Glycobiology and Extracellular Matrices: The Solution Structure of Heparan Sulfate Differs from That of Heparin: IMPLICATIONS FOR FUNCTION

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doi: 10.1074/jbc.M113.492223 originally published online August 6, 2013

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The Solution Structure of Heparan Sulfate Differs from That of Heparin

IMPLICATIONS FOR FUNCTION*

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Background: The polysaccharide heparan sulfate (HS) exhibits key physiological roles.

Results: Analytical ultracentrifugation and x-ray scattering revealed extended but bent HS solution structures.

Conclusion: Scattering fits resulted in molecular models for HS in solution with glycosidic angles in good accord with a HS crystal structure.

Significance: These bent HS models clarify how HS interacts with its ligands.

The highly sulfated polysaccharides heparin and heparan sulfate (HS) play key roles in the regulation of physiological and pathophysiological processes. Despite its importance, no molecular structures of free HS have been reported up to now. By combining analytical ultracentrifugation, small angle x-ray scattering, and constrained scattering modeling recently used for heparin, we have analyzed the solution structures for eight purified HS fragments dp6–dp24 corresponding to the predominantly unsulfated GlcA-GlcNAc domains of heparan sulfate. Unlike heparin, the sedimentation coefficient s20,w of HS dp6–dp24 showed a small rotor speed dependence, where similar s20,w values of 0.82–1.26 S (absorbance optics) and 1.05–1.34 S (interference optics) were determined. The corresponding x-ray scattering measurements of HS dp6–dp24 gave radii of gyration Rg values from 1.03 to 2.82 nm, cross-sectional radii of gyration Rs values from 0.15 to 0.65 nm, and maximum lengths L from 3.0 to 10.0 nm. These data showed that HS has a longer and more bent structure than heparin. Constrained scattering modeling starting from 5,000 to 12,000 conformationally randomized HS structures gave best fit dp6–dp24 molecular structures that were longer and more bent than their equivalents in heparin. Alternative fits were obtained for HS dp18 and dp24, indicating their higher bending and flexibility. We conclude that HS displays bent conformations that are significantly distinct from that for heparin. The difference is attributed to the different predominant monosaccharide sequence and reduced sulfation of HS, indicating that HS may interact differently with proteins compared with heparin.

Heparan sulfate (HS) is a sulfated glycosaminoglycan that is found extensively on animal cell surfaces and other extracellular surfaces (1, 2). HS has key roles in biological recognition processes at the cell-tissue-organ interface through its interactions with a wide range of proteins (3, 4). Specific interactions involving HS include roles in cell growth and development (5), cell adhesion (6), inflammation and wound healing (7), angiogenesis and cancer (8–10), viral invasion (11, 12), and anticoagulation (13). The breadth of these HS-protein interactions offers potential strategies for therapeutic intervention at the cell-tissue-organ interface.

HS is a sulfated polysaccharide composed of uronic acid and D-glucosamine residue pairs linked by (1→4)-glycosidic bonds (Fig. 1) (14, 15). The uronic acid residue is either unmodified β-D-glucuronic acid (β-GlcA), alternating with N-acetylated glucosamine (Fig. 1A), or α-L-iduronic acid (α-IdoA), often 2-O-sulfated, alternating with N-sulfated glucosamine (GlcNS) (Fig. 1B). In the latter, sulfation often occurs at C6 and rarely also at C3 (16, 17). HS has a distinct domain organization that is comprised of short S domains (IdoA2S and GlcNS residues), long N domains with GlcA and GlcNAC residues, and mixed domain regions at the junctions between the S domains and NA domains (15, 16). The S domains and mixed domain regions are termed the hypervariable regions that result in different functional characteristics for HS from different cell types (16).

Three-dimensional structural studies of HS are required to complete an understanding of the physiological significance of HS-protein interactions. Many structural studies have been carried out for heparin, which is an analog for HS but possesses a higher degree of sulfation, being predominantly S region-like in sequence, and for at least 19 heparin-protein co-crystal complexes. This abundance results because of the ease with which heparin is obtained and its strong binding to many of the cell surface proteins whose physiological ligand is HS. An NMR

* This work was supported by the Higher Education Commission of Pakistan, the Biotechnology and Biological Sciences Research Council, the Henry Smith Charity, and the Mercer Fund of the Fight for Sight Charity.
† Author’s Choice—Final version full access.
‡ This article contains supplemental Figs. S1 and S2 and PDB models.
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(A) GlcA-(1,4)-GlcNAc

(B) IdoA-(1,4)-GlcNS

FIGURE 1. Chemical structures of the two disaccharide repeats of HS and heparin. A, the major repeating disaccharide unit of HS (glucuronic acid → N-acetylgalactosamine). The NH-CH₃ group in the second ring is replaced by NH₃SO₃⁻ in 50% of this structure. The resulting molecular mass of this averaged disaccharide is 483 Da. B, the minor repeating unit of HS, which is the major repeating disaccharide unit in 50% of heparin (iduronic acid-2-sulfate → glucosamine-2,6-disulfate). For comparison with this study, heparin is considered to be 50% in the trisulfate form as shown and 50% in a disulfate form where a sulfate group is lost. The resulting molecular mass of this averaged disaccharide is 628 Da.

structure is known for heparin (18). Solution structures are known for six purified fragments dp6-dp36 of heparin from constrained scattering modeling; these forms were shown to be similar in conformation to heparin when observed in heparin-protein crystal structures (19). In distinction, up to now, no molecular structures for free HS are known, and only one crystal structure at 0.21-nm resolution for a dp4 HS oligosaccharide complexed with heparinase II is available (20).

Given the importance of understanding the HS solution structure, we have used a multidisciplinary approach to determine molecular structures for HS based on the combination of three methods, namely analytical ultracentrifugation, small angle x-ray scattering, and constrained scattering modeling (21, 22). This approach is well established for solution structure determinations of large multidomain complement and antibody proteins and was recently applied to small heparin oligosaccharide fragments (19, 23). Here, we apply this approach for the second time for oligosaccharide solution structures, this time for eight HS fragments ranging in sizes from dp6 to dp24, thus permitting detailed comparisons with heparin. The HS fragments exhibited solution structures that were distinct from those of the heparin fragments. In particular, their overall lengths are longer compared with heparin, and their structures display a greater degree of bending with increase in size compared with heparin. Our results are attributed to the difference in monosaccharide sequence between HS and heparin fragments, combined with a much reduced degree of sulfation in the HS fragments, which possessed greater structural flexibility than heparin. These results provided new insight on the potential binding modes of HS to proteins.

Following publication of our original 2011 study, we regretfully discovered an error in the anomeric configuration of our heparan sulfate structural models. This present study supersedes the 2011 study, which has been withdrawn.

