recognition of GP2 was detected in APS3 and

APS9. Taken together, these findings together indicate that a soluble factor(s) contained in plasma from patients with APS, but not in FBS or plasma from healthy individuals, plays an essential role in activation of β_2 GPI-specific T cells in bulk PBMC cultures with β_2 GPI/PS.

IgG anti- β_2 GPI autoantibody as an essential factor for T-cell recognition of β_2 GPI/PS

Since the degree of the β_2 GPI/PS-specific T-cell response appeared to correlate with IgG anti- β_2 GPI antibody titers, we hypothesized that IgG anti- β_2 GPI antibodies in APS plasma are required for peripheral blood T cells to respond to β_2 GPI/PS. To test this hypothesis, we first prepared IgG-depleted APS plasma samples to evaluate the β_2 GPI/PS-induced T-cell response (Figure 3A). Depletion of IgG from APS plasma resulted in complete loss of the β_2 GPI/PS-induced T-cell response, but addition of autologous IgG back to the IgG-depleted APS plasma restored the response in a dose-dependent fashion. In contrast, addition of IgG prepared from healthy plasma had no effect (data not shown). Interestingly, β_2 GPI/PS-induced T-cell response was also detected in medium supplemented with healthy plasma in the presence of IgG derived from APS plasma. This response was abolished when F(ab')₂ was used instead of intact IgG, indicating an important role of the Fc portion of IgG.

We further examined the effects of depletion of β_2 GPI-specific antibody on the β_2 GPI/PS-induced T-cell response in PBMC cultures with APS IgG (Figure 3B). β_2 GPI/PS-induced T-cell

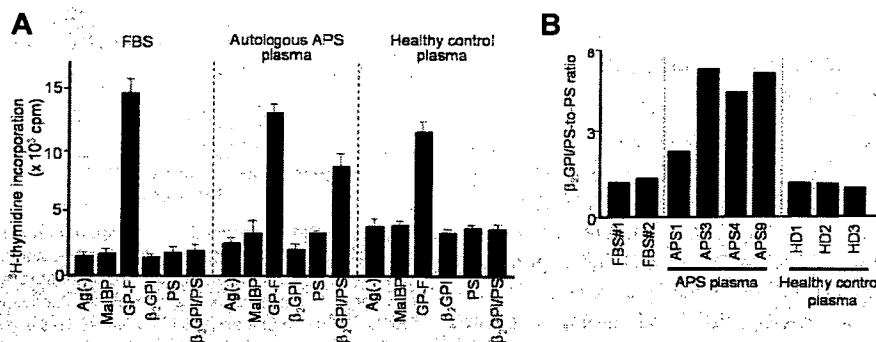


Figure 1. T-cell response to β_2 GPI/PS in bulk PBMC cultures supplemented with FBS, autologous APS plasma, or healthy control plasma. (A) PBMCs from APS4 were cultured in triplicate with or without antigens, including MalBP, GP-F, β_2 GPI, PS, and β_2 GPI/PS, in medium supplemented with FBS, autologous APS plasma, or healthy control plasma. The antigen-induced T-cell proliferative response was assessed by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar) of triplicate measurements. Analogous findings were obtained in 7 independent experiments in PBMCs from all 5 patients with APS. (B) β_2 GPI/PS-specific T-cell response in PBMC cultures of APS4 in medium supplemented with 2 different lots of FBS (no. 1 and no. 2), plasma samples from 4 APS patients (APS1, 3, 4, and 9), or plasma samples from 3 healthy donors (HD1, 2, and 3). β_2 GPI/PS-specific T-cell response was expressed as a β_2 GPI/PS-to-PS ratio, which was the mean cpm incorporated in the triplicate culture with β_2 GPI/PS divided by the mean cpm incorporated in the triplicate culture with PS alone (standard deviations for the individual results were within 20% of the mean in all cases). Similar results were obtained from 3 additional patients with APS (APS1, APS3, and APS9).

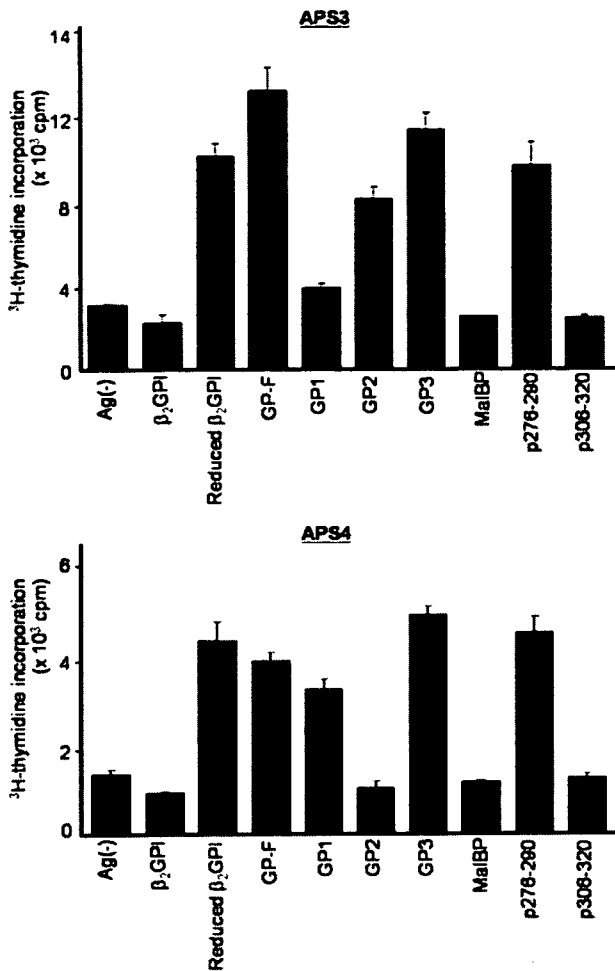


Figure 2. Proliferative responses of β₂GPI/PS-primed T cells to various β₂GPI preparations in secondary cultures. PBMCs from APS3 (top) and APS4 (bottom) were stimulated with β₂GPI/PS for 10 days in medium supplemented with autologous plasma. The viable T cells were then cultured in duplicate with β₂GPI, reduced β₂GPI, GP-F, GP1, GP2, GP3, MalBP, p276-290, or p306-320 in medium containing FBS. After 3 days, ³H-thymidine incorporation was measured. Results are shown as mean (column) and standard deviation (error bar) of duplicate measurements.

response was detected in the presence of mock-treated APS IgG, but completely abolished by depletion of β₂GPI-reactive IgG. These findings indicate that IgG anti-β₂GPI antibodies are required for the T cells of patients with APS to respond to β₂GPI/PS in bulk PBMC cultures.

Roles of β₂GPI/PS-containing immune complex in β₂GPI/PS-induced T-cell response

Since anti-β₂GPI antibodies in sera from patients with APS recognize β₂GPI/PS,²⁰ it is likely that β₂GPI/PS is readily opsonized by IgG anti-β₂GPI antibodies in culture with APS plasma. To evaluate which APCs contained in PBMCs capture this immune complex to induce a specific T-cell response to β₂GPI peptides, we analyzed PBMCs depleted of CD14⁺ monocytes, CD19⁺ B cells, or mock-treated in cultures with β₂GPI/PS and autologous plasma (Figure 4A). The β₂GPI/PS-induced T-cell response was completely inhibited by depletion of monocytes, but was partially suppressed by depletion of B cells, suggesting a primary role of monocytes in our system.

We further evaluated the potential involvement of Fcγ receptors in recognition of the immune complex by monocytes, as the anti-β₂GPI F(ab')₂ was incapable of inducing the T-cell response to

β₂GPI/PS. The β₂GPI/PS-induced T-cell response was completely blocked by anti-FcγRI mAb, but not by control mAb (Figure 4B). Together these findings indicate that efficient β₂GPI/PS-induced T-cell response is achieved by monocytes undergoing FcγRI-mediated uptake of β₂GPI/PS opsonized by IgG anti-β₂GPI autoantibodies.

T-cell response to β₂GPI-treated oxLDL and platelet microparticles

PS liposomes were chemically synthesized, and may not be relevant to patients with APS in vivo. To examine whether anionic substances present in the circulation, such as oxLDL or platelet microparticles, can substitute for PS liposomes in inducing the β₂GPI-specific T-cell response, PBMCs from a representative patient with APS were cultured with various anionic and control substances pretreated with or without β₂GPI in medium supplemented with autologous plasma (Figure 5). OxLDL or activated platelets pretreated with β₂GPI induced a T-cell proliferative

