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## $\beta_2$ -Glycoprotein I – A Protein in Search of Function

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### 1. Introduction

$\beta_2$ -glycoprotein I is a lipid-binding 50-kDa glycoprotein that circulates in plasma at a concentration of approximately 4  $\mu$ M (200  $\mu$ g/ml). The amino acid sequence of human  $\beta_2$ -glycoprotein I was completely determined (1), the cDNAs have been isolated (2, 3) and the crystal structure has been solved (4).  $\beta_2$ -glycoprotein I is a member of the so-called "complement control protein" (CCP) superfamily, whose members are identified by the presence of one or more of a motif containing a characteristic disulfide bond pattern (5). These motifs are called CCP or sushi domains. CCP repeats are units of approximately 60 amino acids with a relatively invariant arrangement of 2 disulfide bonds and a number of other highly conserved residues. Other members of the CCP superfamily include at least 12 complement proteins, the B subunit of blood clotting factor XIII, haptoglobin, the interleukin 2 receptor and selectins.  $\beta_2$ -glycoprotein I is made up entirely of five CCP repeats. CCP5 diverges from the norm for CCPs, including CCPs 1-4 in that it has a relatively unique pattern of 3 disulfide bridges (6), and contains a positively-charged sequence, CKNKEKKC (residues 281-288), that mediates its binding site for anionic phospholipid (7). The crystal structure of  $\beta_2$ -glycoprotein I showed the four CCP domains 1-4 are arranged like a beads on a string and CCP5 folds back giving fishhook-like conformation. The CCP5 contains a central spiral structure with positively charged motif CKNKEKKC close to a hydrophobic patch (LAFW).  $\beta_2$ -glycoprotein I anchors to the anionic phospholipid membrane surface via CCP5 with its hydrophobic loop adjacent to the positively charged lysine rich region in CCP5. Subsequently,  $\beta_2$ -glycoprotein I penetrates the membrane interfacial headgroup region. This binding restricts the mobility of the lipid side chains and aggregates the vesicles without inducing fusion (8-10). In addition to anionic phospholipids,  $\beta_2$ -glycoprotein I binds to sulfatide (11), heparin (12), complement C3 (13), annexin A2 (14), platelet glycoprotein Ib (15), megalin (16), apolipoprotein receptor 2' (17) von Willebrand factor (18, 19) and possibly many others ligands. The solution structure of  $\beta_2$ -glycoprotein I was studied by small angle X-ray scattering (20), the experimentally derived curves fitted poorly to the simulated scattering curves calculated from the crystallographic coordinates of human  $\beta_2$ GPI, suggesting different conformation in solution. Recent studies with negative staining electron microscopic studies showed  $\beta_2$ -glycoprotein I can exist in two different

conformations – a circular conformation due to the interaction of CCP1 with CCP5 and an open elongated conformation consistent with the fishhook-like structure seen in the crystallographic studies (21). In closed conformation  $\beta$ 2-glycoprotein I binds less well to anionic phospholipids or to complement C3 (13). Binding to anionic phospholipids, and possibly other ligands stabilizes the elongated conformation (22). Circulating plasma  $\beta$ 2-glycoprotein I contains free thiols and these moieties are proposed to interact with platelets and endothelium, protecting these cells from oxidative stress (18). Oxidized form  $\beta$ 2-glycoprotein I is increased in patients with thrombosis (23). Oxidized  $\beta$ 2-glycoprotein I induces human dendritic cell maturation and promotes a T helper type I response (24). These studies imply the antibody response to  $\beta$ 2-glycoprotein I are due post translational modifications due to oxidative stress.

$\beta$ 2-glycoprotein I was designated as apolipoprotein H initially as it could be isolated from very low density lipoprotein fractions and had high affinity for triglyceride-rich particles (25). However, recent studies do not suggest an interaction between  $\beta$ 2-glycoprotein I with either high or low density lipoproteins (26).

Despite the extensive physicochemical characterization, the physiological role of  $\beta$ 2-glycoprotein I remains uncertain. Based on several *in vitro* studies, a wide range of functions have been attributed such as regulation of coagulation (27), modulation of complement activity and clearance of apoptotic cells from the circulation (28). In this review, we will summarize newer data on the possible physiological role of  $\beta$ 2-glycoprotein I.

