


The Anticoagulant and Nonanticoagulant Properties of Heparin

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Abstract

Heparins represent one of the most frequently used pharmacotherapeutics. Discovered around 1926, routine clinical anticoagulant use of heparin was initiated only after the publication of several seminal papers in the early 1970s by the group of Kakkar. It was shown that heparin prevents venous thromboembolism and mortality from pulmonary embolism in patients after surgery. With the subsequent development of low-molecular-weight heparins and synthetic heparin derivatives, a family of related drugs was created that continues to prove its clinical value in thromboprophylaxis and in prevention of clotting in extracorporeal devices. Fundamental and applied research has revealed a complex pharmacodynamic profile of heparins that goes beyond its anticoagulant use. Recognition of the complex multifaceted beneficial effects of heparin underscores its therapeutic potential in various clinical situations. In this review we focus on the anticoagulant and nonanticoagulant activities of heparin and, where possible, discuss the underlying molecular mechanisms that explain the diversity of heparin's biological actions.

Keywords

- ▶ heparin
- ▶ anticoagulation
- ▶ metastasis
- ▶ inflammation

Heparin Structure and Biosynthesis

Heparin is a biomolecule that belongs to the class of glycosaminoglycans (GAGs). GAGs exist as large linear polysaccharide structures, composed of repeated structural motifs. Besides heparin, the family of GAGs includes heparan sulfate (HS), dermatan sulfate, chondroitin sulfate (CS), keratan sulfate, and hyaluronan. Each of these GAGs is composed of different repeating disaccharide units and, with the exception of hyaluronan, is sulfated at various positions. These sulfations introduce functionally important negatively charged groups to the GAGs. Both heparin and HS are complex heterogeneous mixtures of repeating uronic acid and D-glucosamine/N-acetyl-D-glucosamine units, with heparin being generally relatively smaller and containing relatively more sulfated groups than HS.¹ The biological

functions of heparin are for a large part dependent on electrostatic interactions and the influence of sulfation on these functions have been well described since decades.²

Heparin biosynthesis occurs in mammalian cells through a complex process that involves many enzymatic steps. After a core tetrasaccharide linker is synthesized, by sequential transfer of monosaccharide units onto a cell-type-specific core protein, serglycin proteoglycan in the case of heparin, heparin synthesis starts. The tetrasaccharide linker is attached to the side-chain oxygen atom of a serine residue and thereby forms an O-linked glycan. The linker then serves as the template onto which heparosan, the precursor molecule of heparin (and of HS), is synthesized by sequential enzymatic transfer of defined disaccharide units. These saccharide units subsequently undergo further posttranslational processing, including sulfations, and will finally result in the mature polyanionic heparin

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proteoglycan molecules that are found exclusively in mast cell granules. Here, the heparin proteoglycans serve to concentrate the various positively charged cytokines, growth factors, and proteases that become secreted upon mast cell stimulation/degranulation. This unique localization is in contrast to the ubiquitously expressed HS proteoglycans (HSPGs), which exist as part of proteoglycans in the extracellular matrix (ECM). Through mild acidic hydrolysis or via random cleavage at glucuronic acid residues by endo- β -D-glucuronidase, a heterogeneous mixture of heparin molecules ranging from 5 to 25 kDa in molar weight can be generated. Of the fully processed heparin molecules, only one in three is able to bind to anti-thrombin (AT).³ The pharmaceutical-grade heparin that is extracted from animal tissues (mostly porcine mucosa) and that contains the mixture of heparin molecules with varying molecular weights is called unfractionated heparin (UFH).

The ubiquitously expressed HSPGs are also heterogeneous with only less than 5% of endothelial HSPG able to bind AT.³ Despite this low percentage, the contribution of HSPGs to anticoagulant homeostasis in the vasculature is considered important due to their presence in the endothelial glycocalyx, thereby contributing to the overall nonthrombogenic nature of the vessel wall under physiological conditions.

The majority of heparins currently being used have been purified from porcine and bovine intestinal mucosa and to a lesser extent from bovine lungs that are collected in the slaughterhouse. These tissues are rich in mast cells that form a first-line defense against invading pathogens. Potential contamination of pharmaceutical heparin together with the vulnerability of the supply chain and increasing demand has led to the consideration of other animal sources and bioengineered heparin.⁴ However, none of these alternative strategies has yet led to a large-scale production of pharmaceutical-grade heparin.

Heparin and Its Derivatives

The discovery of the separability of heparins' anticoagulant effects toward coagulation factor Xa (FXa) and thrombin (see the next section on anticoagulant properties of heparin) led to the development of shorter heparin variants with optimized anticoagulant activity and reduced bleeding risk as compared with UFH. These heparins are obtained by fragmentation of UFH by chemical or enzymological means and are commonly described as low-molecular-weight heparins (LMWHs). While UFH molecules may range in size from 6 to 60 kDa, those for LMWHs vary between 1 and 10 kDa.^{5,6} Many different types of LMWHs exist (e.g., Food and Drug Administration-approved dalteparin, enoxaparin, and tinzaparin) with the more recent generation of second-generation LMWH or ultralow molecular weight heparins such as bempiparin and semuloparin (reviewed in Walenga and Lyman⁷). All these LMWH preparations express different activities toward FXa and have different pharmacokinetic profiles. Consequently, they have different applications in clinical practice.

