Leukocytes and the Natural History of Deep Vein Thrombosis
Current Concepts and Future Directions

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Abstract—Observational studies have shown that inflammatory cells accumulate within the thrombus and surrounding vein wall during the natural history of venous thrombosis. More recent studies have begun to unravel the mechanisms that regulate this interaction and have confirmed that thrombosis and inflammation are intimately linked. This review outlines our current knowledge of the complex relationship between inflammatory cell activity and venous thrombosis and highlights new areas of research in this field. A better understanding of this relationship could lead to the development of novel therapeutic targets that inhibit thrombus formation or promote its resolution. (Arterioscler Thromb Vasc Biol. 2011;31:506-512.)

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Deep vein thrombosis (DVT) is a common condition affecting 1% to 2% of the population, with an annual incidence of 1 in 500.1 DVT can lead to death through pulmonary embolism, and many patients subsequently experience venous reflux, which can lead to the postthrombotic syndrome. This condition is characterized by pain, swelling, and chronic leg ulceration. Around one quarter of patients develop postthrombotic syndrome within 1 year of the episode of thrombosis.2 DVT is therefore potentially fatal, can cause significant patient morbidity, and has become an economic burden for health care services in the developed world.

Treatments for DVT, such as anticoagulation, prevent thrombus propagation and extension but have little effect on existing thrombi, which resolve naturally through a process of organization and vein recanalization.3 Rapid natural resolution is associated with less valvular damage, reduced venous hypertension, and fewer postthrombotic complications.4–6 Treatments that remove thrombus rapidly, however, such as thrombolysis and mechanical removal, are associated with increased morbidity, have significant hemorrhagic side effects, and increase the risk of rethrombosis.7 An organized thrombus (usually those that have been present for more than 14 days), a history of stroke, and a recent operation are contraindications to the use of thrombolytic therapy.8 Alternative forms of treatment that prevent thrombosis or accelerate natural thrombus resolution without hemorrhagic side effects would therefore be attractive. It is likely that these will be developed only from a better understanding of the mechanisms that regulate thrombus formation and its resolution.

There is increasing evidence that inflammatory processes and DVT are intimately linked. This review outlines our current understanding of the complex relationship between inflammatory cell activity and venous thrombosis and highlights new areas of research in this field.

Venous Thrombus Formation
A triad of vessel wall injury, venous stasis, and blood hypercoagulability have historically been considered to predispose to venous thrombosis.9 Venous thrombi arise in both the vein valve pockets and dilated sinuses of the lower limbs, are fibrin and red cell rich, and form on the surface of the endothelium.10–12 They have a laminar structure consisting of layers of platelets, leukocytes and fibrin (lines of Zahn) that encompass the main erythrocyte mass. This is unlike the amorphous structure of a blood clot, which consists predominantly of erythrocytes within a fine fibrin mesh.9

Studies in the 1970s using radiolabeled leukocytes have shown uptake of white blood cells into venous thrombi,13 and accumulation of polymorphonuclear neutrophils (PMNs) on the abluminal side of the endothelium following occlusion of canine veins, led to the speculation that white-cell–induced damage to the endothelium may be a contributing factor to venous thrombosis in humans14 (Figure 1, i). Exposure of the collagen–rich wall is said to lead to platelet aggregation and further leukocyte sequestration, which results in a nidus for thrombus propagation.15,16 C-reactive protein, an inflammatory marker, has been shown to increase in patients experiencing DVT.17 Inflammation is therefore considered an important mechanism for venous thrombus formation.
Inflammation and the Endothelium

The formation, propagation, and dissolution of venous thrombi represent a balance between coagulation and innate protective mechanisms, specifically the circulating inhibitors of coagulation (e.g., tissue factor [TF] pathway inhibitor, thrombomodulin, protein C, plasminogen activator inhibitors) and promoters of fibrinolysis (e.g., plasminogen activators).9

Endothelial microtears containing leukocytes have been demonstrated by electron microscopy in the deep veins following hip surgery in dogs,14 leading to the suggestion that these may be the nidus for venous thrombus formation. Others, however, have found no major overt damage to the endothelium at sites of thrombosis in humans.10 Crushing the vein to cause endothelial damage does not lead to the formation of experimental venous thrombi,19 and scanning electron microscopy reveals minimal endothelial damage immediately after thrombus formation in the rat.20 Disturbance of the endothelium by mechanical (e.g., stretch or surgery) or chemical means (e.g., sepsis) can, however, cause activation rather than overt damage. This results in increased expression of procoagulant proteins, such as tissue factor (TF),21 and cytokines and surface adhesion of molecules that promote leukocyte adhesion and initiate thrombosis22 (Figure 1, i). Genetic knockout of the adhesion molecules E- and P-selectin results in reduced thrombus size, and this is associated with altered leukocyte accumulation in the surrounding vein wall.23 Neutralizing P-selectin glycoprotein ligand-1 also reduces local inflammation and thrombus size and could be a potential treatment for the prevention of DVT in the future.24