## 2. Modulation of hemostasis

Since  $\beta$ 2-glycoprotein I is the target of the majority of antiphospholipid antibodies associated with thrombosis, an anticoagulant function for  $\beta$ 2-glycoprotein I was anticipated. Anionic phospholipid surfaces play an essential role in normal hemostasis by providing a site for the assembly of enzyme-cofactor complexes involved in virtually every step of the enzymatic cascade that results in the generation of fibrin, which polymerizes to form an insoluble fibrin clot. In normal cells, anionic phospholipids such as phosphatidylserine are present only in the inner leaflet of the membrane bilayer. Platelets externalize anionic phospholipid when stimulated by agonists. Binding of  $\beta$ 2-glycoprotein I to anionic phospholipid vesicles (29) and platelets (30, 31) is accompanied by inhibition of phospholipid-dependent coagulation tests (27, 32), suggesting a likely physiological role of  $\beta$ 2-glycoprotein I in the regulation of coagulation, particularly on activated platelets and possibly on other cell surfaces. In addition,  $\beta$ 2-glycoprotein I inhibits the contact activation of the intrinsic coagulation pathway (15, 33).  $\beta$ 2-glycoprotein I binds to factor XI with an affinity equivalent to that of high molecular weight kininogen. The binding inhibits the activation to factor XI by thrombin and FXIIa. This was suggested to be a mechanism, by which  $\beta$ 2-glycoprotein I may modulate thrombin generation.  $\beta$ 2-glycoprotein I also binds to heparin – a fact used in its isolation (12, 29). Heparin binding site had been localized to the positively charged CCP5 (12). Heparin also promotes plasmin cleavage of  $\beta$ 2-glycoprotein I at Lys317-Thr318 bond (34). Plasmin-cleaved  $\beta$ 2-glycoprotein I has markedly decreased affinity for anionic phospholipid. This form of cleaved  $\beta$ 2-glycoprotein I is seen in patients treated with streptokinase and in patients with disseminated intravascular coagulation (35), showing this cleavage reaction can occur *in vivo* during accelerated fibrinolysis.

Several procoagulant effect of  $\beta$ <sub>2</sub>-glycoprotein I have also been described.  $\beta$ <sub>2</sub>-glycoprotein I binds to thrombin and protects it from inactivation by heparin cofactor II/heparin complex (36). Furthermore, Mori et al (37) showed  $\beta$ <sub>2</sub>-glycoprotein inhibited activated protein C inactivation of factor Va – an effect diminished by the addition of phospholipids. At similar concentration,  $\beta$ <sub>2</sub>-glycoprotein I inhibited weakly factor Va- and phospholipid-dependent prothrombinase activity. The depletion of beta  $\beta$ <sub>2</sub>-glycoprotein I from plasma led to only a slight shortening of the diluted Russell's viper venom-dependent clotting time, but to a strong and significant potentiation of the anticoagulant activity of APC. These results suggest that under certain physiological conditions  $\beta$ <sub>2</sub>-glycoprotein I may have procoagulant function.

In contrast to these hemostatic activities demonstrated in vivo, neither the  $\beta$ <sub>2</sub>-glycoprotein I-deficient mice (generated by homologous recombination) nor  $\beta$ <sub>2</sub>-glycoprotein I-deficient individuals exhibit any bleeding manifestations (38-40). On the contrary,  $\beta$ <sub>2</sub>-glycoprotein I-deficient mice have diminished rate of thrombin generation compared with normal or even with heterozygous mice. No significant differences in clotting time were observed in plasma from these three genotypes when measured by dRVVT, dKCT, aPTT, and protein C pathway assays (41). Hereditary deficiency of  $\beta$ <sub>2</sub>-glycoprotein I was reported since 1968 (42), and its potential association with risk of thrombosis had been examined. Bansci *et al.* (43) have described two brothers with total deficiency of  $\beta$ <sub>2</sub>-glycoprotein I, one of whom had experienced recurrent unexplained thrombosis by age 36. However, six other heterozygous individuals (ages 9-73) from this family and the proband's brother with homozygous deficiency were free of thrombosis. Takeuchi et al (39) described two asymptomatic individuals with complete deficiency of  $\beta$ <sub>2</sub>-glycoprotein I. The routine coagulation assays were normal. A slight shortening of the DRVVT was observed in these individuals, which interestingly were not corrected by exogenous addition of  $\beta$ <sub>2</sub>-glycoprotein I.

Thrombosis is a complex multigene phenotype (44). Because of the large number of genes that influence this phenotype teasing out the role of  $\beta$ <sub>2</sub>-glycoprotein I in in this prothrombotic phenotype will be difficult. It is also possible that the thrombosis seen with antiphospholipid antibodies is not related to any of interaction identified above.