The use of LMWH offers several advantages over UFH as the former has a higher bioavailability, allows better predictable dosing, and is associated with less adverse reactions. As

a result, LMWH has become the heparin of choice in many standard clinical situations such as venous thromboembolism (VTE), major surgery, and acute coronary syndrome.^{7,8} UFH, however, remains preferred in situations such as prevention of coagulation in extracorporeal devices and in patients with renal failure due to less dependence on the kidneys for clearance and improved reversibility with protamine.⁹

Considering the drawbacks of heparin isolated from animal origin, a synthetic anticoagulant with a high affinity toward AT was developed.¹⁰ The compound in question, fondaparinux, comprises a specific "pentasaccharide sequence" and will be discussed in the next section in more detail (see Anticoagulant Properties of Heparin). Fondaparinux is approved for the prevention and therapy of VTE after major surgery, and has been successfully applied in patients with, e.g., myocardial infarction.¹¹ Likewise selective modified forms of heparins such as (de)sulfated, deacetylated, or glycol-split heparins are being developed as alternative nonanticoagulant drugs and are currently being tested in preclinical and clinical studies for their application in particular settings.¹²

Anticoagulant Properties of Heparin

Of all its functions, the anticoagulant action of heparin has been described best. It is based on a proposed combined template-based/allosteric mechanism for the interaction between the plasma serine proteinase inhibitor AT and either thrombin or FXa. Although the inhibition of factors VIIa, IXa, XIa, and kallikrein by AT is also stimulated by heparin, these will not be discussed here as these are regarded of overall lesser importance than the inhibition of FXa and thrombin.

Heparin facilitates the interaction between AT and thrombin (or FXa) by binding tightly with a specific site to AT. This site is known as the "pentasaccharide sequence," consisting of a specific sequence of five highly sulfated sugar units in the heparin polymer.^{13,14} **Fig. 1A** shows an experimentally derived structure for the ternary complex composed of a synthetic heparin mimetic binding both thrombin and AT.¹⁵

Moreover, AT undergoes a conformational change upon binding to heparin (**Fig. 1B**). This change makes AT about twice as reactive to thrombin and up to 100 times more reactive to FXa.^{16,17} The remaining 100- to 1,000-fold increase of activity is due to the fact that thrombin binds loosely and aspecifically to the remainder of the heparin chain and is then guided rapidly to the bound AT by one-dimensional approximation along the heparin molecule. FXa inactivation is subject to the same mechanism but binds 10 times less strongly than thrombin and is inactivated three times less efficiently.¹⁸ The interaction between the complex of heparin and AT with FXa or thrombin is thus essentially different. FXa can interact with the pentasaccharide-AT complex as such, whereas inactivation of thrombin requires a "slide" of 17 sugar units at the nonreducing end of the pentasaccharide.¹⁹ Given this requirement in sugar length for the inactivation of thrombin by the heparin-AT complex, the "slide" can best occur with UFH and not with LMWH.⁹

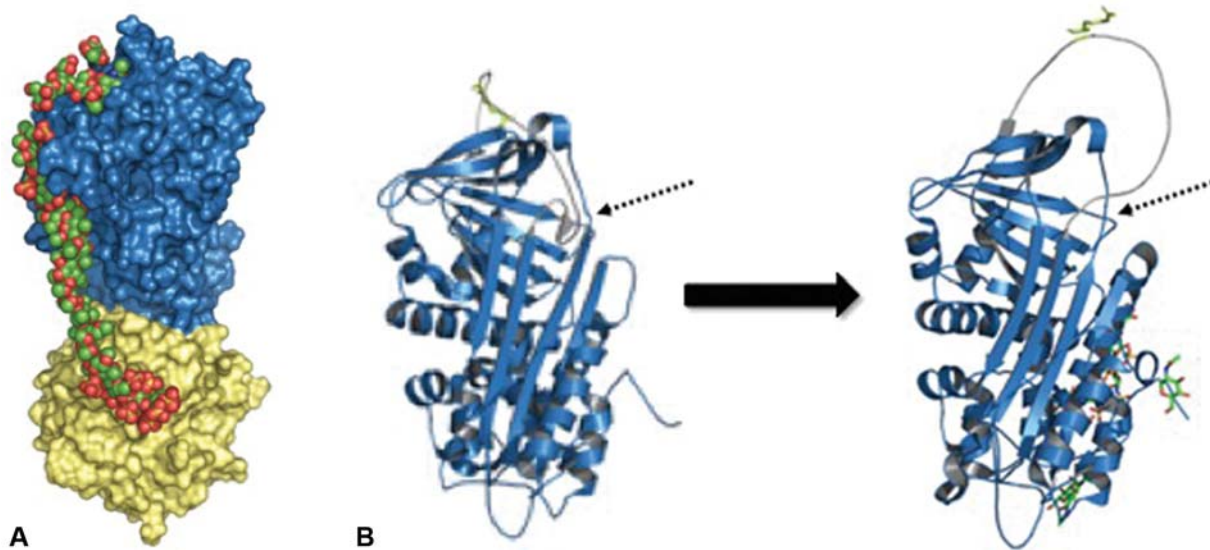


Fig. 1 The interaction of heparin with antithrombin (AT) and thrombin. Three-dimensional structure of the ternary complex between thrombin and AT, bound to the heparin mimetic, SR123781. (A) Shown are the surface structures of thrombin (pale yellow) and AT (blue) that are bound by a heparin fragment (as ball presentation in red and green). This structure represents the 2.5 Å crystal structure as is deposited in the protein data bank by accession number 1TB6.pdb.¹⁵ (B) Conformational change in AT induced by the binding of the pentasaccharide. Left: crystal structure of monomeric AT (1F1T.pdb, in blue) in the nonbound state, with the reactive center loop in gray and P1 residue Arg393 in yellow. Note the hinge region (as indicated by the dashed arrow) in a folded conformation. Right: upon binding of a pentasaccharide (as indicated by the ring structures in green and red) the hinge region assumes a more elongated conformation, causing the reactive center loop (in gray) and the P1 residue (in yellow) to be exposed (from PDB coordination file 2GD4.pdb).

Heparin as a Direct and Indirect Modifier of Coagulation Pathways

The registration and application of heparin as an anticoagulant has spurred the research into mechanistic studies to explain the beneficial effects observed with its clinical use. Interestingly, not only antithrombotic functions were recorded but also a wide variety of nonanticoagulant properties of heparin were elucidated. These nonanticoagulant properties include heparin-mediated modulation of adhesion molecules,^{20,21} heparin-induced release of endothelial-bound tissue factor pathway inhibitor (TFPI),²² modulation of fibrinolytic activators such as tissue plasminogen activator,²³ the binding of chemokines and cytokines,²⁴ and more recently the neutralization of damage-associated molecular patterns (DAMPs).²⁵

Hence, heparin expresses anticoagulant and nonanticoagulant activities (→ Fig. 2) that collectively contribute to the therapeutic effects observed in the wide range of clinical applications of heparin (→ Table 1).

