HEPARAN SULFATE CHAINS OF SYNDECAN-1 REGULATE ECTODOMAIN SHEDDING*

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Running title: Heparan sulfates suppress syndecan shedding

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Background: Syndecan ectodomains shed by cells can enhance progression of cancer, inflammatory disease and pathogen infection. Results: Reducing the amount of heparan sulfate present on syndecan core proteins increases shedding of the ectodomain.

Conclusion: Heparan sulfate chains suppress syndecan shedding.

Significance: Therapeutic inhibition of heparan sulfate degradation could slow disease progression.

SUMMARY

Matrix metalloproteinases (MMPs) release intact syndecan-1 ectodomains from the cell surface giving rise to a soluble, shed form of the proteoglycan. Although it is known that shed syndecan-1 controls diverse pathophysiological responses in cancer, wound healing, inflammation, infection and immunity, the mechanisms regulating shedding remain unclear. We have discovered that the heparan sulfate chains present on syndecan core proteins suppress shedding of the proteoglycan. Syndecan shedding is dramatically enhanced when the heparan sulfate chains are enzymatically degraded or absent from the core protein. Exogenous heparan sulfate or heparin does not inhibit shedding, indicating that heparan sulfate must be attached to the core protein to suppress shedding. Regulation of shedding by heparan sulfate occurs in multiple cell types, for both syndecan-1 and syndecan-4 and in murine and human syndecans. Mechanistically, the loss of heparan sulfate enhances the susceptibility of the core protein to proteolytic cleavage by MMPs. Enhanced shedding of syndecan-1 following loss of heparan sulfate is accompanied by a dramatic increase in core protein synthesis. This suggests that in response to an increase in the rate of shedding, cells attempt to maintain a significant level of syndecan-1 on the cell surface. Together these data indicate that the amount of heparan sulfate present on syndecan core proteins regulates both the rate of syndecan shedding and core protein synthesis. These findings assign new functions to heparan sulfate chains thereby broadening our understanding of their physiological importance and implying that therapeutic inhibition of heparan sulfate degradation could impact the progression of some diseases.

Syndecan-1 is known predominantly as a cell surface proteoglycan that regulates complex cell behaviors such as adhesion, motility, invasion and intracellular signaling. Syndecan-1 performs these diverse functions by acting as a co-receptor for a variety of growth factor receptors or as a binding partner to integrins (1,2). However, syndecan-1 localization is not restricted to the cell surface. It is present in the nucleus where it regulates gene transcription (3), or it can be shed from the cell...
surface and become a soluble or insoluble component of the extracellular matrix (4,5). Shed syndecan-1 retains its glycosaminoglycan chains (both heparan sulfate and chondroitin sulfate) (6) along with their bound ligands which often include growth factors, chemokines and other effectors that can endow the shed proteoglycans with both autocrine and paracrine signaling capacity.

Shed syndecan-1 plays important roles in the pathophysiology of many disease states including inflammation, microbial pathogenesis and cancer (4,5,7). For example, in a murine model of acute lung injury, syndecan-1 mediates the inflammatory response by facilitating the formation of a CXC chemokine gradient to enhance transepithelial migration of neutrophils (8). In pulmonary fibrosis, reactive oxygen species enhance syndecan-1 shedding which leads to induction of neutrophil chemotaxis, impaired epithelial wound healing and increased fibrosis (9).

Shed syndecan-1 also plays an important role in microbial pathogenesis and host defense (4). For example, Pseudomonas aeruginosa induces syndecan-1 shedding through LasA, an enzyme that activates MMP1-mediated shedding of syndecan-1 (10). The shed syndecan-1 inhibits host derived anti-microbial peptides and thus is a component of a virulence mechanism that promotes infection (11).

In cancer, shed syndecan-1 plays an active role in driving tumor progression (5). In a model of breast cancer, syndecan-1 shed from the surface of reactive stromal fibroblasts stimulates signaling and proliferation in adjacent tumor cells (12). Syndecan-1 shed by myeloma cells enhances angiogenesis, osteolysis, growth and spontaneous metastasis of tumor cells in vivo (13-15). Shed syndecan-1 can also be measured in the serum of patients with some cancers including lung cancer and myeloma where high levels of shed syndecan-1 correlates with poor outcome of the patient (16,17).

Although the mechanisms mediating syndecan-1 shedding from the cell surface are not completely understood, shedding occurs via proteolytic cleavage of the core protein close to the cell membrane. Shedding of syndecan-1 occurs constitutively or can be accelerated in response to various agonists, for example, growth factors, chemokines, microbial toxins, insulin, or cellular stress. These diverse stimuli set off a cascade of molecular events that include signal transduction, intracellular regulators and finally, protease activity at the cell surface that cleaves the core protein and releases syndecan-1 (4,5). Intracellularly, activation of protein tyrosine kinases leads to downstream phosphorylation of the syndecan-1 cytoplasmic domain, an event that may enhance syndecan-1 shedding (18,19). In addition, shedding agonists cause dissociation of Rab5 from the syndecan-1 cytoplasmic domain which triggers ectodomain shedding (20). The actual cleavage of syndecan-1 from the cell surface is accomplished by various metalloproteinases including MMPs, MT-MMPs, and ADAMTS (5).

We previously demonstrated that the action of heparanase, an enzyme that cleaves heparan sulfate chains, stimulates increased syndecan-1 shedding (21). We found that increased shedding is due at least in part to heparanase-mediated upregulation of expression of MMP-9 and uPA/uPAR, two enzymes known to be sheddases of syndecan-1 (22). In addition to upregulating proteases, heparanase trims heparan sulfate chains of syndecan-1 resulting in a significantly smaller amount of heparan sulfate present on the proteoglycan. However, a role for such reduction in heparan sulfate content leading to shedding has never been tested. In the present study, we demonstrate that this reduction in heparan sulfate amount dramatically accelerates the rate of syndecan-1 shedding and that this can be blocked by inhibitors of MMPs. Thus, heparan sulfate present on syndecan-1 acts to suppress shedding of the ectodomain possibly by interfering with MMP-mediated cleavage of the core protein. This suppression occurs in multiple cell types, in both
human and murine species and for syndecan-4 as well as syndecan-1. We also find that in response to enhanced shedding, a reduction in syndecan-1 heparan sulfate causes a dramatic increase in synthesis of the core protein of syndecan-1. These data provide new insight into a previously unknown role for the heparan sulfate chains of syndecans.

**EXPERIMENTAL PROCEDURES**

**Cell lines and reagents** - Cell lines CAG, RPMI-8226, ARH-77, and MPC-11 were grown in RPMI-1640 medium supplemented with 10% fetal bovine serum. U266 cells were grown in RPMI-1640 medium supplemented with 15% fetal bovine serum. RPMI-8226, ARH-77, and U266 cells were obtained from American Type Culture Collection (Manassas, VA). Cell lines Panc-1 (a kind gift from Dr. Lacey McNally, University of Alabama at Birmingham), MDA-MB-231 (ATCC) and HeLa (a kind gift from Dr. Selvarangan Ponnazhagan, University of Alabama at Birmingham) were cultured in DMEM supplemented with 10% FBS. Murine mammary epithelial cell line, NMuMG cells were grown in DMEM supplemented with 10% FBS and insulin (23). Bacterial heparinase III (Hep III) and chondroitinase ABC (Chase-ABC) were purchased from Seikagaku (Kogyo, Japan), phorbol 12-myristate 13-acetate, cycloheximide, cytochalasin, heparin and purified bovine kidney heparan sulfate from Sigma-Aldrich (Milwaukee, WI) and inhibitors BB-94, U0126, bisindolylmaleimide, and genistein from Calbiochem (La Jolla, California).

**Constructs and transfections** - Human syndecan-1: The full length human syndecan-1 gene in vector pcDNA3.1(+) (Invitrogen, