Neurology and the blood: haematological abnormalities in ischaemic stroke

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Haematological disorders account for up to 8% of all ischaemic strokes in different series. Table 1 shows the haematological disorders associated with ischaemic stroke. Most studies report them as being more common in younger stroke patients, particularly those who have undetermined stroke aetiology after extensive tests including full cardiac evaluation. Many primary haematological disorders have been associated with ischaemic stroke but in many patients with stroke other aetiological factors are also present making a cause and effect relation difficult to prove. Furthermore, some of these haematological factors, particularly deficiencies of natural anticoagulants, are more potent causes of venous thrombosis. Therefore in such cases paradoxical embolism from the venous system should be considered, and excluded, before arterial thrombosis is implicated.

Normal haemostasis

The haemostatic system is a major defence system of the body. It is the result of interaction of three components: (1) the vessel wall, particularly endothelial cells; (2) platelets; and (3) the coagulation system including the fibrinolytic system.

Keywords: haematological abnormalities; ischaemic stroke

Table 1  Haematological disorders associated with ischaemic stroke

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The aims are to maintain fluidity of the blood and, when there is a break in the integrity of the vessel wall, to rapidly initiate blood coagulation which is maintained locally at the site of vascular damage. The process involves several different proteins. Defects of these, which may be congenital or acquired, will result in disorders of haemostasis which may manifest in clinical syndromes of easy bleeding or bruising (“haemophilias”) or inappropriate thrombosis (“thrombophilia”). A more general breakdown in the initiation and control of haemostasis results in the syndrome of “disseminated intravascular coagulation”, in which there is the apparent paradox of concurrent bleeding and widespread thrombosis which is responsible for much of the organ damage.

The vascular endothelium plays a critical part in maintaining blood fluidity and vascular smooth muscle tone through (a) prostacyclin, a potent vasodilator and platelet antiaggregator synthesised from arachidonic acid through a series of steps, including involvement of cyclooxygenase, which is inhibited by aspirin;

(b) nitric oxide (endothelium derived relaxing factor, EDFR), which is a potent vasodilator and inhibits platelet aggregation. It is not inhibited by aspirin but is inhibited by free haemoglobin.

In addition, thrombomodulin is expressed on the surface of endothelial cells and plays a critical part in the inhibition of fibrin formation through its interaction with thrombin and protein C. Thrombomodulin is reduced on endothelial surfaces in response to hypoxia. Mild hyperhomocysteinaemia may result in inhibition of thrombomodulin and thus explain the increase in thrombosis seen in this condition.

Platelets are anucleate cells derived from megakaryocytes in the bone marrow. Their production is regulated in part by thrombopoietin (TPO), also known as megakaryocyte growth and development factor (MGDF). Activation of platelets brought about by exposure to subendothelial collagen results in changes including shape change, aggregation, and release of intracytoplasmic granule contents.

Functionally these changes result in:
(1) Formation of a primary platelet plug at the site of vascular injury.
Polycythaemia rubra vera is a myeloproliferative disorder resulting from clonal expansion of a transformed haematopoetic stem cell associated with pronounced overproduction of red blood cells and, to a lesser extent, expansion of granulocytic and megakaryocytic elements. It usually begins in late middle age. The increased packed cell volume results in hyperviscosity and reduced cerebral blood flow. This may result in cerebral infarction: transient ischaemic attacks or intracranial venous thrombosis. Stroke rates of about 5% a year have been reported. Other neurological symptoms include headache, dizziness, visual blurring, and confusion resulting from a reduction in cerebral blood flow secondary to hyperviscosity. Treatment may involve phlebotomy, hydroxyurea, and other cytotoxic drugs.

Secondary polycythaemia may be caused by chronic hypoxia, often occurring, for example, in patients with congenital cyanotic heart disease, smoking, cerebellar hemangioblastoma, renal tumours, and patients who smoke. It has been suggested that this is a risk factor for stroke but if this is the case it seems to be only weakly associated. Furthermore the association is confounded by cigarette smoking and blood pressure, both being linked to packed cell volume. Studies have shown no increased risk of stroke in young adults with cyanotic congenital heart disease and secondary polycythaemia or in perioperative thrombotic risk in 100 patients with secondary polycythaemia.