Circulating TF, in the form of microparticles released by activated leukocytes, accumulate in areas of stasis, such as the vein valve pockets.21 Leukocyte microparticles that express P-selectin glycoprotein ligand-1 bind to P-selectin both on platelets and activated endothelial cells,25,26 and this source of TF could sustain the production of thrombin on the forming thrombus, promoting its propagation.27 The relative contribution of leukocyte microparticles to venous thrombosis is, however, not clear, as adaptive transfer of bone marrow (BM) from mice expressing low levels of TF into wild-types does not inhibit thrombus formation.28 It appears that the vessel wall is the most important source of TF that promotes thrombogenesis,28 and therefore activation of the endothelium may be pivotal in the inflammatory processes that lead to thrombosis.

PMNs

PMNs are found in large numbers within the early venous thrombus.30,29 Recent studies on the mechanisms of PMN recruitment to sites of sterile inflammation have revealed that intravascular danger signals, including the activation of the Nlrp3 inflammasome, generation of a chemokine gradient, and release of formyl-peptide signals, act as a guide to these sites.30 PMN adhesion to endothelium is mediated by interactions between the integrin αMβ2 (Mac1) and its endothelial ligand intracellular adhesion molecule-130 (Figure 1, ii). Whether a similar mechanism is involved in PMN accumulation during venous thrombosis, remains to be determined though it appears that the thrombogenic effects of antiphospholipid antibodies are mediated in part by intercellular adhesion molecule-1.31

Aside from the formation of a nidus for thrombus propagation, recent data have emerged that suggest that recruited PMNs may initiate thrombosis through the formation of neutrophil extracellular traps32 (Figure 1, iii). These extracellular DNA fibers, which are made up of histones and neutrophil antimicrobial proteins, form after a cell death program in response to inflammatory stimuli (e.g., interleukin [IL]-8, reactive oxygen species) from cells in vitro33–35 and have been linked to small vessel vasculitis36 and preeclampsia.37 DNA traps appear in the plasma and thrombus following induction of DVT in a baboon model and could provide a scaffold for thrombus formation, although their precise mechanism of action remains to be elucidated.32

The role of PMNs in the natural history of the venous thrombus is complex, however. Selective antibody depletion of PMNs in a rat stasis model led to larger venous thrombi, suggesting that PMNs are not required for thrombus formation but may be important in the removal of forming thrombi38 (Figure 2). However, a similar finding was not demonstrated in a murine model, and CXCR2-dependent thrombus resolution appeared independent of the CXCR2 primary effector leukocyte,
Putative functions of leukocytes in venous thrombus resolution. Leukocytes accumulate in the venous thrombus during its resolution. PMNs predominate in the early stages of resolution, with mononuclear phagocytes (Mo) predominating later. The origin of these cells appears to be from the BM; however, the contribution of tissue-resident macrophages or cells derived from the splenic reservoir remains unknown. Leukocytes signal through a TLR9 mechanism and may be stimulated by fibrinogen and its degradation products. They are speculated to have a number of functions that are important for thrombus resolution.

Red Blood Cells, Inflammation, and Thrombosis

The contribution of red blood cells (RBCs) to venous thrombosis remains poorly understood, despite their abundance in the early venous thrombus and their contribution to platelet rich arterial thrombi. The cytoplasm of RBCs is rich in iron, which when released into the circulation is highly inflammatory because of its oxidative effects on the endothelium. It has been hypothesized that reactive oxygen species, produced by leukocytes and vessel wall at the nidus of thrombosis, oxidizes hemoglobin in RBCs that become trapped by cross-linked fibrin forming at this point. This results in the formation of methemoglobin containing Fe$^{3+}$. The release of Fe$^{2+}$ leads to a cascade of RBC lysis that results in further endothelial dysfunction and thrombus propagation (Figure 1, iv).

