Mosquito Heparan Sulfate and Its Potential Role in Malaria Infection and Transmission*

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Heparan sulfate has been isolated for the first time from the mosquito Anopheles stephensi, a known vector for Plasmodium parasites, the causative agents of malaria. Chondroitin sulfate, but not dermatan sulfate or hyaluronan, was also present in the parasites, the causative agents of malaria. Chondroitin sulfate, sporozoite protein (CSP),2 to heparan sulfate proteoglycans in the binding of the sporozoite major surface protein, the circumsporozoite protein found on the surface of the Plasmodium sporozoite. The heparan sulfate isolated from the whole mosquito binds to circumsporozoite protein, suggesting a role within the mosquito for infection and transmission of the Plasmodium parasite.

Malaria infection is initiated with the bite of an infected anopheline mosquito, which injects Plasmodium sporozoites into the skin of the vertebrate host (Fig. 1A). Sporozoites migrate from the site of injection, enter the blood stream, and are carried to the liver, where they invade hepatocytes and develop into exoerythrocytic forms. Many lines of evidence suggest that clearance of sporozoites by the liver is mediated by the binding of the sporozoite major surface protein, the circumsporozoite protein (CSP),2 to heparan sulfate proteoglycans in the liver (1, 2). Liver heparan sulfate is more highly sulfated than heparan sulfate from other organs (3–6) and frequently contains the fully biosynthetically modified, highly sulfated sequence, →4)β-d-GlcNS6S(1 → 4)-α-L-IdoA2S(1→, that is commonly found in human liver heparan sulfate, which serves as the receptor for apolipoprotein E and is also believed to be responsible for binding to the circumsporozoite protein found on the surface of the Plasmodium sporozoite. The heparan sulfate isolated from the whole mosquito binds to circumsporozoite protein, suggesting a role within the mosquito for infection and transmission of the Plasmodium parasite.

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2 The abbreviations used are: CSP, circumsporozoite protein; apoE, apolipoprotein E; GlcN, 2-deoxy-2 aminoglucopyranose; IdoA, idopyranosyluronic acid; S, sulfate; CHAPS, 3-(β-cholamidopropyl)dimethylammonium)-1-propanesulfonic acid; ΔUA, 4-deoxy-α-L-threo-hexenepyranosyluronic acid; SPR, surface plasmon resonance; HPLC, high performance liquid chromatography; ΔDi, 2-acetamido-2-deoxy-3-O-(β-d-gluco-4-enopyranosyluronic acid)-β-d-galactose(glucose).

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EXPERIMENTAL PROCEDURES

Materials—Actinase E was obtained from Kaken Pharmaceutical Co. Ltd. (Tokyo, Japan). The following reagents were purchased from Sigma: urea, CHAPS, N-hydroxysulfosuccinimide sodium salt, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide, SDS, DNase I, endoglycosidase H (from *Streptococcus plicatus*, recombinant expressed in *Escherichia coli*), α-mannosidase (from *Canavalia ensiformis*), β-mannosidase (from *Helix pomatia*), β-N-acetylgalactosaminidase (from *C. ensiformis*). Bovine serum albumin was from Amresco (Solon, OH). Chondroitinase ABC (EC 4.2.2.4), chondroitinase ACII (EC 4.2.2.5), chondro-4-sulfatase (EC 3.1.6.9), heparin lyase I (EC 4.2.2.7, heparinase I, heparinase II), and heparin lyase III (EC 4.2.2.8, heparinase III, heparitinase I) were obtained from Sigma and Seikagaku Kogyo. Heparin and heparan sulfate, both from porcine intestine, were from Celsus Laboratories. Heparan sulfate from porcine liver was prepared as previously described (4). Recombinant CSP was made in E. coli (DH5α) transfected with pCS27IVC. The plasmid contains the complete *P. falciparum* CSP sequence from the T4 isolate, except that the hydrophobic NH2- and COOH-terminal amino acids 1–26 and 412–424 have been deleted, and 5 histidine residues were added to the COOH terminus to facilitate purification. The protein used in this study was kindly provided by Dr. Bela Takacs (Hoffman-La Roche, Basel, Switzerland) (25).

Standard unsaturated heparan sulfate disaccharides were purchased from Sigma: 2-acetamido-2-deoxy-4-O-(4-deoxy-α-D-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA-GlCNcA), 2-deoxy-2-sulfamido-4-O-(4-deoxy-α-D-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA-GlCNs), 2-acetamido-2-deoxy-4-O-(4-deoxy-α-D-threo-hexenopyranosyluronic acid)-6-O-sulfo-D-glucose (ΔUA-GlCNac6s), 2-acetamido-2-deoxy-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNcA), 2-deoxy-2-sulfamido-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNs), 2-acetamido-2-deoxy-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNac6s), 2-deoxy-2-sulfamido-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNs), 2-acetamido-2-deoxy-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNac6s), 2-deoxy-2-sulfamido-4-O-(4-deoxy-2-O-sulfo-α-L-threo-hexenopyranosyluronic acid)-D-glucose (ΔUA2S-GlCNs), and heparan sulfate (from bovine kidney). Standard unsaturated disaccharides from

FIGURE 1. The life cycle of *Plasmodium* in the mammalian host (A) and the mosquito (B). A, sporozoites are injected during blood feeding by infected anopheline mosquitoes, go to the liver, and invade hepatocytes, where they divide, rupture from the hepatocyte, and enter erythrocytes. The released merozoites invade erythrocytes, mature, divide, rupture from the cell, and enter new erythrocytes. The asexual erythrocytic cycle gives rise to the clinical symptoms associated with malaria. Some merozoites differentiate to gametocytes, which are infective for mosquitoes. This image is reprinted with permission from Ref. 51. B, mosquitoes become infected with *Plasmodium* when they ingest gametocytes (1) during blood feeding. In the midgut (MG) of the mosquito, these cells differentiate (2), fertilization occurs, and the resultant zygote differentiates into an ookinete (3), which traverses the midgut wall (4) and comes to rest extracellularly between the basal lamina and the midgut (5). The ookinete then differentiates into an oocyst, and sporozoites develop (6), and when mature, they emerge into the hemocoe (HC) of the mosquito and invade salivary glands (SG) (7). When the mosquito takes another blood meal, these salivary gland sporozoites are injected into the mammalian host (8), thus continuing the cycle. This image is reprinted with permission from Ref. 52.
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Structure of heparan sulfate glycosaminoglycans. A, unsulfated disaccharide sequence prominent in heparan sulfate. B, fully sulfated disaccharide sequence prominent in heparin. C, structural variants of glucuronic acid containing disaccharide sequences found in heparan sulfate and heparin $\left( X = SO_3^- \right)$. D, structural variants of iduronic acid containing disaccharide sequences found in heparan sulfate and heparin $\left( X = SO_3^- \right)$. E, proposed apoE and CSP binding domain.

chondroitin sulfate were obtained from Seikagaku Kogyo Co. (Tokyo): 2-acetamido-2-deoxy-3-O-(β-D-glucopyranosyluronic acid)-D-glucose (ΔDi-UA2S), 2-acetamido-2-deoxy-3-O-(β-D-glucopyranosyluronic acid)-4-O-sulfo-D-galactose (ΔDi-diS$_3$), 2-acetamido-2-deoxy-3-O-(2-O-sulfo-β-D-glucopyranosyluronic acid)-4-O-sulfo-D-galactose (ΔDi-diS$_4$), 2-acetamido-2-deoxy-3-O-(2-O-sulfo-β-D-glucopyranosyluronic acid)-6-O-sulfo-D-galactose (ΔDi-diS$_5$), 2-acetamido-2-deoxy-3-O-(β-D-glucopyranosyluronic acid)-4,6-di-O-sulfo-D-galactose (ΔDi-diS$_6$), and 2-acetamido-2-deoxy-3-O-(2-O-sulfo-β-D-glucopyranosyluronic acid)-4,6-di-O-sulfo-D-galactose (ΔDi-triS) were obtained from Sigma and Seikagaku Kogyo. M$_r$ cut-off 3,000 dialysis tubing was purchased from Fisher. Bio-Gel P-10 polyacrylamide gels were purchased from Bio-Rad. The Vivapure Q column was purchased from Viva Science (Bedford, MA). All other chemicals used were of analytical reagent grade.

