Probing Structural Selectivity of Synthetic Heparin Binding to Stabilin Protein Receptors

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Abstract

As one of the most widely used drugs worldwide, heparin is an essential anticoagulant required for surgery, dialysis, treatment of thrombosis, cancer, and general circulatory management. Stabilin-2 is a scavenger clearance receptor with high expression in the sinusoidal endothelium of liver. It is believed that Stabilin-2 is the primary receptor for the clearance of unfractionated and low molecular weight heparins in the liver. Here, we identify the modifications and length of the heparin polymer that are required for binding and endocytosis by both human Stabilin receptors: Stabilin-2 and its homolog Stabilin-1 (also found in liver endothelium). Using enzymatically synthesized 35S-labeled heparan sulfate oligomers, we identified that sulfation of the 3-OH position of N-sulfated glucosamine (GlcNS) is the most beneficial modification for binding and endocytosis via both Stabilin receptors. In addition, our data suggest that a decasaccharide is the minimal size for binding to the Stabilin receptors. These findings define the physical parameters of the heparin structure required for efficient clearance from blood circulation. These results will also aid in the design of synthetic heparins with desired clearance rates.

Keywords: Endocytosis, Endothelial Cell, Heparan Sulfate, Heparin, Receptors, Arixtra, Stabilin, Glycosaminoglycan, Liver

Significance: Customized heparin may have therapeutic applications for obtaining the optimal balance between anticoagulation and clearance.

Background: Stabilin-2 is expressed in the liver endothelium and serves as the primary hepatic clearance receptor in mammals.

Results: Increased sulfation and length of heparin increase affinity for Stabilin binding/endocytosis.

Conclusion: The data demonstrate that 3-O-sulfation is not required, but greatly enhances binding to the Stabilin receptors.

Heparin is an anticoagulant drug that is widely prescribed for procedures in which blood is handled, including surgery, blood/plasma donation, coronary and peripheral vascular procedures, thrombosis, dialysis, and in the washing and preparation of medical equipment (1). The current supply of heparin is derived from the pulp of porcine small intestine and is harvested as unfractionated heparin (UFH) with polymeric chain lengths ranging from 3,000 to 30,000 Da. UFH is depolymerized into smaller sizes with chain lengths ranging from 2000 to 8000 Da to yield the products known as low molecular weight heparin (LMWH). Generally, both UFH and LMWH are fast-acting coagulants with different clearance rates. UFH has a circulatory half-life of 30–60 min, in contrast to LMWH, which has a half-life of 2–6 h (2, 3).

The clearance of heparin is primarily mediated by liver sinusoidal endothelial cells (4). The primary receptor responsible for the systemic clearance of heparin is Stabilin-2/HARE (Stab2)(HARE, hyaluronic acid receptor for endocytosis) (5, 6). Thus, the binding affinity of heparin to Stab2 likely relates to the clearance rate of heparin. Stab2 is presented on the cell surface as two isoforms: 315- and 190-kDa type I receptors. A subset of the 315-kDa receptor undergoes proteolytic cleavage to produce the smaller 190-kDa receptor. Both 315-kDa and 190-kDa Stab2 receptors bind to and participate in systemic clearance of heparin (7, 8). The binding affinity of Stab2 to UFH is higher than LMWH, suggesting that interactions between Stab2 and heparin are dependent on the length of the polymer. Stab2 receptors are also responsible for clearing other glycosaminoglycans, including hyaluronic acid and chondroitin sulfates. It is known that the Stab2 binding domain(s) for heparin are distinct from the Link domain for binding hyaluronic acid and chondroitin sulfates (9).