Essential thrombocythaemia

Essential thrombocythaemia is a myeloproliferative disorder in which blood platelet counts above 600 000 cells/ml occur. In addition the platelets are often large and have functional abnormalities. Occasionally such abnormalities of platelet function result in a bleeding tendency but thrombosis is more common. Stroke is a well recognised complication but headache and transient focal and non-focal neurological disturbances are also frequent. Essential thrombocythaemia must be distinguished from secondary thrombocythaemia, which can occur in response to conditions including inflammation, acute bleeding, iron deficiency, splenectomy, and infection. A highly increased platelet count in the absence of an identifiable cause of secondary thrombocythaemia is usually sufficient for a diagnosis of essential thrombocythaemia. Support for the diagnosis may be obtained by in vitro platelet aggregation studies and the documentation of splenomegaly. In essential thrombocythaemia the megakaryocytes are large and hyperploid by contrast with secondary thrombocythaemia when they are usually increased in number and of small diameter and low ploidy.

The management of essential thrombocythaemia requires specialist care and hydroxyurea is often used as an initial treatment. The use of antiplatelet agents such as aspirin is controversial. These may protect against thrombosis but can also increase the risk of haemorrhage. Control of the platelet count is of primary importance but in the occasional patients in whom good control cannot be achieved, or who develop thrombotic complications despite adequate lowering of the platelet...
let count, the addition of low dose aspirin may be warranted.

SICKLE CELL DISEASE

Stroke is a frequent complication of homozygous sickle cell disease, particularly in children.\textsuperscript{11} One study suggested that 75\% of cerebrovascular complications in sickle cell patients occurred in those under 15 years of age\textsuperscript{11} whereas another study found cerebral ischaemia in 15\% of homozygotic Hb6SS patients with a mean age of onset of 15 years.\textsuperscript{3} However, the prevalence of silent infarction on brain imaging is higher.\textsuperscript{13} The mechanism of stroke is often unclear although changes during sickle cell crisis, such as raised whole blood viscosity and red blood cell abnormalities may result in small and large arterial occlusions. In addition, stenosis of large extracranial or intracranial vessels may occur secondary to fibrous proliferation of the intima. Stenoses in the middle cerebral artery can be detected by transcranial Doppler and their presence predicts risk of stroke.\textsuperscript{14} This technique may allow identification of at risk people in whom a programme of exchange transfusion can prevent

Figure 1 The coagulation cascade. Vascular damage initiates coagulation cascade resulting in the explosive generation of thrombin at the site of injury. Thrombin catalyses the conversion of fibrinogen to an insoluble fibrin (clot) matrix, in the presence of factor XIIIa and calcium ions. Critical reactions are closely checked and localised by circulating anticoagulants, such as activated protein C, TFPI, and antithrombin. Fibrinolysis is initiated when fibrin is formed and eventually dissolves the clot. Inappropriate activation of blood coagulation, or depressed fibrinolytic activity, or both may lead to the formation of a thrombus. By contrast, a defect or deficiency in the coagulation process and/or accelerated fibrinolysis is associated with a bleeding tendency. The cascade scheme is organised into the intrinsic (factors XII, XI, IX, VIII, prekallikrein, HMW kininogen), extrinsic (tissue factor, factor VII), and common pathways (factors V, X, XIII, prothrombin, fibrinogen). The extrinsic pathway is initiated when blood is exposed to tissue factor released from damaged endothelium. The intrinsic pathway is initiated when factor XII involving “contact factors” on negatively charged surfaces, such as glass or kaolin in vitro. Feedback activations of factors V, VII, and VIII by factor Xa and the activation of factor XI by thrombin are not shown. (From APC Resistance, version 2.0 1997 Chromogenix AB, Taljegårdsgatan 3, S-431 53 Mölndal, Sweden, with permission.)
stroke. Occlusion of large intracerebral arteries in sickle cell disease may result in a Moya-Moya-like syndrome which can present often in young adults with subarachnoid haemorrhage. The mainstay of treatment of cerebrovascular complications in sickle cell disease is exchange transfusion.