It should be mentioned that also procoagulant/antifibrinolytic properties of heparin have been observed. These include the stimulation of the inhibition of activated protein C by protein C inhibitors,²⁶ the attenuation of fibrinolytic pathways,²⁷ the altered inactivation of coagulation factor Va,²⁸ and the stimulation of clotting of FXa-initiated coagulation of plasma.³ The latter two properties, however, have been tested in the absence of AT. Such conditions may occur in individuals with a consumptive coagulopathy. Treating individuals with heparin may cause depletion of AT.²⁹ Further, of interest here are (1) the observed acceleration of thrombin formation that occurs through inhibition of TFPI function by the nonanticoagulant proteoglycan AV513,³⁰ an effect which promotes the

activation of coagulation factor V³ and (2) the procoagulant properties of polyphosphate, a linear polyanionic polymer of inorganic phosphate,³¹ as well as of extracellular RNA,³² both of which induce autoactivation of coagulation factors XII and XI, but cannot substitute for heparin as an anticoagulant.

Nonanticoagulant Properties of Heparin

Although heparin was discovered and developed as an anticoagulant, it was realized that roughly 70% of the heparin molecules in UFH did not bind to AT,^{33–35} a fraction which was termed “inactive heparin”³⁴ or “low-affinity material.”^{33,35} Hence, over the years patients receiving UFH have been injected also with “inactive” heparin molecules. Given the low frequency and accepted minor side effects of UFH use, the “inactive” heparin molecules may have contributed positively to the overall therapeutic effects of UFH.

Indeed, it has been realized that the earlier termed “inactive” fraction of heparin contains activities that appear to be tissue-protective. In those clinical situations where use of heparin as an anticoagulant is indicated, tissue damage through ischemia and reperfusion is frequent. Therefore, in retrospect nonanticoagulant beneficial properties of heparin may well have contributed to the success of heparin.

Acknowledging the observed nonanticoagulant properties of heparin, both UFH and LMWH have been applied for several other applications besides their use as antithrombotic therapeutics. These include applications such as surface coating of biomedical devices, treatment of hemodynamic disorders, modulation of growth factors, and serving as an adjunct to chemotherapeutic and anti-inflammatory drugs. Currently there are nearly 250 different manually reviewed entries in

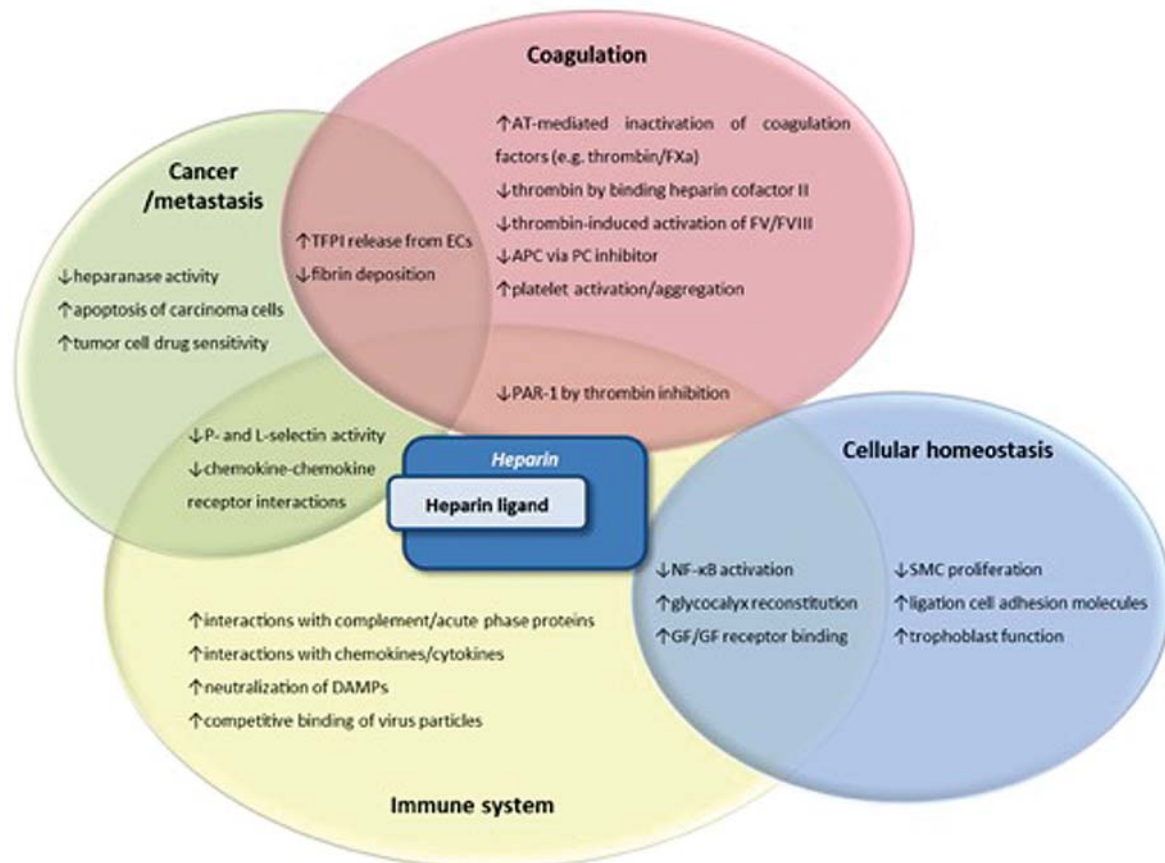


Fig. 2 Heparin–ligand interactions in physiological and pathological states. Schematic illustration of the polypharmacological properties of heparin in cancer/metastasis (green), coagulation (red), cellular homeostasis (blue), the immune system (yellow), or properties able to affect multiple of these processes. Binding of heparin to a plethora of heparin-binding biomolecules may result in alterations of important physiological processes and affect physiological states. (A)PC, (activated) protein C; AT, antithrombin; DAMPs, damage-associated molecular patterns; ECs, endothelial cells; GF, growth factor; PAR-1, protease activated receptor-1; SMC, smooth muscle cell; TFPI, tissue factor pathway inhibitor.