Stabilin-1 (Stab1) is a Stab2 homologous protein expressed in humans, which has not been reported to bind to heparin previously. The topological organization of Stab1 is similar to Stab2 in that they share 21 EGF/EGF-like domains, seven Fasciclin-1 domains, and one X-link domain with an overall 41% amino acid identity and 56% similarity. Stab1, like Stab2, is expressed in the endothelium of liver, lymph node, and spleen,
but is also uniquely expressed in activated macrophages and continuous endothelial vasculature (10, 11), suggesting that this receptor is immune-responsive. Both Stab1 and Stab2 bind an array of ligands. Stab1 binds to SPARC (secreted protein acidic and rich in cysteine) (12) and lactogen (13); Stab2 binds to collagen propeptides (6) in addition to glycosaminoglycans. The common ligands for both receptors are acetylated LDL (5, 14), GDF-15 (15), phosphatidylserine (16–18), and advanced glycation end products (19).

Heparan sulfate (HS) is a glycosaminoglycan consisting of repeating disaccharide units of glucuronic acid (GlcA) or iduronic acid (IdoA) and glucosamine, with each of the saccharide units capable of carrying sulfo groups. The positions of the sulfo groups, the location of the IdoA residues, and the size of the HS chain determine its biological functions. Heparin is a special form of HS, carrying higher levels of sulfation and IdoA. HS is a highly heterogeneous complex mixture. Its biosynthesis involves the preparation of a polysaccharide backbone and a series of sulfations and epimerization modifications. The enzymes that carry out these modification reactions have been expressed and purified (20). Utilizing recombinant HS biosynthetic enzymes, we demonstrated the synthesis of HS polysaccharides with specific sulfation patterns (21, 22) and HS oligosaccharides with defined structures (23, 24). The synthesized polysaccharides and oligosaccharides act as valuable reagents to probe the structural selectivity of HS in a given biological or biochemical experiment (20, 24).

In this study, we used synthetic HS polysaccharides and oligosaccharides to probe the structural requirements for binding to the Stabilin receptors. We report that 3-O-sulfation has a significant impact on binding/endocytosis. The size of polymer also plays a role in endocytosis. We describe the minimal size and sulfations required for endocytosis in recombinant cell lines and in murine liver endothelium. For the first time, we demonstrate that Stab1 is also responsible for the internalization of heparin in cell culture and in binding assays performed in vitro. Our findings advance the understanding of the mechanism for clearing HS and heparin. Our results also provide a molecular basis for designing heparin anticoagulant drugs with different clearance rates.

**Experimental Procedures**

**Materials, Solutions, and Buffers** — Heparin (or unfractionated heparin) was from Sigma. Fondaparinux (Arixtra®) was purchased from a local pharmacy. Flp-In 293 cells, serum, high glucose Dulbecco’s modified Eagle’s medium (DMEM), Hygromycin B, Zeocin, and glutamine were from Invitrogen. Western blot analysis was completed by either colorimetric or chemiluminescence detection of blotted protein. Anti-V5 antibodies and resins were from Bethyl Laboratories (Montgomery, TX). Other materials, reagents, and kits were obtained as described recently (9). Tris-buffered saline with Tween 20 (TBST) contains 20 mm Tris-HCl, pH 7.0, 150 mm NaCl, and 0.1% Tween 20. TBST/BSA is TBST with 1.0% (w/v) bovine serum albumin (BSA). Phosphate-buffered saline (PBS) contains 137 mm NaCl, 8 mm NaHPO₄, 1.5 mm KH₂PO₄, 2.7 mm KCl, pH 7.2. Hanks’ buffered saline solution contains 5 mm KCl, 0.4 mm KH₂PO₄, 0.8 mm MgSO₄, 137 mm NaCl, 0.3 mm Na₂HPO₄, 5.5 mm glucose, 1.26 mm CaCl₂, 0.5 mm MgCl₂, and 28 mm phenol red; at the time of use, 3.5 g/100 ml of NaHCO₃ was added, and the pH was adjusted to 7.2 with HCl. Endocytosis Medium contains DMEM supplemented with 0.05% BSA.
**Preparation of 35S-labeled HS Constructs** — A total of 27 HS constructs were prepared for this study using a chemo-enzymatic approach published previously (Figure 1 and Table 1) (22, 24). Constructs I through 10 (Figure 1A) are polysaccharide constructs differing in sulfation types and IdooA content, whereas constructs 11 through 23 are oligosaccharide constructs ranging from hepta- to nonadecasaccharides. A representative structure of a decasaccharide (15b) is shown in Figure 1B. For the synthesis of polysaccharide constructs (1 through 10), N-sulfo heparosan was used as a starting material and incubated with the appropriate enzymes and the sulfo donor 3′-phosphoadenosine 5′-phosphosulfate (PAPS) (22). The polysaccharide products were analyzed by disaccharide analysis to confirm the anticipated sulfations (25). To prepare the oligosaccharide constructs (11 through 23), both elongation and modification steps were involved. During the elongation step, a disaccharide starting material (GlC-AnMan) was first elongated to the desirable size with KfiA (N-acetyl glucosaminyl transferase of *Escherichia coli* K5 strain) and pmHS2 (*Pasteurella multocida* heparosan synthase 2) in the presence of UDP-GlcA and UDP-GlcNAc or UDP-GlcNTFA. The elongated products were confirmed by electrospray ionization mass spectrometry (ESI-MS). The oligosaccharides were then converted to N-sulfo oligosaccharides by treating with triethylamine followed by N-sulfotransferase modification. The products were demonstrated to have the anticipated molecular size and purity by ESI-MS. The ESI-MS spectra of the octa- to decasaccharides are shown in Supplemental Figure S1. For those oligosaccharides larger than dodecasaccharides, the ESI-MS spectra are shown in separate publications (23).*

