Evidence for Inhibitory Interaction of Hyaluronan-binding Protein 1 (HABP1/p32/gC1qR) with Streptococcus pneumoniae Hyaluronidase*

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Bacterial hyaluronidase enzymes are the major virulence factors that enable greater microbial ingress by cleaving hyaluronan (HA) polymers present predominantly in extracellular space of vertebrates. Based on the premise that effective inhibitors may bind to and stabilize HA thereby protecting it from degradation, we here investigated inhibitory activity of human hyaluronan-binding protein 1 (HABP1) on bacterial hyaluronidase because it is highly specific to HA and localized on the cell surface. Biochemical characterization revealed that HABP1 is a competitive inhibitor of Streptococcus pneumoniae hyaluronate lyase (SpnHL) with an IC50 value of 22 μM. This is thus the first report of an endogenous protein inhibitor that may be used during natural antibacterial defense. Our findings also support a novel multipronged mechanism for the high efficacy of HABP1-mediated inhibition based on structural modeling of enzyme, substrate, and inhibitor. Evidence from docking simulations and contact interface interactions showed that the inherent charge asymmetry of HABP1 plays a key role in the inhibitory activity. This novel role of HABP1 may pave the way for peptide inhibitors as alternatives to synthetic chemicals in antibacterial research.

Bacterial hyaluronidases have long been known as the “spreading factors” or virulence factors responsible for a wide range of acute and invasive infections in humans causing life-threatening diseases, such as pneumonia, meningitis, and septicemia among many others (1–3). These enzymes directly facilitate the spread of infection by degrading the HA matrix in all vertebrates, acting as a diffusion barrier to the active site of hyaluronidases. Bacterial hyaluronidases, isolated from strains of various microorganisms such as Clostridium, Micrococcus, Streptococcus, or Strep-tomyces, degrade HA at very fast rates via a β-elimination reaction that cleaves the alternate β-1,4-glycosidic linkage between the repeating units, resulting in unsaturated disaccharide products (8, 9). Although the mechanism of hyaluronidase-mediated degradation of HA has been very well elucidated through biochemical as well as structural investigations (10, 11), relatively little is known about the defense mechanisms used by the host during infection or about the role of HA and hyaluronidase in pathophysiological processes. It is believed that the balance of HA biosynthesis and intrinsic hyaluronidase levels can be regulated by invoking specific inhibitors (12–14). As a result, potent and specific inhibitors of bacterial hyaluronate lyase would have enormous therapeutic potential as drugs in the treatment of various bacterial infections.

The search for effective hyaluronidase inhibitors began more than half a century ago (2, 15–18) and continues to this day. Exogenous small molecule inhibitors have been identified, but they have weak potency and low selectivity, and there are many unanswered questions with regard to their mechanism of inhibition and whether they would be active against bacterial hyaluronidases as well. For example, heparins and derivatives of alginic acids (19, 20) have structural similarity to HA, whereas polyphenols, flavone analogues (21), saponins, norlignans, or antiallergic drugs are among other reported inhibitors of hyaluronidases (22, 23). Vitamin C, 1-arginine derivatives, and fatty acids have been reported to inhibit streptococcal hyaluronidases at submillimolar concentrations, but a comparison of percent inhibition and IC50 values between these and previous reports is difficult because of differences in the experimental conditions (24–26). In view of the enormous medical applications of effective inhibitors, a systematic search has continued, using computer-aided drug design strategies. A rational analysis based on structural interactions led to the identification of a hydrophobic derivative of vitamin C as a strong inhibitor of