Apparatus—The chromatographic equipment included a gradient pump (L-7000), a chromato-integrator (D-7500) from Hitachi Instruments (Tokyo, Japan), a double plunger pump for the reagent solution and reaction heater (Shimamura Instruments, Tokyo, Japan), a sample injector with a 20-μl loop (model 7125; Reodyne, Rohnert Park, CA), and a fluorescence spectrophotometer (FP-1580; JASCO, Tokyo, Japan). SPR measurements were performed on a BIAcore 3000 operated using BIAevaluation software (version 4.1). 1H NMR spectra were acquired on a Varian 500 MHz spectrometer at 25 °C in D$_2$O.

Insects—Adult A. stephensi were raised as previously described (26). Our analysis was performed with 10–14-day-old adult female mosquitoes that had never had a blood meal. For organ harvest, mosquitoes were placed on ice, rinsed briefly in 70% ethanol, and transferred to buffered saline (150 mM NaCl, 4 mM KCl, 3 mM CaCl$_2$, 1.8 mM NaHCO$_3$, 0.6 mM MgCl$_2$, 25 mM Hepes, pH 7). Dissections were performed in buffered saline, and dissected organs were placed into glass tubes and kept on ice. After dissection, tubes were spun at 200 × g for 5 min to pellet the dissected organs, excess buffer was removed, and the organs were frozen at −80 °C until use. Whole mosquitoes were counted, washed in ethanol as above, rinsed in buffered saline, and then frozen at −80 °C. Weights of midguts and salivary glands were measured using a Cahn microbalance (model C-34; ATI). Organs were dissected dry and then transferred for weighing using siliconized capillary pipettes. Midguts were weighed in groups of two, and pairs of salivary glands were weighed in groups of five.

Preparation of Crude Glycosaminoglycans for Microanalysis—Mosquito samples for disaccharide analysis were first lyophilized to dryness. Lyophilized sample (5 mg) was homogenized with acetone (1 ml), and the homogenate was washed with excess acetone and dried. The pellet was extracted with a solution containing SDS (1.0 ml 0.5%), NaOH (0.1 ml), NaBH$_4$ (0.8%) with constant stirring for 16 h at room temperature. Sodium acetate (200 μl of 1 M solution) and HCl (300 μl of 1 M solution) were then added, the solution was filtered, and HCl obtained from Millipore Corp. (Bedford, MA).
(200 µl, 1 m) was added to the filtrate. Insoluble material was removed by centrifugation at 2,500 × g for 10 min at 4 °C. Ethanol (7 ml) was added to the supernatant and chilled for 2 h at 0 °C, and the crude glycosaminoglycan fraction was collected by centrifugation at 2,500 × g for 10 min at 4 °C. The resulting precipitate was dissolved in 250 µl of water, and the crude glycosaminoglycan solution was stored frozen (24).

Disaccharide Microanalysis—Crude glycosaminoglycan solution (20 µl) was diluted to 100 µl with water and used as the sample solution for chondroitin disaccharide microanalysis. Chondroitinase digestion was performed using Tris-acetate buffer (5 µl, 0.2 m, pH 8.0), and 10 µl of an aqueous solution containing chondroitinase ABC (50 mIU) and chondroitinase ACII (50 mIU) were added to a 20-µl portion of the sample solution and incubated at 37 °C for 3 h. An 8-µl portion of this mixture was analyzed by high performance liquid chromatography. For heparan sulfate microanalysis, crude glycosaminoglycan sample (230 µl) and sodium phosphate buffer (50 µl, 0.3 m, pH 6.0) were added, and the solution was applied on an Ultrafree-MC DEAE membrane, which had been equilibrated with sodium phosphate buffer (pH 6.0) containing NaCl (0.15 m). The fractions eluted with NaCl (1.0 M) were collected, desalted with Biomax-5, evaporated, and resuspended in 12 µl of water in preparation of heparin lyase digestion. Next, sodium acetate buffer (5 µl, 0.1 m, pH 7.0) containing calcium acetate (10 m.mx) and 15 µl of an aqueous solution containing heparin lyase mixture (Seikagaku), heparin lyase I (1 mIU), heparin lyase II (1 mIU), and heparin lyase III (1 mIU) were added to a 5-µl portion of sample. The mixture was incubated at 37 °C for 16 h, and an 8-µl aliquot was loaded onto the high performance liquid chromatograph (24).