The oligosaccharides were then modified by C5-epimerase, 2-O-sulfotransferase (2-OST), 6-O-sulfotransferase 1 and 3 (6-Ost-1 and 6-Ost-3), and 3-O-sulfotransferase 1 (3-Ost-1). After the modifications, a mixture of oligosaccharides with different levels of sulfation was obtained as determined by DEAE (diethylaminoethyl)-HPLC (Supplemental Figure S2). To introduce a 35S label to the polysaccharides or oligosaccharides, [35S]PAPS replaced unlabeled PAPS. 35S-labeled 3-O-sulfated heparin was prepared by incubating heparin with 3-Ost-5 enzyme and [35S]PAPS, and the product was purified by ESI-MS chromatography.

The procedures for preparing the N-sulfo heparosan, PAPS and [35S]PAPS, UDP-GlcNTFA, and disaccharide (GlC-An-Man) starting materials are described elsewhere (22, 23, 27).

**Expression Plasmids** — The cDNA for human Stab1 (a kind gift of J. Kzhyshkowska, University of Heidelberg) and Stab2/315-HARE were ligated into the multi-cloning site of pSecTag/FRT/V5–6×HIS-TOPO, which provides a secretion signal for the 190-HARE protein (8). Plasmids encoding the desired ectodomain were expressed in *E. coli* and incubated with the appropriate enzymes and the sulfo donor 3′-phosphoadenosine 5′-phosphosulfate (PAPS) (22). The polysaccharide products were analyzed by disaccharide analysis to confirm the anticipated sulfations (25). To prepare the oligosaccharide constructs (11 through 23), both elongation and modification steps were involved. During the elongation step, a disaccharide starting material (GlC-AnMan) was first elongated to the desirable size with KfiA (N-acetyl glucosaminyl transferase of *Escherichia coli* K5 strain) and pmHS2 (*Pasteurella multocida* heparosan synthase 2) in the presence of UDP-GlcA and UDP-GlcNAc or UDP-GlcNTFA. The elongated products were confirmed by electrospray ionization mass spectrometry (ESI-MS). The oligosaccharides were then converted to N-sulfo oligosaccharides by treating with triethylamine followed by N-sulfotransferase modification. The products were demonstrated to have the anticipated molecular size and purity by ESI-MS. The ESI-MS spectra of the octa- to decasaccharides are shown in Supplemental Figure S1. For those oligosaccharides larger than dodecasaccharides, the ESI-MS spectra are shown in separate publications (23).*