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3 The abbreviations used are: HA, hyaluronan; HABP1, hyaluronan-binding protein 1; SpnHL, Streptococcus pneumoniae hyaluronate lyase; a.s.a., accessible surface area.
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Streptococcus agalactiae hyaluronidase, but it was weakly active against Streptococcus pneumoniae, a major human Gram-positive pathogen (27). Computer-aided drug design-based strategies mainly explore new leads by virtue of structural similarity to HA or, in some cases, by rational de novo approaches. An alternative strategy may also exist wherein an inhibitor may bind to and stabilize HA thereby protecting it from degradation. Such an alternative is plausible because HA has been known to bind and interact with receptors and a large number of HA-binding proteins. Hyaladherins are a very well characterized and highly selective family of proteins reported to interact non-covalently with HA and have a wide array of functions, including receptor-mediated activation of intracellular signaling and structural aggregations in the extracellular matrix (28–30). We therefore began this work with the surmise that a potent hyaluronidase inhibitor may be supplied endogenously by one of these ubiquitous proteins. HABP1 (synonym human p32 or gC1qR), a member of the multifunctional hyaladherin family, is a 68-kDa protein originally purified and cloned from human fibroblast as a novel HA receptor by our group (31). In this work, HABP1 was tested as an inhibitor of bacterial hyaluronidase because we had already confirmed its localization on the cell surface (32). According to the crystal structure of HABP1, it exists as a homotrimer having N-terminal α-helices and asymmetric charge distribution with structural plasticity (33).

Here we report the role of HABP1 as an inhibitor of S. pneumoniae hyaluronidase (SpnHL) and the elucidation of its competitive nature of inhibition. We also propose a unique multi-pronged mechanism of action based upon evidence from structural modeling of the interactions among substrate, enzyme, and inhibitor. Thus we present biochemical evidence for a novel role of HABP1 in pathophysiological processes as a highly potent inhibitor of SpnHL, offering a variety of avenues for optimization of selectivity and potency.

Experimental Procedures

General Materials and Reagents—Substrate from human umbilical cord HA was purchased from Sigma Co. All other chemicals were of analytical grade and obtained from Sigma.

Purification and Estimation of HABP1—HABP1 (amino acids 74–282) was overexpressed in Escherichia coli and purified following the method described earlier (34) with modifications using the 55–90% ammonium sulfate cut followed by ion exchange chromatography on Mono Q® HR (10/10) column (Amersham Biosciences) using a linear gradient of 0–1 M NaCl in 20 mM Tris, pH 7.2, after washing the column with 150 mM NaCl in 20 mM Tris, pH 7.2. The pooled peak fractions were dialyzed in 150 mM NaCl, 20 mM Tris, pH 7.2, and quantified for further use.

Assay of SpnHL Enzymatic Activity—The activity of SpnHL was determined following the method described earlier (35) and is defined as its ability to break down HA to unsaturated disaccharide units (36). In a 1-ml solution of 0.2 mg/ml HA in 50 mM sodium acetate buffer, 10 mM CaCl₂, pH 6.5, 10 μl of enzyme sample (3 ng/μl, diluted just before taking measurement) was added. The reaction mixture was incubated for 5 min during which the measurements were carried out by monitoring the increase in absorbance at 232 nm at 25 °C in a Cary 100 Bio UV-visible double beam spectrophotometer (Varian Inc.).

Determination of Inhibitory Activity on Hyaluronidase—In vitro enzyme inhibition experiments were performed with varying concentrations of HABP1. The inhibitory effect of HABP1 on S. pneumoniae hyaluronidase was measured by using two different enzymatic inhibition assays. In the first assay 0.2 mg/ml HA was preincubated with varying concentrations (0–100 μM) of HABP1 for 1 h at 4 °C in a 1-ml reaction volume of 50 mM sodium acetate buffer, 10 mM CaCl₂, pH 6.5. The reaction was initiated with 30 ng of enzyme, and activity was measured. In the second assay, 30 ng of SpnHL was preincubated in the above mentioned 1-ml reaction volume of buffer with varying concentrations of HABP1 (0–100 μM) for 1 h at 4 °C. The reaction was initiated with 0.2 mg/ml HA, and enzymatic activity was measured. In both assays, the reaction was initiated exactly at 60 s after the addition of either enzyme or HA, respectively, to the reaction mixture. The progress of the reaction was monitored by detecting the appearance of HA oligosaccharide at 232 nm absorbance.