### 3. $\beta$ <sub>2</sub>-glycoprotein I as an opsonin

The term opsonins is used to refer molecules that target a cell for phagocytosis. A number of observations suggest  $\beta$ <sub>2</sub>-glycoprotein I can be an opsonin for clearance of anionic phospholipid vesicles containing surfaces from the circulation. In normal cells, anionic phospholipids such as phosphatidylserine are present only in the inner leaflet of membrane bilayer. There is transbilayer movement of phosphatidylserine during apoptosis and phosphatidylserine exposed can be a tag for their clearance by macrophages (45-47). In artificial membranes, the phosphatidylserine content has to be at least 5-10% before a significant binding of  $\beta$ <sub>2</sub>-glycoprotein I could be observed (48). Nevertheless, the binding of  $\beta$ <sub>2</sub>-glycoprotein I to phosphatidylserine containing surfaces such as apoptotic cells and platelet microvesicles have been shown (49, 50). In addition to the anionic phospholipids,  $\beta$ <sub>2</sub>-glycoprotein I is also shown to bind the Ro/SSA, a 60 kDa a nuclear antigen and target of autoantibodies in primary Sjogren syndrome (19). Ro/SSA translocates to cell surface during apoptosis and can serve as additional binding site. The complex of anionic

phospholipid and  $\beta$ 2-glycoprotein I are taken into a receptor-mediated pathway by macrophages and possibly endothelial cells also. The phagocytic receptors mediating the uptake have been shown to be toll-like receptor 4 in macrophages (51) and lipoprotein receptor related family members (49). In endothelial cells toll-like receptor 2 and 4 (52, 53), annexin A2 (14), and apolipoprotein E receptor 2 (54) have been implicated. Deficiencies of factors, implicated in the removal of apoptotic cells such as lactadherin and Gas6 receptors, are associated with systemic lupus erythematosus and autoimmunity (55). However, no immunological dysfunction is reported in  $\beta$ 2-glycoprotein I deficiency.

$\beta$ 2-glycoprotein I may also have a role in the clearance of exogenous liposomes. Liposomes have been used extensively as vehicles for drug delivery and following in vivo infusions, liposomes are preferentially taken up by the mononuclear phagocytic cells of the reticuloendothelial system (56). In 1982, Wurm et al (57) showed that infusion of  $\beta$ 2-glycoprotein I in rats results in an accelerated clearance of triglyceride-rich vesicles from the circulation. The clearance of liposomes by the phagocytic cells, is markedly affected lipid composition of the liposomes and anionic phospholipid containing are cleared very rapidly from blood (56). By analyzing the proteins that associate with the liposomes in blood, Chonn et al. have identified  $\beta$ 2-glycoprotein I as a major protein associated with rapidly cleared liposomes and noted that pretreating the mice with anti- $\beta$ 2-glycoprotein I antibodies markedly increased the circulating half-life of the liposomes (58). It is interesting to note that in 1982, Wurm et al (57) showed that infusion of  $\beta$ 2-glycoprotein I in rats results in an accelerated clearance of triglyceride-rich vesicles from the circulation.

The complement system is involved in the clearance of dead cells and debris from the circulation and recently a role for  $\beta$ 2-glycoprotein I its regulation has been identified (13). The elongated and open conformation of  $\beta$ 2-glycoprotein I binds to C3 and induces a conformational changes so that the regulator factor H binds. As factor H promotes factor I-induced the cleavage of C3,  $\beta$ 2-glycoprotein I acts as special cofactor for factor H and factor I. The enhanced the degradation of C3 limits further complement amplification. Deficiencies of complement factor H and I are associated atypical hemolytic uremic syndrome and no such association has been described for  $\beta$ 2-glycoprotein I.

#### **4. A role in gestation**

Because of the association with fetal loss and anti- $\beta$ 2-glycoprotein I antibodies, a role in gestation has been proposed. Infusion of cyanine labeled  $\beta$ 2-glycoprotein I in mice show preferential localization on the endothelium of uterine vessels and at the implantation sites in pregnant mice (59), suggesting a role in early gestation. However, the  $\beta$ 2-glycoprotein I null mice were fertile and carried viable fetuses to term and there were no thrombosis in placental vessels (60). Nevertheless, there was an 18% reduction in the number of viable implantation sites and reduced fetal weight and fetal:placental weight ratio in late gestation in  $\beta$ 2-glycoprotein I null mice.