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**Endocytosis Assays** — Stably transfected cells expressing Stab1 or Stab2 receptors or only Hygromycin B-resistant (empty vector) were plated in 24-well dishes and grown in DMEM with 8% FBS and 50 μg/ml Hygromycin B for at least 2 days prior to the experiments. The cells were incubated at 37 °C for 3 h with fresh Endocytosis Medium supplemented with labeled [35S]HS constructs (2.0 × 104 cpm/ml). For those experiments utilizing antithrombin III (ATIII), the [35S]HS constructs were preincubated with ATIII (0.2 mg/ml) for 30 min prior to diluting 10-fold in endocytosis medium. Specific binding or endocytosis was assessed in the presence of excess unlabeled heparin (0.1 mg/ml) to determine background count per minute (cpm) values. These values were subtracted from all data points to determine the specific [35S]HS endocytosis. At the termination times, cells were washed three times with ice-cold Hank’s buffered saline solution and lysed in 0.3 n NaOH, and radioactivity and protein content were determined and expressed as cpm/μg of protein ± S.D. 35S radioactivity of all samples was measured by a Beckman-Coulter LS6500 scintillation counter. Non-specific binding/background radiation levels were consistently between 16 and 20 cpm for all experiments.

**Direct Binding Assays** — To assess direct protein-HS binding, ectodomains of each receptor were expressed and secreted in stable cell lines. The ectodomains were immunoprecipitated with a goat anti-V5 resin (Bethyl Laboratories), washed with TBS, and then incubated with 4.0 × 107 cpm of each [35S]HS construct for 1.5 h under rotation. The resin was centrifuged, washed three times with TBS, and then placed in scintillation fluid and quantified by a Beckman-Coulter LS6500 scintillation counter. The amount of protein on the resin was quantified by separation with 5% SDS-PAGE, blotted, and probed with rabbit anti-V5 antibody (Bethyl Laboratories), and images were captured on film.

**Assessment of Liver Clearance** — All animal procedures were approved by the Institutional Animal Care and Use Committee (IACUC) of the University of Nebraska under the Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC) guidelines. BALB/c mice were placed under general anesthesia (isoflurane) on a warming platform during the entire procedure. Once unconsciousness was confirmed, the mice were injected via the lateral tail vein using a 27-gauge × 1/2 inch needle with 0.053 μCi of 35S-labeled HS construct. The radiolabeled material was allowed to circulate for 10 min followed by abdominal exposure and severance of the descending aorta abdominallis for bleed out. The liver was collected, briefly washed to get rid of residual blood, cut into smaller (~0.1 g) pieces, weighed, and then homogenized with a PowerGen 125 (Fisher) tissue homogenizer in 0.75 ml of 1% Nonidet P-40. Homogenized tissue was then spun at 12,000 × g for 2 min to clear out insoluble material, and supernatants were mixed with 4 ml of scintillation fluid and counted.

**Results**

**35S-labeled Heparin Binds Stabilin-1 and Stabilin-2 Receptors** — Cell lines stably expressing Stab1 and empty vector were created as described previously (7, 8). The cell line stably expressing human Stab1 cDNA was also constructed with identical methodology. The expressions of Stab1 and Stab2 are shown by a representative Western blot of the cell lysates.

* Y. Xu, E. H. Pempe, and J. Liu, submitted for publication.
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Figure 2. Internalization of 3-O-sulfated heparin via Stab1 and Stab2. A, cell lysates (20 μg) were separated by 5% SDS-PAGE, blotted to nitrocellulose, and probed with anti-V5 antibody. Lane 1, Stab2/190-HARE; lane 2, Stab2/315-HARE; lane 3, Stab1. B, stable cell lines expressing Stab1 (white bar), both Stab2 isoforms (gray bars), and empty vector (EV, black bar) were incubated with 35S-labeled 3-O-sulfated heparin for 3 h. The dotted line represents nonspecific binding values and is subtracted from the data to determine receptor-specific endocytosis. Endocytosis was evaluated by cpm/μg of cell lysate protein for each cell line, mean ± S.D., n = 3.