Enzyme Activity in Absence and Presence of Inhibitor—To study the effect of varying the concentration of substrate in the presence of inhibitor, HA concentration in solution was expressed as moles of hexasaccharide based on a hexasaccharide molecular weight of 1203.9 (37). In the absence of inhibitor, varying concentrations of HA (0.0625–0.750 mM) in 150 mM sodium acetate buffer, 10 mM CaCl₂, pH 6.5, was added to the enzyme (30 ng), and the activity was measured. In the presence of inhibitor, 20 μM HABP1 was preincubated with the enzyme (30 ng) for 1 h at 4 °C in a 1-ml reaction volume of buffer, increasing concentrations of HA (0.0625–0.750 mM) were added, and enzymatic activity was measured exactly 60 s after the HA was added to the reaction mixture. The progress of the reaction was monitored by detecting the appearance of HA oligosaccharide at 232 nm absorbance.

Initial Velocity Measurement and Data Analysis—The reaction progress curves for the enzymatic activity and enzyme inhibition were obtained by measurement of the absorbance at 232 nm due to the unsaturated disaccharide product. The reaction was carried out in a 1-ml solution of 50 mM sodium acetate buffer, 10 mM CaCl₂, pH 6.5, with substrate concentration ranging from 0.2072 to 0.3469 mM HA in the above buffer. The reaction was initiated by the addition of 10 μl of enzyme in the same buffer. The product absorbance was measured every 60 s for 5 min at 25 °C. The reaction progress was linear over the time of measurement within the precision of the instrument. The initial velocity of the reaction was determined from the increase in absorbance over the first 5 min of the reaction. The slope was divided by the molar absorption coefficient for the disaccharide product, 5.5 × 10⁴ mol⁻¹ cm⁻¹ (36). The values were calculated and expressed in μM/min HA degradation by SpnHL in the presence of HABP1.

The data for initial velocity, V̇, from each experiment in which the concentration of substrate, S, was varied, were fit into the Michaelis-Menten equation with a nonlinear regression program (GraphPad Prism, version 5, GraphPad Software Inc., San Diego, CA). For all experiments, goodness-of-fit statistics showed that R² and correlation values were greater than 0.987
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In Vitro Effect of HABP1 on the Activity of SpnHL

The in vitro enzyme inhibition experiments demonstrated that highly purified HABP1 inhibits SpnHL activity in a concentration-dependent manner as shown in Fig. 1. The activity of purified SpnHL gradually decreased in the presence of increasing concentrations of purified HABP1 either when HABP1 was preincubated with HA and the reaction was initiated with SpnHL or when it was preincubated with the enzyme SpnHL, and the reaction was initiated with HA. This can be seen much better in Fig. 1B, which shows a plot of percent inhibition of SpnHL activity versus inhibitor concentration. HABP1 was found to inhibit SpnHL with an IC50 value of 22 μM.
It has been reported that the alternate $\beta-1,4$-glycosidic linkage within HA gets cleaved by SpnHL (37), and we were interested in finding out whether HABP1 mediates SpnHL inhibition by protecting this bond during its interaction with HA. To test this, we structurally modeled the HABP1 and HA interaction. The results showed that binding of HA to HABP1 indeed involves the $\beta-1,4$-glycosidic linkage site on the HA oligomer, explaining the observed reduction in enzyme activity. Docking simulations depicted a steric fit between substrate and inhibitor. Interestingly when the top ranking orientations were analyzed to elucidate the molecular details of binding, the HA hexasaccharide was predicted to bind with an energy of $7.59$ kcal/mol, whereas the tetrasaccharide and disaccharide HA oligomers showed binding energies of $6.56$ and $6.28$ kcal/mol, respectively. These observations are in agreement with previous experimental reports stating that the HA hexasaccharide shows better binding affinity than its shorter oligomers (42). Fig. 2 shows the top ranking confirmation between the hexasaccharide substrate and inhibitor after refinement as described under “Experimental Procedures.” A total of 24 contacts were observed between the sugar and HABP1 of which 17 were from chain A and seven were from chain B. Thus, only two of the three polypeptide chains of HABP1 participate in this interaction, and binding occurs at the junction of these two chains. The hexasaccharide sugar was found to bind with HABP1 strongly via hydrophobic as well as hydrogen bond interactions. Interestingly the two $\beta-1,4$-glycosidic linkages of the HA polymer (one between HA1 and HA2 and the other between HA2 and HA3) are both entirely buried in this complex; they are tethered on both sides by hydrogen bonds as well as hydrophobic interactions. For example, the C-1 and C-4 carbon atoms that mediate the $\beta-1,4$-glycosidic linkages are bound by multiple hydrophobic interactions with HABP1 residues His-75 and Thr-76 for the first linkage and Asp-245, Arg-246, and Gly-247 for the second linkage. Furthermore residue Arg-122 of the HA binding motif plays a key role in the first linkage, making hydrogen bonds with the glucuronate moiety on the sugar preceding the linkage and the secondary alcohol group on the sugar succeeding the linkage. Motif residues Arg-122 and Ala-125 also participate in hydrophobic interactions with the ligand. The binding between HABP1 and the HA hexasaccharide is further strengthened by side chains of Lys-80, Ser-106, and Arg-244 that make hydrogen bonds with the sugar and Met-74, Gly-107, Asp-249, and Thr-251 that are involved in hydrophobic interactions. Atomic details of all these interactions have been provided in Table 1.