#### **5. $\beta$ 2-glycoprotein I and angiogenesis**

$\beta$ 2-glycoprotein I is enzymatically cleaved by plasmin at the peptide bond between Lys317-Thr318 to form a cleaved form  $\beta$ 2-glycoprotein I (61, 62). This form is seen in the circulation in patients with increased fibrinolysis. The cleaved form of  $\beta$ 2-glycoprotein I binds to

plasminogen and inhibits plasmin generation. In addition to modulating fibrinolysis, a role in angiogenesis had been proposed for the cleaved form of  $\beta_2$ -glycoprotein I. The cleaved form of  $\beta_2$ -glycoprotein I inhibits endothelial cell proliferation *in vitro*, inhibits neovascularization into subcutaneously implanted angiogenic matrices and the growth of orthotopic prostate cancer in C57BL/6 mice (63, 64). The cleaved  $\beta_2$ -glycoprotein I strongly reduced HUVEC growth and proliferation as evidenced by the MTT and BrdU assay and delayed cell cycle progression arresting endothelial cells in the S- and G2/M-phase (65). However, the cleaved form of  $\beta_2$ -glycoprotein I can also promote angiogenesis as it binds angiostatin 4.5 (plasminogen kringle 1-5) and attenuates its antiangiogenic property (66). The murine  $\beta$  *in vivo* apparently displayed only mild anti-angiogenic properties.  $\beta_2$ -glycoprotein I deficient mice developed larger tumors with more vessels than  $\beta_2$ -glycoprotein I replete mice but no survival benefit is conferred to tumor bearing animals regardless of  $\beta_2$ GPI status raising questions about its pathophysiological role in tumorigenesis(66).

## 6. Conclusion

Since its discovery in the sixties and following the recognition that it is the antigenic target for antiphospholipid antibodies in nineties, several structural and functional studies have been described. However, there is no convincing pathogenetic mechanism or theoretical framework for the hypercoagulable state associated with antibodies to this protein. Many hypotheses have been proposed based on *in vitro* findings and most of them revolve on the anionic phospholipid binding properties of  $\beta_2$ -glycoprotein I. At least two patients are described with antiphospholipid syndrome who had mutations in  $\beta_2$ -glycoprotein I rendering it incapable of binding phospholipids (67, 68), questioning its phospholipid binding in pathogenesis. These findings underscore the importance of finding its physiological function to elucidate the mechanism of thrombosis seen with antibody to this molecule.

## 7. Acknowledgment

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## 8. References

- [1] Lozier, J., Takahashi, N., and Putnam, F.W. 1984. Complete amino acid sequence of human plasma beta 2-glycoprotein I. *Proc Natl Acad Sci U S A* 81:3640-3644.
- [2] Steinkasserer, A., Estaller, C., Weiss, E.H., Sim, R.B., and Day, A.J. 1991. Complete nucleotide and deduced amino acid sequence of human beta 2-glycoprotein I. *Biochem J* 277 ( Pt 2):387-391.
- [3] Day, J.R., O'Hara, P.J., Grant, F.J., Lofton-Day, C., Berkaw, M.N., Werner, P., and Arnaud, P. 1992. Molecular cloning and sequence analysis of the cDNA encoding human apolipoprotein H (beta 2-glycoprotein I). *Int J Clin Lab Res* 21:256-263.
- [4] Schwarzenbacher, R., Zeth, K., Diederichs, K., Gries, A., Kostner, G.M., Laggner, P., and Prassl, R. 1999. Crystal structure of human beta2-glycoprotein I: implications for phospholipid binding and the antiphospholipid syndrome. *EMBO J* 18:6228-6239.
- [5] Reid, K.B., and Day, A.J. 1989. Structure-function relationships of the complement components. *Immunol Today* 10:177-180.