Determination of Unsaturated Disaccharides from Heparan Sulfate and Chondroitin Sulfate—Unsaturated disaccharides produced enzymatically from heparan sulfate and chondroitin sulfate were analyzed by a reversed-phase ion pair chromatography with sensitive and specific postcolumn detection. A gradient was applied at a flow rate of 1.1 ml/min using a Senshu Pak Docosil column (4.6 × 150 mm) at 55 °C. The eluents used were as follows: A, H2O; B, 0.2 m sodium chloride; C, 10 mM tetra-n-butylammonium hydrogen sulfate; D, 50% acetonitrile. The gradient program was as follows: 0–10 min, 70–67% eluent A, 1–4% eluent B; 10–11 min, 67–56% eluent A, 4–15% eluent B; 11–20 min, 56–46% eluent A, 15–25% eluent B; 20–22 min, 46–18% eluent A, 25–53% eluent B; 22–29 min, 18% eluent A, 53% eluent B; equilibration with 70% A and 1% B for 20 min. The proportions of eluents C and D were constant at 12 and 17%, respectively. A solution of 2-cyanoacetamide (aqueous, 0.5% (w/v)) and sodium hydroxide (0.25 m) was added at the same flow rate (0.35 ml/min) to the effluent using a double plunger pump. The mixture passed through a reaction coil (internal diameter, 0.5 mm; length, 10 m) set in a dry reaction temperature-controlled bath at 125 °C and a following cooling coil (internal diameter, 0.25 mm; length, 3 m). The effluent was monitored fluorometrically (excitation, 346 nm; emission, 410 nm) (24).