EXPERIMENTAL PROCEDURES

Purification of HS Fragments—HS oligosaccharide fragments were prepared according to a similar method to that previously used for heparin oligosacharides (19, 24–26). Exhaustive heparinase digestion was used to minimize the content of fully sulfated sequences. Approximately 100 mg of HS (prepared from a crude glycosaminoglycan mixture, the kind gift of Laboratori Derivati Organici, Italy: a mixture of HS-I and HS-II as described in Ref. 27) was weighed out and dissolved in phosphate buffer, pH 7. An aliquot of 200 μl of heparinase I stock solution was added and left to digest at room temperature for at least 2 h, long enough for the reaction to run to completion. The reaction mixture was evaporated to dryness, using a rotary evaporator at 50 °C.

To isolate the HS fragments, the dried digest was dissolved in 1.5 ml of 2% ammonium bicarbonate solution and filtered through a 0.45-micron syringe filter before injection onto the preparative gel filtration column. The filtered digested HS was then applied to a preparative gel permeation chromatography column (100 × 1.6 cm; packed with Biogel P10 (Bio-Rad). The HS fragments were eluted using 2% ammonium bicarbonate at a flow rate of 0.2 ml/min in 2-ml fractions. The absorbance of the fractions was measured at 234 nm, and the top fractions corresponding to each individual resolved peak were pooled. The HS oligosaccharides larger than dp12 were not completely resolved (Fig. 2). The pooled fractions were evaporated under reduced pressure and lyophilized before assessment of their sizes by analytical gel permeation chromatography (25). Like heparin, gel permeation chromatography was carried out using two columns (TSK G3000 SW-XL, 30 cm; TSK G2000 SW-XL, 30 cm; Anachem) connected in series. The eluant was 0.1 M ammonium acetate solution at a flow rate of 0.5 ml/min, and HS was detected with a refractive index detector (RI-1530; Jasco). The chromatography system was calibrated using the First International Reference Reagent Low Molecular Weight Heparin for Molecular Weight Calibration (NIBSC 90/686). HS quantization was achieved by integration of the area under each refractive index peak and comparison with a standard curve prepared using known concentrations of low molecular weight
heparin. An absorption coefficient of 5500 M⁻¹ cm⁻¹ at 232 nm was used for HS experiments (28).

**PAGE of HS Fragments**—The HS fragments were analyzed by PAGE to determine the level of purity of each one according to a previously described method (29). Each HS fragment (5 μg) was mixed with 20% glycerol up to a maximum volume of 10–15 μl and then loaded into separate wells (Fig. 2, inset). Phenol red in 20% glycerol in a maximum volume of 10 μl was also applied to a separate well as a marker. Initially samples were run through a stacking gel (5% acrylamide, 0.5% bisacrylamide) at 150 V for 20–30 min until the phenol red started to enter the resolving gel. In the resolving gel (25% acrylamide, 1% bisacrylamide), samples were run at a constant current of 18 mA until the phenol red reached the bottom of the gel. The discontinuous buffer system of Laemmli (30) consisted of 0.125 M Tris/HCl, pH 6.8, in the stacking gel and 0.375 M Tris/HCl, pH 8.8, in the resolving gel. The gel running buffer was 25 mM Tris, 0.192 mM glycine, pH 8.3. The gel was stained with 0.08% aqueous Azure A for 10 min to visualize HS bands. The gel was then destained in water to remove excess dye and clear the gel background.

**Analytical Ultracentrifugation of HS Fragments**—Sedimentation velocity data for eight HS fragments (dp6, dp8, dp10, dp12, dp14, dp16, dp18, and dp24) were obtained on two Beckman XL-I analytical ultracentrifuges (Beckman-Coulter Inc., Palo Alto, CA) using both absorbance and interference optics. Experiments with the dp6 – dp24 fragments were performed at concentrations of 0.5 mg/ml in 10 mM HEPES and 137 mM NaCl (pH 7.4). The buffer density was measured at 20 °C using an Anton-Paar DMA5000 density meter to be 1.00480 g/ml. A partial specific volume of 0.467 ml/g determined for heparin (31) was used for HS. An alternative higher value of 0.55 ml/g for HS has been reported elsewhere and was used for data processing only when required to confirm that the partial specific volume has no effect on the outcome of this study (32). Analytical ultracentrifugation runs were carried out in an eight hole AnTi50 rotor with standard double-sector cells with column heights of 12 mm at 20 °C using absorbance optics at 234 nm and interference optics. Sedimentation velocity data were collected at 40,000, 50,000, and 60,000 rpm using absorbance optics and interference optics. The continuous c(s) analysis method was used to determine the sedimentation coefficients s₂₀,w of the eight HS fragments using SEDFIT software (version 9.4) (33, 34). The c(s) analysis directly fits the experimental sedimentation boundaries using the Lamm equation, the algorithm for which assumes that all species have the same frictional ratio f/f₀ in each fit. The final SEDFIT analyses used a fixed resolution of 200, and optimized the c(s) fit by floating the meniscus and cell bottom when required and holding the f/f₀ value, base line, and cell bottom fixed until the overall root mean square deviations and visual appearance of the fits were satisfactory (Fig. 3). The individual f/f₀ values calculated previously for the heparin fragments were used for the equivalent HS fragments (19).

**Synchrotron X-ray Scattering of HS Fragments**—X-ray solution scattering of the above eight HS fragments dp6 – dp24 were performed on the Beamline ID02 at the European Synchrotron Radiation Facility at Grenoble, France, in two sessions with a...
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ring energy of 6.0 GeV (35). In the first session, data were collected for six HS fragments in 16-bunch mode using beam currents from 63 to 89 mA. In the second session, data were collected for all eight HS fragments in 16-bunch mode using beam currents from 65 to 78 mA. Data were acquired using an improved fiber optically coupled high sensitivity and dynamic range CCD detector (FReLoN) with a smaller beamstop. The sample to detector distance was 3.0 m. Experiments used the same HS concentrations of 0.5 mg/ml and buffers used in the sedimentation velocity experiments. For each HS fragment, the samples were measured in a flow cell, which moved the sample continuously during beam exposure in 10 time frames with different exposure times of 0.1, 0.25, 0.5, and 1 s to check for the absence of radiation damage effects. This exposure was optimized using on-line checks for the absence of radiation damage to show that this was not detectable.