We also examined the direct binding between the polysaccharide constructs and different Stabilin receptor ectodomains (Figure 4). The Stabilin receptors were captured from the medium of cells secreting the corresponding receptor (Figure 4A) using anti-V5 antibody coupled to Sepharose resin and exposed to 35S-labeled HS constructs. As anticipated, a similar trend was observed for direct binding, namely that higher sulfation levels for a given construct lead to greater binding to the Stabilin receptors (Figure 4, B–D). In these experiments, the empty vector negative control was used as a baseline for nonspecific polysaccharide binding due to the inherent “stickiness” of the high charge on the HS to the antibody resin. Again, the polysaccharides carrying 3-O-sulfation (constructs 8 and 9) exhibited the highest binding toward the Stabilin receptors. We also observed that the binding levels of 8 and 9 to the 190-HARE and Stab1 receptors were comparable, suggesting that the receptor makes no distinction between IdoA and GlcA (Figure 4, B and D). Although slightly higher binding was observed for construct 9 with 315-HARE (Figure 4C), we cannot conclude that IdoA decreases the binding of HS to the receptor.

Antithrombin III Competes with Stabilins for HS Binding — ATIII is a serine protease inhibitor that regulates blood coagulation by targeting thrombin and factor Xa (29). Inhibition of these targets occurs via the formation of a 1:1 complex between ATIII and HS that contains a specific heparin pentasaccharide sequence for ATIII binding (30, 31). The 3-O-sulfated glucosamine residue within the pentasaccharide is the last modification step during heparin biosynthesis and is carried out by 3-OST-1 (32). Because the 3-O-sulfated polysaccharides exhibited increased Stabilin-mediated endocytosis, we asked whether Stabilin and ATIII bind to the same site in HS. First, we demonstrated the inhibition of 35S-heparin endocytosis with Stab1 cells in the presence of increasing concentrations of ATIII (Figure 5A). We utilized both ATIII-binding polysaccharide constructs, constructs 8 and 9 containing the 3-O-sulfation, and ATIII-nonbinding polysaccharide constructs, constructs 6 and 7. As expected, constructs 6–9 displayed distinct Stabilin-mediated internalization effects in the presence and absence of ATIII. ATIII did not interfere with the endocytosis of constructs 6 and 7 in all three

Stabilin receptors, we chemoenzymatically produced a series of HS polysaccharides with specific modifications ranging from N-sulfation alone on the glucosamine residue (construct 1) to the much more complex sulfation patterns commonly found in heparin and HS (constructs 8 and 9) (Table 1 and Figure 1). We observed that the polysaccharides carrying N-sulfation only or a combination of N-sulfation and a single O-sulfation type display low internalization via the Stabilin receptors (Figure 3, constructs 1–5). Increasing the O-sulfation types (or combining N-sulfation with 2-O- and 6-O-sulfation) elevated the internalization via the Stabilin receptors (Figure 3, constructs 6 and 7). Interestingly, the addition of 3-O-sulfation showed the highest internalization rate (Figure 3, constructs 8 and 9). Given the fact that 3-O-sulfation represents a rare modification in HS (28), our results suggest that Stabilin receptors recognize unique sulfation patterns independent of the charge density. This conclusion was further strengthened by the competitive inhibition using antithrombin (see Figure 5) as well as by employing the oligosaccharide substrates as described below (see Figure 7). Constructs 6 and 8 (with IdoA residues) and 7 and 9 (without IdoA residues) exhibited similar binding to the receptors, suggesting that the presence of IdoA has no significant effect on endocytosis.