Thus the collective strength of a substantial number of non-covalent interactions at the contact interface enhances the binding of HA oligomers to HABP1, thereby protecting HA from degradation. It may be noted that the inhibitor is in the homotrimeric form, and the interaction causes a very small loss of accessible surface area (479 Å²; roughly 2% of the entire molecule) at one of the three interchain junctions (see Fig. 2). Therefore it is plausible that the remaining two interchain junctions of HABP1 could potentially bind more HA molecules, or a much longer HA polymer may utilize all three binding sites at various regions along its length, ensuring protection.

**Competitive Nature of SpnHL Inhibition by HABP1**

To determine the nature of inhibition observed during the preincubation of SpnHL and HABP1 (Fig. 1), enzyme assays were carried out either in the absence or in the presence of purified HABP1 (20 μM). Fig. 3 depicts the plot of substrate concentrations against the enzyme activity in the presence and absence of inhibitor. An inhibition of 20–30% was observed in the presence of the inhibitor. The plot shows that reaction was linear over the time of measurement, and the initial velocity and
TABLE 1
Interactions and surface accessibility at the HABP1-HA hexasaccharide contact interface

For each HA unit, UA and NAc denote β-D-glucuronic acid and N-acetyl-β-D-glucosamine, respectively. Stars (**) mark the C-1 and C-4 atoms that participate in the two crucial β-1,4-glycosidic linkages present in the HA hexasaccharide. Distance, distance in Å between interacting atoms from sugar and HABP1. RSA δ (HA) is the loss in residue surface accessibility (RSA) of HABP1 residues upon complexation with HA measured in Å². "Hyp" denotes a hydrophobic interaction. RSA δ (HL) shows the partial a.s.a. lost from three residues of one chain of the HABP1 trimer during SpnHL inhibition. The last three rows depict residues that lose accessibility but do not directly interact with the sugar. HA1, HA2, and HA3 denote the three hyaluronan units of the hexasaccharide. chn, chain; res, residue; id, identity; na, not applicable.