Stroke may also complicate haemoglobin sickle cell disease. 15

PAROXYSMAL NOCTURNAL HAEMOGLOBINURIA
Paroxysmal nocturnal haemoglobinuria is a rare disorder which is an acquired clonal disease in which red cells show increased sensitivity to lysis by complement. 16 The complement activation indirectly stimulates platelet aggregation and hypercoagulability, which is probably responsible for the tendency to thrombosis. Patients present with a haemolytic anaemia, and often mild lymphopenia and thrombocytopenia. Haemoglobinuria may occur. The diagnosis can be made by the Ham test in which sensitivity of the patient’s cells to lysis by complement can be shown or with CD59 assessment by flow cytometry. Cerebral venous thrombosis may occur; stroke is occasionally part of the syndrome. 17

THROMBOCYTOPENIA
Thrombotic thrombocytopenic purpura is a rare but often fatal disorder characterised by thrombocytopenia, a microangiopathic haemolytic anaemia, renal failure, fever, and neurological symptoms. It may be initiated by endothelial injury and subsequent release of Von Willebrand factor and other procoagulant materials from the endothelial cells. In addition in some patients circulating protein may induce platelet aggregation. 16 Many of the symptoms are due to widespread small platelet microthrombi which cause infarction in many organs including the brain. Neurological symptoms include a fluctuating encephalopathic picture with confusion and seizures and this can be accompanied by focal symptoms and signs. 16-18 However, it can occasionally present with isolated stroke or transient ischaemic attack. The thrombocytopenia on full blood count will point to the diagnosis. Brain CT may be normal or show infarction and occasionally intracerebral haemorrhage. 19

Thrombotic thrombocytopenic purpura is similar to haemolytic-uraemic syndrome, which usually occurs in children less than 5 years old. This multisystem disorder presents with fever, thrombocytopenia, a microangiopathic haemolytic anaemia, hypertension, and varying degrees of renal failure. Haemolytic thrombi are particularly seen in the afferent arterioles and glomerular capillaries of the kidneys and neurological symptoms, other than those associated with uraemia, are less common but can still occur. 20

Treatment for thrombotic thrombocytopenic purpura involves exchange transfusion or extensive plasmapheresis coupled with infusion of fresh frozen plasma. This therapeutic approach has led to a considerable reduction in the overall mortality with over half of the patients with thrombotic thrombocytopenic purpura recovering.

Heparin induced thrombocytopenia
It has been estimated that as many as 10%-15% of patients receiving therapeutic doses of heparin may develop a degree of
thrombocytopoenia. This may occur due to drug-antibody binding to platelets or occasionally secondary to direct platelet agglutination by heparin. It may lead to severe bleeding or intravascular platelet aggregation and paradoxical thrombosis. Stroke may occur.

LEUKAEMIA
Leukaemia more often causes intracerebral haemorrhage due to thrombocytopoenia or direct leukaemic infiltration of the CNS, than arterial occlusion. When stroke does occur it is thought to be due to increased blood viscosity.

INTRAVASCULAR LYMPHOMA
Intravascular lymphoma is an uncommon malignancy, defined pathologically by neoplastic proliferation of lymphoid cells within the lumens of capillaries, small veins, and arteries with little or no adjacent parenchymal involvement. It used to be called malignant angioendotheliosis but recent immunohistochemical studies have demonstrated that the tumours are neoplastic lymphoid cells more commonly of B-cell origin and therefore it is now referred to as intravascular lymphoma or angiotrophic large cell lymphoma. Most commonly symptoms are confined to the skin or peripheral nervous system findings may mimic neurological manifestations without abnormalities on bone marrow biopsy, chest and abdominal CT, or CSF examination. One neurological presentation is with recurrent stroke-like episodes. It may also present with a dementia with or without focal neurological signs, a spinal cord syndrome, and peripheral or cranial neuropathies. It may produce an identical clinical picture to primary angitis of the CNS, including similar angiographic appearances, and distinction may only be possible on brain biopsy or postmortem. Similarly the peripheral nervous system findings may mimic systemic vasculitis and again only be differentiated on biopsy. Autoantibodies may occur which can make distinction from vasculitis even more difficult. Most of the cases of CNS involvement have been diagnosed at postmortem. In some cases an improvement has been made after corticosteroid therapy although this may only be partial or transient. Chemotherapy has resulted in remission in a few case reports.