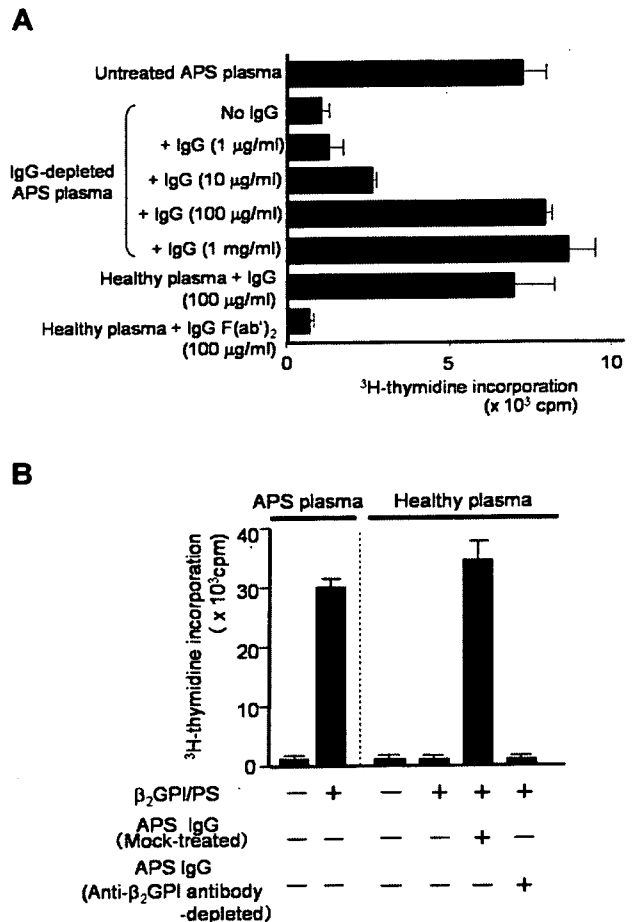


Figure 3. β₂GPI/PS-induced T-cell response in PBMC cultures with or without IgG derived from APS plasma. (A) PBMCs obtained from APS3 were cultured in triplicate with β₂GPI/PS in medium supplemented with untreated or IgG-depleted autologous APS plasma, or healthy plasma. Purified IgG (1 μg/mL-1 mg/mL) or IgG F(ab')₂ (100 μg/mL) from APS3 was added to the cultures. After 7 days, the T-cell proliferative response induced by β₂GPI/PS was measured by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar). Concordant results were obtained with a sample from APS4. (B) PBMCs derived from APS3 were cultured in triplicate with or without β₂GPI/PS in medium supplemented with autologous APS plasma or healthy plasma. An anti-β₂GPI antibody-depleted or mock-treated autologous IgG fraction was added to the initiation of cultures. After 7 days, the T-cell proliferative response was measured by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar). Concordant results were obtained with a sample from APS4.

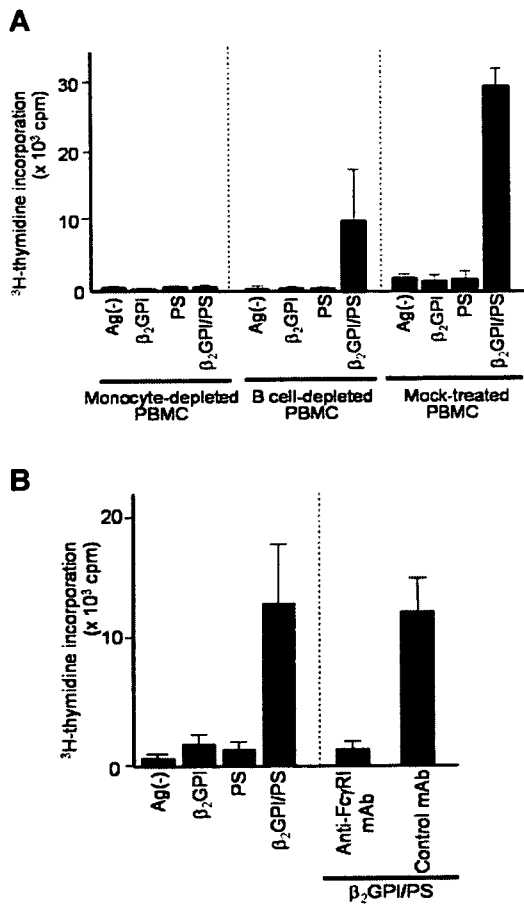


Figure 4. Effects of APC depletion or anti-Fc γ RI mAb on β_2 GPI/PS-induced T-cell response. (A) CD14⁺ monocyte-depleted, CD19⁺ B-cell-depleted, and mock-treated PBMCs derived from APS3 were cultured for 7 days with or without β_2 GPI, PS, or β_2 GPI/PS in medium supplemented with autologous APS plasma, and the T-cell proliferative response was measured by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar) of duplicate measurements. Analogous results were obtained in a total of 4 independent experiments using samples from 3 patients with APS (APS1, APS3, and APS4). (B) PBMCs from APS2 were cultured for 7 days with or without β_2 GPI, PS, or β_2 GPI/PS in medium supplemented with autologous APS plasma. Anti-Fc γ RI or isotype-matched control mAb was added to the initiation of cultures. The T-cell proliferative response was evaluated by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar) of duplicate measurements. Concordant results were obtained with samples from 3 patients with APS (APS2, APS3, and APS9).

response, as observed in cultures with β_2 GPI/PS. These responses were specifically inhibited by anti-HLA-DR mAb (data not shown). Thus, oxLDL and activated platelets can be in vivo sources of anionic surfaces that bind β_2 GPI and promote the efficient presentation of β_2 GPI cryptic peptides by APCs.

Discussion

This study evaluated the potential cellular and molecular mechanisms that induce sustained presentation of the dominant cryptic β_2 GPI determinant that activates β_2 GPI-reactive T cells to subsequently produce pathogenic anti- β_2 GPI antibodies in patients with APS. Here we demonstrate that efficient presentation of cryptic determinants recognized by β_2 GPI-reactive T cells is achieved by monocytes undergoing Fc γ RI-mediated uptake of β_2 GPI/PS opsonized by IgG anti- β_2 GPI antibodies. High avidity IgG anti- β_2 GPI antibodies, which were reported to possess high pathogenicity,²⁴ would also have enhanced capacity to promote this process. We

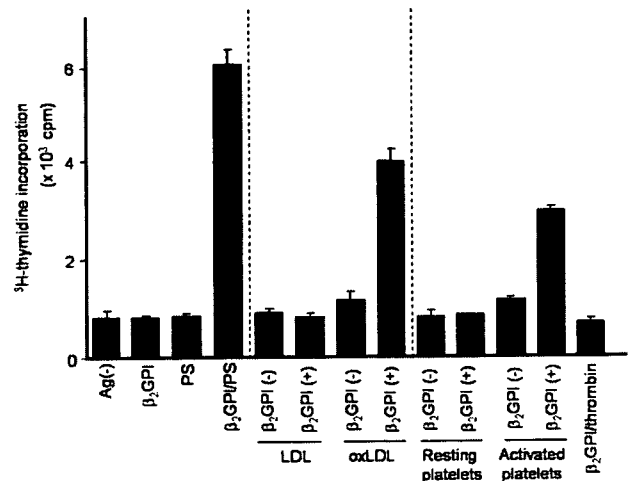


Figure 5. T-cell responses to β_2 GPI-treated anionic substances present in circulation. PBMCs from APS4 were cultured with or without various antigen preparations in medium supplemented with autologous APS plasma. Antigens used included β_2 GPI alone, as well as PS, LDL, oxLDL, resting platelets, and activated platelets, which were treated either with or without β_2 GPI. Thrombin, which was used to activate platelets, in combination with β_2 GPI served as a control. T-cell proliferative response was measured by ³H-thymidine incorporation. Results are shown as mean (column) and standard deviation (error bar) of duplicate measurements. Analogous results were obtained in samples from all 5 patients with APS.

propose a model by which a pathogenic loop maintains sustained anti- β_2 GPI autoantibody production in patients with APS (Figure 6). This model consists of 3 major players: β_2 GPI-reactive CD4⁺ T cells, anti- β_2 GPI antibody-producing B cells, and macrophages. Upon recognition of β_2 GPI cryptic peptides, such as p276-290, presented by macrophages in the context of HLA-DR, β_2 GPI-reactive CD4⁺ T cells are activated and exert helper activity that induces IgG anti- β_2 GPI antibody production from B cells. This process can be achieved by T-B cell collaboration through CD40-CD154 engagement and T cell-derived IL-6.¹¹ IgG anti- β_2 GPI antibodies subsequently recognize β_2 GPI-bound anionic surfaces in circulation, resulting in enhanced phagocytosis of this immune

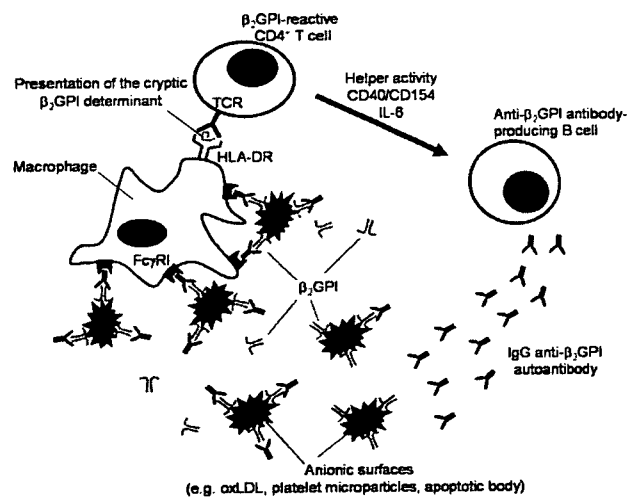


Figure 6. A schematic model representing a continuous autoimmune loop carried out by macrophage, β_2 GPI-reactive CD4⁺ T cell, and anti- β_2 GPI antibody-producing B cell. The macrophage efficiently presents the cryptic β_2 GPI determinant in the context of HLA-DR. The β_2 GPI-reactive CD4⁺ T cell is activated by recognition of the cryptic β_2 GPI peptide and exerts helper activity that induces production of IgG anti- β_2 GPI autoantibodies from the specific B cell. The immune complex consisting of anionic surfaces, β_2 GPI, and IgG anti- β_2 GPI antibodies were captured by macrophages via Fc γ RI.

complex by macrophages through Fc γ RI. In this regard, it has been shown that anti- β_2 GPI antibodies in APS sera are predominantly of IgG2 subclass,^{25,26} which has low affinity to Fc γ RI. However, anti- β_2 GPI antibodies of IgG1 or IgG3 subclass were also detected in many patients with APS. These low levels of anti- β_2 GPI antibodies with high binding affinity to Fc γ RI may be sufficient to drive the pathogenic loop. We have previously shown that β_2 GPI binding to anionic substances promotes the generation of β_2 GPI cryptic peptides by protecting the major PL-binding site from protease attack during antigen-processing by dendritic cells or macrophages.¹⁵ Since it has been shown that antibody binding to the antigen boosts the generation of some minor epitopes,²⁷ binding of IgG anti- β_2 GPI antibodies to the β_2 GPI-anionic substance complex may further amplify generation of previously cryptic β_2 GPI peptides. Moreover, this immune complex is likely to stimulate monocytes via Fc γ RI to secrete tissue factor, which is shown to play an important role in thrombus formation in patients with APS.²⁸ Partial suppression of the β_2 GPI/PS-induced T-cell response by depletion of B cells in our system suggests that presentation of cryptic β_2 GPI peptides could be mediated through B cells that capture β_2 GPI/PS via specific B-cell receptors. This process, however, might have less of an impact on the T-cell response, due likely to low abundance of specific B cells recognizing β_2 GPI/PS. The mechanism that triggers anti- β_2 GPI antibody response in patients with APS remains unclear, but once this autoimmune loop is established, pathogenic anti- β_2 GPI antibodies are continuously produced.