Guinier analyses give the radius of gyration, \( R_G \), which measures the degree of structural elongation in solution if the internal inhomogeneity of scattering within the macromolecules has no effect. Guinier plots at low Q values (where \( Q = 4 \pi \sin \theta / \lambda \), \( 2\theta \) is the scattering angle, and \( \lambda \) is the wavelength) gives the \( R_G \) and the forward scattering at zero angle \( I(0) \) (36).

\[
\ln I(Q) = \ln I(0) - R_G^2 Q^2 / 3
\]  
(Eq. 1)

This expression is valid in a \( Q-R_G \) range up to 1.5. If the structure is elongated (i.e., rod-shaped), the radius of gyration of the cross-sectional structure \( R_{XS} \) and the mean cross-sectional intensity at zero angle \( I(Q)Q_{Q \rightarrow 0} \) parameters are obtained from fits in a higher \( Q \) range.

\[
\ln[I(Q) \cdot Q] = \ln[I(0) \cdot Q_{Q \rightarrow 0} - R_{gX}^2 Q^2 / 2]
\]  
(Eq. 2)

The \( R_G \) and \( R_{XS} \) analyses were performed using an interactive PERL script program SCTPL73 on Silicon Graphics OCTANE workstations. Indirect Fourier transformation of the full scattering curve \( I(Q) \) in reciprocal space gives the distance distribution function \( P(r) \) in real space. This yields the maximum dimension of the macromolecule \( L \) and its most commonly occurring distance vector \( M \) in real space.

\[
P(r) = \frac{1}{2\pi} \int_0^\infty I(Q)Qr \sin(Qr) dQ
\]  
(Eq. 3)

The transformation was carried out using GNOM software (37). For dp6–dp16, the full x-ray \( I(Q) \) curves contained 295–343 data points in similar \( Q \) ranges between 0.29 and 1.80 nm\(^{-1}\).

Molecular Modeling of HS Fragments—Linear HS models were created using the crystal structure of the HS tetrasaccharide (dp4) observed in its complex with heparinase II (PDB code 3E7J) with Discovery Studio (version 2.5) molecular graphics software (Accelrys, San Diego, CA). The monosaccharide residues in the HS tetrasaccharide of 3E7J were N-acetyl \( \alpha-D \)-glucosamine (internal), \( \beta-D \)-GlcNAc (reducing terminal), \( \beta-D \)-GlcA (internal), and 4,5-dehydro-\( \beta-D \)-GlcA (\( \Delta \)UA; nonreducing terminal). Unfortunately the PDB three-letter abbreviations used in 3E7J do not conform to the PDB conventions. The abbreviation NAG (correctly used only for \( \beta-D \)-GlcNAc) is used for both \( \alpha \)- and \( \beta \)-anomers; the abbreviation GCU (correctly used only for \( \alpha-D \)-GlcA) is used for \( \beta-D \)-GlcA. In addition, we also point out that the dp4 structure was written out in the original crystallography paper (20) as NAG-GCU-NAG-GCD (where GCD is the PDB code for \( \Delta \)UA). This order, with the reducing end to the left, is unconventional. In the current study, the disaccharide \( \alpha-D \)-GlcNAc-(1→4)-\( \beta-D \)-GlcA, from the two internal monosaccharides in the 3E7J tetrasaccharide, was taken to be the base HS structure, and these disaccharide units (PDB code NDG-BDP) were joined by glycosidic linkages to generate a fully extended linear HS dp30 structure. In this, the \( \Phi \) and psi (\( \Psi \)) angles were maintained at similar values observed in the starting dp4 structure. Linear HS dp6–dp24 models were created from this extended dp30 starting model by the removal of nonrequired disaccharides.

Totals of 5,000 conformationally randomized models for each of dp6, dp8, dp10 and dp12, 8,000 similar models for each of dp14 and dp16, and 12,000 similar models for each of dp18 and dp24 were created starting from each linear model. In the original HS dp4 structure, the \( \Phi \) and \( \Psi \) angles were ~90° and 124° respectively for the GlcA-GlcNAc (BDP-NDG) disaccharide and 85° and 95° respectively for the GlcNAc-GlcA (NDG-BDP) disaccharide. These \( \Phi \) and \( \Psi \) angles were randomized to take any value in a maximum range of ±45° starting from the preceding values using the TorsionKick function in a PERL script that was modified from the ExtractAngle.pl script provided with the Discovery Studio software. For example, in the case of dp16, a total of eight \( \Phi \) and \( \Psi \) angles for GlcA-GlcNAc and seven \( \Phi \) and \( \Psi \) angles for GlcNAc-GlcA were randomized in this way. To avoid steric clashes between the dp16 atoms in each randomized structure, a constant force field termed DREIDING minimation provided in Discovery Studio was used to correct this. DREIDING minimation was useful in generating structures by providing accurate geometries and reasonably accurate steric barrier for organic, biological and inorganic main groups (38).

Constrained Scattering and Sedimentation Coefficient Modeling—Each HS model was used to calculate the x-ray scattering curve for comparison with the experimental curve using Debye sphere models (39–41). A cube side length of 0.520 nm in combination with a cutoff of 4 atoms was used to create the spheres for the HS dp6–dp16 models. The hydration shell corresponding to 0.3 g/g H\(_2\)O was created by adding spheres to the unhydrated sphere models using HYPRO (42), where the optimal total of hydrated spheres is listed in Table 1. The x-ray scattering curve \( I(Q) \) was calculated using the Debye equation as adapted to spheres and assuming a uniform scattering density for the spheres (43). Other details are given elsewhere (39–41, 44). X-ray curves were calculated without instrumental corrections as these were considered to be negligible for the pinhole optics used in synchrotron x-ray instruments. First, the number of spheres \( N \) in the dry and hydrated models after grid transformation was used to assess steric overlap between the HS disaccharides, where models showing less than 95% of the optimal totals (see Table 1) were discarded. This procedure was found to be insensitive to steric overlap in the case of oligosaccharides,
and was discontinued in favor of the DREIDING minimization procedure (above). Next, the models were assessed by calculation of the x-ray $R_G$ values from Guinier fits of the modeled curves using the same $Q$ ranges used for the experimental Guinier fits to allow for any approximations inherent in the use of the $Q R_G$ range up to 1.5. Models that passed the $N$ and $R_G$ filters were then ranked using a goodness-of-fit $R$ factor to identify the best fit eight models for each HS fragment.

Sedimentation coefficients $s_{20,w}$ for each of the eight best fit HS scattering models were calculated directly from molecular structures using the HYDROPRO shell modeling program (45). The default value of 0.31 nm for the atomic element radius for all structures using the HYDROPRO shell modeling program (45).

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given that the four smallest HS fragments dp6–dp12 were eluted as well resolved peaks, whereas the four larger fragments dp14–dp24 were less well resolved (Fig. 2). Analytical high performance size exclusion chromatography of the HS oligosaccharide fractions as described (25), and PAGE was performed to show that all eight peak fractions showed altered sizes as expected and were relatively homogenous in size. Proton NMR spectroscopy (not shown) showed that the GlcA-GlcNAc disaccharide was the predominant structure between the 77 and 86% values would correspond to a 10% predicted reduction in $s_{20,w}$ value. If real, the difference between the 77 and 86% values would correspond to a 10% smaller frictional coefficient for HS compared with heparin, i.e., HS may have a slightly more compact solution structure than heparin.