- [6] Kato, H., and Enjyoji, K. 1991. Amino acid sequence and location of the disulfide bonds in bovine beta 2 glycoprotein I: the presence of five Sushi domains. *Biochemistry* 30:11687-11694.
- [7] Hunt, J., and Krilis, S. 1994. The fifth domain of beta 2-glycoprotein I contains a phospholipid binding site (Cys281-Cys288) and a region recognized by anticardiolipin antibodies. *J Immunol* 152:653-659.
- [8] Willems, G.M., Janssen, M.P., Pelsers, M.M., Comfurius, P., Galli, M., Zwaal, R.F., and Bevers, E.M. 1996. Role of divalency in the high-affinity binding of anticardiolipin antibody-beta 2-glycoprotein I complexes to lipid membranes. *Biochemistry* 35:13833-13842.
- [9] Hammel, M., Schwarzenbacher, R., Gries, A., Kostner, G.M., Laggner, P., and Prassl, R. 2001. Mechanism of the interaction of beta(2)-glycoprotein I with negatively charged phospholipid membranes. *Biochemistry* 40:14173-14181.
- [10] Gushiken, F.C., Le, A., Arnett, F.C., and Thiagarajan, P. 2002. Polymorphisms beta2-glycoprotein I: phospholipid binding and multimeric structure. *Thromb Res* 108:175-180.
- [11] Merten, M., Motamedy, S., Ramamurthy, S., Arnett, F.C., and Thiagarajan, P. 2003. Sulfatides: targets for anti-phospholipid antibodies. *Circulation* 108:2082-2087.
- [12] Guerin, J., Sheng, Y., Reddel, S., Iverson, G.M., Chapman, M.G., and Krilis, S.A. 2002. Heparin inhibits the binding of beta 2-glycoprotein I to phospholipids and promotes the plasmin-mediated inactivation of this blood protein. Elucidation of the consequences of the two biological events in patients with the anti-phospholipid syndrome. *J Biol Chem* 277:2644-2649.
- [13] Gropp, K., Weber, N., Reuter, M., Micklisch, S., Kopka, I., Hallstrom, T., and Skerka, C. {beta}2 glycoprotein 1 ({beta}2GPI), the major target in anti phospholipid syndrome (APS), is a special human complement regulator. *Blood*.
- [14] Ma, K., Simantov, R., Zhang, J.C., Silverstein, R., Hajjar, K.A., and McCrae, K.R. 2000. High affinity binding of beta 2-glycoprotein I to human endothelial cells is mediated by annexin II. *J Biol Chem* 275:15541-15548.
- [15] Shi, T., Iverson, G.M., Qi, J.C., Cockerill, K.A., Linnik, M.D., Konecny, P., and Krilis, S.A. 2004. Beta 2-Glycoprotein I binds factor XI and inhibits its activation by thrombin and factor XIIa: loss of inhibition by clipped beta 2-glycoprotein I. *Proc Natl Acad Sci U S A* 101:3939-3944.
- [16] Moestrup, S.K., Schousboe, I., Jacobsen, C., Leheste, J.R., Christensen, E.I., and Willnow, T.E. 1998. beta2-glycoprotein-I (apolipoprotein H) and beta2-glycoprotein-I-phospholipid complex harbor a recognition site for the endocytic receptor megalin. *J Clin Invest* 102:902-909.
- [17] van Lummel, M., Pennings, M.T., Derksen, R.H., Urbanus, R.T., Lutters, B.C., Kaldenhoven, N., and de Groot, P.G. 2005. The binding site in {beta}2-glycoprotein I for ApoER2' on platelets is located in domain V. *J Biol Chem* 280:36729-36736.
- [18] Passam, F.H., Rahgozar, S., Qi, M., Raftery, M.J., Wong, J.W., Tanaka, K., Ioannou, Y., Zhang, J.Y., Gemmell, R., Qi, J.C., et al. Redox control of beta2-glycoprotein I-von Willebrand factor interaction by thioredoxin-1. *J Thromb Haemost* 8:1754-1762.
- [19] Reed, J.H., Giannakopoulos, B., Jackson, M.W., Krilis, S.A., and Gordon, T.P. 2009. Ro 60 functions as a receptor for beta(2)-glycoprotein I on apoptotic cells. *Arthritis Rheum* 60:860-869.