The presence of anionic substances with the capacity to bind β_2 GPI is essential to drive the pathogenic loop inducing continuous anti- β_2 GPI antibody production in patients with APS. Potential anionic substances in the circulation include apoptotic bodies, microparticles derived from activated platelets and endothelial cells, and oxLDL. Since β_2 GPI is abundantly present in the circulation (~200 μ g/mL), excessive exposure to anionic substances would result in the immediate formation of a complex with β_2 GPI. In the present study, we have clearly demonstrated that microparticles derived from activated platelets and oxLDL can function as a substitute for the PS liposome that binds to β_2 GPI and facilitates presentation of the cryptic epitopes of β_2 GPI as a consequence of antigen processing. In addition, some of our group (E.M. and K.K.) reported that stable and nondissociable β_2 GPI-oxLDL complexes were frequently detected in sera from patients with APS and/or systemic lupus erythematosus, but not in healthy individuals.²⁹ In addition, β_2 GPI is known to have antiatherosclerosis activity by preventing oxLDL uptake by macrophages via scavenger receptor, but binding of IgG anti- β_2 GPI antibodies to β_2 GPI-oxLDL complexes mediates atherosclerosis by promoting phagocytosis of macrophages via Fc γ receptor.²⁹⁻³¹ Furthermore,

elevated levels of procoagulant microparticles were detected in patients with APS in association with anti- β_2 GPI antibodies and lupus anticoagulant.³²⁻³⁴ The presence of a large quantity of anionic substances in circulation in patients with APS supports our proposed model.

Based on our model, therapeutic strategies that inhibit pathogenic anti- β_2 GPI antibody production should target interrupting the continuous autoimmune loop carried out by macrophages and β_2 GPI-reactive CD4⁺ T cells and B cells. These immune cells are already targets of therapies under consideration, such as the anti-CD20 chimeric antibody rituximab.³⁵ Another potential therapeutic approach includes the removal of immune complexes consisting of β_2 GPI, anionic substance, and anti- β_2 GPI antibodies. Accordingly, plasma exchange and double filtration plasmapheresis, which theoretically remove such immune complexes, are shown to be effective for patients with intractable APS, including catastrophic APS.^{36,37} Alternatively, small molecules that inhibit Fc receptor downstream signaling would have beneficial effects in patients with APS by suppressing the generation of β_2 GPI cryptic peptides.³⁸

In summary, excessive exposure to anionic surfaces may play a key role in maintaining the pathogenic anti- β_2 GPI antibody response in patients with APS. Further studies should focus on mechanisms that prime the autoimmune loop and development of novel therapeutic strategies targeting the pathogenic process.

Acknowledgments

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Authorship

Contribution: Y.Y., N.S., J.K., K.K., and E.M. performed experiments; Y.Y. and M.K. analyzed results and made the figures; Y.Y. and M.K. designed the research and wrote the paper.

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Blood Coagulation, Fibrinolysis and Cellular Haemostasis

A large deletion of the *PROS1* gene in a deep vein thrombosis patient with protein S deficiency

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Summary

Inherited deficiency of protein S encoded by the *PROS1* gene constitutes an important risk factor for deep vein thrombosis (DVT). Nevertheless, although more than 200 deleterious genetic variations in *PROS1* have been identified, causative point mutations of *PROS1* gene are not detected in approximately half of protein S-deficient families. The present study investigated whether there may exist a large deletion in *PROS1* that constitutes a genetic risk factor for Japanese DVT patients. A multiplex ligation-dependent probe amplification analysis was employed to identify the deletions in *PROS1* in 163 Japanese patients with DVT. A large gene deletion was identified in one patient who

showed 16% protein S activity and did not carry point mutations in *PROS1* by DNA sequencing and it was validated by the quantitative PCR method. The deletion spanned at least the whole *PROS1* gene (107 kb) and at most from the centromere located downstream of *PROS1*, to before the D3S3619 marker, the first heterozygous marker in the upstream of *PROS1* in chromosome 3. In conclusion, a large deletion in *PROS1* was shown to partly account for DVT with protein S deficiency. Screening for large deletions in *PROS1* might be warranted in *PROS1* causative point mutation-negative DVT patients with protein S deficiency.

Keywords

Gene deletion, deep vein thrombosis, protein S deficiency, *PROS1*, MLPA

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Introduction

Imbalance between procoagulant and anticoagulant potencies triggers coagulation. Protein S (PS) is a cofactor for the anticoagulant protease, activated protein C (APC), which proteolytically inactivates activated coagulation factors V and VIII. Therefore, inherited deficiency of PS constitutes an important risk factor for venous thrombosis (1). In the general Japanese population, PS deficiency is relatively common, and its estimated prevalence, 1.12%, is much higher than that in the Caucasian population (0.16–0.21%) (2–4). Recently, we and others ident-

ified a PS K196E mutation (PS Tokushima) as a genetic risk factor for the development of deep vein thrombosis (DVT) (5–8).

The gene for PS (*PROS1*) is located near the centromere on chromosome 3. It spans approximately 101 kb of genomic DNA and comprises 15 exons and 14 introns (9–12). Human DNA contains the active *PROS1* gene at 3q11.2 and a closely linked pseudogene (*PROSP*) at 3p21-cen that shows 96.5% homology to exons 2 to 15 of the *PROS1* gene. To date, more than 200 deleterious mutations in the *PROS1* gene have been reported (13). Most of the gene defects are small point mutations, such as missense, nonsense, frameshift, or splice site mutations, which are

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scattered throughout the coding region of the *PROS1* gene. Some studies have reported that causative point mutations in *PROS1* were found only in approximately 50% of the cases with functional low PS activity (14). One possible cause of such PS deficiency is the existence of large deletions or inversions covering all or part of the *PROS1* gene, which are difficult to identify by current PCR-based sequencing methods. Several studies in PS-deficient families with *PROS1* causative point mutation-negative members indicated that large deletions of *PROS1* have been identified in patients with PS deficiency and screening for large deletions in *PROS1* causative point mutation-negative individuals is therefore warranted (15–18).

A wide range of methods are available for gene dosage measurement to detect large deletions or insertions, but no one method offers overwhelming advantages. Recently developed technique, multiplex ligation-dependent probe amplification (MLPA), based on PCR amplification of ligated probes hybridized to the target DNA (19, 20), would be beneficial for detecting large cryptic genetic variations. This method has proven to be accurate and reliable for identifying deletions, duplications, and amplifications in several diseases (21–23). In this study, we utilized the MLPA method to determine whether there might be large deletions in *PROS1* that constitute genetic risk factors for DVT in Japanese patients.

Materials and methods

Patients

One hundred sixty-three DVT patients, 78 men and 85 women, were registered by the Study Group of Research on Measures for Intractable Diseases, working under the auspices of the Ministry of Health, Labor, and Welfare of Japan. Six centers (one each in Tochigi, Tokyo, Nagoya, and Kyoto, and two in Osaka) participated in this study (5). The protocol for the study was approved by the ethical review committee, and only those subjects who provided written informed consent for genetic analysis were included.

Clinical profile of a patient with a large *PROS1* gene deletion

A previously healthy 22-year-old female was admitted to our hospital due to swelling of her left limb for the past two months. She had no family history of thrombotic diseases. On physical examination, she showed no abnormal findings except swelling of the left thigh and calf.

Laboratory examination showed that the levels of free PS antigen and PS activity had decreased to 2.1 µg/ml (normal range: 6.5–9.8 µg/ml) and 16% (normal range: 65–105%), respectively, while antithrombin (112.7%; normal range: 80–120%), PC (119%; normal range: 75–125%) and plasminogen (104.8%; normal range: 70–120%) were within the normal ranges. ^{99m}Tc-MAA scintigraphy revealed obstruction of the left common iliac and bilateral popliteal veins with no evidence of pulmonary embolisms. Because of the presence of DVT, decreased level of PS, and exclusion of other underlying diseases causing thrombosis, the patient was diagnosed with PS deficiency.

Laboratory values measurement

Some of the patients' plasma samples were stored at –80°C. We measured the PC amidolytic activity (24) and plasma PS activity using Staclot protein S (Diagnostica Stago, Asnieres, France) (3) in 34 samples of DVT patients.