The sedimentation coefficient $s_{20,w}$ values were calculated using HYDROPRO software from molecular models of HS. For this, 13 linear HS models (dp6 – dp30) were computed starting from the HS dp4 crystal structure seen in its complex with heparinase II (20). Even though the rate of increase of the $s_{20,w}$ values with size was predicted correctly, the theoretical $s_{20,w}$ values for HS were consistently lower than those seen experimentally (Fig. 4, A and B). The theoretical values for the HS fragments were consistently lower when compared with their experimental values and therefore show that HS sediments more rapidly than predicted, i.e., the overall solution structures are more compact through bending than the linear HS structures. Below, this difference in Fig. 4 is explained by the best fit bent HS structures that were modeled from the scattering data. It is noteworthy that this difference between experimental and theoretical values was not seen for heparin (19); this shows that the HS and heparin solution structures are different. In addition, the linear HS and heparin structures differ in their degree
of elongation. The theoretical $s_{20,w}$ value for a linear HS structure is 70% lower than the corresponding theoretical value for a linear heparin structure, in reflection of a longer linear HS structure.

X-ray Solution Scattering Data for Eight HS Fragments—Solution scattering is a diffraction technique that provides structural information on biological macromolecules in random orientations in solution (21, 22). To complement the analytical ultracentrifugation data in more detail, the solution structures of the same eight HS fragments dp6–dp24 at 0.5 mg/ml were characterized by synchrotron x-ray scattering. The scattering experiments reports the scattering curve $I(Q)$ as a function of scattering angle $Q$. Tests for possible radiation damage effects ("Experimental Procedures") showed that they were not detectable; accordingly data were acquired with the longest exposure time of 1 s to maximize signal:noise ratios. Guinier analyses of $\ln I(Q)$ versus $Q^2$ at low $Q$ values gives the radius of gyration, $R_G$, which monitors the degree of macromolecular elongation. Because of the very different sizes of the HS fragments, different $Q$ ranges were used for each fragment to work within acceptable linear $Q$-$R_G$ ranges (Fig. 5A). Thus the Guinier fit $Q$ range of 0.4 to 0.8 nm$^{-1}$ for dp6 was successively reduced in stages to that of 0.28–0.55 nm$^{-1}$ for dp24 (Fig. 5A). The mean Guinier $R_G$ values increased from 1.03 ± 0.08 nm for dp6 up to 2.82 ± 0.10 nm for dp24 (Table 1). These $R_G$ values for the eight HS fragments do not show a linear relationship with the size of the HS fragments, unlike the $R_G$ values calculated from linear models, showing reduced values instead (Fig. 6A). Thus the comparison of the $R_G$ values between the linear models and the experimental data also showed that the HS fragments were bent in solution. In addition, the experimental HS $R_G$ values are larger for the dp18–dp36 fragments than those seen for the heparin dp18 and dp24 fragments (Fig. 6A). This shows that HS has a more elongated structure than that for heparin.

Macromolecules that are sufficiently elongated in shape will show a cross-sectional radius of gyration $R_{XS}$ value. The $R_{XS}$ value monitors the degree of bending within the macromolecular length. As for the $R_G$ analyses, different $Q$ ranges were used for the $R_{XS}$ fits for the different HS fragments depending on the size of the HS fragment, all of which were larger than the $Q$ ranges used for the $R_G$ analyses (Fig. 5B). In all cases, despite the worsened signal:noise ratios of the $I(Q)$ data, linear fit ranges were identified in the plots of $\ln I(Q)$-$Q$ versus $Q^2$. These ranges gave experimental $R_{XS}$ values of 0.31 nm for dp6 that increased
up to 0.65 nm for dp24 (Fig. 6B and Table 1). This increase in the \( R_{XX} \) values correlated with the deviation of the \( R_G \) values from linearity (Fig. 6A). They were larger than the calculated \( R_{XX} \) values of 0.40 nm for the linear HS dp6-dp30 models, thus supporting the conclusion that the HS fragments become progressively more bent with an increase in size. Combination of the \( R_G \) and \( R_{XX} \) values according to the relationship \( L^2 = 12 \ (R_G^2 - R_{XX}^2) \) for an elliptical cylinder (36) showed that HS dp6, dp8, dp10, dp12, dp14, dp16, dp18, and dp24 have approximate lengths of 3.4, 3.9, 4.6, 5.5, 5.8, 6.8, 7.8, and 9.5 nm in that order. Similar lengths of 7.0, 9.1, 9.6, and 10.7 nm were observed for the heparin dp18, dp24, dp30, and dp36 fragments (19). In conclusion, the comparison of the dp18 and dp24 lengths showed that HS has a longer structure than that of heparin, in addition to being more bent than heparin.

The distance distribution function \( P(r) \) is calculated from the full Q range of the \( I(Q) \) scattering curve (“Experimental Procedures”). The \( P(r) \) curve represents the distribution of all the distances between the atoms within the macromolecule. This provides \( R_G \) values and model-independent determinations of the overall length, \( L \), following an assumption of the maximum dimension, \( D_{max} \) (Fig. 6C); note that \( L \) is not a contour length. The mean \( R_G \) values obtained from the \( P(r) \) curves increase from 0.98 ± 0.05 nm for dp6 to 3.0 ± 0.05 nm for dp24 (Table 1). These \( P(r) \) \( R_G \) values are in excellent accord with the corresponding Guinier \( R_G \) values from the low \( Q \) values and follow the same trends with size (Table 1 and Fig. 6A). Model-independent \( L \) values are determined from the \( r \) value where the \( P(r) \) curve reaches zero at large \( r \). These experimental \( L \) values were 3.0 nm for dp6, 3.5 nm for dp8, 4.5 nm for dp10, 5.5 nm for dp12, 6.0 nm for dp14, 7.0 nm for dp16, 8.5 nm for dp18, and 10.0 nm for dp24. These values show increasing deviation with size from the longer lengths measured for the linear HS dp6–dp24 models (i.e., 3.0 nm for dp6, 3.9 nm for dp8, 4.4 nm for dp10, 5.7 nm for dp12, 7.0 nm for dp14, 8.5 nm for dp16, 8.9 nm for dp18, and 11.0 nm for dp24), noting that a hydration shell of thickness 0.6 nm (2 × 0.3 nm) is added to these linear model lengths (44). These experimental \( L \) values from the \( P(r) \) curves are more accurate compared with the approximate \( L \) values calculated from the \( R_G \) and \( R_{XX} \) values that assumed an elliptical
FIGURE 5. Experimental Guinier x-ray scattering analyses of eight HS dp6 – dp24 fragments. A, Guinier $R_G$ plots for dp6 – dp24 at concentrations of 0.5 mg/ml. The filled circles were used to determine the radius of gyration $R_G$, based on the best fit lines as shown. The $Q$ ranges used for the $R_G$ analyses were 0.40–0.8 nm$^{-1}$ for dp6, 0.42–0.78 nm$^{-1}$ for dp8, 0.34–0.74 nm$^{-1}$ for dp10, 0.30–0.66 nm$^{-1}$ for dp12, 0.30–0.64 nm$^{-1}$ for dp14, 0.29–0.55 nm$^{-1}$ for dp16, 0.28–0.55 nm$^{-1}$ for dp18, and 0.28–0.55 nm$^{-1}$ for dp24. B, Guinier $R_{XS}$ plots for dp6 – dp24. The filled circles represent the $Q$ ranges used to determine the cross-sectional radius of gyration $R_{XS}$, based on the best fit lines as shown. The $Q$ ranges used for $R_{XS}$ analyses were 1.2–1.6 nm$^{-1}$ for dp6, 1.1–1.6 nm$^{-1}$ for dp8, 1.0–1.54 nm$^{-1}$ for dp10, 1.0–1.44 nm$^{-1}$ for dp12, 0.82–1.3 nm$^{-1}$ for dp14, and 1.0–1.4 nm$^{-1}$ for dp16, dp18, and dp24.