Disorders of coagulation/fibrin
CONGENITAL
Natural anticoagulation disorders
The natural anticoagulants (heparin cofactor 2, antithrombin III, protein C, and protein S) inhibit thrombosis in the normal subject. Deficiencies of these anticoagulants may be hereditary or acquired. Such deficiencies may be responsible for as many as 20% of nontraumatic venous embolisms but their role in arterial thrombosis remains unclear. Many case reports have suggested an association but more controlled studies have often failed to prove this. Heparin cofactor 2 has probably only a minor role in venous thrombosis, and there are no convincing data linking it with arterial thrombosis.

Protein C
Protein C is a vitamin K dependent protein which binds to the endothelial cell surface protein thrombomodulin and is converted to an active protease by thrombin. Activated protein C, in conjunction with protein S proteolyses factor Va and factor VIIIa, which reduces thrombin formation. Activated protein C may also promote fibrinolysis and accelerate clot lysis. Protein C is synthesised in hepatocytes and its synthesis is encoded by a single gene located on chromosome 2. Hereditary protein C deficiency is usually an autosomal dominant disorder although dysfunctional molecules have also been identified in some patients with thrombosis. Heterozygotes with 25%-50% of protein C concentrations occur in about 1 in 300 to 1 in 3000 unselected blood donors, but many of these people are asymptomatic. Symptomatic deficiency is much less common, occurring in perhaps as few as 1 in 36 000 people.

There are many reports of protein C deficiency associated with ischaemic stroke. In young patients with stroke and protein C deficiency, who also have a strong family history of premature thrombosis, it is likely that the relation between protein C deficiency and the stroke is causal. However, often in older patients with moderate degrees of protein C deficiency, determining whether the association is causal can be difficult. The correlation between protein C and protein S concentrations and the risk of thrombosis is not as precise as for antithrombin III deficiency, and protein C concentrations in asymptomatic deficient people overlap with those seen in patients with recurrent thromboembolism. Furthermore, in the acute stage of stroke low concentrations of protein C are fairly common and may reflect consumption. The degree of reduction in protein C concentration has been associated with the severity of stroke. Serial sampling has shown that protein C concentrations may return to normal some months after stroke. Therefore, if low protein C concentrations are found in the acute phase of stroke measurement should be repeated after three months and family members should be tested. Apart from patients with extensive thrombosis or stroke, secondarily low protein C concentrations may occur in severe liver disease, the nephrotic syndrome, disseminated intravascular coagulation, in the postoperative period, and in patients receiving warfarin. It is generally recommended that patients with primary protein C deficiency and stroke should be heparinised and treated with oral anticoagulants. Whether these should be continued lifelong remains uncertain. Warfarin therapy itself will reduce protein C concentrations making determinations of concentrations while on therapy difficult.

An association has been reported between warfarin induced skin necrosis and protein C deficiency. This is a rare but serious complica-
tion of warfarin. It usually presents with localized pain followed by a petechial rash and ecchymoses. This can progress to widespread full thickness skin necrosis. If these symptoms appear during warfarin therapy the drug should be discontinued and vitamin K injected parenterally. Heparin can be given safely.

**Protein S deficiency**

Protein S is a vitamin K dependent plasma glycoprotein which serves as a cofactor for activated protein C. It is synthesised primarily in the liver and encoded on chromosome 6. In plasma about 60% is bound to a C4b-binding protein and 40% is in an active free form. Therefore for full assessment both total and active or “free” protein S need to be measured. It has been estimated that protein S deficiency occurs in 1 in 3000 to 1 in 15 000 people. Hereditary protein S deficiency is an important cause of idiopathic venous thrombosis and may account for 5% or more cases. Some case reports and small series have reported an association between protein S deficiency and ischaemic stroke. However, the same difficulties occur as in interpreting the association between protein C and stroke. Protein S concentrations may fall in ill patients both with stroke and admitted to hospital for other reasons. Therefore as for protein C, follow up concentrations after three months and screening of family members is necessary. In the presence of persistent protein S deficiency in stroke the recommended treatment is anticoagulation. Low protein S concentrations may also occur with pregnancy, warfarin therapy, and acute illness, whereas women have lower concentrations of protein S than men.