- [20] Hammel, M., Kriechbaum, M., Gries, A., Kostner, G.M., Laggner, P., and Prassl, R. 2002. Solution structure of human and bovine beta(2)-glycoprotein I revealed by small-angle X-ray scattering. *J Mol Biol* 321:85-97.
- [21] Agar, C., van Os, G.M., Morgelin, M., Sprenger, R.R., Marquart, J.A., Urbanus, R.T., Derksen, R.H., Meijers, J.C., and de Groot, P.G. Beta2-glycoprotein I can exist in 2 conformations: implications for our understanding of the antiphospholipid syndrome. *Blood* 116:1336-1343.
- [22] Pengo, V., Biasiolo, A., and Fior, M.G. 1995. Autoimmune antiphospholipid antibodies are directed against a cryptic epitope expressed when beta 2-glycoprotein I is bound to a suitable surface. *Thromb Haemost* 73:29-34.
- [23] Ioannou, Y., Zhang, J.Y., Qi, M., Gao, L., Qi, J.C., Yu, D.M., Lau, H., Sturgess, A.D., Vlachoyiannopoulos, P.G., Moutsopoulos, H.M., et al. Novel assays of thrombogenic pathogenicity for the antiphospholipid syndrome based on the detection of molecular oxidative modification of the major autoantigen ss2-glycoprotein I. *Arthritis Rheum*.
- [24] Buttari, B., Profumo, E., Mattei, V., Siracusano, A., Ortona, E., Margutti, P., Salvati, B., Sorice, M., and Rigano, R. 2005. Oxidized beta2-glycoprotein I induces human dendritic cell maturation and promotes a T helper type 1 response. *Blood* 106:3880-3887.
- [25] Lee, N.S., Brewer, H.B., Jr., and Osborne, J.C., Jr. 1983. beta 2-Glycoprotein I. Molecular properties of an unusual apolipoprotein, apolipoprotein H. *J Biol Chem* 258:4765-4770.
- [26] Agar, C., de Groot, P.G., Levels, J.H., Marquart, J.A., and Meijers, J.C. 2009. Beta2-glycoprotein I is incorrectly named apolipoprotein H. *J Thromb Haemost* 7:235-236.
- [27] Nimpf, J., Bevers, E.M., Bomans, P.H., Till, U., Wurm, H., Kostner, G.M., and Zwaal, R.F. 1986. Prothrombinase activity of human platelets is inhibited by beta 2-glycoprotein-I. *Biochim Biophys Acta* 884:142-149.
- [28] Balasubramanian, K., Chandra, J., and Schroit, A.J. 1997. Immune clearance of phosphatidylserine-expressing cells by phagocytes. The role of beta2-glycoprotein I in macrophage recognition. *J Biol Chem* 272:31113-31117.
- [29] Wurm, H. 1984. beta 2-Glycoprotein-I (apolipoprotein H) interactions with phospholipid vesicles. *Int J Biochem* 16:511-515.
- [30] Schousboe, I. 1980. Binding of beta 2-glycoprotein I to platelets: effect of adenylate cyclase activity. *Thromb Res* 19:225-237.
- [31] Nimpf, J., Wurm, H., and Kostner, G.M. 1985. Interaction of beta 2-glycoprotein-I with human blood platelets: influence upon the ADP-induced aggregation. *Thromb Haemost* 54:397-401.
- [32] Bevers, E.M., Janssen, M.P., Comfurius, P., Balasubramanian, K., Schroit, A.J., Zwaal, R.F., and Willems, G.M. 2005. Quantitative determination of the binding of beta2-glycoprotein I and prothrombin to phosphatidylserine-exposing blood platelets. *Biochem J* 386:271-279.
- [33] Schousboe, I., and Rasmussen, M.S. 1995. Synchronized inhibition of the phospholipid mediated autoactivation of factor XII in plasma by beta 2-glycoprotein I and anti-beta 2-glycoprotein I. *Thromb Haemost* 73:798-804.
- [34] Ohkura, N., Hagihara, Y., Yoshimura, T., Goto, Y., and Kato, H. 1998. Plasmin can reduce the function of human beta2 glycoprotein I by cleaving domain V into a nicked form. *Blood* 91:4173-4179.