Table 1 Physiological/pathological states and medical procedures for which heparin is being applied

Targets and applications of heparin			
Acute myocardial infarction	Deep vein thrombosis	Ischemic stroke	Surgery
Alzheimer's disease	Drug delivery	Pregnancy	Extracorporeal circulation and hemodialysis
Atrial fibrillation	Infectious disease	Pulmonary embolism	Transplantation

Note: Heparin is a multifunctional biomolecule which is used in many different types of medical applications and for various physiological and pathological states. Some examples of heparin's targets and applications, as extracted from references throughout this text, are given.

the Swiss-Prot database for proteins that bind heparin and whose biological properties can be modulated by heparin. Modulating effects of heparin not necessarily depend on its anticoagulant activity and an increasing number of described effects can be contributed to heparin's nonanticoagulant properties. The following sections provide an overview of the most studied nonanticoagulant properties of heparin, supported, if possible, by molecular mechanisms that explain the observed effects of heparin.

Growth Factor Modification

Binding of growth factors to heparins has already been described in the 1980s. In fact, the preparation of purified

growth factors can proceed efficiently through the use of affinity chromatography using immobilized heparin.

Several observations may help to explain the effects that heparins have on growth factor function and availability. HSPG-bound fibroblast growth factor-2 (FGF-2) was shown to enter cells via nonclathrin-mediated lipid raft-dependent pathways, thus preventing the FGF-2 to be directed toward the lysosomes.³⁶ Notably, heparin needs charged molecules to be able to translocate to the cell nucleus and compete with DNA for binding to growth factors.³⁷ It is known that many growth factors (e.g., vascular endothelial growth factor [VEGF], FGF, and platelet-derived growth factor [PDGF]) bind to GAGs present on the cell surface and in the ECM

where they are deposited in the basement membrane.³⁸ As such, these growth factors are protected from proteolytic inactivation or physical denaturation.³⁹ At the same time, they can reside in the endothelial GAG layer and serve as a pool of ready-to-use growth factors that can for instance respond to vascular damage. When the GAG chains are broken down by physiological or pathological stimuli, the bound growth factors are released and may bind (likely in complex with the released GAG) to their respective cellular receptors.⁴⁰ Since heparin can interact directly with both growth factors and their respective receptors, it acts as a bridging or coreceptor molecule which can mediate the formation of high-affinity growth factor-receptor complexes. Addition of exogenous heparin may thus compete with the cell-bound GAGs for binding to the immobilized growth factor pool and stabilize these growth factor-receptor complexes. This stabilization by heparin can result in signaling events and has been observed for heparin sequences exceeding a tetramer polysaccharide length of appropriate composition and structure.^{41,42} Interestingly, for the promotion of VEGF activity, soluble heparin with a chain length of >22 saccharides is required, while a chain length of <18 sugar units results in the inhibition of VEGF binding to its receptor.⁴³ Therefore, LMWH appears to be a potent inhibitor of both angiogenesis and fibrosis.^{43,44}

In recent years, novel applications of heparin-growth factor modifications have been investigated in the field of nano- and regenerative medicine.⁴⁵ Heparin-tailored biomaterials/scaffolds, such as hydrogels, nanoparticles, polymers, and liposomes, have been studied in tissue engineering and drug-delivery applications.⁴⁶ Using high-affinity delivery systems, heparin, growth factors, and/or drugs can site-specifically be delivered to the tissue of interest and exert effects depending on the delivery system.

Antimetastatic Properties of Heparin

Use of heparin as a potential chemotherapeutic adjunct has been under investigation for many years.

Interest in the antimetastatic properties of heparin was sparked by several clinical observations indicating that cancer patients who were treated with heparin or heparin derivatives for cancer-associated thromboembolic disease appeared to have prolonged survival.^{47,48} Several preclinical studies investigating predominantly melanoma or mammary carcinoma tumors aimed to elaborate on the anticancer and antimetastatic properties of heparin (reviewed in Borsig⁴⁹).

Besides heparin's anti-FXa and antithrombin effects, an additional anticoagulant mechanism was suggested to be involved in metastatic mechanisms, namely the induction of TFPI release from cells,^{22,50} a phenomenon that is observed in cancer patients receiving heparin.⁵¹ TFPI exhibits antiangiogenic and antimetastatic effects in vitro and in vivo and its release is induced by both anticoagulant and nonanticoagulant LMWHs.⁵² Furthermore, heparin is thought to downregulate the expression and activity of tissue factor through modulation of growth factor receptor-mediated activation of nuclear factor-kappa B (NF- κ B).⁵³ Another potential antimetastatic mechanism of heparin is hypothesized to be the inhibition

of fibrin deposition around tumor cells, which lifts the protection from cancer cells against immune cell attack.⁵⁴

Besides these anticoagulant mechanisms through which heparin is able to exert anticancer properties, nonanticoagulant mechanisms are assumed to be involved in these effects as well. One of the contributing nonanticoagulant activities was proposed to be the modulation of selectin activity (see the Anti-inflammatory Properties section). Heparin is known to block P- and L-selectin binding to natural and tumor cell ligands. This implies that heparin is able to modify selectin-mediated events, thereby influencing cell-cell interactions that drive tissue growth and differentiation. Interestingly, there appears to be specificity in the effects that glycans have on selectin binding, with, e.g., heparin and HS showing differential interaction with selectins.⁵⁵ UFH and LMWH were shown to inhibit metastasis of carcinoma cells to an extent comparable to that of P-selectin deficient cells.^{49,56} It is however difficult to describe this selectin-blocking heparin property as being specifically antimetastatic since the exact mechanisms still remain under investigation.^{57,58}