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### Table 1. Summary of the polysaccharide and oligosaccharide constructs

<table>
<thead>
<tr>
<th>Constructs</th>
<th>Anticipated disaccharide repeating unit</th>
<th>Size of the construct</th>
<th>Modification steps involved</th>
<th>$^{35}$S labeling site</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>(-GlcA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>2</td>
<td>(-GlcA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST and 6-OST</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>3</td>
<td>(-IdoA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST, C₂-epi, and 6-OST</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>4</td>
<td>(-GlcA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST and 2-OST</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>5</td>
<td>(-IdoA-GlcNAc6S)₂⁻ tech</td>
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<tr>
<td>6</td>
<td>(-GlcA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST, C₂-epi, 2-OST and 6-OST</td>
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</tr>
<tr>
<td>7</td>
<td>(-GlcA-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST, 2-OST and 6-OST</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>8</td>
<td>(-GlcA-GlcNS6S ± 6S ± 3S)⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST, C₂-epi, 2-OST, 6-OST, and 3-OST-1</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>9</td>
<td>Heparin, (-IdoA2S-GlcNAc6S)₂⁻ tech</td>
<td>Polysaccharide, &gt;6,000 Da</td>
<td>NST, 2-OST, 6-OST, and 3-OST-1</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>10</td>
<td>GlcNAc6S-GlcA-GlcNAc6S-GlcA-AnMan</td>
<td>Heptasaccharide (7-mer)</td>
<td>Synthesis started from pure N-sulfo, N-acetylated heptasaccharide</td>
<td>3-O-[³⁵S]sulfation</td>
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<tr>
<td>12a</td>
<td>GlcA-GlcNAc6S-GlcA-GlcNAc6S-GlcA-AnMan</td>
<td>Octasaccharide (8-mer with 3-O-sulfation)</td>
<td>Synthesis started from pure N-sulfo octasaccharide</td>
<td>N$^{35}$S sulfation</td>
</tr>
<tr>
<td>15b</td>
<td>GlcA-GlcNAc6S-GlcA-GlcNAc6S-GlcA-AnMan</td>
<td>Decasaccharide (10-mer with 3-O-sulfation)</td>
<td>Synthesis started from pure N-sulfo decasaccharide</td>
<td>6-O-[³⁵S] sulfation</td>
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<tr>
<td>17b</td>
<td>GlcA-GlcNAc6S-GlcA-GlcNAc6S-GlcA-AnMan</td>
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<td>Synthesis started from pure N-sulfo decasaccharide</td>
<td>6-O-[³⁵S] sulfation</td>
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cell lines (Figure 5, B and C). In contrast, endocytosis of constructs 8 and 9 was significantly decreased, but not entirely inhibited, in all three cell lines (Figure 5, D and E). It is also interesting to note that there is a significant amount of internalization of $^{35}$S-polysaccharide after ATIII inhibition (constructs 8 and 9). The internalization levels that could not be inhibited by ATIII for constructs 8 and 9 were similar to the ones observed for constructs 6 and 7. These data suggest that ATIII and membrane-bound Stabilin receptors bind to the same site in HS for internalization. Our data also suggest that Stabs bind to HS in two modes. One mode is that it binds to the site that also interacts with ATIII, which may be the more significant interaction site; another mode is that it binds to highly negatively charged polysaccharides without structural selectivity.

The Size of the Saccharide Versus Endocytosis in Stab1 and Stab2 Cell Lines — We next investigated the length of heparin required for endocytosis. First, we used heparin oligosaccharides of different lengths ranging from a 10-mer to a 19-mer (constructs 14-23). None of these oligosaccharides were taken up by the empty vector cells including those without IdOA residues (data not shown). Next, we tested all of the constructs 14-23 with both Stab1 and Stab2 (315-HARE) cell lines. As summarized in Figure 6B, both the 7-mer (11) and 8-mer (12) exhibit very little endocytosis, but endocytosis is slightly increased in the 9-mer (13b), especially for the Stab1 cells. The biggest jump in the incidence of endocytosis occurs when one additional sugar is added to make a 10-mer (15b). The amount of internalized oligosaccharides remained steady up to a nonadecasaccharide (19-mer, 23). From these data, the minimal size for significant endocytosis is a 10-mer for both Stab1 and Stab2 receptors. Similar results were observed for the 190-HARE receptor.

