Heritable Thrombophilia and Hypofibrinolysis

Possible Causes of Retinal Vein Occlusion

Charles J. Glueck, MD; Howard Bell, MD; Lou Vadlamani, MD; Arun Gupta, PhD; Robert N. Fontaine, PhD; Ping Wang, PhD; Davis Stroop, MS; Ralph Gruppo, MD

Objective: To determine whether heritable thrombophilia and hypofibrinolysis were risk factors for retinal vein occlusion.

Design: Measures of thrombophilia (increased likelihood of thrombus formation) included anticardiolipin antibodies (IgG and IgM), the lupus anticoagulant (including dilute Russell viper venom clotting time), antithrombin C and S, and homocysteine. Polymerase chain reaction assays were performed for 3 thrombophilic gene mutations (factor V Leiden, methylenetetrahydrofolate reductase, and prothrombin gene). Measures of hypofibrinolysis (reduced ability to lyse thrombi) included lipoprotein Lp(a), plasminogen activator inhibitor activity, and polymerase chain reaction analysis of the hypofibrinolytic 4G/5G polymorphism of the PAI1 gene. These coagulation measures were performed in 17 patients with retinal vein occlusions with comparison with serologic coagulation measures and polymerase chain reaction assays in 40 and 234 healthy normal volunteers as controls, respectively.

Results: Of 14 patients with retinal vein occlusion with measures of dilute Russell viper venom clotting time, a thrombophilic antiphospholipid antibody, 6 (43%) had abnormal results (>38.8 seconds) compared with 1 (3%) of 30 controls (P = .002). Of 17 patients with vein occlusion, 3 (18%) were heterozygous for the thrombophilic factor V Leiden G1691A mutation compared with 7 (3%) of 233 controls (P = .02). Of 17 patients with vein occlusion, 2 (12%) had normal alleles (5G/5G) for the plasminogen activator inhibitor gene promoter; the other 15 (88%) were heterozygous or homozygous for the 4G polymorphism, which is associated with hypofibrinolysis. Of 234 controls, 85 (36.3%) had the 5G/5G allele; 149 (63.7%) were heterozygous or homozygous for the 4G polymorphism (P = .03). Patients with vein occlusion were more likely to have high levels of the major determinant of hypofibrinolysis, plasminogen activator inhibitor activity. These levels were high (>22 U/L) in 6 (38%) of 16 patients with vein occlusion compared with 1 (2%) of 40 controls (χ² = 12.8; P = .001). Patients with vein occlusion were more likely (8/16 [50%]) to have high levels of hypofibrinolytic Lp(a) (>35 mg/dL) than controls (5/40 [13%]; χ² = 9; P = .003). The median Lp(a) level in patients with vein occlusion who had the 4G/4G genotype was 62 mg/dL compared with 5.3 mg/dL in controls with the 4G/4G genotype (P = .05).

Conclusion: Thrombophilia and hypofibrinolysis are possible causes of retinal vein occlusion.

PATIENTS AND METHODS

STUDY PROTOCOL

The study was approved by the Institutional Review Board of the Jewish Hospital, Cincinnati, Ohio, and was carried out with informed consent.

Blood specimens for measures of thrombophilia, hypofibrinolysis, and lipid profiles were obtained between 8 and 10 AM after an overnight fast from seated patients with vein occlusion. Retinal vein occlusion was diagnosed by the referring ophthalmologists based on either or both the results of fluorescein angiography and characteristic fundus features.

cDNA PCR ASSAYS

A full range of cDNA PCR assays and measures of thrombophilia and hypofibrinolysis were systematically performed in patients with vein occlusion and in the healthy control groups. The PCR analyses of 3 thrombophilic gene mutations (factor V Leiden G1691A mutation, the MTHFR C677T mutation, and the 20210A allele of the prothrombin gene) and the hypofibrinolytic 4G/5G polymorphism of the PAI1 gene promoter were performed at the Coagulation Research Laboratory, Children’s Hospital Medical Center, Cincinnati.

Genomic DNA for each PCR assay was obtained by a salting-out procedure. A PCR assay was performed for factor V Leiden to determine the presence or absence of a single point mutation at nucleotide 1691 leading to a glutamine substitution for an arginine residue at amino acid 506 in the factor V molecule. The PCR product was digested with the restriction enzyme MnlI. A PCR assay was performed for the enzyme MTHFR to determine the presence of a point mutation at nucleotide 677 leading to a valine substitution for an alanine residue. The PCR product was digested with the restriction enzyme Hinfl. A PCR assay for the prothrombin gene was performed to determine the presence or absence of a point mutation at nucleotide 20210 leading to a G-to-A transition. The PCR product was digested with the restriction enzyme HindIII. A PCR assay was performed for the enzyme PAI1 gene promoter region of the PAI1 gene 675 base pairs upstream from the transcriptional starting site, resulting in 2 alleles containing either 4 or 5 guanosines.

MEASURES OF THROMBOPHILIA

Measures of thrombophilia included antithrombin III, anticoagulant proteins C and S (Coagulation Research Laboratory, Children’s Hospital Medical Center), the lupus anticoagulant, and homocysteine (Alliance Hospital Laboratories, Cincinnati, Ohio). Established, previously published methods were used.

MEASURES OF HYPOFIBRINOLYSIS

The major hypofibrinolytic factors measured by coagulation techniques included Lp(a) (Alliance Hospital Laboratories) and PAI-Fx (Coagulation Research Laboratory, Children’s Hospital Medical Center).

PATIENTS

Of the 17 patients with retinal vein occlusion, 15 were referred from the practice of a single ophthalmologist and 2 from other ophthalmology practices. The diagnosis of retinal vein occlusion was made by ophthalmoscopic fundus examination revealing disc swelling, venous dilation or tortuosity, retinal hemorrhages, and cotton-wool spots and by fluorescein angiography demonstrating extensive areas of capillary closure, venous filling defects, and increased venous transit time.

CONTROL SUBJECTS

For comparison with the cDNA polymorphisms of the patients with vein occlusion, 234 healthy controls (194 children and 40 adults) were used; for comparison with their coagulation measures, 40 healthy adult controls were used.

STATISTICAL METHODS

Proportions of patients with retinal vein occlusion and those of controls having abnormalities of thrombophilia or hypofibrinolysis were compared between patients and controls by fitting regression lines for patients and controls and then comparing slopes (Figure 5). To determine whether the 3 patients with retinal vein occlusion who also had osteonecrosis disproportionately accounted for patient-control differences, all of the statistical analyses were repeated after excluding these 3 patients. Data are expressed as the mean ± SD.

After the discovery that resistance to activated protein C (mediated by the thrombophilic mutant factor V Leiden gene) is a major risk factor for venous thrombosis, Williamson, Larsson, and Scat and their colleagues reported a higher prevalence of resistance to activated protein C in patients with retinal vein occlusion than in controls. A 1996 study by Graham et al, however, found no significant association between the factor V Leiden G1691A mutation and the presence of retinal vein occlusion. Furthermore, a recent study by Gottlieb et al in patients younger than 50 years with central retinal vein occlusion reported protein C resistance in
The body maintains an elegant balance between thrombosis and fibrinolysis (Figure 1).14 Thromboplastin (tissue factor) arises from endothelial cells and macrophages, the extrinsic pathway. Tissue factor acts through factor VII on factor X, the final common pathway for thrombosis initiation. Factor X, in turn, acts on prothrombin. Mutations in the prothrombin gene lead to high prothrombin levels13 and increase the risk of thrombosis. Platelets, exposed collagen, and bacterial endotoxins activate the intrinsic pathway, acting through factors VIII and IX on factor X. Factor Xa (the activated form) binds to factor Va and mediates the conversion of prothrombin to thrombin, which then acts on fibrinogen to form a fibrin clot. Thrombin and factor Xa are inhibited by antithrombin III. Protein C is activated by thrombin, which is bound at the endothelium to thrombomodulin, and once activated, protein C inactivates factors VIIa and Va. The factor V Leiden mutation affects factor V, rendering it resistant to inactivation by protein C, a thrombophilic effect.16 Protein S enhances protein C’s inactivation of factors Va and VIIa. Proteins C and S and antithrombin III are thus endogenous anticoagulants; deficiencies of proteins C and S have been associated with osteonecrosis and with arterial thrombosis.14 After the fibrin clot is formed, it is lysed through the action of plasmin (fibrinolysis), which arises from plasminogen. The conversion of plasminogen to plasmin is stimulated by tissue plasminogen activator, which in turn is inhibited by plasminogen activator inhibitor 1 (PAI-1). The 4G/4G polymorphism of the PAI1 gene is hypofibrinolytic, associated with high levels of plasminogen activator inhibitor activity (PAI-Fx).17 Low tissue plasminogen activator levels, high PAI-1 levels, or both, major causes of hypofibrinolysis, have been associated with osteonecrosis and are also risk factors for arterial thrombosis. Not depicted in Figure 1 because it has multiple postulated interactions with the coagulation cascade is the common mutation in the methylenetetrahydrofolate reductase (MTHFR) gene, associated with increased levels of the thrombophilic amino acid, homocysteine.

Further confirmation of pathogenic relationships between hypofibrinolysis and thrombophilia and retinal vein occlusion,3-5,7-10,13 with better understanding of the molecular genetic cause of venous thrombosis,14-20 should promote further studies of the surveillance, prevention, and treatment of central retinal vein thrombosis.

Our specific aim was to systematically assess measures of thrombophilia and hypofibrinolysis in patients with retinal vein occlusion, including newly developed complementary DNA (cDNA) polymerase chain reaction (PCR) assays for mutant genes that affect coagulation.14-21

### RESULTS

Retinal vein occlusion was bilateral in 3 patients. There were 9 women and 8 men, 1 of whom was African American, and the rest were white. The age of patients was 52 ± 10 years (median, 51 years; range, 33-69 years) and of controls was 37 ± 7 years (median, 37 years; range, 24-54 years). Of the 40 adult controls, 23 were female; 36 were white, 2 African American, and 2 “other.” Six patients (35%) had hypertension that was well controlled with antihypertensive agents, 3 (18%) ingested 7 or more alcoholic beverages per week, 2 (12%) had type 2 (mature-onset) diabetes mellitus, and 1 (6%) smoked 1 pack or more of cigarettes per day.

Three (18%) of the patients had a history of osteonecrosis, 4 patients (24%) had other thrombotic events, primarily deep venous thrombosis not including osteonecrosis. No patients had been given high-dose corticosteroid therapy (prednisone, >20 mg/d, for ≥ week) before the development of retinal vein thrombosis. One patient received high-dose corticosteroid treatment of his retinal vein thrombosis for 2 months; this antedated and contributed to his subsequent development of osteonecrosis of the hips, requiring bilateral hip replacement.

Exclusion of the 3 patients who had both osteonecrosis and retinal vein thrombosis from the patient-control comparisons did not alter any of the statistical comparisons. One of these 3 patients was heterozygous...
for the mutant factor V Leiden gene, and 1 was homozygous for the 4G/4G polymorphism of the PAI1 gene.

THROMBOPHILIC AND HYPOFIBRINOLYTIC GENE MUTATIONS (PATIENTS VS CONTROLS)

Of the 17 patients with vein occlusion, 3 (18%) were heterozygous for the thrombophilic factor V Leiden G1691A mutation compared with 7 (3%) of 233 controls ($P = .02$, Fisher exact test) (Figure 2). Of the 3 patients heterozygous for the factor V Leiden G1691A mutation, 2 were also homozygous for the 4G/4G polymorphism of the PAI1 gene, and 1 of these 2 patients also had high Lp(a) levels (70 mg/dL). One patient was heterozygous for the mutant PAI1 gene and also had high Lp(a) levels (40 mg/dL).

The distribution of the MTHFR C677T mutation did not differ between patients and controls ($P = .41$) (Figure 2).

All 17 patients with measurement of the 20210*A allele of the prothrombin gene had the normal "wild-type" gene, which was not different ($P > .41$) from 9 (3.8%) of 234 controls (Figure 2).

The distribution of polymorphism in the promoter region of the PAI1 gene was skewed toward the 4G/4G and 4G/5G genotypes in patients with vein occlusion ($\chi^2 = 4.8; P = .09$) (Figure 2). Controls were 3 times more likely than patients (64% vs 88%) to be heterozygous or homozygous for the 4G mutation ($P = .03$, Fisher exact test) (Figure 3). The frequency of the 4G allele was 21 (0.62) of 34 in patients vs 196 (0.42) of 468 in controls ($\chi^2 = 5.1; P = .02$).