Modulation of chemokine-chemokine receptor interactions constitutes another action of heparin affecting cell migration and metastasis, as was shown for the chemokine receptor CXCR4 and its ligand CXCL12 through in vitro studies.⁵⁹ Also the interaction of the integrin VLA-4 with its ligand VCAM-1 was found to be inhibited by heparin, reducing melanoma cell adhesion and experimental melanoma metastasis.⁶⁰ CXCR1/2 and CXCL8 are other potential targets for heparin.⁶¹ Heparanase is an endoglycosidase that degrades HS in the ECM and on cell surfaces. Inhibition of heparanase activity by heparin was shown to inhibit tumor invasion and metastasis.⁶² The inhibition is abolished by using a totally desulfated type of heparin.⁶³

Other ways through which heparin is thought to exert anticancer effects are the direct induction of apoptosis of carcinoma cells through cell-cycle arrest⁶⁴ and the enhancement of tumor cell sensitivity to chemotherapy by increasing drug uptake and reversing the downregulation of tumor suppressor genes.^{65,66} The Wnt signaling pathway is thought to be involved in this latter process.⁶⁶ Recently it was also shown that galectin-3, a β -galactoside-binding protein which is commonly overexpressed in most types of cancers, can be inhibited by heparins, independent of their anticoagulant properties. The exact mechanism via which this inhibition results in reduced metastasis is unclear but ligation by heparin has clearly inhibitory effects on galectin-3-mediated cancer cell behavior.⁶⁷

In view of the close interplay between the hemostatic system and those mechanisms that underlie processes of cancer cell growth and metastasis, it is likely that the observed therapeutic effects of heparin in cancer patients can be attributed to multiple effects of the drug. It appears that prescription of heparin (mainly LMWH or UFH) in cancer patients without associated thrombosis results in increased survival, in the absence of an increased bleeding risk, as was recently shown by means of a meta-analysis for particularly small cell lung cancer patients⁶⁸ or in a randomized phase III study in non-small cell lung cancer patients.⁶⁹ It should be noted, however,

that a recent multicenter randomized trial evaluating the prophylactic use of LMWH in >2,000 lung cancer patients found no difference in overall or metastasis-free survival between heparin-treated and untreated groups, while there was an increase in clinically relevant nonmajor bleeding events.^{70,71} Other trials indicated that patients with a “better prognosis” might benefit most from heparin treatment.^{72–74}

Of note, the aforementioned studies have used registered anticoagulant heparin formulations and it has been observed before that the antimetastatic properties do not depend on the anticoagulant functions of heparin.^{75,76} Potentially contributing to this is the activity of heparin cofactor II, a plasma serpin with anticoagulant activities, which is known to act through both pentasaccharide-containing heparin (anticoagulant heparin) as well as through nonanticoagulant heparin. Via both *in vitro* study and animal studies it was shown that heparin cofactor II enhances metastasis in nonsmall cell lung cancer, which supported an observed correlation between high pretreatment plasma levels of heparin cofactor II and a reduced survival.⁷⁷ As such, nonanticoagulant heparins are being probed for their usefulness as adjuncts in preclinical antimetastatic models. Several studies suggest that such heparin derivatives could indeed provide the sought after anticancer effects, without affecting the coagulation system.^{76,78} The fact that these heparin forms can be dosed to higher concentrations without risking bleeding in patients may prove beneficial to their potential application.

Also, the effects of heparin on the inhibition of metastasis were not observed for fondaparinux,⁵⁶ suggesting that the earlier listed anticoagulant properties are not per se the most significant contribution to the overall activity. Likewise, the mentioned inhibitory interaction of heparins with selectins is independent of the anticoagulant function of heparin and was described also for chemically modified forms of heparin *in vitro*.⁷⁹ As the hemostatic system is involved in cancer development, it remains to be proven what the true therapeutic value of inclusion of nonanticoagulant heparin in cancer treatment is and what the best treatment strategy is in current clinical practice. Hence, the possibility that different types of heparin have different therapeutic effects in various types of cancer should be considered. Therefore, clinical studies are needed that are designed to determine anticancer effects of a specified heparin type in a specified patient population.

Anti-inflammatory Properties of Heparin

Given the fact that endogenously produced heparin is stored in the granules of mast cells, it may not be surprising that pharmaceutical-grade heparin has immunomodulatory properties. Mast cells play a major role in inflammatory and allergic diseases as they contribute to increased vascular permeability and to allergic and anaphylactic reactions. It has been reported that heparin from mast cells, which is structurally different from clinical-grade heparin, has proinflammatory properties via the stimulation of bradykinin,⁸⁰ similar to what has been reported for oversulphated CS.⁸¹ However, endogenous heparin-like GAGs present on the endothelial cell surface and administered heparin appear to have quite the opposite effect. Patients with pathologies

having a distinct immunological component seem to benefit from administration of heparin because of its anti-inflammatory properties, even though the evidence is not always equally convincing. These pathologies include conditions such as asthma, allergic rhinitis, inflammatory bowel disease, ocular disorders, cystic fibrosis, and burns.^{82–86} Also in several cardiovascular conditions with a clear inflammatory component such as acute coronary syndrome, cardiopulmonary bypass, and thrombophlebitis,^{87–89} the use of heparin is beneficial. In organ preservation and transplantation, in which processes of both ischemia and reperfusion are apparent, heparin is used to reduce vascular thrombosis and ischemia-reperfusion injury.^{90,91} However, the exact benefit or risk of heparin treatment during different stages of these processes is still a subject of debate.

The mechanisms by which heparin and its derivatives are able to express their anti-inflammatory properties reflect their multifaceted effects in biological processes. The anti-inflammatory properties can be roughly divided into two modes of action (→Fig. 3): (1) modulation through binding to soluble plasma ligands and (2) modulation through binding to cell-surface-bound receptors or macromolecules, with potential effects on downstream signaling pathways. In this way, heparin is able to interfere with several (if not all) stages of leukocyte transmigration and extravasation into the target tissue.