| HA of HA1 | UA of HA1 | HA Sugar | HABP1 Residues | Distance (Å) | RSA δ (HA) | Interaction | RSA δ (HL) |
|-----------|-----------|----------|----------------|--------------|------------|-------------|------------|
|           |           |          | chn, res id, atom |              |            |             |            |
| C         | C         | glucurionate CO₂ | A 106 SER CB | 3.53 | 31.92 | hyp | none |
| C2        | A         | 251 THR CB | 3.69 | 33.21 | hyp | none |
| C3        | A         | 251 THR CB | 3.986 | " | hyp | none |
| C35       | A         | 249 ASP CG | 3.697 | 38.49 | 2.81 |
| H41       | A         | 249 ASP OD1 | 1.789 | " | H-bond | |
|           | NAc of HA1 |          |                |              |            |             |            |
| 8H2       | A         | CH₂OH group | 122 ARG NH2 | 2.052 | 34.29 | H-bond | none |
| C26       | A         | Acetyl group | 244 ALA C | 3.774 | 25.76 | hyp | 6.72 |
| C27       | A         | CH₂OH group | 107 GLY CA | 3.396 | 21.76 | hyp | none |
| C28       | A         | 122 ARG CZ | 3.282 | 34.29 | hyp | none |
| C29       | A         | 247 GLY CA | 3.171 | " | hyp | none |
| C30       | A         | C3 atom | 247 GLY CA | 3.538 | " | hyp | |
| C31       | A         | 247 GLY CA | 3.802 | " | hyp | none |
| C32       | A         | C1 atom ** | 245 ASP C | 3.648 | 38.68 | 11.45 |
| C32       | A         | " ** | 246 ARG C | 3.61 | na | none |
| C33       | A         | 244 ALA C | 3.07 | 23.42 | hyp | none |
| C34       | A         | 244 ALA CB | 3.626 | 25.76 | 6.72 |
| H38       | A         | NH₂ group | 244 ALA O | 1.283 | " | H-bond | none |
| H52       | A         | 247 GLY N | 1.685 | 23.42 | H-bond | none |
|           | UA of HA2 |          |                |              |            |             |            |
| C38       | A         | 245 ASP C | 3.989 | 38.68 | hyp | 11.45 |
| C38       | A         | 245 ASP CA | 3.804 | " | hyp | none |
| C39       | A         | C4 atom ** | 245 ASP C | 3.642 | " | hyp | |
| C40       | A         | 245 ASP C | 3.666 | " | hyp | |
| O6A       | A         | glucurionate CO₂ | 245 ASP O | 1.644 | " | H-bond | |
| O6A       | "       | " | 106 SER HG | 2.244 | 31.92 | H-bond | none |
| O6A       | "       | " | 122 ARG HHH2 | 1.863 | 34.29 | H-bond | none |
| O6B       | "       | " | 122 ARG HHH1 | 2.127 | " | H-bond | none |
|           | NAc of HA2 |          |                |              |            |             |            |
| C10       | B         | HA2      | 74 MET CG | 3.9 | 39.09 | hyp | none |
| C11       | B         | C1 atom ** | 76 THR CA | 3.858 | 36.01 | hyp | |
| C11       | B         | " ** | 76 THR CG2 | 3.585 | " | hyp | |
| C12       | B         | "       | 76 THR CG2 | 3.724 | " | hyp | |
| C13       | B         | C3 atom | 76 THR CG2 | 3.782 | " | hyp | |
| C7        | B         | Acetyl group | 76 THR CA | 3.704 | " | hyp | |
| C7        | B         | "       | 76 THR CG2 | 3.903 | " | hyp | |
| C8        | B         | CH₂OH group | 74 MET CG | 3.318 | 39.09 | hyp | |
| C9        | B         | 74 MET CB | 3.643 | 39.09 | hyp | |
| C9        | B         | 74 MET CG | 3.102 | " | hyp | |

|           | UA of HA3 |          |                |              |            |             |            |
| C14       | B         | His      | 75 HIS C | 3.875 | 49 | hyp | none |
| C23       | B         | His      | 75 HIS C | 3.997 | " | hyp | |
| C23       | B         | His      | 75 HIS CB | 3.618 | " | hyp | |
| C4        | B         | Acetyl group | 76 THR C | 3.904 | 36.01 | hyp | |
| C4        | B         | "       | 76 THR CA | 3.944 | " | hyp | |
| C6        | B         | C4 atom ** | 75 HIS C | 3.162 | 49 | hyp | |
| C6        | B         | " ** | 76 THR CA | 3.723 | 36.01 | hyp | |
| H4        | B         | OH group | 75 HIS H | 2.182 | " | H-bond | |
| O6        | B         | "       | 75 HIS H | 1.488 | 49 | hyp | |
| na        | B         | "       | 77 ASP na | 4.389 | " | hyp | |
| na        | A         | "       | 119 THR na | 4.389 | " | hyp | |
| na        | A         | "       | 124 VAL na | 4.43 | " | hyp | |
other parameters were calculated as described under “Experimental Procedures.” The $K_m$ was calculated to be 0.2765 and 0.41129 mM in the absence and presence of the inhibitor, respectively. A Lineweaver-Burke plot (double reciprocal) of the data obtained was indicative of competitive inhibition. The value for $K_i$ was obtained from plots of $1/V$ versus $1/[S]$. It was found that HABP1 inhibits the enzyme at a $K_i$ value of 34.24 μM.