MLPA method

MLPA was performed as described by Schouten et al. (19) using SALSA MLPA KIT P112 *PROS1* (MRC-Holland, Amsterdam, the Netherlands), a kit for screening deletions or duplications in the *PROS1* gene. The kit contains probes for 13 out of 15 exons in *PROS1*, one probe for the promoter located 6.5 kb upstream of *PROS1*, and one probe for exon 4 in *PROSP*. It also contains 17 control probes located on different chromosomes, 1p36, 2q14, 2q24, 3p22, 5q35, 7q31, 7q, 9p21, 10p12, 12p13, 12q14, 14q22, 15q21, 16q22, 18q21, 19p13, and 21q11. The ligation products were amplified by PCR using the common primer set with the 6-FAM label distributed by the supplier. Approximately 100 ng of genomic DNA was utilized for one MLPA reaction. Amplification products were run on an ABI PRISM 3130 DNA Sequencer with the GeneScan 500 LIZ size standard (Applied Biosystems, Foster City, CA, USA) and analyzed by GeneMapper Software 5.0 (Applied Biosystems).

Dosage analyses based on a comparison between deleted and reference wild-type DNA samples were performed on an Excel software Coffalyser V3 (MRC-Holland). Briefly, all *PROS1* peak areas were normalized by dividing each peak area by the combined areas of all the control probe peaks in the same sample. Then, these normalized peak areas were divided by the corresponding wild-type normalized peak areas (average of five independent normal DNA samples) of that probe amplification product, in order to obtain a series of patient-to-normal DNA copy number ratios. The MLPA analysis was repeated three times for all samples.

Q-PCR analysis

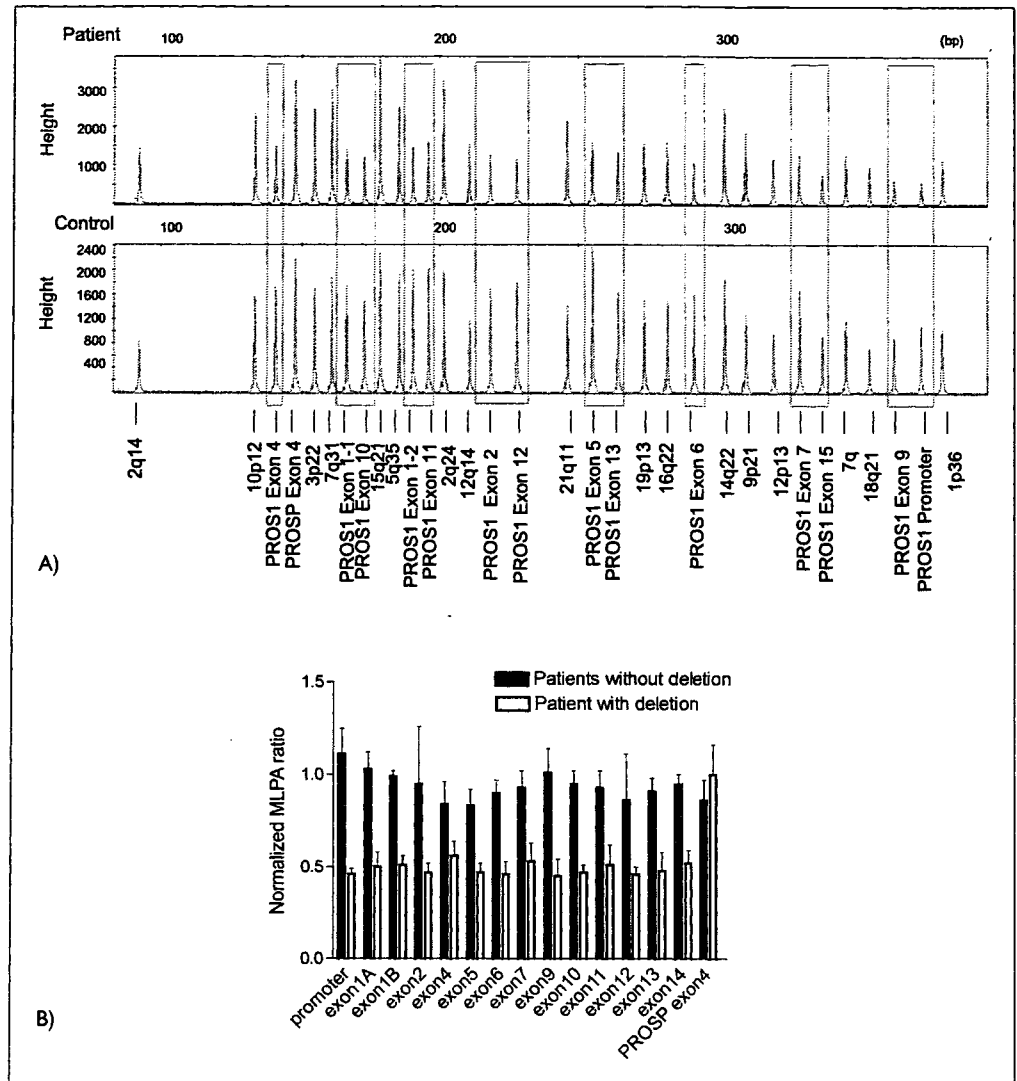
DNA copy numbers for 14 exons in *PROS1* and exon 4 in *PROSP* were determined by Q-PCR. Primers for each sequence (except for the sequence of exon 1) were designed to avoid concurrent amplification of the *PROSP* sequence. The amplicon length ranged from 67 to 198 bps. The PCR reaction and the reference genes were as described previously (25). Each assay included a no-template control (in duplex), 20 ng of normal calibrator human genomic DNA (Roche Applied Science, Indianapolis, IN, USA) (in quadruplicate), and 20 ng of sample DNA (in quadruplicate). Calculation of the gene copy number was performed using the comparative ($\Delta\Delta C_T$) method (26) and the Q-PCR automated analysis software package "qBase" (<http://medgen.ugent.be/qbase/>). A haploid copy number of 1 is expected for a normal sample and a value of ≤ 0.5 for a sample with deletion.

PROS1 sequence analysis

A genomic DNA sample from the DVT patient with the deletion in *PROS1* was sequenced for all exons in *PROS1* using an ABI 3730 sequencer (Applied Biosystems) (27). Genetic variations were identified with NAMIHEI (Mitsui Knowledge Industry Co., Ltd., Tokyo, Japan) and Sequencher (Gene Codes Corpor-

Figure 1: MLPA analysis of the deletion in the *PROS1* gene.

A) The representative MLPA electropherograms in one patient with the deletion in *PROS1* (top) and in a normal control DNA (bottom). Deleted exons are marked by squares. This is one representative of three rounds of MLPA analyses. Probes are 13 out of 15 exons in *PROS1* (two probes for exon 1), one probe for promoter located 6.5 kb upstream of *PROS1*, one probe for exon 4 in *PROSP* and 17 controls located on different chromosomes. B) The representative histograms showing a deletion of 14 probes in *PROS1* in one patient (white bars) and the average normalized MLPA ratio in the patients without deletion (black bars). The probe for exon 4 in *PROSP* is shown as the control probe. The data are presented as the means \pm SD from three rounds of MLPA.



ation, Ann Arbor, MI, USA) software, followed by visual inspection (28).

SNPs genotyping and microsatellite analysis

Five SNP markers located within a 252 kb region centered on *PROS1* were chosen on the basis of the available data from NCBI Map Viewer (Build 36.2) and the genotyping of the markers was performed by the TaqMan-PCR genotyping system (29).

Four microsatellite markers (D3S3681, D3S1276, D3S3634 and D3S1271) covering a 22 Mbp region centered on the *PROS1* locus were obtained from the ABI Linkage Mapping Set Version 2.5 (Applied Biosystems), and the microsatellite marker analyses were performed according to the manufacturer's instructions. For the other five microsatellite markers (D3S3556, D3S1251, D3S3619, D3S1752 and D3S3716) in the region, we designed PCR primer sets. Primer sequences were available on request. PCR products were detected by Agilent Bioanalyzer 2100 (Agilent Technologies, Boeblingen, Germany) or by electrophoresis through the 12% polyacrylamide gel stained with

SYBR Green I (Nucleic Acid Gel Stain (Molecular Probes, Eugene, OR, USA). The products were also directly sequenced to reveal the allele zygosity.

Results

Deletion detection using MLPA analysis

To identify the large deletions in *PROS1* in patients with DVT, we utilized MLPA analysis. This method has the advantage of using a relatively small amount of DNA – i.e. only 100 ng of genomic DNA are required for one MLPA analysis to reveal a *PROS1* gene deletion. A deletion of one copy of a probe target sequence is usually apparent by a 35–55% reduction in relative peak area. A gain in copy number from two to three copies is usually apparent by a 30–55% increase in relative peak area.

We identified reduced peak areas in the *PROS1* regions in one DVT patient, which indicates a heterozygous deletion encompassing the whole *PROS1* gene (Fig. 1). No significant changes were found in the peak areas of the control probes in-

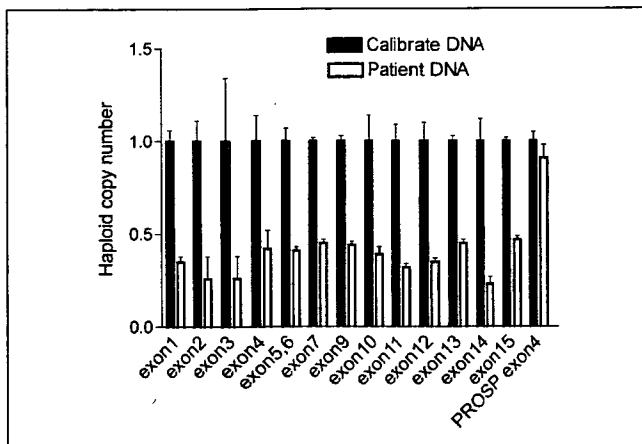


Figure 2: Haploid copy number status of *PROS1* determined by Q-PCR. Representative histogram showing the haploid copy numbers of *PROS1* exons in one patient with the *PROS1* deletion (white bars) and in the normally calibrated DNA (black bars). Data are presented as the means \pm SD from Q-PCR reaction in quadruplicate. The copy number of *PROSP* exon 4 is plotted as no deletion control.

cluding *PROSP* exon 4 in this patient. For other patients, no reduced or increased peak areas were detected. The results of the MLPA analysis were reproducible, with three rounds of MLPA yielding identical findings in all patients. Based on the MLPA analysis, a minimum distance of *PROS1* deletion in this patient was estimated to be 107 kb between the *PROS1* promoter and exon 15 MLPA probes.