Modulation by Heparin through Binding of Soluble Plasma Ligands

Heparin is known to interact with a wide variety of ligands, in particular proteins, and these interactions confer upon heparin its anti-inflammatory functions. Through binding of inflammatory mediators and enzymes, heparin can inhibit activation of inflammatory cells and subsequent propagation of the inflammatory response and tissue damage.⁹² Examples of such ligands are complement proteins which have been described to express changed functional properties upon heparin binding.⁹³ Heparin interferes with both the classical and alternative complement pathways by binding and inhibiting the formation of several complement factors (e.g., active C1 complex, C3 convertase) as well as the membrane attack complex, interfering with terminal cell lysis.⁹⁴

Likewise, proinflammatory molecules like chemokines and cytokines are able to bind to heparin, in which electrostatic forces and interaction sites seem to determine the relative affinity toward a particular molecule.^{95,96} Binding of cytokines to cell-surface GAGs and binding to mast-cell-derived heparins at the site of inflammation result in the concentration and protection of cytokines against proteolytic inactivation and rapid clearance from the circulation. Given that these cytokines and chemokines exert their proinflammatory functions through receptor-mediated events, the administration of exogenous heparin will result in competitive binding between the inflammatory mediators and either the endogenous GAGs/heparins or the administered heparin. Consequently, administered heparin can dissociate cytokines and other proteins from their stationary binding, as it is observed during an acute-phase reaction, in which heparin-binding proteins are elevated in post-heparin plasma.⁹⁷

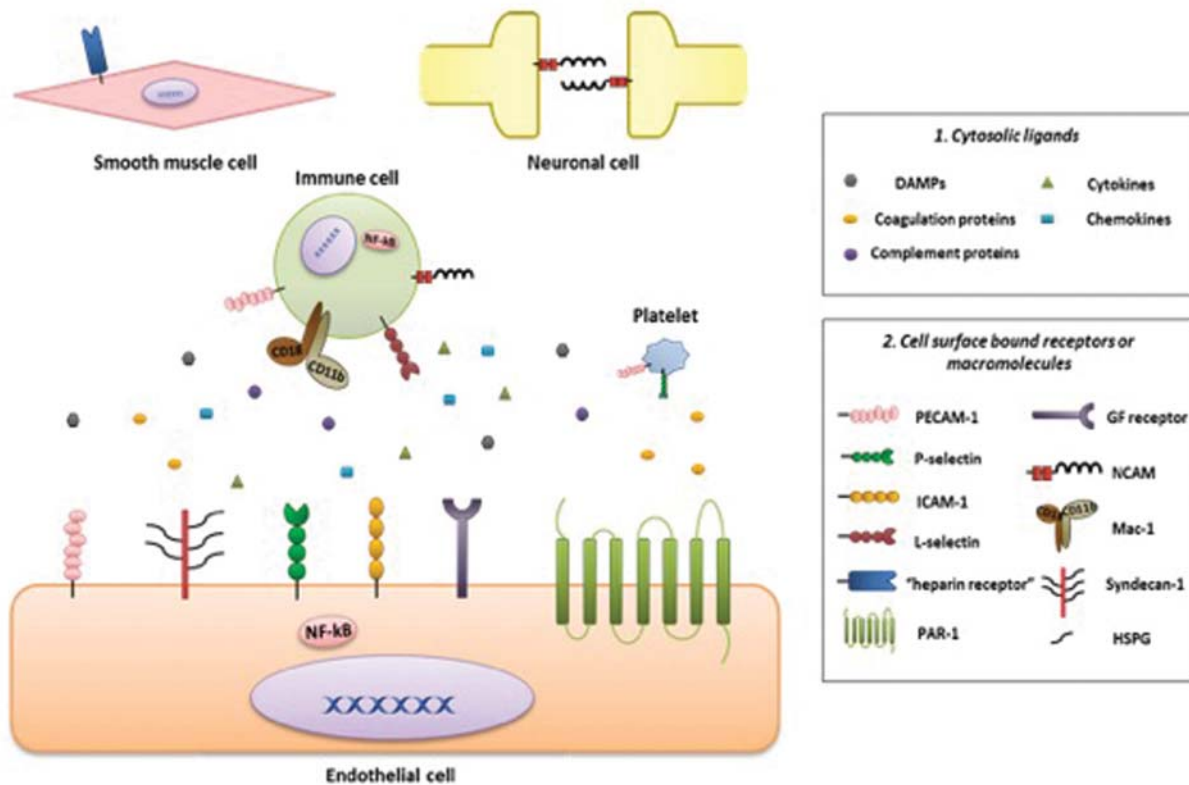


Fig. 3 The anti-inflammatory nature of heparin–ligand interactions. Illustration of the anti-inflammatory polypharmacology of heparin. Besides its use as an anticoagulant, the anti-inflammatory properties of heparin are widely recognized. These properties depend on the overall effect of heparin affecting many different ligands. These ligands can be either found in plasma, are surface-bound, or are present in the intracellular compartment.

In 2004 it was shown that nuclear proteins, although not present in the circulation under normal physiological conditions, may appear in plasma as a result of NETosis.⁹⁸ NETosis is the process in which decondensed chromatin composed of nuclear DNA–histone scaffolds, designated neutrophil extracellular traps (NETs), are formed in response to a trigger, which is commonly a pathogen. Besides the DNA–histone network, NETs contain several granule proteins, oxidases, and elastases.⁹⁸ Histones are known to be cytotoxic when present extracellularly.⁹⁹ In vitro and in vivo experiments by Wildhagen and coworkers have shown that heparin is able to neutralize the cytotoxicity of extracellular histones through a mechanism that is independent of its anticoagulant properties,²⁵ findings that were later confirmed.^{100,101} Furthermore, both heparin and HS were able to prevent accumulation of histones in the lungs of rabbits when administered prior to histone injection.¹⁰² Other granular proteins from activated immune cells such as elastase and cathepsin G have also been shown to be inhibited by heparin.¹⁰³ NETs are a major component of arterial and venous thrombi, as demonstrated by several in vivo models and in patients, whereby LMWH was shown to prevent NETosis and thereby the extent of deep vein thrombosis in mice.¹⁰⁴ This effect could demonstrate the strongest anticoagulant function of heparin.