These results confirmed that the inhibition observed in Fig. 1 is competitive in nature.

**Structural Modeling: Inhibition of Hyaluronidase Mediated by HABP1**

To comprehend the molecular details and structural implications of the observed competitive inhibition, HABP1 was docked with SpnHL as described under “Experimental Procedures.” Analysis of the top ranking docking complex confirmed that HABP1 binds to the enzyme in the same region where substrate HA has been reported to bind (37); the data thus support the competitive inhibition observed in the experimental assays.

Fig. 4A depicts the overall mode of binding with the inset (Fig. 4B) showing the molecular details of the interaction. An accessible surface area (a.s.a.) of $\sim 2938 \text{ Å}^2$ (roughly 11% of the total a.s.a.) is buried on the negatively charged face of HABP1 trimer upon binding to SpnHL, occluding an a.s.a. of $5489 \text{ Å}^2$ (roughly 20% of total a.s.a.) from the enzyme around its active site cleft. The majority of interactions at the contact interface emerge from the juxtaposition of these two large and oppositely charged surfaces. All three monomers of the inhibitor participate in binding. The refined model of binding revealed a total of 207 contacts between the two structures mediated by 49 residues from HABP1 and 81 residues from SpnHL. Of these, 25 are
ionic interactions. The inhibitory activity of HABP1 is evidently due to the combined effect of the considerable number of non-covalent interactions it forms with catalytically essential residues in the active site of the enzyme.

As can be seen in Fig. 4, the binding potency of the inhibitor is a virtue of electrostatic, hydrophobic, and hydrogen bonding interactions all along the contact interface mainly in the catalytic pocket region and a few involving the α-domain of the enzyme. Several interesting features of the complex become evident because the orientation of inhibitor obstructs the region on the enzyme where substrate must enter and bind, thereby rendering the cleft inaccessible. The majority of interactions at the interface are electrostatic in nature. It may be noted that the enzyme active site, which is in the form of a cleft between the two structural domains of SpnHL, has a highly positive center, and the negatively charged projection of HABP1 plays a key role in facilitating the attachment, thereby effecting the high charge complementarity at the interface. The maximum ionic contacts are observed in the product release region, which has previously been shown to be an important target site for hyaluronidase inhibitors (27). Small molecule-based inhibitor design strategies also target specific residues within the active site that are known to be crucial for catalysis. Targeting of these sites in the enzyme has been reported to add to the potency of inhibition (43). These reports add strength to the notion that the residues of the HA binding motif (of HABP1) are not involved in any of its interactions with SpnHL. Of the three potential HA binding sites on a given HABP1 trimer, two are completely accessible during the inhibitory interaction. The third binding site shows a small overlap as three of the 24 HABP1 residues required for sugar binding lose partial side chain accessibility during the inhibitory interaction. These three residues are shown in Table 1. A perusal of this partially buried region reveals that the lost a.s.a. is relatively insignificant (roughly 6%) as a result of which all three binding sites of HABP1 retain sufficient accessibility when it mediates the inhibition of SpnHL. The significance of this observation for the mechanism of action is discussed below.

**DISCUSSION**

In the present study, we have characterized human HABP1 as the only known endogenous protein inhibitor of SpnHL, thus implying indirectly its functional role in defense against bacterial invasion. The mechanistic/structural aspects of this inhibition have been elucidated with the help of docking simulations. HABP1 with an IC_{50} value of 22 μM is one of the most potent inhibitors of SpnHL reported to date. Ascorbic acid 6-hexadecanoate has previously been reported as an inhibitor of *S. agalactiae* hyaluronate lyase with an IC_{50} of 4 μM, but it showed much weaker inhibition of SpnHL with an IC_{50} of 100 μM (27). We have analyzed enzyme substrate interaction in context with enzyme-inhibitor and substrate-inhibitor interactions to elucidate the structural details of the observed inhibition.