**Activated protein C resistance**

Functional resistance to the anticoagulant effects of activated protein C seems to be the most common inherited prothrombotic state. The genetic basis for this functional abnormality has recently been defined as a point mutation in factor V at the exact site (Arg 506) where activated protein C normally cleaves and inactivates the Va procoagulant; this is referred to as the Leiden factor V mutation. Some small studies have suggested an association between activated protein C resistance or the Leiden factor V mutation and stroke but larger case-control studies have failed to confirm the association. The situation is complicated by a recent report associating activated protein C resistance with cerebrovascular disease independent of the factor V mutation. In view of the frequency of asymptomatic heterozygotes for the factor V Leiden mutation in the normal population the optimal treatment of patients with the mutation and stroke is uncertain. However, in young patients with no obvious risk of stroke anticoagulation with warfarin seems a reasonable approach. Paradoxical venous embolism should be sought in view of the strong association with venous thrombosis.

**Antithrombin III**

Antithrombin III is a plasma glycoprotein synthesised in hepatocytes and endothelial cells, and encoded for by a gene located on chromosome 1. It inhibits thrombin and other activated serine proteases including factors Ixa, Xa, XIa, XIIa, and calicrin. Congenital antithrombin III deficiency is inherited as an autosomal dominant trait. The most common defect is mild (heterozygous) antithrombin deficiency which occurs in 1 in 2000 people. In addition, dysfunctional antithrombin molecules with mutations affecting either the serine protease binding site or the heparin binding site have been described. It has been estimated that between 1 in 2000 and 1 in 5000 of the general population have antithrombin III deficiency. Antithrombin III deficiency has been associated with a high risk of developing venous thrombosis with as many as 85% of patients developing thrombosis by the age of 50. There are case reports linking hereditary antithrombin III deficiency and ischaemic stroke and in some of these stroke occurred in family members, suggesting a causative association. An acquired antithrombin III state can be caused by severe hepatic failure leading to reduced synthesis of antithrombin III, nephrotic syndrome, oral contraceptive use, heparin therapy, disseminated intravascular coagulation, leukaemia, malnutrition, and diabetes. Heparin therapy increases antithrombin III activity and rapidly decreases antithrombin III plasma concentrations. The plasma concentrations normalise two to three days after stopping therapy. Because heparin requires antithrombin to exert its anticoagulant action, treatment of antithrombin III deficiency with heparin alone is inadequate. However, patients with antithrombin III who develop acute thrombosis or embolism can be treated with intravenous heparin initially as there is usually sufficient normal antithrombin to act as a heparin cofactor. However, after this they should be placed on long term warfarin therapy. It is not certain whether lifelong warfarin treatment is required and currently decisions should be made according to the severity of the condition, the family history, and the number of recurrences. Family studies should be conducted when an antithrombin deficient person is discovered as up to half the members of a kindred may be affected. Asymptomatic subjects with antithrombin III deficiency should receive prophylactic anticoagulation with heparin or antithrombin III concentrate infusions to raise their antithrombin III concentration before medical or surgical procedures which may increase their risk of thrombosis. Chronic oral anticoagulation is not recommended until those at risk have a clinical thrombotic episode. A recent alternative treatment is antithrombin III replacement with antithrombin III concentrates.
Fibrinolytic system disorders

CONGENITAL

Plasminogen deficiency
Families have been described with recurrent venous thrombosis and embolism due to defects in fibrinogen or plasminogen, or with decreased synthesis or release of tissue plasminogen activator. There are a few case reports associating low functional levels of plasminogen activity in young people with stroke.70 71

Hereditary dysfibrinogenaemia is characterised by abnormal fibrinogen molecules that are resistant to cleavage by plasmin. This is a rare disorder which has been linked to thrombosis, including strokes.72

ACQUIRED

Disseminated intravascular coagulation
Disseminated intravascular coagulation is a rare disease characterised by fibrin thrombi in small vessels and haemorrhagic lesions. It more commonly causes an encephalopathic picture rather than stroke-like episodes although stroke-like episodes may occur. Pathology shows widespread haemorrhagic cerebral infarcts and intracranial haemorrhages. The diagnosis is confirmed by a low platelet count accompanied by low fibrinogen and raised fibrin degradation products.

Lupus anticoagulant/anticardiolipin syndrome
The lupus anticoagulant and anticycardolipin antibodies are closely related autoantibodies belonging to a group of antibodies which react with proteins associated with phospholipid. Anticardiolipin antibodies seem to be directed against the plasma protein β2-glycoprotein whereas thrombin is probably the target protein for the lupus anticoagulant. They are most commonly found in patients with systemic lupus erythematosus but may also occur in patients without the disease and are associated with arterial and venous thrombosis.