FIGURE 6. Experimental Guinier and $P(r)$ x-ray data analyses of eight HS dp6 – dp24 fragments. A, comparison of the experimental $R_G$ values from Guinier plots (△) and $P(r)$ curves (●) with the predicted $R_G$ values calculated from the linear models of Fig. 4 (○). The six corresponding values for heparin from Ref. 19 are denoted by ○ and ★, respectively, and fitted to a dotted line. B, comparison of the experimental cross-sectional $R_{XS}$ values (●) with the predicted $R_{XS}$ values calculated from the linear models of Fig. 4 (○). The corresponding four values for heparin dp18 – dp36 from Ref. 19 are denoted by ○ and fitted to a dotted line. C, the distance distribution function $P(r)$ analyses for dp6 – dp24. The $r$ values of the maximum at $M$ were 1.02 nm (dp6), 1.15 nm (dp8), 1.30 nm (dp10), 1.43 nm (dp12), 1.44 nm (dp14), 1.61 nm (dp16), 1.87 nm (dp18), and 1.90 nm (dp24). The eight fragments are denoted by continuous, dashed, and dotted lines in alternation. D, comparison of the $P(r)$ analyses for HS dp6 – dp24 with heparin dp6 – dp24. The curves corresponding to the four HS fragments dp6, dp12, dp18, and dp24 are denoted by dashed lines, whereas the corresponding four curves for heparin are denoted by continuous lines. The heparin $P(r)$ data are from Ref. 19.
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Constrained Modeling of Eight HS Fragments—The experimental x-ray $R_G$ and $R_{XS}$ values showed that the solution structures for HS is longer and more bent than those of heparin. Here, constrained scattering modeling was performed with HS to provide a molecular explanation of these scattering data. The linear models created from the HS dp4 crystal structure were the starting constraint. All eight HS fragments dp6 – dp24 were subjected to modeling. They were considered as belonging to a structurally homologous series. As illustrated in the previous modeling of heparin, the linkage connectivity between the oligosaccharide rings was maintained (Fig. 1A), whereas the $\Phi$ and $\Psi$ rotational angles at each glycosidic linkage was varied randomly in a range of up to $\pm 45^\circ$ from their preceding values. In all, 5,000 – 12,000 models for each of the eight HS fragments were generated. For each fragment, x-ray scattering curves were calculated from these randomized models and fitted to the experimental curve. The $R_G$, $R_{XS}$, and $R$ factor values were calculated for each modeled curve, where the $R_G$ and $R_{XS}$ values were calculated using the same $Q$ range used for the experimental Guinier fits (Fig. 5, A and B), and the $R$ factor is a measurement of goodness of fit. The $R$ factor distributions (supplemental Fig. S1, A–H) showed that all the dp6 – dp24 models (yellow circles) encompassed the experimental $R_G$ values (dashed lines) and that the lowest $R$ factor values were close to the 5 – 8% level usually expected for excellent curve fits (47).

Totals of 5,000 models for dp6 – dp10 and 8,000 models for dp12 – dp16 provided enough randomized conformers to be able to determine best fit HS solution structures. Typically the lowest $R$ factors showed the best agreement with the experimental x-ray curves. Near these minima, the best fit $R_G$ values for dp6 – dp16 (one red and seven cyan circles) agreed well with the experimental $R_G$ values. The best fit $R_{XS}$ values for dp6 – dp16 also showed good agreement with the experimental $R_{XS}$ values (supplemental Fig. S1, I–P). In distinction, the linear models for the dp6 – dp16 HS fragments showed greater deviations from the experimental $R_G$ and $R_{XS}$ values (green circles). These modeling analyses (Table 1 and Fig. 7) confirmed that a nearly linear structure with slight bending accounted for the x-ray and analytical ultracentrifugation data for dp6 – dp16.

For HS dp6, the eight best fit models gave $R$ factors of 4.4 – 4.6%, $R_G$ values of 1.01 – 1.03 nm, $R_{XS}$ values of 0.30 – 0.32 nm, and maximum lengths, $L$, of 3.0 – 3.1 nm. These agree well with the experimental $R_G$ value of 1.03 ± 0.08 nm, $R_{XS}$ value of 0.31 ± 0.06 nm, and the $P(r)$ length of 3.0 nm (Table 1). In distinction to these agreements, the linear dp6 model gave an $R$ factor of 4.1%, an $R_G$ value of 0.98 nm, and a $L$ value of 3.0 nm. The visual agreement between the experimental and modeled $l(Q)$ curves and $P(r)$ curves was excellent (Fig. 7A). The calculated $s^0_{20,w}$ values from the eight best fit models gave 0.47 – 0.53 S. These values are lower but comparable with the experimental $s_{20,w}$ values of 0.82 ± 0.05 and 1.05 ± 0.04 S, given that the typical accuracy of the $s^0_{20,w}$ calculation is ± 0.21 S (22).