DIFFERENCES BETWEEN PATIENTS AND CONTROLS IN MAJOR COAGULATION MEASURES

Thrombophilic Factors

Measurements of levels of antigenic proteins C and S, anticardiolipin antibodies IgG and IgM, and homocysteine did not differ between patients and controls ($P > .10$; data not shown). The major component of the lupus anticoagulant, however, dilute Russell viper venom clotting time, differed between patients and controls. Of 14 patients with dilute Russell viper venom clotting times, 6 (43%) had prolonged (abnormal) times (>38.8 seconds) compared with 1 (3%) of 30 controls ($P = .002$, Fisher exact test) (Figure 4).

Hypofibrinolytic Factors

Patients with vein occlusion were more likely to have high levels of hypofibrinolytic Lp(a) (>35 mg/dL) ($\chi^2 = 8/16 [50%]$) than controls (5/40 [13%]) ($\chi^2 = 9; P = .003$) (Figure 4). Patients were more likely to have high levels of the major determinant of fibrinolysis, PAI-Fx. Levels of PAI-Fx were high (>22 U/L) in 6 (38%) of 16 patients.
compared with 1 (2%) of 40 controls ($P = .001$, Fisher exact test) (Figure 4).

Hypofibrinolytic disorders often occurred in clusters that were present with and without inclusion of the 3 patients who also had osteonecrosis. Of the 6 patients with high PAI-Fx levels, 4 (67%) also had high Lp(a) levels, and 2 (33%) also had prolonged dilute Russell viper venom clotting times. Of the 8 patients with high Lp(a) levels, 4 (50%) had high PAI-Fx values.

Of the 6 patients homozygous for the 4G/4G polymorphism in the promoter sequence of the PAI1 gene, 5 had measures of Lp(a). Of these 5 patients, 4 (80%) also had high Lp(a) levels; of 9 patients heterozygous for the 4G allele, 4 (44%) also had high Lp(a) levels, whereas the 2 patients with the 5G/5G genotype had normal Lp(a) levels.

When classifying the patients and controls by the PAI1 gene polymorphism, patients homozygous for the 4G allele had much higher median Lp(a) levels than controls (62 vs 5.3 mg/dL; $P = .05$) (Figure 5). Patients heterozygous for the 4G/5G trait also had higher median Lp(a) levels than controls heterozygous for the 4G/5G allele (16 vs 3.3 mg/dL; $P = .06$) (Figure 5). Patients with the 5G/5G genotype had much higher median Lp(a) levels than controls with the same genotype (17.5 vs 6.7 mg/dL), but this difference was not significant ($P = .62$) (Figure 5).

When the relationships between the PAI1 gene and Lp(a) levels were compared in patients vs controls, the slopes of the 2 regression lines differed ($P = .04$) (Figure 6). In patients, but not in controls, increased 4G alleles for the PAI1 gene were associated with increased Lp(a) levels (Figure 6).

**Lipids and Lipoprotein-Cholesterol Levels**

Of 16 patients with vein occlusion having lipid profiles in the fasting state, 10 (63%) had high total cholesterol levels (>5.17 mmol/L [>200 mg/dL]), 2 (13%) had high triglyceride levels (>2.82 mmol/L [>250 mg/dL]), 1 (6%) had low-high-density-lipoprotein-cholesterol levels (<0.90 mmol/L [<35 mg/dL]), and 8 (50%) had high low-density-lipoprotein–cholesterol levels (>3.36 mmol/L [>130 mg/dL]). Of the patients with high low-density-lipoprotein–cholesterol levels, 5 (63%) also had high Lp(a) levels (>35 mg/dL). Low-density-lipoprotein–cholesterol levels correlated with Lp(a) levels ($r = 0.48$; $P = .06$).