In the absence of systemic lupus erythematosus they may form one part of the antiphospholipid antibody syndrome which can present with recurrent miscarriages, arterial and venous thrombosis in any size vessel, livedo reticularis, cardiac valve vegetations, and thrombocytopenia.73 However, anticardiolipin antibodies are not specific to the antiphospholipid syndrome and may occur in normal subjects, patients with other autoimmune disorders, malignancy, and HIV infection, and after the use of various drugs including phenytoin, sodium valproate, procainamide, hydralazine, and quinidine.

Studies have found widely varying frequencies of antiphospholipid antibodies in patients with stroke with a prevalence from 1 to about 50%.74 75 A recent study in an unselected stroke population found no evidence to support the hypothesis that anticardiolipin antibodies are an independent risk factor for stroke in young people.76 There was an increase in IgG titre with age and number of vascular risk factors in patients with stroke, but the authors interpreted this as suggesting that it may be a non-specific accompaniment of vascular disease. By contrast, a recent study by the Antiphospholipid Antibodies and Stroke Study Group found that a single anticardiolipin antibody value > 10 GPL units at the time of an initial ischaemic stroke was a significant independent risk factor for stroke but when patients were followed up, anticardiolipin antibody positivity did not confer a significantly increased risk for subsequent thrombo-occlusive events or death, including that secondary to stroke.77 Therefore the overall contribution of antiphospholipid antibodies to risk of stroke remains uncertain. Such cases are relatively rare and therefore no association may be detected in small studies of unselected stroke patients. The symptoms of cerebral ischaemia may be atypical, sometimes with atypical amaurosis fugax in the absence of carotid artery disease. Nevertheless, in those with persistently high titres of anticardiolipin antibody, or a persistently positive lupus anticoagulant and some features of the antiphospholipid syndrome there does seem to be an association between the antiphospholipid antibodies and stroke. In such patients, anticoagulation with an international normalised ratio (INR) ≥ 3 seems to be more effective than low intensity warfarin or aspirin in preventing recurrent thrombosis.78 However, in patients with raised anticardiolipin antibodies and stroke, in the absence of other features of the antiphospholipid syndrome, the association is more tenuous. In such patients it is sensible to repeat the titre and look for other causes of stroke before treating with anticoagulation.

Pregnancy and the puerperium

In developed countries stroke complicating pregnancy or the puerperium is rare with a frequency of perhaps only 1 or 2 per 10 000 deliveries.79 It is more common in India.80 A proportion of these cases is caused by a prothrombotic state that may result in acute middle cerebral artery or other large cerebral artery occlusion, perhaps due to paradoxical embolism from the pelvic or leg veins, or cerebral venous thrombosis. This most often occurs in the puerperium.81

Oral contraceptives

Studies have shown an increased stroke risk in women taking the oral contraceptive pill.82 83 Studies assessing the risk of stroke in women on the second and third generation combined oestrogen/progesterone oral contraceptives are only now being published. A recent hospital based case-control study assessed the risk of ischaemic and haemorrhagic stroke in 20–44 year olds in 21 centres around the world.84 85 Six hundred and ninety seven cases of cerebral infarction confirmed by CT were compared with 1962 age matched hospital controls and the overall odds of ischaemic stroke were 2.99 (95% confidence interval (95% CI) 1.65–5.4) in Europe, and 2.93 (95% CI 2.15–4.00) in developing countries. Odds ratios were lower in non-smokers, younger women, and those without hypertension. In Europe the odds ratio associated with low dose oral contraceptives was 1.53 (95% CI 0.71–3.31) compared with 5.30 (95% CI 2.56–11.0) for higher dose oral contraceptives but no such difference was
found in developing countries. For haemorrhagic stroke 1068 cases and 2910 controls were studied and overall the use of combined oral contraceptives was associated with a slightly increased risk, but only in developing countries (odds ratio 1.76, 95% CI 1.35–2.30). Current oral contraceptive users and hypertensive women had substantially increased risk. Overall in this study about 13% of all strokes in women aged 20–44 in Europe are attributed to the use of oral contraceptives and 8% of strokes in a similar age group in women in developing countries. Significantly increased risks are seen in older women, those who smoke, and those with a history of hypertension. Recently it has been suggested that persons heterozygous for the Leiden factor V mutation are at increased risk of having ischaemic stroke while on the contraceptive pill.