Deletion detection using Q-PCR

To confirm the deletions detected by MPLA in the patient, the patient’s genomic DNA and a normally calibrated DNA were

subjected to Q-PCR analysis to determine the copy numbers of *PROS1*. The analysis confirmed that all exons of *PROS1* were deleted in this patient (Fig. 2). No deletion was detected in *PROSP* exon 4.

Direct sequencing of the coding regions of *PROS1*

As described in *Methods*, PS antigen and PS activity of the patient with the deletion in *PROS1* decreased to 2.1 μ g/ml (normal range: 6.5–9.8 μ g/ml) and 16% (normal range: 65–105%), respectively, while antithrombin, PC and plasminogen were within normal ranges. This suggested that the patient might be a compound heterozygote for PS deficiency. To identify additional genetic variations in *PROS1* in the patient, we sequenced all exons of *PROS1* in this patient, but no point mutations were found.

Distribution of microsatellites and SNP genotypes

To define the range of deletions detected by the MLPA analysis, five SNP markers and nine microsatellite markers covering a 22 Mbp window centered on the *PROS1* locus were analyzed for their allele zygosity (Table 1). When the allele carries a deletion, it will be hemizygous for all markers within the deleted region, but will appear homozygous. Thus, if a specific marker is homozygous, there might be a deletion covering that locus, but if a specific marker is heterozygous, there must be no deletion covering that locus.

The five SNP markers flanking *PROS1* were homozygous. The marker D3S3619, which was located 1,949 kb upstream of *PROS1*, was detected as the first microsatellite heterozygosity in the upstream region of *PROS1* (Table 1). As a result, the boundary of the deletion in this patient must be within the range of D3S3619 to D3S1276. Since the centromere is located between *PROSP* and exon 15 of *PROS1*, one breakpoint of the deletion must be closer to exon 15 than to *PROSP* (Fig. 3). Therefore, the

Marker	Location, kbp	Allele or repeats	Minor allele frequency	Heterozygosity, %	Zygosity in the patient
D3S3681	79894	CA	—	—	heterozygous
D3S1276	85338	CA	—	—	heterozygous
rs9289536	95031	C/T	0.11	0.20	homozygous
rs9713061	95074	G/T	0.10	0.15	homozygous
rs6786905	95213	C/A	0.10	0.18	homozygous
rs13100168	95220	C/T	0.06	0.12	homozygous
rs10433405	95283	G/A	0.14	0.24	homozygous
D3S3556	95525.66	CA	—	—	homozygous
D3S3634	95525.74	CA	—	—	homozygous
D3S1251	95760	CA	—	—	homozygous
D3S3619	97124	CA	—	—	heterozygous
D3S1752	99228	GAT	—	—	heterozygous
D3S3716	100219	CA	—	—	heterozygous
D3S1271	102217	CA	—	—	heterozygous

Minor allele frequencies and heterozygosities of SNPs were calculated from 44 unrelated normal Japanese studied in the International HapMap project. Locations are from NCBI Map Viewer build 36.2. The zygosity analysis was performed for a patient with a deletion in *PROS1*.

Table 1: Description of microsatellite and SNP markers.

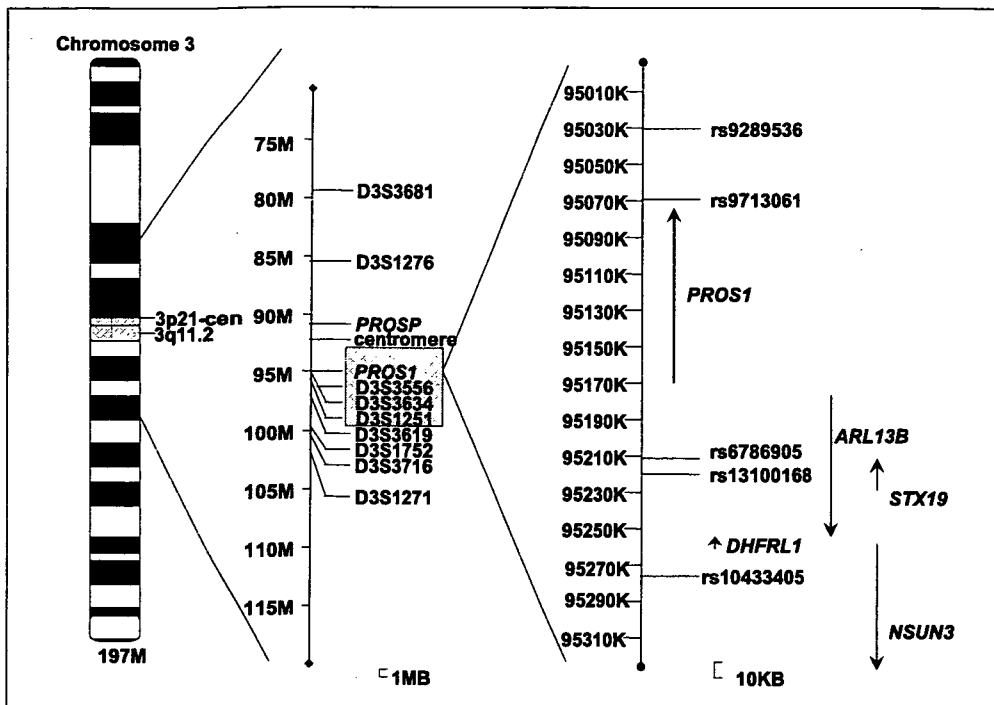


Figure 3: Schematic presentation of the large *PROS1* deletion on chromosome 3. The distributions of *PROS1* (3q11.2), *PROSP* (3p21-cen), *PROS1* flanking SNP and microsatellite markers are mapped on chromosome 3. Possible maximum range of the deletion identified in the patient with DVT is boxed with a grey square. Transcriptional orientations of the genes are shown by arrows. Nucleotide positions are according to NCBI Map Viewer (Build 36.2).

deletion ranged at a maximum from the centromere to before the D3S3619 marker, the first heterozygous marker in the upstream of *PROS1* in chromosome 3. If we take this maximum deletion into account, the large deletion could also include the genes *ARL13B* (ADP-ribosylation factor-like 13B), *STX19* (syntaxin 19), *DHFRL1* (dihydrofolate reductase-like 1), and *NSUN3* (NOL1/NOP2/Sun domain family, member 3), because they are located between *PROS1* and the marker D3S3556 (Fig. 3).

***PROS1* deletion carrier in *PROS1* causative point mutation-negative Japanese DVT patients with low PS activity**

We measured PC and PS activity in 34 out of 163 DVT patients. We found that 11 patients showed PS activity of less than 50% with PC activity of more than 90%, suggesting functional low PS activity, where the patient with the large *PROS1* deletion was included. Among 11 patients with low PS activity, five carried causative point mutations including missense or splice site mutations (manuscript in preparation by YS and TM). Therefore, in our Japanese DVT patient group, one patient with a large *PROS1* deletion was identified in six DVT patients with low PS activity and no causative point mutations in *PROS1*.

Discussion

With a few exceptions, gene deletion measurements are not routine in most mutation screening studies, and thus alterations of gene deletion on the kilobase scale are thought to be underestimated. By MLPA, Q-PCR, and direct sequencing analysis of the *PROS1* coding regions, in addition to genotyping of the microsatellites and SNP markers flanking *PROS1*, we detected a large

deletion encompassing at least the whole *PROS1* coding regions in 1 out of 163 Japanese patients with DVT.

Evaluation of the deletion detection methods

The presence of the highly homologous pseudogene (>96%) *PROSP* hampered the analysis of the gene deletion in *PROS1* by the conventional techniques. In the present study, therefore, we used the MLPA method to detect the deletions in *PROS1*. The accuracy of this method was confirmed by three rounds of MLPA and validated directly by Q-PCR. The presence and possible range of the deletion was further confirmed by direct sequencing of *PROS1* as well as microsatellite analysis and SNP linkage analysis of the flanking regions of *PROS1*. With the sequencing of *PROS1* exon regions, the loss of heterozygosity of all the SNPs of *PROS1* indicates the existence of a hemizygous deletion encompassing the whole *PROS1* gene in one DVT patient. The boundaries of the deletion were deduced by allele zygosity analysis of the SNPs and microsatellite markers centered on *PROS1*. The marker density selected for analysis also has an impact on the accuracy of discriminating the range of the deletion. In the present study, the breakpoints of the deletion could only be limited to the range between D3S3619, which was detected as the first heterozygous microsatellite in the region upstream of *PROS1*, and the centromere located downstream of *PROS1*. To further define the range of the deletion, it will be necessary to determine the segregation pattern of the investigated SNP and microsatellite markers in other affected individuals of the same family (18).