It is worthwhile noting that heparin accelerates inhibition of thrombin by AT and, thereby, will affect also downstream thrombin targets/substrates, such as the protease activated

receptor-1 (PAR-1).¹⁰⁵ PAR receptors contribute to the proinflammatory response. Consequently, heparin dampens inflammation auxiliary to activated coagulation. An illustration of this intertwining of mechanisms is seen in the fact that in vivo thrombin formation can lead to inflammation, but also vice versa, in which a proinflammatory state may lead to thromboembolic events.¹⁰⁶ In as much atherosclerosis can be seen as an inflammatory condition, it has been known for several decades that heparin use is associated with a lowered risk for atherosclerosis. Support for this association is given by the interplay between hemostatic and atherogenic processes,¹⁰⁷ both of which are directly affected by the anticoagulant properties of heparin but also through the binding of heparin to other plasma and vessel-wall components (see also next paragraph). Collectively these effects result in a state in which the endothelial barrier function is preserved and thrombin generation and platelet adhesion are inhibited. Likewise, studies from the 1960s already showed that lipoprotein lipase activity in the blood is increased in the presence of heparin, resulting in reduced lipoprotein uptake by the vessel wall and an overall improved lipid profile by a lowering of low-density lipoproteins.¹⁰⁸

Heparin-Dependent Modulation of Cell-Surface-Bound Receptors or Macromolecules

Several studies have shown that heparin is able to modify interactions between cells. Heparin-dependent binding to cell adhesion molecules as well as heparin-independent altered

expression of adhesion molecules appears to mediate these effects.^{55,109} Particularly, the inhibition of interactions between endothelial cells and blood cells (e.g., leukocytes, platelets) by heparin results in a more anti-inflammatory state, whereby heparin reduces tumor necrosis factor- α -induced leukocyte rolling *in vivo*.¹¹⁰ This effect is likely caused by the binding of heparin to P-selectin that is present on the surface of activated endothelial cells and activated platelets.¹¹¹ Heparin also binds L-selectin on the surface of leukocytes and has anti-inflammatory activity *in vivo*.¹¹² HS that is present on activated endothelial cells is known to be of importance for the adherence and rolling of leukocytes to the endothelium, and exogenously added heparins can interfere with and prevent this interaction.^{113,114} Interestingly, despite its structural similarity to P- and L-selectin, E-selectin does not bind heparin.⁵⁵ This difference relies on two specific amino acid residues in the EGF-like domain, meaning that E-selectin would be able to bind heparin if these are altered.¹¹⁵ The reported down-regulation of E-selectin and other adhesion molecules in heparin-treated endothelial cells is thought to occur (in part) through NF- κ B inhibition.¹¹⁶ Intercellular adhesion molecule-1 (ICAM-1), a cell adhesion molecule of the immunoglobulin superfamily of proteins which binds strongly to the integrins CD11a/CD18 or CD11b/CD18, is present in the membranes of both activated endothelial cells and leukocytes. ICAM-1 interactions are important for the maintenance of the barrier function of endothelial tissue and for leukocyte endothelial transmigration. Heparin was shown to attenuate ICAM-1-mediated interactions by binding to this adhesion molecule¹¹⁷ and to suppress increased ICAM-1 expression during endothelial cell activation by reducing gene expression.¹¹⁸ Platelet endothelial cell adhesion molecule or PECAM-1, expressed on the surface of platelets and immune cells, which is involved in transmigration of immune cells, is thought to bind to heparin under preferable mild acidosis conditions.^{119,120} Also neuronal cell adhesion molecule (NCAM)¹²¹ and the integrin macrophage-1 antigen (Mac-1)¹²² have been shown to bind heparin. Mac-1 is present on the surface of several immune cells, including neutrophils, macrophages, and natural killer cells. This pattern recognition receptor consists of protein subunits CD11b/CD18 and binds to several ligands including the complement proteins iC3b and C4b and several bacterial surface epitopes. Heparin binding to Mac-1 was shown to inhibit the binding of Mac-1 ligands, resulting in modulation of inflammation and cell proliferation.

Heparin was also shown to influence interaction of viruses with cells and heparin can compete with cell-surface-bound HS for binding to several types of viruses. This attenuates the functional interaction between the virus and its target cells and thus can result in inhibition of infection.^{123–125}

The above-mentioned receptor-mediated anti-inflammatory effects of heparin can be considered as in direct contrast to the effects resulting from the heparin-mediated inhibition of thrombin and FXa. Through the enhancement of inhibition of thrombin/FXa activity *in vivo*, heparin thus has a twofold indirect anti-inflammatory property since it is known to dampen the procoagulant responses of endothelial cells and platelets by limiting their activation by coagulation factors.

In addition it was shown that both heparin and low anticoagulant heparin act in a PAR-1-dependent manner through a mechanism that does not depend on thrombin binding and can thus directly interfere with cell signaling.¹²⁶ The dampening effect of heparin on platelet reactivity appears in contrast to effects recorded for use of dabigatran, an oral anticoagulant (direct oral anticoagulant [DOAC]) for which it was shown that its use in a cohort of atrial fibrillation resulted in enhanced platelet reactivity and increased platelet PAR-1/PAR4 expression.¹²⁷

In addition to endothelial cells, heparin also targets vascular smooth muscle cells (VSMCs) and modulates their biology. Heparin induces switching from VSMC synthetic to contractile phenotype and inhibits VSMC proliferation by interfering with cell-cycle progression.¹²⁸ This property of heparin can be employed in stent technology to prepare coatings that control neointimal cell growth.¹²⁹ The modulation of VSMCs appears to depend neither on the anionic charge¹³⁰ nor on the anticoagulant properties of heparin.¹³¹

Many experimental and clinical studies have illustrated in a more integrative approach the overall beneficial *in vivo* effects of heparin in inflammation. More recent studies confirmed that both UFH and LMWH reduce neutrophil sequestration and lung permeability with concomitant tissue protection in a rat model of lipopolysaccharide (LPS)-induced acute lung injury.^{132,133} The protective effect was likely independent of heparin's anticoagulant properties since heparins devoid of anticoagulant activity reduce neutrophil infiltration and protect lungs in a mouse model of LPS-induced sepsis.²⁵