Putative Functions of Inflammatory Cells During Thrombus Resolution

The role of inflammation, and specifically leukocytes, such as PMNs and the monocyte/macrophage, during thrombus res-
olution is not completely understood (Figure 2). Studies in mice lacking the ets transcription factor Pu.1 suggest that inflammatory cell activity may not be a prerequisite for tissue repair.57 Nevertheless, leukocytes make up a significant proportion of the cells in the thrombus, and interventions that lead to alterations in their accumulation have significant effects on subsequent thrombus resolution.

There is increased plasminogen activator content, both tissue-type and urokinase-type (uPA) that colocalizes with macrophages in thrombus formed in the rat.58,59 This has led to the speculation that fibrinolysis is important for thrombus resolution. Deletion of the tissue-type plasminogen activator gene (tPA−/−), however, has no effect on this process.51 By contrast, deletion of the urokinase-type plasminogen activator gene (uPA−/−) prevents resolution and is associated with reduced macrophage numbers in the thrombus.51 Adoptive transfer of wild-type BM into uPA−/− mice rescues normal resolution,51,60–62 and upregulation of uPA in macrophages enhances this process.56 Binding of urokinase to its receptor (uPAR) on the cell membrane leads to the generation of plasmin at the cell surface. Plasmin activates other proteases, such as matrix metalloproteinases (including matrix metalloproteinases 2 and 9), that degrade extracellular matrix, and facilitate cell migration.60–62 These data lead us to speculate that monocyte-associated urokinase activity is important for venous thrombus resolution.

Neovascular channels appear around the thrombus wall junction and within the thrombus as resolution proceeds.3,10,11 Histological studies in rodent models suggest that these channels are derived from the vein wall, and they may also be derived from cells residing in the thrombus.63 We have found that enhancing the levels of vascular endothelial growth factor either alone or simultaneously with a number of other angiogenic factors through upregulation of HIF1α within the thrombus enhances its resolution.64–67 These outcomes are linked to accumulation of macrophages within the thrombus, which perhaps act as cellular chaperones, as recently described in the development of the vascular network.68

Leukocyte Signaling During Venous Thrombus Resolution

Fibrin(ogen) and its degradation products are present in abundance in the thrombus.51,69 These molecules are able to stimulate recruitment and activation of leukocytes, produce cytokines (tumor necrosis factor α and IL-1β) and chemokines (IL-8 and monocyte chemotactic protein-1) in inflammatory settings,70,71 and promote phagocytosis and cell migration in vitro.72 Interaction of the fibrinogen γ chain residues 390 to 396 with Mac1 is thought to be an important pathway by which these molecules influence leukocyte activity.73 Mononuclear cells may also interact with fibrinogen to produce chemokines through a TLR4-dependent mechanism.73 Examination of thrombus formation and resolution in mice that carry mutations, such as Fibγ 390 to 396A, may provide new insights into the role of fibrin(ogen) and its degradation products in the natural history of venous thrombi.

More recently, data have emerged that have provided new insights into the signaling mechanisms that regulate the functions of leukocytes during venous thrombus resolution. Deletion of TLR9−/− gene impairs resolution assessed at days 2 and 8 after induction, despite an increase in the numbers of both PMNs and Mac2+ macrophages in the thrombus.74 These data suggest that TLR9 is important for leukocyte function during thrombus resolution. This effect was independent of MyD88 signaling (a major TLR signaling pathway) and was related to NOTCH ligand δ-like 4 pathways. TLR9−/− mice have reduced thrombus neovascularization and decreased levels of the Th1 inflammatory cytokines interferon-α, IL-1α, and IL-2 in the vein wall, which appear to be important for venous thrombus resolution.52 Further investigation is required to elucidate the role of TLR9 and MyD88 in later phases of thrombus resolution (beyond 8 days) and whether other TLRs are involved.

Mononuclear Phagocytes

Macrophage accumulation within the thrombus is a hallmark of resolving thrombus in both humans and experimental models.59,70 Adoptive transfer experiments suggest that these cells are derived from the BM; however, macrophages that are not of BM origin may also have a role in thrombus resolution.51 Macrophages that accumulate in the thrombus could be derived from cells that are resident within the vein wall or may even arise from the splenic reservoir that has recently been described.26 The relative contribution of these sources for cells implicated in the resolution of venous thrombi remains unknown.

Although it is generally thought that the accumulation of macrophages is dependent on the recruitment of circulating monocytes, direct visualization of these cells entering the thrombus has yet to be demonstrated. Renewal of certain mononuclear cells (microglia and Langerhans cells) in the steady state appears to be independent of BM.77,78 Adult Langerhans cells self-renew in situ and proliferate during inflammation.79 A demonstration that macrophages proliferate in the thrombus would change the current paradigm regarding the nature of their accumulation.