In our study, patients with retinal vein occlusion were more likely than healthy normal controls to have the heritable thrombophilic factor V Leiden G1691A mutation and the thrombophilic antiphospholipid lupus anticoagulant. They were also more likely to have homozgyosity or heterozygosity for the heritable 4G PAI1 allele and, accompanying this, were much more likely to have high levels of the hypofibrinolytic PAI-Fx. Patients with retinal vein occlusion who were homozygous for the 4G PAI1 allele were more likely than controls to have high Lp(a) levels, a double dose of hypofibrinolytic factors that, we postulate, contribute to retinal vein occlusion. Patients were also more likely than controls to have high levels of the heritable hypofibrinolytic Lp(a). The lipoprotein Lp(a) is postulated to be hypofibrinolytic by virtue of its sequence homology with plasminogen, thus competing with plasminogen for fibrin binding and hindering fibrin digestion. It can also interact with cellular plasminogen receptors with a plasminogen-like affinity. Most relevant to the present study, where 4G/4G homozygosity of the PAI1 gene was associated with high Lp(a) levels, Lp(a) enhanced PAI1 transcription and expression on cultured endothelial cells. As noted by Rosendaal, venous thrombosis is a multicausal disease; usually more than 1 coagulation abnormality needs to be present before thrombosis occurs. “The younger an individual, the more risk factors are required to precipitate thrombosis.” This is relevant to retinal vein occlusion because systemic workup for the cause of the disease is usually limited to younger
patients. The common concurrence of other venous thrombosis and central retinal vein occlusion should promote studies of the pathogenesis of this type of conjoint thrombosis.

Whether synergism between the patients’ high low-density-lipoprotein–cholesterol and high Lp(a) levels, known to cause arterial occlusive disease,26 could have played any role in their ophthalmic disease is unknown. However, 2 earlier studies1,2 implicated hyperlipidemia as a risk factor for retinal vein occlusion, consistent with the findings of the present study.

In agreement with previous studies of retinal vein occlusion, we found a high prevalence of the hypofibrinolytic Lp(a),4 the thrombophilic lupus antibody,5 and thrombophilia had been identified as causes of retinal vein occlusion.1,7,13 Other retrospective studies14-19 of the causes of central retinal vein occlusion in young patients were inconclusive. The newly discovered, most common heritable thrombophilia, however, resistance to activated protein C, appears to be 4 to 5 times more common than other known inherited thrombophilias or hypofibrinolyses in young patients with central retinal vein occlusion.8-10 The PCR assay for polymorphism of the PAI1 gene has only recently become available17,25 and, to our knowledge, has not been used previously in the evaluation of patients with retinal vein occlusion. Major recent advances have been made in coagulation measurements that allow the diagnosis of the most common coagulation disorders, with the recognition of resistance to activated protein C10,12 and with cDNA PCR measurements of the mutant factor V Leiden gene12,16,19 and the 4G/4G polymorphism of the PAI1 gene.17,25 By enabling the diagnosis of common, heritable coagulation disorders, these cDNA PCR methodological advances8,10,16,19,25 provide a higher level of certainty, unaffected by age,19,16,25 that thrombophilia and hypofibrinolysis are causes of retinal vein occlusion. Furthermore, the diagnosis of heritable thrombophilia, hypofibrinolysis, or both, as putative causes of retinal vein occlusion provides important prospective diagnostic insight into the high likelihood of other thrombotic disorders in patients with central retinal vein occlusion. Thus, 18% of our patients had a history of osteonecrosis,19,10,22 and 24% had other thrombotic events exclusive of osteonecrosis, primarily deep venous thrombosis. The recognition of resistance to activated protein C in patients with retinal vein occlusion also has important ramifications for their first-degree relatives.39 In 177 subjects from 34 kindreds in which 1 family member had resistance to activated protein C, Svensson and Dahlback39 reported that 27% had a history of thrombosis. At age 45 years, the likelihood that a subject with resistance to activated protein C would be free of thrombosis was only 59% compared with 97% in subjects without resistance to activated protein C.

The recognition of frequent hypofibrinolysis and thrombophilia as possible causes of retinal vein occlusion1-13 calls for further prospective analyses and study. Patients with retinal vein occlusion, particularly those with a family history of thrombosis,8 should have tests designed to diagnose heritable thrombophilias and hypofibrinolysis, including assessment of the factor V Leiden G1691A mutation. These analyses should include studies of the 4G/5G polymorphism of the PAI1 gene promoter, the prothrombin gene, the MTHR C677T mutation, PAI-Fx, Lp(a), the lupus anticoagulant antibody, and anticardiolipin antibodies. Patients with these predominantly heritable abnormalities are at increased risk of both venous and arterial thrombi.

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Corresponding author: Charles J. Glueck, MD, Cholesterol Center, Jewish Hospital, 3200 Burnet Ave, Cincinnati, OH 45229 (e-mail: glueckch@healthall.com).