Large Scale Preparation and Purification of Heparan Sulfate from Whole Mosquitoes for Surface Plasmon Resonance (SPR) Studies—Approximately 8000 mosquitoes were ground with dry ice, washed with acetone three times to remove lipids, and then dried in vacuo. Mosquito acetone powder was dissolved in 100 ml of water, and 20 ml of 0.05 m Tris-acetate buffer (pH 8.0) containing 1% actinase E was added and incubated at 45 °C overnight. After actinase E digestion, 2 ml of 100 mM acetic acid was added and boiled for 5 min in a water bath and centrifuged at 1,000 × g for 20 min. The supernatant was dialyzed against water in M, cut-off 3,000 dialysis tubing overnight and freeze-dried. The resulting peptidoglycan (glycosaminoglycan linked to residual core peptide) from mosquitoes was dissolved in 8 m urea, 2% CHAPS buffer (pH 8.3) and loaded onto the Vivapure Q spin column and centrifuged at 500 × g for 15 min. The column was then washed with 8 m urea, 2% CHAPS buffer and then washed five times with 50 mM NaCl. The bound peptidoglycan was released by washing the spin column three times with 2.7 m (16%) NaCl at 500 × g. Methanol (final concentration 80%) was added to the eluent, and the peptidoglycan component was allowed to precipitate overnight at 4 °C. The peptidoglycan fraction was collected by centrifugation at 2,500 × g for 20 min. Core peptide was removed by β-elimination, by dissolving the isolated peptidoglycan in 2 ml of 0.5% SDS, 0.1 m NaOH, 0.8% NaBH4, and incubated at room temperature for 16 h with constant stirring. Sodium acetate (400 µl of 1 m) and HCl (600 µl of 1 m) were then added, the solution was filtered, and HCl (400 µl of 1 m) was added to the filtrate. Finally, the solution was dialyzed against M, cut-off 3,000 dialysis tubing overnight and lyophilized. To remove chondroitin sulfate, the resulting glycosaminoglycan sample was redissolved in 0.2 m Tris-HCl buffer and incubated with a mixture of chondroitinase ABC and AC II overnight at 37 °C. After digestion, the sample was dialyzed against M, cut-off 3,000 dialysis tubing overnight to remove the chondroitin disaccharide that had been formed and then freeze-dried. Residual DNA was removed by dissolving the sample in DNase I reaction buffer (10 mm Tris-HCl, 2.5 mM MgCl2, 0.5 mM CaCl2, pH 7.6), incubating with DNase I at 37 °C overnight. Chromatography on a P-10 column (1 × 50 cm) eluted with water removed DNase and nucleotides, and intact glycosaminoglycan was recovered and lyophilized. 1H NMR spectroscopy (not shown) suggested the presence of residual contaminating mannose-containing glycans. These contaminating glycans were removed by dissolving the sample in 250 mM sodium phosphate buffer (pH 5.5) and incubating with endoglycosidase H at 37 °C for 48 h followed by α-mannosidase, β-mannosidase, and β-N-acetylgalactosami- nidade for 24 h at 25 °C. The sample was dialyzed against water in M, cut-off 3,000 dialysis tubing overnight and lyophilized. All of the enzymes (proteins) that had been added were then removed by dissolving the sample in 8 m urea, 2% CHAPS buffer (pH 8.3) and passing through a Vivapure Q spin column at 500 × g for 15 min. The column was washed with 8 m urea, 2% CHAPS buffer, followed by washing five times with 50 mM NaCl. The heparan sulfate was released from the column with three washes of 2.7 m (16%) NaCl at 500 × g. The heparan sulfate was precipitated at 4 °C overnight by adding absolute methanol (final concentration 80%). The pure heparan sulfate was collected by centrifugation at 2,500 × g for 20 min. A carbazole assay (27) showed that 532 µg of heparan sulfate was obtained.
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*Surface Plasmon Resonance Studies—HBS-N buffer was used for immobilization and interaction studies. CM5 chips were activated with 50 mM N-hydroxysuccinimide sodium salt and 200 mM N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide mixture for 7 min at a flow rate of 5 µl/min. CSP was immobilized at a concentration of ~700 nM. The surfaces were deactivated with 1 M ethanolamine hydrochloride, pH 8.5. The final response units after ethanolamine was 8,000–10,000. The control surface was prepared the same way, except ~6000 response units of bovine serum albumin was immobilized. At a flow rate of 5 µl/min and temperature of 25 °C, the interaction between CSP and various concentrations (1, 5, 10, 20, and 50 mM) of heparan sulfate was studied, selecting an interaction with an association time of 3 min and dissociation time of 4 min. Between each injection, the surface was regenerated with 2 M NaCl. The interaction was monitored by plotting the increase in response over time (sensorgram), and the control flow cell was subtracted from the experimental flow cell. The curves were evaluated with BIAevaluation version 4.1.*

**RESULTS**

Preliminary studies on salivary glands, prepared from adult female *A. stephensi* mosquitoes, showed that they contained both chondroitin sulfate and heparan sulfate. The amount of chondroitin sulfate in the mosquito, however, appeared to be considerably lower than that reported in the whole body of *D. melanogaster* third stage larvae (24). Based on these preliminary findings, hundreds of adult female *A. stephensi* mosquitoes and selected tissues from these mosquitoes were collected. Salivary glands, midguts, and malpighian tubules with ovaries were collected in buffered saline, centrifuged to remove excess buffer, frozen, and then ground into a fine powder for analysis. The total dry mass, prior to removing salts and lipids, of each sample is presented in Table 1. The dry mass of individual mosquitoes or organs was obtained by either dividing the total mass by the number of individuals or by obtaining wet weights on a Cahn microbalance and correcting for water content. Salivary glands (in pairs) and midguts corresponded to 0.6 and 1.2% of the total insect body mass, respectively.