Nature of the large deletion of *PROS1*

The large deletion in the present study was defined as ranging at a minimum distance of 107 kb between the *PROS1* promoter and

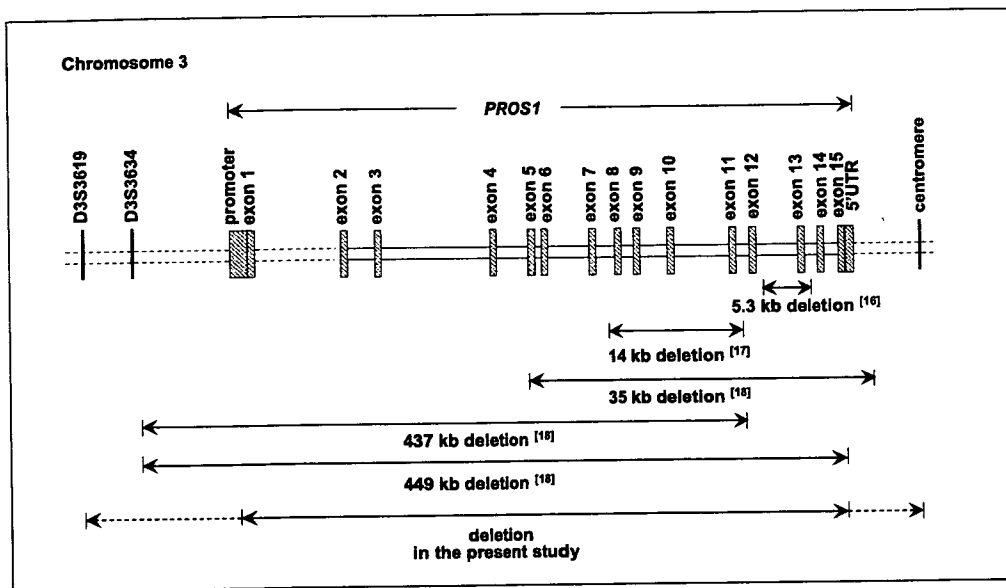


Figure 4: Outline of reported large deletions encompassing *PROS1* on chromosome 3. The location of centromere, *PROS1*, and two upstream microsatellite markers (D3S3619 and D3S3634) are sketched on chromosome 3. The relative long distance between exons 1 and 2 is represented by the dotted line. The distance of the regions flanking *PROS1* is represented by the dotted line. The minimum range of the deletion in the present study is marked with the solid line and the flanking maximum range is marked with the dotted line. The reported deletions are taken from the references (16–18).

exon 15 MLPA probes and at a maximum distance from the centromere to before the D3S3619 marker, the first heterozygous marker in the upstream of *PROS1* in chromosome 3. To date, a total of five large *PROS1* deletions have been described (Fig. 4) (16–18). A 5.3 kb deletion spanning over 90% of intron 12, the entire exon 13, and about a quarter of intron 13, was detected in two *PROS1*-deficient families by Southern hybridization (16). A long-PCR-based technique identified a deletion from intron 7 to 11 that removed approximately 14 kb (17). With a dense set of SNP and microsatellite markers, three large deletions that encompassed at least 35 kb including exons 5–15, 437 kb including exons 1–11, and 449 kb including the whole *PROS1* gene were identified (18). The deletion in the present study was another one with the definite minimum range encompassing the whole *PROS1* gene.

Large deletion of *PROS1* and PS activity

The free PS antigen level and PS activity of the patient with the deletion in *PROS1* were 2.1 $\mu\text{g/ml}$ (normal range: 6.5–9.8 $\mu\text{g/ml}$) and 16% (normal range: 65–105%), respectively, so that, in addition to the hemi-deletion of the *PROS1* gene, other genetic variations could influence the levels of the PS antigen and activity. However, no causative point mutations were found in this patient by DNA sequencing. If we take the deletion as the maximum, the deletion may encompass four other genes, *ARL13B*, *STX19*, *DHFR1L1*, and *NSUN3* (Fig. 3). To our knowledge, none of these genes are known to be associated with thrombosis, and thus their absence is unlikely to affect the PS deficiency phenotype. The four genes were also deleted in other patients with PS deficiency (18). Since the characteristics of these genes and the corresponding phenotypes have not yet been clarified, quantitative analysis of these genes was not considered in the present study. Another genetic variation that was missing by our analysis was the existence of chromosomal rearrangements such as an inversion of a large segment of DNA which has been found to be a major genetic defect in the factor VIII gene in patients with se-

vere hemophilia A (30). Mutations in the promoter region might be another possible explanation for the low PS activity. The *PROS1* promoter region has recently been characterized and liver-specific cis-acting DNA elements have been identified, but the role of this region in PS deficiency is still unknown (31).

Mechanism of the large deletion

We speculated that the deletion might occur by non-allelic homologous recombination which can generate rearrangements as a result of recombination between highly similar duplicated sequences, such as segmental duplications (32). It has been confirmed that 24% of the 1,447 copy-number variations in the human genome are associated with segmental duplications. Segmental duplications were more frequently associated with the long copy-number variations (33). However, the segmental duplications were not present in the vicinity of the *PROS1* gene at the pericentromeric region in chromosome 3 (34). Regardless of the recombination mechanism, genomic architectural features have been associated with many rearrangement breakpoints (35). This suggests that the large deletion in *PROS1* is not a random event, but might result from a predisposition to rearrangement due to a complex genomic architecture that may create instability in the genome.

Large *PROS1* deletion in Japanese DVT patients

All of the large deletions of *PROS1* so far reported were detected in affected individuals in PS deficiency families (15–18). In the present study, a *PROS1* deletion was identified in a patient with DVT, but unfortunately, the family members were not available for analysis. The presence of a large deletion in one out of the 163 DVT patients suggested that a large deletion of *PROS1* is relatively rare in DVT patients. However, if the deletion was obtained in specified patients with low PS activity (<50%) and normal PC activity (>90%), it was found in one out of 11 patients. When we looked at DVT patients with low protein S activity who did not carry the causative point mutations in *PROS1*, one out of

six patients had a large *PROS1* deletion. Large deletions in *PROS1* have been previously identified by segregation analysis in three out of eight *PROS1* causative point mutation-negative families with PS deficiency (18). Large deletions of *PROS1* might be more common in familial PS deficiency than previously thought, and MLPA analysis is an effective tool to detect the deletion in *PROS1*.

In conclusion, a large deletion encompassing the whole *PROS1* gene was detected in one DVT patient with 16% PS ac-

tivity. The large deletion in *PROS1* partly accounted for the low PS activity in patients with DVT. Screening for large deletions in *PROS1* might be warranted in causative point mutation-negative DVT patients with PS deficiency.

Acknowledgements

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ADAMTS13 assays and ADAMTS13-deficient mice

Toshiyuki Miyata, Koichi Kokame, Fumiaki Banno, Yongchol Shin and Masashi Akiyama

Purpose of review

Thrombotic thrombocytopenic purpura can be induced by acquired or congenital deficiency of the plasma von Willebrand factor-cleaving protease, ADAMTS13.

Measurement of ADAMTS13 activity is important for the diagnosis and treatment of microangiopathies including thrombotic thrombocytopenic purpura. Phenotypic analysis of mice lacking the *Adamts13* gene is valuable for understanding the pathogenesis of microangiopathies.

Recent findings

The minimum substrate for ADAMTS13 activity was identified as 73 amino acid residues in the A2 domain of von Willebrand factor, called VWF73. Several new assays have been developed using this sequence. The VWF73-based assays are rapid, quantitative, and easy to handle, and are well correlated with the measures from previous assays. Mice lacking the *Adamts13* gene were produced. The mice were viable and fertile. They showed a prothrombotic state but no symptoms of spontaneous thrombocytopenia, hemolytic anemia, or microvascular thrombosis were observed.

Summary

VWF73-based ADAMTS13 assays will significantly facilitate the accurate diagnosis of microangiopathies and contribute to the improved clinical treatment of these diseases. Accumulated clinical information on patients with ADAMTS13 deficiency and mice lacking the *Adamts13* gene indicates that additional environmental or genetic susceptibility factors are required to trigger thrombotic thrombocytopenic purpura.

Keywords

ADAMTS13, microangiopathy, thrombotic thrombocytopenic purpura, von Willebrand factor

Abbreviations

CUB	complement components C1r/C1s, Uegf (epidermal growth factor-related sea urchin protein), and bone morphogenetic protein-1
HUS	hemolytic uremic syndrome
TSP-1	thrombospondin type-1
TTP	thrombotic thrombocytopenic purpura
ULVWF	ultralarge von Willebrand factor
VWF	von Willebrand factor

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Introduction

Thrombotic thrombocytopenic purpura (TTP) is characterized by thrombocytopenia and microangiopathic hemolytic anemia accompanied by variable-penetrance of neurologic dysfunction, renal failure, and fever. In the microvasculature of patients with TTP, systemic platelet thrombi are developed, largely resulting from the accumulation of ultralarge von Willebrand factor (ULVWF) multimers [1]. ULVWF can be accumulated by acquired or congenital deficiency of the von Willebrand factor (VWF)-cleaving protease, ADAMTS13 (a disintegrin-like and metalloprotease with thrombospondin type 1 motif, 13) [2,3**]. TTP caused by congenital deficiency of ADAMTS13 is also called Upshaw-Schulman syndrome.

Since the cloning of its cDNA in 2001, this new antithrombotic factor has been intensively studied [4-6,7*,8**,9**,10*]. Here we summarize the recent progress on ADAMTS13, focusing on assays for ADAMTS13 and mice lacking the *Adamts13* gene.

Genetic mutations in congenital ADAMTS13 deficiency

ADAMTS13 consists of 1427 amino acid residues with a calculated molecular mass of 145 kDa. It is composed of multiple discrete domains, as shown in Fig. 1 [4-6,7*,8**,9**]. Unlike other ADAMTS family members, the ADAMTS13 sequence has a short pro-sequence and two C-terminal CUB [complement components C1r/C1s, Uegf (epidermal growth factor-related sea urchin protein), and bone morphogenetic protein-1] domains. The human ADAMTS13 gene comprises 29 exons, encompassing 37 kb on chromosome 9q34. It is expressed mainly in the liver; primarily in stellate cells [11,12]. Platelets and endothelial cells also express ADAMTS13 [13,14,15*,16**]. The CUB domains are required for apical sorting of ADAMTS13 in endothelial cells [16**].