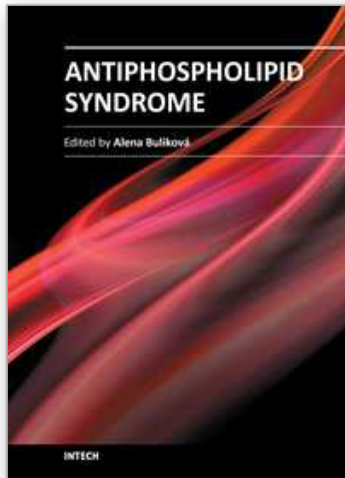


- [35] Horbach, D.A., van Oort, E., Lisman, T., Meijers, J.C., Derksen, R.H., and de Groot, P.G. 1999. Beta2-glycoprotein I is proteolytically cleaved in vivo upon activation of fibrinolysis. *Thromb Haemost* 81:87-95.
- [36] Rahgozar, S., Giannakopoulos, B., Yan, X., Wei, J., Cheng Qi, J., Gemmell, R., and Krilis, S.A. 2008. Beta2-glycoprotein I protects thrombin from inhibition by heparin cofactor II: potentiation of this effect in the presence of anti-beta2-glycoprotein I autoantibodies. *Arthritis Rheum* 58:1146-1155.
- [37] Mori, T., Takeya, H., Nishioka, J., Gabazza, E.C., and Suzuki, K. 1996. beta 2-Glycoprotein I modulates the anticoagulant activity of activated protein C on the phospholipid surface. *Thromb Haemost* 75:49-55.
- [38] Sheng, Y., Reddel, S.W., Herzog, H., Wang, Y.X., Brighton, T., France, M.P., Robertson, S.A., and Krilis, S.A. 2001. Impaired thrombin generation in beta 2-glycoprotein I null mice. *J Biol Chem* 276:13817-13821.
- [39] Takeuchi, R., Atsumi, T., Ieko, M., Takeya, H., Yasuda, S., Ichikawa, K., Tsutsumi, A., Suzuki, K., and Koike, T. 2000. Coagulation and fibrinolytic activities in 2 siblings with beta(2)-glycoprotein I deficiency. *Blood* 96:1594-1595.
- [40] Yasuda, S., Tsutsumi, A., Chiba, H., Yanai, H., Miyoshi, Y., Takeuchi, R., Horita, T., Atsumi, T., Ichikawa, K., Matsuura, E., et al. 2000. beta(2)-glycoprotein I deficiency: prevalence, genetic background and effects on plasma lipoprotein metabolism and hemostasis. *Atherosclerosis* 152:337-346.
- [41] Miyakis, S., Robertson, S.A., and Krilis, S.A. 2004. Beta-2 glycoprotein I and its role in antiphospholipid syndrome-lessons from knockout mice. *Clin Immunol* 112:136-143.
- [42] Cleve, H. 1968. [Genetic studies on the deficiency of beta 2-glycoprotein I of human serum]. *Humangenetik* 5:294-304.
- [43] Bancsi, L.F., van der Linden, I.K., and Bertina, R.M. 1992. Beta 2-glycoprotein I deficiency and the risk of thrombosis. *Thromb Haemost* 67:649-653.
- [44] Zoller, B., Garcia de Frutos, P., Hillarp, A., and Dahlback, B. 1999. Thrombophilia as a multigenic disease. *Haematologica* 84:59-70.
- [45] Navratil, J.S., and Ahearn, J.M. 2001. Apoptosis, clearance mechanisms, and the development of systemic lupus erythematosus. *Curr Rheumatol Rep* 3:191-198.
- [46] Ravichandran, K.S., and Lorenz, U. 2007. Engulfment of apoptotic cells: signals for a good meal. *Nat Rev Immunol* 7:964-974.
- [47] Casciola-Rosen, L., Rosen, A., Petri, M., and Schliessel, M. 1996. Surface blebs on apoptotic cells are sites of enhanced procoagulant activity: implications for coagulation events and antigenic spread in systemic lupus erythematosus. *Proc Natl Acad Sci U S A* 93:1624-1629.
- [48] Thiagarajan, P., Le, A., and Benedict, C.R. 1999. Beta(2)-glycoprotein I promotes the binding of anionic phospholipid vesicles by macrophages. *Arterioscler Thromb Vasc Biol* 19:2807-2811.
- [49] Maiti, S.N., Balasubramanian, K., Ramoth, J.A., and Schroit, A.J. 2008. Beta-2-glycoprotein 1-dependent macrophage uptake of apoptotic cells. Binding to lipoprotein receptor-related protein receptor family members. *J Biol Chem* 283:3761-3766.
- [50] Nomura, S., Komiyama, Y., Matsuura, E., Kokawa, T., Takahashi, H., and Koike, T. 1993. Binding of beta 2-glycoprotein I to platelet-derived microparticles. *Br J Haematol* 85:639-640.

- [51] Lambrianides, A., Carroll, C.J., Pierangeli, S.S., Pericleous, C., Branch, W., Rice, J., Latchman, D.S., Townsend, P., Isenberg, D.A., Rahman, A., et al. Effects of polyclonal IgG derived from patients with different clinical types of the antiphospholipid syndrome on monocyte signaling pathways. *J Immunol* 184:6622-6628.
- [52] Alard, J.E., Gaillard, F., Daridon, C., Shoenfeld, Y., Jamin, C., and Youinou, P. TLR2 is one of the endothelial receptors for beta 2-glycoprotein I. *J Immunol* 185:1550-1557.
- [53] Pierangeli, S.S., Vega-Ostertag, M.E., Raschi, E., Liu, X., Romay-Penabad, Z., De Micheli, V., Galli, M., Moia, M., Tincani, A., Borghi, M.O., et al. 2007. Toll-like receptor and antiphospholipid mediated thrombosis: in vivo studies. *Ann Rheum Dis* 66:1327-1333.
- [54] Romay-Penabad, Z., Aguilar-Valenzuela, R., Urbanus, R.T., Derksen, R.H., Pennings, M.T., Papalardo, E., Shilagard, T., Vargas, G., Hwang, Y., de Groot, P.G., et al. Apolipoprotein E receptor 2 is involved in the thrombotic complications in a murine model of the antiphospholipid syndrome. *Blood* 117:1408-1414.
- [55] Nagata, S., Hanayama, R., and Kawane, K. Autoimmunity and the clearance of dead cells. *Cell* 140:619-630.
- [56] Senior, J.H. 1987. Fate and behavior of liposomes in vivo: a review of controlling factors. *Crit Rev Ther Drug Carrier Syst* 3:123-193.
- [57] Wurm, H., Beubler, E., Polz, E., Holasek, A., and Kostner, G. 1982. Studies on the possible function of beta 2-glycoprotein-I: influence in the triglyceride metabolism in the rat. *Metabolism* 31:484-486.
- [58] Chonn, A., Semple, S.C., and Cullis, P.R. 1995. Beta 2 glycoprotein I is a major protein associated with very rapidly cleared liposomes in vivo, suggesting a significant role in the immune clearance of "non-self" particles. *J Biol Chem* 270:25845-25849.
- [59] Agostinis, C., Biffi, S., Garrovo, C., Durigutto, P., Lorenzon, A., Bek, A., Bulla, R., Grossi, C., Borghi, M.O., Meroni, P., et al. In vivo distribution of {beta}2 glycoprotein I under various pathophysiological conditions. *Blood*.
- [60] Robertson, S.A., Roberts, C.T., van Beijering, E., Pensa, K., Sheng, Y., Shi, T., and Krilis, S.A. 2004. Effect of beta2-glycoprotein I null mutation on reproductive outcome and antiphospholipid antibody-mediated pregnancy pathology in mice. *Mol Hum Reprod* 10:409-416.
- [61] Hunt, J.E., Simpson, R.J., and Krilis, S.A. 1993. Identification of a region of beta 2-glycoprotein I critical for lipid binding and anti-cardiolipin antibody cofactor activity. *Proc Natl Acad Sci U S A* 90:2141-2145.
- [62] Yasuda, S., Atsumi, T., Ieko, M., Matsuura, E., Kobayashi, K., Inagaki, J., Kato, H., Tanaka, H., Yamakado, M., Akino, M., et al. 2004. Nicked beta2-glycoprotein I: a marker of cerebral infarct and a novel role in the negative feedback pathway of extrinsic fibrinolysis. *Blood* 103:3766-3772.
- [63] Beecken, W.D., Engl, T., Ringel, E.M., Camphausen, K., Michaelis, M., Jonas, D., Folkman, J., Shing, Y., and Blaheta, R.A. 2006. An endogenous inhibitor of angiogenesis derived from a transitional cell carcinoma: clipped beta2-glycoprotein-I. *Ann Surg Oncol* 13:1241-1251.
- [64] Sakai, T., Balasubramanian, K., Maiti, S., Halder, J.B., and Schroit, A.J. 2007. Plasmin-cleaved beta-2-glycoprotein 1 is an inhibitor of angiogenesis. *Am J Pathol* 171:1659-1669.

- [65] Beecken, W.D., Ringel, E.M., Babica, J., Oppermann, E., Jonas, D., and Blaheta, R.A. Plasmin-clipped beta(2)-glycoprotein-I inhibits endothelial cell growth by down-regulating cyclin A, B and D1 and up-regulating p21 and p27. *Cancer Lett* 296:160-167.
- [66] Nakagawa, H., Yasuda, S., Matsuura, E., Kobayashi, K., Ieko, M., Kataoka, H., Horita, T., Atsumi, T., Koike, T. 2009. Nicked  $\beta$ 2-glycoprotein I binds angiostatin 4.5 (plasminogen kringle 1-5) and attenuates its antiangiogenic property. *Blood* 114:2553-2559.
- [67] Passam, F.H., Qi, J.C., Tanaka, K., Matthaei, K.I., and Krilis, S.A. In vivo modulation of angiogenesis by beta 2 glycoprotein I. *J Autoimmun* 35:232-240.
- [68] Nash, M.J., Camilleri, R.S., Liesner, R., Mackie, I.J., Machin, S.J., and Cohen, H. 2003. Paradoxical association between the 316 Trp to Ser beta 2-glycoprotein I (Beta2GPI) polymorphism and anti-Beta2GPI antibodies. *Br J Haematol* 120:529-531.
- [69] Gushiken, F.C., Arnett, F.C., and Thiagarajan, P. 2000. Primary antiphospholipid antibody syndrome with mutations in the phospholipid binding domain of beta(2)-glycoprotein I. *Am J Hematol* 65:160-165.

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The antiphospholipid syndrome has been described for the first time by Graham Hughes in 1983 as a condition connected with thromboses or foetal losses and antiphospholipid antibodies presence. From that time there has been a great progress in knowledge, including antiphospholipid antibodies characterisation, their probable and also possible action, clinical manifestations, laboratory detection and treatment possibilities. This book provides a wide spectrum of clinical manifestations through Chapters written by well known researchers and clinicians with a great practical experience in management of diagnostics or treatment of antiphospholipid antibodies' presence.

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