The whole insect and these isolated organs were extracted by a method previously developed for use with *D. melanogaster* (24) to recover total glycosaminoglycans. A single mosquito was found to contain 6.0 ng of chondroitin sulfate and 11.3 ng of heparan sulfate (Tables 2 and 3). We used mosquitoes that had taken a blood meal 1 week prior to the analysis, because blood feeding initiates changes in the mosquito’s transcriptional repertoire (28), which could, in turn, alter the composition of the cell barriers encountered by malaria parasites (15). Analysis of whole rabbit blood, however, showed that it contained 3–4 ng/µl glycosaminoglycans, making it theoretically possible that the blood meal contributed to the chondroitin sulfate and heparan sulfate detected in the mosquitoes. To control for this possibility, we com-

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**TABLE 1**

Samples prepared for analysis

| Sample                        | Total mass | Number | Mass/insect or organ | Percentage of insect body mass |
|-------------------------------|------------|--------|----------------------|--------------------------------|
| Whole *A. stephensi* mosquitoes | 71         | 150    | 340                  | 100                            |
| Salivary glands               | 4.3        | 486    | 2.0                  | 0.6                            |
| Midguts                       | 3.7        | 247    | 4.1                  | 1.2                            |
| Malpighian tubules and pairs of ovaries | 10 | 320     | ND                  | ND                             |

* Based on the dry weight of 1,000 insects multiplied by 0.001.
* Calculated from six independent determinations of the wet weight of either two midguts or five pairs of salivary glands. The dry wet weight of each organ was calculated by multiplying the wet weight by 0.25 (corresponding to a water content of 75%).
* ND, not determined.

**TABLE 2**

Unsaturated disaccharides from chondroitin sulfate (CS)

| Unsatuated disaccharide | Sulfo group/disaccharide | Amount of chondroitin sulfate |
|-------------------------|--------------------------|-------------------------------|
|                         | mol %                    | ng of CS/mg of tissue         | ng of CS/insect or organ      |
|                         |                          | mg/g                          | mg                             |
| Female *A. stephensi* mosquitoes | 73.4 | 26.6 | 0.3 | 17.6 | 6.0 |
| Mosquito salivary glands | 78.1 | 21.9 | 0.3 | 21.6 | 0.043 |
| Mosquito midguts        | 69.2 | 30.8 | 0.3 | 6.4 | 0.026 |
| Mosquito malpighian tubules and ovaries | 54.1 | 45.9 | 0.5 | 25.2 | ND |
| *D. melanogaster* (adult)* | 70.8 | 29.2 | 0.3 | 126 | 25 |

* ND, not determined.
* The mass of a dry fruit fly is 200 µg (see Ref. 24).

**TABLE 3**

Unsaturated disaccharides from heparan sulfate

| Heparan sulfate source | Amount of heparan-sulfated unsaturated disaccharide | Sulfo group/disaccharide | Dry tissue | HS/insect or organ |
|------------------------|-----------------------------------------------|--------------------------|------------|-------------------|
|                        | mol %                                        |                         |            |                   |
| Female *A. stephensi* mosquitoes | 38.9 | 24.4 | 4.8 | ND | ND | 3.3 | 1.0 | 33.1 | 11.3 |
| Mosquito salivary glands | 41.3 | 22.8 | 3.2 | ND | 19.4 | 10.3 | 3.0 | 0.9 | 23.0 | 0.045 |
| Mosquito midguts        | 35.3 | 37.8 | 1.1 | ND | 11.1 | 12.7 | 2.0 | 0.9 | 50.4 | 0.2 |
| Mosquito malpighian tubules and ovaries | 40.0 | 29.6 | 3.0 | ND | 16.6 | 9.1 | ND | 1.7 | 0.9 | 42.3 | ND |
| *D. melanogaster* (adult)* | 42.6 | 18.8 | 4.4 | ND | 15.4 | 14.5 | ND | 4.3 | 1.0 | 19.7 | 3.9 |
| Human liver              | 37.3 | 15.5 | 9.8 | 1.2 | 7.8 | 5.9 | 0.7 | 21.8 | 1.2 | ~8 |

* ND, not detected.
* See Ref. 24.
* See Ref. 5.
pared the chondroitin sulfate and heparan sulfate content of blood-fed mosquitoes (1 week after blood feeding) with age-matched naive mosquitoes and found no difference in disaccharide composition between the two groups (data not shown).

Next we performed a disaccharide analysis of whole mosquitoes and mosquito organs. Treatment of total purified intact glycosaminoglycans with chondroitin ABC lyase afforded chondroitin disaccharides. The resulting disaccharide mixture was analyzed by reversed-phase ion pairing high performance liquid chromatography (HPLC) using ultrasensitive detection following postcolumn fluorescence derivatization (24, 29) (Fig. 3 and Table 2). The whole insects and insect organs contained 54–78% ΔUA (1 → 3) GalNAc (OS) and 22–46% ΔUA (1 → 3) GalNAc4S (4S) disaccharide. Six of the eight possible chondroitin disaccharides were not present, since no traces of 6-O-sulfo, 2-O-sulfo, disulfated, or trisulfated chondroitin disaccharides could be observed. The chondroitinase-sensitive glycosaminoglycans obtained from mosquitoes contained no hyaluronic acid or dermatan sulfate. The presence of hyaluronic was ruled out using a graphite column that can discriminate between glucosaminoglycans and galactosaminoglycans (29). The presence of dermatan sulfate was ruled out, since the disaccharide analysis with chondroitinase AC was identical to that with chondroitinase ABC (24). These data on mosquito and mosquito organs are consistent with the known absence of dermatan sulfate and hyaluronan in D. melanogaster (24).

Interestingly, the whole mosquito contained only one-fourth as much chondroitin sulfate as the fly on a per insect basis. In contrast, the whole mosquito contained ~3 times as much heparan sulfate as the fly on a per insect basis. The same glycosaminoglycan samples from whole mosquito and isolated organs were collected and treated with a mixture of heparin lyase I, II, and III, to convert the heparan sulfate into disaccharides. Disaccharide analysis (Fig. 4 and Table 3) revealed that all samples contained six of the eight common heparan sulfate disaccharides, with ΔUA2S (1 → 4) GlcNAc (2SNAc) and ΔUA2S (1 → 4) GlcNAc6S (2SNAc6S) conspicuously absent. The ΔUA2S (1 → 4) GlcNS6S trisulfated disaccharide, probably arising from heparin lyase cleavage of the (→3)-GlcNS6S(1 → 4)IdoA2S(1 → 4) sequence, was found in all of the mosquito samples. No clear patterns or differences in disaccharide composition between different insect organs could be discerned. Comparison of the composition of these samples with human liver heparan sulfate clearly showed a lower degree of sulfation, 0.9–1.0 sulfo groups/disaccharide for the insect samples compared with 1.2 and 1.0 sulfo groups/disaccharide for the human and porcine liver heparan sulfate (5).

Domain mapping of whole mosquito heparan sulfate was undertaken by treatment with heparin lyase III, followed by polyacrylamide gel electrophoresis (30) (not shown). The results of this analysis demonstrated the presence of heparin lyase III-resistant (high sulfate) domains ranging in size from hexasaccharide to hexadecasaccharide.

Next, ~8,000 mosquitoes were collected and used to prepare ~0.5 mg of heparan sulfate required for CSP binding studies. A number of additional steps were added to ensure the purity of the heparan sulfate and the absence of contaminating nucleic acids, proteins, and non-heparan sulfate glycans. The interaction of the purified mosquito heparan sulfate with immobilized CSP was studied using SPR and compared with previous results obtained on the interaction of CSP with porcine heparin and heparan sulfate (10, 11). The resulting sensorgrams all afforded excellent fitting to a 1:1 conformational change model with compensation for mass transfer limitations. The sensorgrams
for mosquito heparan sulfate binding to immobilized CSP are shown in Fig. 5. A single concentration, corresponding to 10 μM, was selected to compare the binding to CSP of mosquito heparan sulfate with that of heparin and other heparan sulfates (Table 4). Heparin bound with a much higher affinity than all other ligands, with porcine liver heparan sulfate binding with the second highest affinity. Mosquito heparan sulfate showed a CSP binding affinity comparable with porcine intestinal heparan sulfate. It is noteworthy that the heparan sulfate from mosquitoes, while fitting the 1:1 conformational change model quite well, gave very different binding kinetics from the samples prepared from mammalian tissues. The kinetics of mosquito heparan sulfate was dominated by its slow on-rate and off-rate, followed by a very slow conformational change.