Paraproteinaemias
Paraproteinaemias such as Waldenstrom’s macroglobulinaemia and multiple myeloma can result in a hyperviscosity syndrome which usually presents with an encephalopathic picture with symptoms such as headache, ataxia, lethargy, poor concentration, visual blurring, and drowsiness and coma. Occasionally focal stroke-like episodes may occur and these may also be secondary to occlusion of vessels with acidophilic material thought to result from the abnormal plasma proteins.65

General management considerations
It is difficult to determine the importance of the various deficiencies of natural anticoagulants in the pathogenesis of stroke in view of the conflicting data that have been published. However, most well controlled studies suggest that their contribution to overall stroke risk is low in the general population of patients with stroke. They are likely to be a more important cause in young patients without other obvious causes for stroke. On current evidence it would seem reasonable to screen for protein C, protein S, activated protein C resistance, and possibly antithrombin III deficiency in such patients aged 55 or under. These tests can be screened for by assays of protein concentrations or functional testing. Interpretation of the results is complicated both by the acute phase changes seen with protein C and S and the roughly 5% prevalence of heterozygotes with the factor V Leiden mutation in the normal population. These factors have to be carefully considered before embarking on long term anticoagulant treatment in such patients. In general this should only be instituted if there are no other obvious causes for stroke. However, in young patients including children with stroke associated with hereditary deficiencies and a strong family history of stroke in the presence of anticoagulation abnormalities, anticoagulation is usually appropriate. The situation is further complicated by the coexistence of more than one abnormality in some patients which may possibly confer increased stroke risk.65

| Table 2 Screening for haematological disorders in patients with stroke |
|----------------------------------|-------------------------------------------------------------|
| In all cases | Full blood count, erythrocyte sedimentation rate, plasma viscosity |
| In young stroke (age ≤55) | Protein C, Protein S, Antithrombin III, Lupus anticoagulant, Anticardiolipin antibodies, APC resistance/Leiden factor V mutation, Haemoglobin electrophoresis |

*In subjects of African or Mediterranean origin.

Haematological evaluation in patients with ischaemic stroke
From the above discussion it is clear that haematological causes for stroke are uncommon and that the association between some of these and stroke remains uncertain. All patients with stroke should have a full blood count. A detailed family history should be taken. More detailed haematological assessment should be confined to patients in whom the likelihood of detecting a significant abnormality is greater. This includes younger patients, patients without any other obvious cause for stroke, and those with a strong family history of nonatheromatous stroke. In such patients it is reasonable to screen for protein C, protein S (and total protein S if abnormal), and antithrombin III deficiencies, activated protein C resistance, anticardiolipin antibodies, and the lupus anticoagulant. If activated protein C resistance is abnormal factor V Leiden polymorphism should be determined. When such tests are performed in the acute phase of stroke, particularly in patients with larger strokes, if abnormalities are found the tests should be repeated some weeks later before any long term decisions on management are made. It should be remembered that concurrent anticoagulation therapy (for example, heparin or oral coumarin) will reduce the concentration of the anticoagulant proteins such as protein C and S. All young black and eastern Mediterranean people with stroke should have haemoglobin electrophoresis for sickle cell disease. Table 2 summarises this screening procedure.

Summary
Haematological disorders account for 0%-8% of ischaemic strokes in different series. These include cellular disorders such as polycythaemia rubra vera, essential thrombocythaemia, sickle cell disease, thrombocytopenia, and other disorders. They also include disorders of coagulation, and recently particular interest has centred on protein C and S deficiency and activated protein C resistance. Antiphospholipid antibodies represent an acquired disorder of coagulation. A prothrombotic state induced by more common factors including the contraceptive pill, pregnancy, and the puerperium, and neoplasia also seems to increase stroke risk. Haematological causes of ischaemic stroke were reviewed in this chapter and a protocol for exclusion of such disorders in patients with ischaemic stroke was discussed.

1 Marcus AJ, Webster BB, Jaffe EA, et al. Synthesis of prosta-
Further reading