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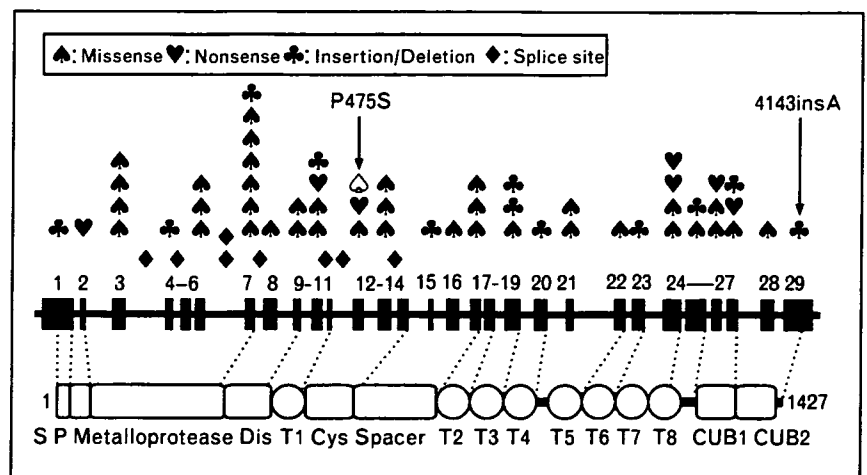
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Figure 1 Genomic structure and domain organization of human *ADAMTS13* and nonsynonymous mutations identified in patients with congenital thrombotic thrombocytopenic purpura

Missense mutations are indicated by black spades. Nonsense mutations, insertion/deletion mutations, and splice site mutations are shown by hearts, clubs, and diamonds, respectively. The P475S mutation commonly observed in the Japanese population and the A insertion at nucleotide number 4143 found in multiple European populations are shown. S, signal peptide; P, propeptide; Dis, disintegrin-like domain; T (numbered 1–8), thrombospondin type 1 motifs; CUB, complement components C1r/C1s, Uegf (epidermal growth factor-related sea urchin protein), and bone morphogenetic protein-1; Cys, cysteine-rich domain.



More than 60 different mutations of the *ADAMTS13* gene have been reported [6,17–19,20,21]. More than 50% are missense mutations, as well as frame-shift mutations such as insertions and deletions, and nonsense mutations and splice-site mutations. These mutations are distributed throughout the various domains of *ADAMTS13* (Fig. 1). The correlation of genotype and protease activity in some of these mutations has been examined by expression analysis *in vitro* [18,20,21]. Most of the mutations have been identified in a single family but there are at least six recurrent mutations in unrelated patients. Among them, the 4143insA mutation has been identified as a genetic background for TTP in multiple families and is frequent among patients with congenital *ADAMTS13* deficiency in Northern and Central European countries [22^{**}]. Interestingly, several missense mutations can interact and alter the phenotype of *ADAMTS13* deficiency [23^{**}].

There are at least nine missense polymorphisms in the *ADAMTS13* gene. The Q448E mutation was shown not to affect the protease activity, whereas the P475S mutation decreased the activity [18]. This mutation is present in the Asian population but not in the white population [19,24,25]. It has been shown not to be a genetic risk factor for deep-vein thrombosis [26^{*}].

Information regarding the wide variation of phenotypes in TTP patients with congenital *ADAMTS13* deficiency has accumulated. Most patients with congenital *ADAMTS13* deficiency had their first episode as newborn babies or in early infancy [10^{*}]. After this period, the clinical manifestations of congenital *ADAMTS13* deficiency vary from patient to patient, and patients are often incorrectly diagnosed with idiopathic thrombotic purpura or Evans' syndrome during childhood. Seven women with congenital *ADAMTS13* deficiency

who exhibited TTP at 5–6 months of pregnancy have been reported [27].

The factor V Leiden mutation is a well characterized and the most prevalent genetic risk factor for venous thrombosis. A previous study suggested that the factor V Leiden mutation may be a pathogenic risk factor in patients with thrombotic microangiopathy who have normal *ADAMTS13* activity [28]. A recent study did not support the link between the factor V Leiden mutation and thrombotic microangiopathy [29].

Hemolytic uremic syndrome (HUS) is a thrombotic microangiopathy with manifestations of hemolytic anemia, thrombocytopenia, and renal impairment [1,8^{**},9^{**},30]. In most cases, typical HUS is triggered by Shiga toxin-producing *Escherichia coli* and manifests diarrhea (D+HUS). Atypical non-Shiga toxin-associated HUS is not associated with diarrhea (D–HUS), and deficiencies of complement factor H, membrane cofactor protein, and complement factor I have been reported in such atypical HUS [9^{**},31].

Epitopes of *ADAMTS13* autoantibodies

Inhibitory autoantibodies for *ADAMTS13* cause a deficiency of *ADAMTS13* among patients with autoimmune TTP [3^{**},8^{**}]. The prevalence of *ADAMTS13* deficiency among patients with TTP varies from 13 to 100% depending on the criteria of the study. TTP may develop within 2–6 weeks after the antiplatelet agent ticlopidine is administered. TTP may develop in patients with HIV infection. No other apparent etiologies of inhibitory autoantibodies have been identified.

Several studies using recombinant *ADAMTS13* and its truncated forms have identified the epitopes of

autoantibodies in patients with TTP. Many inhibitory autoantibodies recognized the spacer domain as the target epitope [32–34,35*]. Some reacted with the C-terminal CUB domains and the first thrombospondin type-1 (TSP-1) repeat [33]. Multiple B-cell clones producing antibodies directed against the spacer domain have been reported [36].

Assays of ADAMTS13 activity

ADAMTS13 assays such as multimer analysis by SDS/agarose-gel electrophoresis and residual collagen-binding analysis have been utilized for plasma ADAMTS13 activity in thrombotic microangiopathies and other pathophysiological conditions [2,4,6]. These assays quantify ADAMTS13 activity by measuring residual VWF multimers or their activity, suggesting that more accurate and simpler assays are in demand.

New assays using VWF73 peptide as the ADAMTS13 substrate

In 2004, we identified a 73-amino acid sequence spanning residues D1596–R1668 of the VWF A2 domain, VWF73, as the minimum region for ADAMTS13 cleavage [37]. A shorter peptide, VWF64, from D1596 to R1659, was not a good substrate for ADAMTS13. Since the development of VWF73 as the substrate for ADAMTS13, several new assays based on the VWF73 sequence have been developed [37–39,40*–43*]. So far, seven assays have been published for the VWF73-based measurements of ADAMTS13 activity. The characteristics of these assays are summarized in Table 1 [37–39,40*–43*]. The assays have the advantages of being simple, rapid, accurate, and quantitative. They do not require denaturing conditions; therefore, the incubation time of the substrates and plasma samples is reduced to less than 1 h. They give rise to quantitative measures. In addition, most of the assays are compatible with the 96-well

microplates that clinical-laboratory workers are familiar with, so they have the potential to be widely used in clinical settings.

Of the seven assays, FRET-VWF73, a fluorogenic substrate for ADAMTS13, is well characterized. The advantage of using FRET-VWF73 is that the ADAMTS13 activity can be determined by the initial velocity of the increase in fluorescence. Therefore, the assay is highly quantitative. ADAMTS13 cleaves VWF with a K_{mapp} of $3.7 \pm 1.4 \mu\text{g/ml}$ or 15 nM in VWF subunits, which is comparable with the plasma VWF concentration of 5–10 $\mu\text{g/ml}$, and with a value for k_{cat} of $\sim 0.83 \text{ min}^{-1}$ [44*]. ADAMTS13 cleaves FRET-VWF73 with a K_{mapp} of $3.2 \pm 1.1 \mu\text{M}$ and a k_{cat} of $\sim 58 \text{ min}^{-1}$. Thus, the affinity of ADAMTS13 to FRET-VWF73 was decreased ~ 200 -fold compared with VWF, but the catalytic efficiency was ~ 70 -fold greater than VWF. Therefore, ADAMTS13 cleaves VWF and FRET-VWF73 with roughly comparable catalytic efficiencies of 55 and $18 \mu\text{M}^{-1} \text{ min}^{-1}$, respectively [44*].

FRET-VWF73 was evaluated by three different research groups [45–47]. Although the definitive evaluation remains to be determined in a large cohort of patients diagnosed with acquired or congenital TTP, ADAMTS13 activity determined by FRET-VWF73 assay was in good accordance with that measured by conventional assays. FRET-VWF73 is now commercially available (Table 2).