For HS dp8, the modeling analyses indicated slightly bent structures similar to those seen for dp6. The eight best fit models gave $R$ factors of 4.5%, $R_G$ values of 1.18 – 1.20 nm, $R_{XS}$ values of 0.40 nm, and $L$ values of 3.5 – 3.8 nm. These values agree well with the experimental $R_G$ value of 1.19 ± 0.08 nm, $R_{XS}$ value of 0.40 ± 0.03 nm, and the $P(r)$ length of 3.5 nm (Table 1). In distinction to these agreements, the linear dp8 model again showed a higher $R$ factor of 4.8% and an $R_G$ value of 1.21 nm, an $R_{XS}$ value of 0.43 nm, and a larger $L$ value of 3.9 nm. The visual agreement of the experimental and modeled $l(Q)$ and $P(r)$ curves was excellent (Fig. 7B). The eight calculated $s^0_{20,w}$ values of 0.61 – 0.62 S are smaller but comparable with the experimental $s_{20,w}$ values of 0.94 ± 0.06 and 1.06 ± 0.08 S.

For HS dp10, the modeling analyses showed good agreements with slightly bent structures, in which the deviation from a linear dp10 structure for dp10 was slightly increased. The eight best fit dp10 models gave $R$ factors of 4.4%, $R_G$ values of 1.40 – 1.42 nm, $R_{XS}$ values of 0.43 – 0.44 nm, and $L$ values of 4.5 – 5.0 nm. These correspond well with the experimental $R_G$ value of 1.41 ± 0.07 nm, $R_{XS}$ value of 0.44 ± 0.04 nm, and $L$ value of 4.5 nm (Table 1). The deviations from a linear dp10 model are larger, for which the $R$ factor is 4.4%, the $R_G$ value is 1.44 nm, the $R_{XS}$ value is 0.47 nm, and the $L$ value is 4.4 nm. The visual agreement of the experimental and modeled $l(Q)$ and $P(r)$ curves was again excellent (Fig. 7C). The eight modeled $s^0_{20,w}$ values of 0.67 – 0.68 S are again comparable with the experimental $s_{20,w}$ values of 0.95 ± 0.09 and 1.09 ± 0.06 S.

For HS dp12, the modeling analyses were also successful, in which the deviation from a linear dp12 structure was greater. The eight best fit models gave $R$ factors of 4.2 – 4.5%, $R_G$ values of 1.63 – 1.65 nm, $R_{XS}$ values of 0.48 – 0.50 nm, and $L$ values of 5.5 – 5.7 nm. These agree well with the experimental $R_G$ value of 1.65 ± 0.09 nm, the $R_{XS}$ value of 0.49 ± 0.04 nm, and the $L$ value of 5.7 nm (Table 1). The linear dp12 model showed a higher $R$ factor of 4.5%, a higher $R_G$ value of 1.68 nm, a reduced $R_{XS}$ value of 0.42 nm, and a longer $L$ value of 5.7 nm. The visual agreement of the experimental and modeled $l(Q)$ and $P(r)$ curves was excellent (Fig. 7D). The eight modeled $s^0_{20,w}$ values of 0.74 S are comparable with the experimental $s_{20,w}$ values of 1.08 ± 0.09 and 1.16 ± 0.05 S.

For HS dp14, good agreements between the models and experimental data were obtained, whereas the deviation from a linear dp14 structure was larger. The eight best fit dp14 models gave $R$ factors of 4.3%, $R_G$ values of 1.73 – 1.76 nm, $R_{XS}$ values of 0.48 – 0.49 nm, and $L$ values of 6.0 – 6.3 nm. These values agree well with the experimental $R_G$ value of 1.76 ± 0.07 nm, the $R_{XS}$ value of 0.51 ± 0.02 nm, and the $L$ value of 6.0 nm (Table 1). These values deviated from the linear dp14 model, which had an $R$ factor of 4.7%, a higher $R_G$ value of 1.91 nm, a lower $R_{XS}$ value of 0.38 nm, and an $L$ value of 7.0 nm. Again the visual
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(A) dp6

(B) dp8

(C) dp10

(D) dp12

(E) dp14

(F) dp16

(G) dp18

(H) dp24

\( \ln(I(Q)) \) vs. \( Q \) (nm\(^{-1}\))

\( L = 3.0 \text{ nm} \)
\( L = 3.1 \text{ nm} \)

\( L = 3.9 \text{ nm} \)
\( L = 3.8 \text{ nm} \)

\( L = 4.4 \text{ nm} \)
\( L = 4.6 \text{ nm} \)

\( L = 5.7 \text{ nm} \)
\( L = 5.5 \text{ nm} \)

\( L = 7.0 \text{ nm} \)
\( L = 6.1 \text{ nm} \)

\( L = 8.5 \text{ nm} \)
\( L = 7.2 \text{ nm} \)

\( L = 8.9 \text{ nm} \)
\( L = 8.5 \text{ nm} \)

\( L = 9.0 \text{ nm} \)
\( L = 8.5 \text{ nm} \)
agreement of the experimental and modeled \(I(Q)\) and \(P(r)\) curves was excellent (Fig. 7E). The eight modeled \(s_{20\text{,w}}^0\) values of 0.82–0.83 S compare well with the experimental \(s_{20\text{,w}}^0\) value of 1.07 ± 0.07 and 1.18 ± 0.04 S, respectively.

For HS dp16, the outcome of the modeling analyses was similar to that of dp14. The eight best fit models gave \(R\) factors of 6.5–6.6%, \(R_G\) values of 1.93–2.00 nm, \(R_{XS}\) values of 0.49–0.51 nm, and \(L\) values of 6.4–7.2 nm. These agree well with the experimental \(R_G\) value of 2.03 ± 0.07 nm, the \(R_{XS}\) value of 0.52 ± 0.01 nm, and the \(L\) value of 7.0 nm (Table 1). The linear dp16 model gave a poorer fit with an \(R\) factor of 6.7%, an \(R_G\) value of 2.15 nm, an \(R_{XS}\) value of 0.38 nm, and an \(L\) value of 8.5 nm. The experimental and modeled \(I(Q)\) and \(P(r)\) curves showed excellent agreement (Fig. 7F). The eight best fit models gave two values of 0.92 S. For dp24, the experimental \(R\) values were less than the experimental \(R_{XS}\) value of 0.43 nm, and an \(L\) value of 11.0 nm. Notably, \(R_G\) values of 2.22 nm, \(R_{XS}\) value of 0.31 nm, and an \(L\) value of 8.9 nm. For extended dp24, modeled \(R_G\) values of 2.68–2.78 nm, \(R_{XS}\) values of 0.52–0.58 nm, and \(L\) values of 9.0–10.0 nm were obtained. The modeled \(R_G\) and \(R_{XS}\) values were less than the experimental \(R_G\) value of 2.34 ± 0.03 nm and the \(R_{XS}\) value of 0.61 ± 0.05 nm (Table 1).

As expected, the linear model gave a poorer fit with an \(R\) factor of 8.2%, \(R_G\) value of 2.22 nm, \(R_{XS}\) value of 0.31 nm, and an \(L\) value of 8.9 nm. For extended dp24, modeled \(R_G\) values of 2.68–2.78 nm, \(R_{XS}\) values of 0.52–0.58 nm, and \(L\) values of 9.0–10.0 nm were obtained. The modeled \(R_G\) and \(R_{XS}\) values were less than the experimental \(R_G\) value of 2.82 ± 0.10 nm and the \(R_{XS}\) value of 0.65 ± 0.05 nm (Table 1). The linear model also gave a poorer fit with an \(R\) factor of 11.9%, an \(R_G\) value of 2.72 nm, an \(R_{XS}\) value of 0.43 nm, and an \(L\) value of 11.0 nm. Notably, the \(R_G\) values of 2.13 nm (dp18) and 2.47 nm (dp24) at the \(R\) factor minima of supplemental Fig. S1 (G and H) (Table 1) did not coincide with the experimental \(R_G\) values of 2.34 nm (dp18) and 2.82 nm (dp24). Better agreements were observed for more bent models (Table 1). These differences in \(R_G\) values suggested that conformational heterogeneity between extended and bent structures was present, i.e., dp18 and dp24 exhibited multiple conformations in solution. The experimental and modeled \(I(Q)\) and \(P(r)\) curves showed good visual agreements for either conformation (Fig. 7, G and H). For dp18, the modeled \(s_{20\text{,w}}^0\) values of 0.90–0.92 S (extended) or 0.90–0.94 S (bent) were similar to the experimental \(s_{20\text{,w}}^0\) values of 1.12 ± 0.06 and 1.25 ± 0.07 S, with the best fit models giving two values of 0.92 S. For dp24, the modeled \(s_{20\text{,w}}^0\) values of 0.91–1.0 S (extended) or 0.78–1.07 S (bent) were similar to the experimental \(s_{20\text{,w}}^0\) values of 1.26 ± 0.06 S and 1.34 ± 0.06 S, with the two best fit models giving a value of 0.96 or 1.07 S.

**DISCUSSION**

The size and spacing of \(S\) domains in HS are proposed to be as important to its biologically significant interactions with proteins as are the detailed sequences of the \(S\) domains themselves (49). Heparin, a commonly used model compound for HS, consists of lengthy \(S\) domains, made up largely of the repeating trisulfated disaccharide shown in Fig. 1B, separated by much smaller, unsulfated \(NA\) domains. In HS the position is reversed, and long \(NA\) domain sequences (Fig. 1A) act as spacers to separate the short \(S\) domains. Although the \(S\) domain conformation, exemplified by heparin, has been the subject of numerous studies (50), the \(NA\) domain has not. It has been proposed that the \(NA\) domain repeating sequences are both less flexible (51) and more flexible (52) than the \(S\) domains.

The application of constrained scattering modeling has proved to be as effective for the HS fragments as it was for heparin previously. Usually scattering fits are performed for protein structures of size 20–100 kDa and higher (21, 22). The HS fragments dp6–dp24 and the heparin fragments dp6–dp36 are notably smaller in size with masses of 1–7 and 2–11 kDa, respectively. The ability to measure their scattering curves was attributed to the high x-ray beam intensity and low background at the instrument, together with improved detector technology. Constrained scattering modeling determines a three-dimensional molecular structure that best accounts for the observed scattering curve through trial and error searches that rule out structures that are incompatible with the observed scattering curves. By fixing the analyses to what is already known about the macromolecule, namely the carbohydrate rings, and varying only the \(\Phi\) and \(\Psi\) angles of each glycosidic linker, relatively few modeling variables are involved in the scattering fits. Through the variation of \(\Phi\) and \(\Psi\), the resulting 5,000–12,000 models provided sufficient statistical detail to result in clear V-shaped graphs of \(R\) factor versus \(R_G\) and \(R\) factor versus \(R_{XS}\) values. The best fit models were identified by the lowest \(R\) factors, and they were verified by the agreement of the modeled and experimental \(R_G\) and \(R_{XS}\) values at this point. The quality of the HS dp6–dp16 scattering fits was similar to those of the heparin dp18–dp36 fits (19). Interestingly, different fits were obtained for the HS dp18 and dp24 structures. The monodispersity of these two fragments had been established by the single peaks seen in the ultracentrifugation (c) analyses (Fig. 3); thus the potential contribution of sample heterogeneity in dp18 and dp24 can be ruled out as an explanation for the different fits. The ability to fit either extended or bent HS dp18 and dp24 structures to the scattering curves resulted in the conclusion of multiple conformations for HS dp18 and dp24. This modeling outcome is distinct from that for heparin dp6–dp36, where only single conformations were
required for good fits. This outcome suggested that the heparin structures show greater rigidity than the HS structures. The HS fragments used in this study were produced by extensive depolymerization using heparinase 1, an enzyme that cleaves only within the S domains (28) leaving NA domains untouched. It is therefore likely that some minor degree of sulfation remains at the reducing and nonreducing end of our fragments, but that internal disaccharides are unsulfated. Such fragments bear a closer resemblance to NA domains of intact HS than the most commonly used model compound for this sequence, the capsular polysaccharide from Escherichia coli K5 (51, 52). These structures provide novel comparative insight into the structures of HS and heparin, and the likely manner that these two polyanionic oligosaccharides interact with their protein ligands. HS and heparin both share similar covalent structures (Fig. 1). The comparison of our two sets of structures for HS and heparin becomes essentially that between NA and S domains. The greater bending and flexibility of HS compared with heparin (see Fig. 9) may be attributed to the difference in uronic acid residue, in which GlcNAc alternates with GlcA in HS, causing HS to adopt a distinct conformation from that of fully sulfated heparin (53).

More detailed inspection of the best fit models for HS dp6–dp16 clarifies their progressively more bent structures in solution with increase in HS size. This outcome is visible from the superimposition of the eight best fit models for each HS fragment (Fig. 8). When comparing the solution structures of heparin dp18 and dp24 (Fig. 8 of Ref. 19) with those of HS dp18 and dp24, the HS structures were visibly more bent than those of heparin (Fig. 9A). Crystal structures containing HS or heparin showed that the glycosidic linkage in HS has a similar length to that in heparin (Fig. 9B). In HS, the separation between the C1–C4 atoms of GlcA-GlcNAc is 0.237 ± 0.003 nm and that between GlcNAc-GlcA is 0.235 ± 0.002 nm. For heparin, analyses of five crystal structures containing dp6 showed that very similar separations were seen between IdoA-GlcNS of 0.241 ± 0.004 nm and between GlcNS-IdoA of 0.243 ± 0.006 nm. Thus the increased length of HS in solution compared with heparin is mostly the consequence of altered Φ and Ψ angles. In terms of rotational bend, it is already known that the crystallographic Φ and Ψ angles for smaller HS and heparin structures agree with each other within error (Table 2 and supplemental Fig. S2). The Φ and Ψ angles for HS dp6–dp24 were all similar, including...
those in the small crystal structures (Table 2). The linkage geometries of our HS models fall approximately into the low energy regions for \( ^1H_9251 \) (13) and \( ^1H_9252 \) (13) linkages between glucose residues, as determined for maltose and cellobiose by experimental and theoretical methods (48). Thus the mean \( ^1H_9021 \) and \( ^1H_9023 \) angles for the GlcA-GlcNAc linkage were 85 and 127° in HS, being similar but not identical with the corresponding IdoA-GlcNS values of 61 and 132° in heparin. The mean \( ^1H_9021 \) and \( ^1H_9023 \) angles for the GlcNAc-GlcA linkage were 85 and 91° in HS, close to the corresponding values of 98 and 86° for GlcNS-IdoA in heparin (Table 2). The distribution of the \( ^1H_9021 \) and \( ^1H_9023 \) values in supplemental Fig. S2 (A and B) showed low rotational variability at both glycosidic linkages in HS. These distributions were in good accord with those for heparin in supplemental Fig. S2 (C and D).

In conclusion, the more bent structure in solution for HS than for heparin (Fig. 9A) is attributed to small but reproducible differences of 24 and 13° for the \( ^1H_9021 \) angles of heparin and HS (Table 2). In contrast, the two \( ^1H_9023 \) angles of heparin and HS differ by only 5°.

The solution structures of the HS dp6–dp24 fragments exhibited bending and flexibility (Figs. 8 and 9A). In addition, the HS structures are slightly longer for reason of alterations in the glycosidic \( \Phi \) and \( \Psi \) angles (supplemental Fig. S2). The physical basis for these changes in HS is likely to arise from the GlcA-GlcNAc sequences (as opposed to the IdoA-GlcNS sequences in heparin). Unlike HS, heparin will be influenced by greater repulsion between regular repeats of sulfate-sulfate, sulfate-carboxylate, and carboxylate-carboxylate groups. The combination of the \( \alpha \) and \( \beta \) domain structures within the parent HS structure suggests that different parent HS structures with greater or lesser bending may arise through variations of the ratio in sizes of the \( \alpha \) and \( \beta \) domains. These variations are likely to be as important as the fine structure of the individual domains in the physiological functions of these complex glycosaminoglycans (49, 55).

In terms of biological function, the greater degree of bending in HS accounts for the ability of HS to bind to a diverse range of protein ligands in all orientations. This will facilitate the assembly of large multipartner complexes on cell surfaces such as those involving the 20-domain structure of complement factor H (56). The outcome of 19 protein-heparin crystal structures has been discussed previously (19), whereas only one protein-HS crystal structure is known (20). It was of interest that the \( \Phi \) and \( \Psi \) angles for the HS crystal structure are similar to those seen in the 19 heparin-protein crystal structures (Table 2). In addition, the mean \( \Phi \) and \( \Psi \) angles for HS in solution are similar and close to those seen by crystallography for HS dp4. Several studies of HS-protein interaction have noted that SAS sequences, in which two short \( \beta \) domains are separated by an \( \alpha \) sequence, are preferred for optimum binding (48). This is particularly true for oligomeric proteins.

**TABLE 2**

<table>
<thead>
<tr>
<th>Summary of the ( \Phi ) and ( \Psi ) angles in the crystal and solution structures of HS and heparin</th>
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<tr>
<td>( \Phi (\degree) )</td>
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<td><strong>GlcA-GlcNAc</strong></td>
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<td>HS crystal structure *</td>
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<td>Heparin dp6</td>
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<td>Average (5/54 values) *</td>
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</table>

* The mean value from two HS dp4 molecules seen in the crystal structure (PDB code 3E7J).

* The average is calculated from all the \( \Phi \) and \( \Psi \) angles in the eight best fit models.

* The average is calculated from the heparin models of Ref. 19.
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such as, for example, MIP-1α or platelet factor 4 (7, 57). In these cases, the multiple heparin-binding sites on the oligomer are not always arranged in a linear way, so that a single long S domain cannot readily bind to more than one site on the same multimer. It has been reasonable to suppose that NA domains, composed of alternating α(1→4)- and β(1→4)-linked hexopyranoses, would be more flexible than the unusually stiff heparin structure of S domains, allowing an SAS domain to bend to present two S domains to heparin-binding sites on opposite sides of a protein complex. The conclusion from our present study of HS NA fragments supports this intuitive reasoning.

Acknowledgments—We thank Dr. Azubuike I. Okemefuna for excellent computational help and Dr. Anne Imberty (CNRS Grenoble) for useful discussions. We thank the National Institute for Biological Standards and Control for the provision of laboratory facilities.

Addendum—In our originally reported HS best fit structures (58), the anomeric configurations of the GlcA and GlcNAc residues should have alternated between α and β. Instead, the anomeric configurations were all β in our original HS models. The present study replaces our original study (58), which has now been withdrawn. The error was traced back to a misunderstanding of the starting HS structure in the Protein Data Bank, and we take full responsibility for our mistake. First, the dp4 structure had been written out in the original crystallography paper (20) as NAG-GCU-NAG-GCD (where NAG is the PDB code for GlcNAc, GCU is the code for GlcA, and GCD is the code for ΔUA). This order, with the reducing end to the left, is unconventional and was a contributing factor to our misunderstanding. Second, the use of the terms NAG and GCU corresponded to the β-GlcNAc and α-GlcA anomers, respectively, in the PDB dictionary. In fact, HS is comprised of the α-GlcNAc and β-GlcA anomers and should have been described as NDG and BDP in the HS dp4 crystal structure (PDB code 3E7)). Inspection of PDB depositions containing GCU showed the existence of other terminological confusions. Several relatively recent structures in the PDB use GCU for internal β-GlcA residues, possibly following the example of the classic hyaluronate and chondroitin sulfate studies (PDB codes 1HYA, 2HYA, 4HYA, 1C4S, and 2C4S). In addition to the PDB code 3E7) of the present study, the structures of bee venom hyaluronidase with a hyaluronate tetrasaccharide (PDB code 1FCV) and CD44 with a hyaluronate octamer (PDB code 2CJQ) use GCU and not BDP. The PDB structure 2CJQ (from the same study as PDB code 2CJQ) uses the correct term BDP. Related discrepancies in the PDB have also been reported by others (54, 59).

REFERENCES
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