DISCUSSION

The proteoglycans chondroitin sulfate and heparan sulfate, as well as the structurally related heparin (31), have been observed in a wide variety of animals, ranging from mammals (3–6, 32, 33), to birds (34), clams (35), fruit flies, and worms (24). Nothing, however, is known about these molecules in insects of medical importance, where they may play a role in the transmission of pathogens. The recent genome sequencing of two medically important mosquito species, A. gambiae and Aedes aegypti demonstrates that the genes encoding the biosynthetic machinery necessary for synthesis of heparan and chondroitin sulfates are present in these species. Our current work in A. stephensi, a species closely related to A. gambiae, demonstrates that these molecules are indeed biosynthesized and elucidates their fine structure.

Heparan sulfate is found linked to a number of core proteins, including members of the syndecan family (integral membrane proteins), the glypican family (GPI-anchored proteins), and the perlecan proteoglycans (extracellular matrix proteins) (36). The heparan sulfate glycosaminoglycan side chains of these proteoglycans serve an essential role in cell-cell interactions and cellular communication (37, 38). Of relevance to our findings, heparan sulfate glycosaminoglycan chains are involved in cell-cell adhesion by binding to a variety of different proteins, such as selectins, fibronectin, laminin, and vitronectin (38, 39). The interaction of heparan sulfate with heparin-binding proteins is believed to be specific and related to the sequence within heparan sulfate (40). The sequences within a given heparan sulfate chain are the result of its biosynthesis in the Golgi through the sequential action of N-deacetylase, N-sulfotransferase, C5 epimerase, and O-sulfotransferases. Interestingly, some of these enzymes have multiple isoforms that are differentially expressed in a tissue-dependent and a time-dependent fashion, leading to a complex and little understood mixture of glycotopes.

A number of infectious agents are known to use the negative charge of cell surface glycosaminoglycans to localize, enter, and infect animal cells (41, 42). Viruses, including dengue virus (43), herpes simplex virus 1 (44), hepatitis C virus (45, 46), and human immunodeficiency virus-1 (47), are known to utilize heparan sulfate as a receptor. Herpes simplex virus 1 targets heparan sulfate having a unique sequence found in nervous tissue (44) and requires 3-O-sulfation of specific
TABLE 4
SPR data for a 1:1 conformational change model with mass transport limitation at 10 μm analyte glycosaminoglycan concentration binding to immobilized CSP

| Sample              | $k_1$  | $k_2$  | $k_{-1}$ | $k_{-2}$ | $R_{max}$ | $x^2$ | $K_D$ |
|---------------------|--------|--------|----------|----------|------------|-------|-------|
| Heparin             | $2.3 \times 10^5$ | $5.0 \times 10^{-2}$ | $1.5 \times 10^{-2}$ | $3.4 \times 10^{-4}$ | 216 | 0.356 | 0.05 |
| Porcine liver HS    | $1.1 \times 10^4$ | $5.1 \times 10^{-2}$ | $1.2 \times 10^{-2}$ | $5.7 \times 10^{-4}$ | 149 | 0.876 | 0.2  |
| Porcine intestinal HS | $2.9 \times 10^2$ | $2.8 \times 10^{-2}$ | $1.1 \times 10^{-2}$ | $1.9 \times 10^{-3}$ | 102 | 0.211 | 2    |
| Mosquito HS         | $5.0 \times 10^2$ | $5.8 \times 10^{-4}$ | $5.7 \times 10^{-6}$ | $1.0 \times 10^{-5}$ | 131 | 0.277 | 2    |

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