There may be limitations of the VWF73-based ADAMTS13 assays. VWF73, a small fragment of the A2 domain of VWF, may lack additional sites on VWF that interact with ADAMTS13. The A1 domain of VWF binds cofactors such as platelet glycoprotein Ib and heparin to regulate cleavage [48], and the A3 domain may be a docking site for ADAMTS13 [49]. VWF73 lacks

Table 1 VWF73-based ADAMTS13 activity assays

Substrate	Principle	Reference
GST–VWF73 fusion protein with the C-terminal 6×His tag	Western-blot detection using anti-GST antibody	[37]
FRET-VWF73, synthetic VWF73 peptide with a fluorophore at the P7 position and a quencher at the P5' position	Fluorescence resonance energy transfer, initial-velocity method	[39]
Immobilized GST–VWF73 fusion protein with the C-terminal 6×His tag	Enzyme immunoassay, the amount of 6×His remaining was assayed with anti-6×His IgG conjugated with HRP, end point method	[38]
Immobilized His- and biotin-labeled VWF73 conjugated with HRP	Enzyme-linked assay, endpoint method	[40*]
Immobilized GST–VWF73 fusion protein with the C-terminal 6×His tag	Mass-spectrometry analysis of the products, endpoint method	[41*]
Immobilized GST–VWF73 fusion protein with the C-terminal 6×His tag	Enzyme immunoassay, the amount of products were assayed with anti-N10 mAbs conjugated with HRP, endpoint method	[42*]
Recombinant 6×His-tagged VWF73 peptide labeled with fluorescein at both the P7 and P6' positions	Fluorescence resonance energy transfer, initial-velocity method	[43*]

GST, glutathione-S-transferase; HRP, horseradish peroxidase; mAb, monoclonal antibody; VWF73, 73 amino acid residues of von Willebrand factor (VWF) from D1596 to R1668.

Table 2 Commercially available kits for assaying ADAMTS13 activity and antigen

Kit	Maker/supplier	Objectives	Time
FRETS-VWF73	Peptide Institute, Peptides International	Activity	1 h
ATS-13 ADAMTS-13 Activity	GTI	Activity	30 min
ADAMTS13 ELISA kit	Mitsubishi Kagaku Iatron	Antigen	3.5 h
ADAMTS13 activity ELISA kit	KAINOS LABORATORIES	Activity	3.5 h
TECHNOZYM ADAMTS-13	Technoclone GmbH	Activity/antigen	2.5 h/4 h
TECHNOZYM ADAMTS-13 INH	Technoclone GmbH	Autoantibody	2.5 h
ACTIFLUOR ADAMTS13 Activity Assay kit	American Diagnostica	Activity	
IMUBIND ADAMTS13 ELISA	American Diagnostica	Antigen	5 h
IMUBIND ADAMTS13/rXI Complex ELISA	American Diagnostica	ADAMTS13/rXI complex	4 h
IMUBIND ADAMTS13 Autoantibody ELISA	American Diagnostica	Autoantibody	4 h

these domains. Therefore, if enzyme defects in patients with TTP affect the ADAMTS13-binding site for these domains, cleavage of VWF73 will not reflect these defects.

Measurements of ADAMTS13 autoantibodies

Autoantibodies neutralizing ADAMTS13 activity are a major cause of acquired TTP. The presence or absence of inhibitory autoantibodies is important in discriminating acquired from congenital TTP. An inhibitor assay is generally carried out using mixtures of heat-inactivated plasma from patients and normal plasma at a 1:1 dilution or several dilutions. Assays for ADAMTS13 activity so far developed, including VWF73-based assays, are compatible for the inhibitor assay. It should be noted that nonneutralizing autoantibodies may reduce the circulating ADAMTS13 levels by antibody-mediated clearance.

ELISA has been developed to detect autoantibodies against ADAMTS13. In this process, immobilized ADAMTS13 in the plate wells captures both inhibitory and noninhibitory autoantibodies in plasma samples; then secondary detection antibodies, such as goat antihuman IgG or IgM antibodies labeled with horseradish peroxidase, are added and the levels of ADAMTS13-binding IgGs are determined [50]. Using this assay, low titers of IgG antibodies were detected in four out of 111 healthy control donors who lacked anti-ADAMTS13 inhibitory activity by inhibitor assays. IgG autoantibodies were found in 97% of untreated patients with acute acquired thrombotic microangiopathies who had plasma ADAMTS13 activity levels below 10% [51]. This assay was more sensitive than the standard functional inhibitor assay for detecting autoantibodies against ADAMTS13. The ELISA kit utilizing this principle is now commercially available (Table 2), and has been validated to be useful [52].

Antigen assays for plasma ADAMTS13

ELISA for measuring plasma ADAMTS13 antigen levels has also been developed by several research groups. ADAMTS13 antigen ELISA kits are also commercially available (Table 2).

Healthy plasma ADAMTS13 levels

The ELISA assay to detect plasma ADAMTS13 levels can estimate the plasma ADAMTS13 concentration when the ADAMTS13 standard can be obtained from recombinant full-length ADAMTS13 protein. The ADAMTS13 antigen concentration in normal human plasma pooled from white donors was $1.03 \pm 0.15 \mu\text{g/ml}$ of plasma [53^{*}]. Interestingly, normal Chinese donors have significantly lower antigen levels ($0.62 \pm 0.13 \mu\text{g/ml}$). In another study of 99 healthy Austrian donors, the median plasma ADAMTS13 level was $1.08 \mu\text{g/ml}$ using recombinant ADAMTS13 as the standard [54^{*}]. The plasma ADAMTS13 level in Japanese donors was reported to be 0.82 ± 0.15 and $0.70 \pm 0.13 \mu\text{g/ml}$ using two different ELISA systems when recombinant ADAMTS13 was used as the standard [55^{*}].

Phenotype of mice lacking *Adamts13* gene

The mouse is a promising animal model for seeking genetic or environmental susceptibility factor(s) for a certain disease phenotype. Two types of mouse *Adamts13* cDNA have been isolated and characterized [56]. cDNA isolated from the 129/Sv strain showed a domain organization identical to the human one. The other cDNA lacked the C-terminal two TSP-1 motifs and two CUB domains due to the insertion of an intracisternal A particle retrotransposon in intron 23, which creates a premature stop codon. Both recombinant proteins showed VWF-cleaving activity *in vitro*.

Mice lacking the *Adamts13* gene have been recently developed by us and another group [57,58^{**},59^{**}]. We generated mice lacking the *Adamts13* gene by replacing exons 3–6 encoding the catalytic domain by a neomycin-resistant cassette and analyzed phenotypes on a 129/Sv genetic background of the ADAMTS13-deficient mice [58^{**}]. The ADAMTS13-deficient mice were born in the expected Mendelian distribution. Plasma from homozygous mice showed no ADAMTS13 activity. The mice were viable and fertile. Hematologic and histologic examinations failed to detect any evidence of thrombocytopenia, hemolytic anemia, or microvascular thrombosis. However, ULVWF multimers were observed in the plasma of homozygotes. Thrombus formation on immobilized

collagen under flow was significantly elevated in homozygotes in comparison with wild-type mice. Thrombocytopenia was more severely induced in homozygotes than in wild-type mice after intravenous injection of a mixture of collagen and epinephrine. Therefore, a complete lack of ADAMTS13 in mice caused a prothrombotic state, but it alone was not sufficient to cause TTP. Factors in addition to ADAMTS13 deficiency may be necessary for development of TTP.

Mice lacking the *Adamts13* gene have also been generated with replacement of exons 1–6 by a neomycin cassette [57]. The ADAMTS13-deficient mice were born in the expected Mendelian distribution and homozygous mice were viable and fertile. When the VWF multimer analysis was examined in the ADAMTS13-deficient mice on a mixed-strain C57BL/6J and 129X1/SvJ genetic background, the multimers of wild-type mice and ADAMTS13-deficient mice were indistinguishable. However, the ADAMTS13-deficient mice, after two generations of backcrossing to the CASA/Rk strain (a mouse strain with elevated plasma VWF), showed ULVWF multimers compared with wild-type littermates. Mice with a mixed CASA/Rk background showed a significant decrease in platelet count and a fraction of the deficient mice exhibited severe thrombocytopenia and significantly decreased survival compared with wild-type or heterozygous controls. These mice showed a TTP-like phenotype such as severe microangiopathic changes in the peripheral blood and VWF-rich and fibrin-poor hyaline thrombi in the small vessels. Deficient mice showed prolongation of VWF-mediated platelet-endothelial interactions, indicating that ADAMTS13 regulates VWF-mediated platelet adhesion *in vivo*. When Shiga toxin was infused intravenously, TTP-like symptoms were observed in ADAMTS13-deficient mice with a mixed CASA/Rk background, but not in mice with a mixed C57BL/6J background. Shiga toxin is known to induce HUS through endothelial dysfunction. Thus, TTP can be induced in ADAMTS13-deficient mice by agents causing endothelial dysfunction. This strain-specific difference of TTP pathogenesis in mice may indicate the contribution of additional genetic factors.

Further characterizations of events *in vivo* in ADAMTS13-deficient mice on a mixed-strain C57BL/6J and 129X1/SvJ genetic background have been examined [59**]. When the microvenule endothelium in ADAMTS13-deficient mice was activated with calcium ionophore, ULVWF multimers were secreted from Weibel–Palade body, and platelet aggregation resulting in spontaneous thrombus formation was observed using intravital microscopy. In wild-type littermates, platelet strings and very small aggregation could be seen attached to the endothelium, but thrombi did not form. A ferric chloride injury model on arterioles

exhibited that ADAMTS13 downregulates both platelet adhesion to the exposed subendothelium and thrombus formation. Infusion of recombinant ADAMTS13 into ADAMTS13-deficient or wild-type mice inhibited similar thrombus growth. These findings revealed that ADAMTS13 is a natural anticoagulant.

Conclusion

A highly accurate and quantitative assay method for measuring ADAMTS13 activity has been developed. These assays are now commercially available and will be widely utilized for a clinical diagnosis in patients with microangiopathy to discriminate TTP from HUS or other thrombocytopenia. Mice lacking the *Adamts13* gene were viable and fertile. They did not show the TTP-like phenotype such as spontaneous thrombocytopenia, but intensive analyses revealed that they were prothrombotic. They are useful models to reveal how ADAMTS13 deficiency interacts with other genetic and environmental factors.

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References and recommended reading

- Papers of particular interest, published within the annual period of review, have been highlighted as:
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- Additional references related to this topic can also be found in the Current World Literature section in this issue (p. 296).
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