We next compared the internalization of the 7-mer (11) and 19-mer (23) for internalization in full-length Stab1 and Stab2 cell lines. We did not observe any internalization of the 7-mer in cells expressing either Stab1 (white bars) or Stab2/315-HARE (gray bars) (Figure 6B). Furthermore, we did not observe a sig-
significant decrease in the internalization of the 19-mer (23) by competition with the synthetic pentasaccharide, Arixtra, at 50-fold molar excess. These results demonstrate that the minimal length of a heparin oligosaccharide cleared by these cell lines is at least 10 sugars in length and that the synthetic pentasaccharide is not cleared by cells expressing the Stabilin receptors.

**3-O-Sulfation Is Required for Decasaccharides to Bind to Stab Receptors and for Endocytosis** — To further establish the role of 3-O-sulfation and the size requirement for the binding to Stab receptors, we utilized a series of oligosaccharides with and without 3-O-sulfation. These experiments also utilized labeled heparin as a positive control (black bars) and a structurally defined heptasaccharide (7-mer, 11) (24) as a negative control. Both the Stab1 (Figure 7A) and the Stab2 (Figure 7B) cells show robust endocytosis with heparin and no endocytosis with 11. The oligosaccharides composed of eight or nine sugars without 3-O-sulfation (12a and 13a) did not show endocytosis in contrast to their 3-O-sulfated counterparts that exhibited marginal to very low endocytosis (12b and 13b, Figure 7, gray bars). This is in contrast with the 3-O-sulfated decasaccharide (15b) and dodecasaccharide (17b), which exhibit robust endocytosis with the dodecasaccharide that is almost comparable with the heparin control (Figure 7, A and B). Furthermore, the deca- and dodecasaccharide without 3-O-sulfation (15a and 17a, white bars) were not internalized near the extent as their sulfated counterparts (gray bars). Taken together, our data are consistent with the conclusions that 3-O-sulfation plays a significant role in binding to Stab receptors in the endocytosis of heparin and that the minimum size for binding to the Stab receptors is a decasaccharide.
Structural Selectivity of Synthetic Heparin Binding to Stabilin Protein Receptors

It is known that UFH is retained by liver to display a fast clearance rate, whereas Fondaparinux (a pentasaccharide) is not and displays a slow clearance rate in vivo (33). Therefore, these size and sulfation parameters should hold true if the Stabilin receptors are responsible for the bulk clearance of systemic heparin. To test this, we individually injected mice via the tail vein with four 35S-labeled constructs: heparin (positive control); 7-mer (21, negative control); the decasaccharide without 3-O-sulfation (12a); and the decasaccharide with 3-O-sulfation (12b) (Figure 8). Mice under general anesthesia were injected with equal amounts of radioactive material that was allowed to circulate; the mice were then bled out, and the livers were collected and processed for scintillation counting. We found that the 7-mer (21, white bar) was not retained in the liver. The decasaccharide with 3-O-sulfation (12b, dark gray bar) was retained in the liver to a similar degree as heparin (black bar). In contrast, the retention of the decasaccharide without 3-O-sulfation (12a) in the liver was significantly reduced. To this end, our data suggest that the 3-O-sulfation contributed to the retention in liver in vivo.