The function of mononuclear cells in the thrombus also remains to be fully elucidated. As they are phagocytic by definition, it seems reasonable to speculate that they contribute to the clearance of cells, nucleotides, and matrix proteins within the thrombus. These cells may also promote fibrinolysis, are associated with angiogenesis, and could regulate tissue remodeling—processes seemingly beneficial for thrombus resolution. To add further complexity, both monocytes and macrophages consist of heterogeneous populations of cells, which appear to have distinct functions.85

Circulating monocyte subsets can be distinguished on the basis of their expression of surface receptors.84 Circulating inflammatory monocytes express Ly6C in the mouse and are recruited into tissue, where they undergo activation in a pathogen-dependent response.84,85 In a model of spinal cord injury, recruitment of Ly6C+ monocytes appears important for tissue repair,86 and these monocytes may also contribute to the fraction of myeloid-derived suppressor cells that promote tumor-driven angiogenesis.87 The contribution of Ly6C+ monocytes in the formation and resolution of venous thrombosis has yet to be established, but our previous studies suggest that recruitment of this subset may be important for...
thrombus resolution. Impaired resolution occurs in Ccr2−/− mice, and CCR2 is required for the exit of Ly6C+ monocytes from BM.

Ly6C− monocytes exhibit long-range crawling over the endothelium of both arteries and veins. It has been hypothesized that they are involved in the surveying of the vasculature and sensing of tissue damage such as dying or infected cells. In a model of myocardial infarction, Ly6C− monocytes initially accumulate in the healing myocardium and may digest damaged tissue. In a later reparative phase, Ly6C− monocytes predominate and are suggested to be involved in tissue repair by inducing myofibroblast accumulation, angiogenesis, and collagen deposition. The human equivalent of Ly6C− murine monocytes (CD14dim monocytes) have recently been implicated in the pathogenesis of autoimmune diseases, such as lupus, and respond to viruses and nucleic acid–containing immune complexes via a proinflammatory TLR7-TLR8-MyD88-mitogen-activated protein kinase pathway. Whether these patrolling cells have a role in venous thrombosis (initiation or resolution) remains to be determined, although we speculate that their patrolling function makes them an ideal candidate for the detection of endothelial dysfunction and possible initiation of coagulation.

When monocytes enter tissue, they differentiate into macrophages. Based on in vitro studies, these cells have been tentatively classified by some into 2 main phenotypes: those that promote inflammatory responses (M1 or classically activated, expressing inflammatory mediators, such as tumor necrosis factor α and nitric oxide synthase 2) and those that attenuate inflammatory responses (M2a-c or alternatively activated, expressing arginase, mannose receptor, and the transcription factors Fizz1 and Ym1/2). Others consider dividing macrophage populations based on their immunologic or trophic roles in response to granulocyte/macrophage colony stimulating factor (CSF) (also known as CSF2) or macrophage CSF (also known as CSF1) respectively. A rigid classification of macrophages probably represents the extremes of a continuous spectrum and may be too simplistic, as these cells may exhibit characteristics of more than 1 phenotype. M2-like macrophages have, however, been reported in the healing myocardium and injured skeletal muscle, where they are considered to be involved in tissue repair and wound resolution.

Clinical trials involving the therapeutic targeting of macrophages in other vascular diseases, such as atherosclerosis, has been largely unsuccessful. This is in part because of a lack of understanding of their function. Their roles in venous thrombosis require investigation, especially because different monocyte and macrophage phenotypes may have complementary and contrasting functions. This could be achieved through the use of functional reporter mice that express fluorescent proteins linked to cell specific genes. We are currently developing these tools to examine cellular functions in thrombus resolution.

Inflammation is a central mechanism in both the genesis and resolution of venous thrombi. The temporal accumulation of leukocytes in the forming (PMMs) and resolving thrombus (macrophages) is part of a dynamic intravascular wound healing process that results in either the early lysis of the thrombus or its stabilization and subsequent resolution. Enhancing our understanding of the cellular and molecular pathways that mediate sterile inflammation in the context of venous thrombosis could lead to the development of novel therapeutic targets for (1) prevention of DVT in a manner that does not promote pathological bleeding and (2) acceleration of natural thrombus resolution to reduce the incidence of postthrombotic complications. These may be achieved through developments in molecular and cellular imaging capable of delineating specific inflammatory processes, that are currently on the horizon.

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Disclosures

None.

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