Finally, heparin acts positively on vascular endothelium by restoring glycocalyx function after inflammation-induced glycocalyx shedding.^{134,135} It is hypothesized that heparin can replenish the cell-surface proteoglycan network by mobilizing an intercellular pool of syndecan-1¹³⁶ and that heparin can take over partly the functions of syndecan-1.¹³⁷

As discussed above heparin can influence heparin-independent intracellular signaling through modulation of receptor-ligand interactions. Heparin was shown to inhibit p38 MAPK and NF- κ B activation in an *in vitro* LPS-induced inflammatory response on endothelial cells¹³⁸ as well as in an *in vivo* experimental model of LPS-induced inflammation.¹³² Likewise, heparin was able to alleviate endothelial barrier dysfunction through regulation of p38 MAPK in an inflammatory model induced by the DAMP high mobility group box 1 (HMGB1).¹³⁹ While some report heparin does not affect the translocation of NF- κ B into the nucleus,¹⁴⁰ other have proposed that it may compete with DNA for NF- κ B binding sites.^{138,141}

Another process in which heparin likely acts through NF- κ B is the decreased degranulation of mast cells and other immune cells after exogenous heparin administration. This decrease subsequently leads to a reduction in lysosomal enzyme activation and reactive oxygen species (ROS) generation.¹⁴² As heparin was found to be an antioxidant, heparin is possibly exerting its effects through NF- κ B given the crosstalk between both ROS and NF- κ B.¹⁴³

An important indication of heparin's anti-inflammatory activities in the clinical setting is provided by recently published meta-analyses interrogating the effect of heparin use in

patients with sepsis.^{144–146} The general outcome of these analyses is that the use of heparin and LMWH results in reduced 28-day mortality in these critically ill patients, despite the recording of an increased number of bleeding events. Septic patients often have a compromised coagulation system. This awareness has prevented a more widespread clinical use of heparin as an anti-inflammatory agent in sepsis.⁹⁷ Since a significant part of heparin's anti-inflammatory actions does not depend on anticoagulant activity, it is hypothesized that heparins with low anticoagulant activity have beneficial therapeutic effects without increasing the risk of bleeding in patients with sepsis. Clinical trials are needed to address this hypothesis.

Adverse Effects of Heparin

The extensive experience that was gained over the many years of heparin use has revealed several side effects of heparins that may limit their use and should always be considered when heparins are being prescribed. A foremost side effect of anticoagulant heparins is the occurrence of bleeding. For several reasons, depending on the clinical context, a dose of heparin may completely disturb the hemostatic balance such that bleeding occurs. Given the relative short plasma half-time of heparin, this effect is relatively transient as compared with bleeding caused by vitamin K antagonist use. Moreover in an emergency situation, heparin can be reversed with protamine, a positively charged protein that binds to the negatively charged heparin, upon which the complex is removed from circulation by the reticuloendothelial system. A further well-described and potentially dangerous side effect is the occurrence of heparin-induced thrombocytopenia (HIT), which is estimated to occur in 0.1 to 5% of patients who receive therapeutic doses of heparin (for a recent review see Sanford et al¹⁴⁷). HIT can be induced by various types of heparins, but the incidence of HIT is lower with the use of more fractionated heparins, as LMWH. Two types of HIT exist, with the first type, which is less significant in clinical practice, being triggered by the exposure of patients to UFH or high heparin doses for the first time. This a nonimmune thrombocytopenic response caused by binding of heparin to platelet-exposed PF4, resulting in ultralarge complexes (ULCs) that trigger platelet activation or via a fibrinogen receptor mediated lowering of platelet counts. HIT type 2, is an immune-mediated response where antibodies are generated against ULCs. This type of HIT causes life-threatening thromboses and thrombocytopenias and is seen usually 5 to 10 days after the onset of heparin therapy. Immediate cessation of heparin administration is essential to allow increase of platelet counts. A recent guideline by the American Society of Hematology recommends the use of a nonheparin anticoagulant (such as argatroban, bivalirudin, danaparoid, fondaparinux, or DOACs) for treatment of acute HIT.¹⁴⁸

Data on other adverse effects of heparin use are more scarce. These include the association of heparin use and osteoporosis¹⁴⁹ and the occurrence of skin lesions.¹⁵⁰ It is generally accepted that long-term use of UFH is associated with a 2.2 to 5% incidence of heparin-induced osteoporotic fracture, an

effect that is far less clear for LMWH. The molecular mechanism through which heparin induces bone loss is not completely understood, but it likely involves the resorption of bone by osteoclasts, an effect that is stimulated by heparin. At the same time osteoblast function is suppressed, together leading to decreased bone mass. Few data are available on the incidence of heparin-induced skin lesions, but a study from 2009¹⁵⁰ estimated that 7.5% of patients receiving subcutaneous heparin developed skin lesions, which makes heparin-induced skin lesions relatively common. In all these patients a delayed-type hypersensitivity reaction was the cause of the lesion which occurred after prolonged heparin use.

Conclusion and Outlook

Discovered about a century ago as a water-soluble anticoagulant obtained from animal tissues, heparin has ever since been used in the clinic because of its anticoagulant properties. Clinical use became widespread after Kakkar showed that systematic administration after surgical interventions dramatically decreased postoperative thrombosis with acceptable bleeding risk. Research over the past decades has unveiled diverse nonanticoagulant properties of heparin that potentially have therapeutic value in a variety of diseases with inflammatory components. UFH is a mixture of polysaccharides with variable saccharide compositions and molecular weights. So far it has not been demonstrated that all the polysaccharides in the mixture contribute to the nonanticoagulant properties. Further research is required to delineate the structure–function relationship of the various nonanticoagulant properties, in stark contrast to the structure–function knowledge available in the anticoagulant properties of heparin. Nevertheless, the anticoagulant property of heparin will impact its use in nonanticoagulant applications by limiting the dose due to bleeding risk. One strategy to exploit fully the nonanticoagulant properties therapeutically encompasses the continued formulation of heparins that have no or limited anticoagulant activity.

Conflict of Interest

C.P.R., H.C.H., and G.A.F.N. are inventors of a patent, owned by the Maastricht University, on the use of nonanticoagulant heparin to treat systemic inflammation and sepsis.

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