Discussion

Understanding the clearance rate of heparin has significant clinical implications for improving the safety of heparin-based drugs. For surgical applications, a fast-clearing heparin drug is preferred, allowing the anticoagulant effect to rapidly disappear after the operation and thus reducing the risk for bleeding side effects. In contrast, a slow-clearing heparin is more desirable for patients with ongoing use, avoiding repetitive dosing. Indeed, UFH is widely used in surgical procedures and kidney dialysis, whereas LMWH and Fondaparinux are more commonly used as prophylactic agents among high-risk patients prone to thrombosis (34, 35). Although it is known that the size of the heparin chain plays a role in the rate of clearance in patients, the precise structural requirements for regulating clearance remain uncharacterized. Previous studies have demonstrated that a liver endothelial cell receptor, known as Stab, is primarily responsible for heparin clearance (5). By taking advantage of our recent success in synthesizing heparins (22, 24), we investigated the contribution of the sulfation and the size of heparin to the binding to Stab proteins. Our results provide a
This is the first study demonstrating the structural selectivity of heparin binding to Stab1. The Stab1 receptor is expressed in the liver sinusoidal endothelium as well as in alternatively activated macrophages (11) and other physiological niches (38) and may play a role in both systemic and localized clearance of heparinoid molecules. Activated macrophages may modulate the amount of dermatan sulfate in wound fluids that promote FGF-2 activity (39, 40). It is possible that Stab 1 plays a role in regulating the amount of anticoagulant HS in wound fluid.

It should be noted that the rate of endocytosis also depends on the concentration of Stab receptors on the cell surface. The 190-HARE Stab2 isoform always showed the highest increase in endocytosis because these cells produce more receptor per microgram of cell lysate than the other two cell lines. Not surprisingly, the liver endothelial cells exhibit a much higher ratio of 190/315-HARE Stab2 than the recombinant cell lines, which may account for the rapid uptake of heparin within the liver (41). The consistently lower internalization rate of heparin in the Stab1 cells may be reflective of the amount of total surface receptor available to bind and internalize heparin, a lower binding affinity, or the unavailability to bind ligand due to the very short transient time on the cell surface. Others have reported the very short transient time of Stab1 on the cell surface (11, 42). In these experiments, we noticed a discrepancy in binding versus endocytosis. Our in vitro binding assays by immunoprecipitation of the ectodomain of Stab1 revealed that the binding of the HS oligomers was qualitatively about as high as that of the Stab2 ectodomains. This contrasts with the cell-based assays and reveals that a combination of surface availability and rapid turnover, not binding affinity, may be responsible for lower endocytic rates in Stab1 cells. In addition, the expression of Stab receptors can be regulated by other cellular mechanisms. A genetic screen in which human umbilical vein endothelial cells activated with VEGF revealed an increase of Stab1 expression reveals that growth factors and cytokines are able to alter endocytosis profiles of tissues (26).

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References


Probing the structural selectivity of synthetic heparin binding to the Stabilin receptors

(Supplementary Information)

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Suppl. Fig. S1. ESI-MS analysis of pure N-sulfated oligosaccharides. A, octasaccharide, measured MW: 1592.5 ± 0.5. B, nonasaccharide, measured MW: 1833.9 ± 0.8. C, decasaccharide, measured MW: 2009.9 ± 0.7. D, dodecasaccharide, measured MW: 2427.4 ± 0.5.

Suppl. Fig. S2. DEAE-HPLC chromatograms of HS oligosaccharides. A, Octasaccharide (12a); B, Nonasaccharide (13a); C, Decasaccharide without 3-0-sulfation (15a); D, Decasaccharide with 3-0-sulfation (15b); E, Dodecasaccharide without 3-O-sulfation (17a); F, Dodecasaccharide with 3-O-sulfation (17b). Multiple or broad peaks indicates a mixture.
**A.**

C_{46}H_{77}N_{3}O_{50}S_{3} Calculated MW: 1592.3

**B.**

C_{54}H_{88}N_{4}O_{57}S_{4} Calculated MW: 1833.5

**C.**

C_{60}H_{96}N_{4}O_{63}S_{4} Calculated MW: 2009.7

**D.**

C_{72}H_{115}N_{5}O_{76}S_{5} Calculated MW: 2427.0