Preincubation of HABP1 and HA was able to inhibit the enzyme activity, and there was no further degradation of HA upon addition of SpnHL to the reaction solution. It was observed that HA degradation and product release are totally dependent on the ratio of HABP1 and HA molecules present in the reaction solution. With the increase in HABP1 concentration in the reaction solution, a decrease in enzymatic activity was observed, whereas upon increasing the concentration of HA, the enzymatic activity was retained. For a given amount of HA, however, low HABP1 was not able to block enzyme or sequester substrate, but with high concentration of HABP1, 95% inhibition was observed. Furthermore there was no change in enzymatic activity when the concentration of HA was kept equal to that of HABP1. It is therefore plausible that during the preincubation binding occurred between HABP1 and its ligand of the inhibitor. For example, arginines 462 and 466 of this patch lose 27 and 47 Å² of their residue surface accessibility, respectively, to ionic interactions with glutamates 198 and 200 of HABP1, amounting to a 77.4% loss of accessibility. Other residues comprising the positive patch, namely Lys-250, Arg-300, Arg-355, and Arg-480, also get buried during the inhibitory interaction, resulting in an overall loss of 71% surface accessible area for this patch. The negative patch comprising residues Glu-388, Asp-398, and Thr-400 has been implicated in the product release process of SpnHL. The above mentioned interactions involving the positive patch of the enzyme also lead to the occlusion of the residues in the negative patch because the two patches lie in close vicinity of each other. In particular, Glu-388 was observed to lose 12.5-Å² (roughly 26%) surface area during the interaction with HABP1.

It may be noted that the residues of the HA binding motif (of HABP1) are not involved in any of its interactions with SpnHL. Of the three potential HA binding sites on a given HABP1 trimer, two are completely accessible during the inhibitory interaction. The third binding site shows a small overlap as three of the 24 HABP1 residues required for sugar binding lose partial side chain accessibility during the inhibitory interaction. These three residues are shown in Table 1. A perusal of this partially buried region reveals that the lost a.s.a. is relatively insignificant (roughly 6%) as a result of which all three binding sites of HABP1 retain sufficient accessibility when it mediates the inhibition of SpnHL. The significance of this observation for the mechanism of action is discussed below.

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HA, thereby protecting the latter from the enzyme. This deduction was structurally supported by docking experiments between HABP1 and HA that revealed a tight binding with the β-1,4-glycosidic linkage of HA via hydrogen bonding as well as hydrophobic interactions (shown in Fig. 2 and Table 1).

Having thus confirmed the protection of HA by HABP1, we tested whether this was the exclusive mode of inhibition used by the inhibitor. As shown in Fig. 1A, the inhibition of enzyme observed upon preincubating substrate and inhibitor was found to be comparable to the inhibition observed upon preincubating the enzyme and inhibitor. Subsequently assays were carried out to determine the nature of this inhibition. As shown in Fig. 3, Lineweaver-Burke plot analysis confirmed that HABP1 inhibits SpnHL activity in a competitive manner. Docking experiments between SpnHL and HABP1 further provided the molecular details of the contact interface between HABP1 and the active site cleft of SpnHL (Fig. 4). It was also observed that HABP1 uses only the negative side/face of its trimeric (roughly flat) structure during the inhibition of SpnHL. These observations were taken together to build a novel mechanism of HABP1-mediated inhibition of SpnHL.

A Novel Mechanism for HABP1-mediated Inhibition of SpnHL.—The fascinating aspect of our study is that the sugar is capable of binding to both enzyme and inhibitor, and in both cases, it is the β-1,4-glycosidic linkage of the sugar, which is buried in the receptors. Rationalization of the competitive nature of inhibition has further revealed that the enzyme SpnHL binds to the substrate and inhibitor in the same region, namely the active site cleft. It is remarkable that the doughnut-shaped HABP1 acting as an inhibitor uses only its negative face/side for blocking the enzyme, whereas the opposite side/face is used for binding to the relatively negatively charged HA. Evidently the inhibitor uses two distinct and spatially separate sites for binding to the enzyme and the substrate. Based on these observations from experiments as well as computational simulations, we hypothesize that HABP1 may use a novel and unique multipronged mechanism/model for lowering hyaluronidase activity. According to this model, on one hand, HABP1 obstructs the active site of the enzyme to inhibit its activity, thereby following the nature of competitive inhibition. On the other hand, it uses its large free side/face to bind and sequester up to three additional HA polymers from the matrix, thereby protecting HA from SpnHL-mediated degradation during infection. This model has been structurally depicted in Fig. 5. Although one of the three inter chain junctions (or HA binding sites) shows a slight loss of residue surface area during the inhibitory interaction, 21 of 24 residues depicted in Table 1 remain fully accessible and free to bind HA oligomers. Alternatively a single long polymer of HA may bind to HABP1 using all three of its binding sites to circumvent degradation.

Biological Implications—Our findings reveal that HABP1 interacts with and blocks a large number of crucial/conserved groups of residues in the active site cleft of SpnHL. A majority of bacterial hyaluronate lyases are known to have a high degree of sequence conservation in the cleft region, and consequently, we speculate that they may also be inhibited by HABP1 in a similar manner. In contrast, mammalian and bacteriophage hyaluronidase enzymes have very different sequence as well as structural architectures and are therefore not likely to be inhibited by HABP1. It may be conjectured that this strategy of selective defense against bacterial hyaluronidases may have evolved in mammalian hosts to protect their own hyaluronidases from inhibition. It is encouraging to note that there have been reports of up-regulation and involvement of HABP1 in inflammation, apoptosis, and bacterial infections like Listeria (44,45). Based on the crystal structure it has been suggested that terminal α-helices of HABP1 are critical for maintaining trimeric assembly as well as protein-protein interactions (33,46). This is also evident from the Trp-Glu signature motif at the N-terminal region of HABP1 that is homologous to the WD-40 family of proteins that play predominant roles in interprotein interactions (47). A few viral proteins and bacterial proteins have already been reported to interact with HABP1 (48). However, although its multifunctional nature has been characterized, its role in hyaluronan protection has never been explored before. Apart from the ability to competitively inhibit bacterial hyaluronidases, HABP1 has the added advantage of being able to bind and sequester multiple HA oligomers by virtue of its unique structural asymmetry. Our observations thus also explain the inherent charge asymmetry of the HABP1 structure that had not been accounted for until now. It may be speculated that, at least to some extent, this characteristic may be used to
Inhibition of SpnHL by HABP1

In conclusion, the identification of HABP1 as a competitive inhibitor of SpnHL constitutes a new line of defense against bacterial invasion and offers novel opportunities for therapeutic intervention, making it a highly suitable candidate to be investigated further for optimization of potency and specificity against bacterial virulence factors. Furthermore based on the insights obtained in this study, it will be possible to design short inhibitory peptides for binding and protecting HA polymers. Efforts in this direction present an alternative to synthetic chemicals and are currently underway in our laboratory.

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protect the host HA matrix from endogenous hyaluronidas as well.

Competitive inhibition conventionally refers to structural analogy between substrate and inhibitor leading to competition of both for the active site of the enzyme. It must be noted that although we demonstrate the phenomenon of competitive inhibition here the inhibitor is structurally distinct from the substrate, and there is also a simultaneous blocking interaction of HABP1 with its substrate HA. Such a dual mode of inhibition is a unique phenomenon by itself, not previously demonstrated by any other enzyme inhibitor. It is true that free unbridled hyaluronidase activity would create much havoc, and nature must have created a great number of mechanisms for keeping these enzymes in check, mechanisms that must function at multiple levels. We believe that the data presented in this work may describe how remarkable such checking mechanisms can be and that they may indeed function at multiple levels simultaneously.

In conclusion, the identification of HABP1 as a competitive inhibitor of SpnHL constitutes a new line of defense against bacterial invasion and offers novel opportunities for therapeutic intervention, making it a highly suitable candidate to be investigated further for optimization of potency and specificity against bacterial virulence factors. Furthermore based on the insights obtained in this study, it will be possible to design short inhibitory peptides for binding and protecting HA polymers. Efforts in this direction present an alternative to synthetic chemicals and are currently underway in our laboratory.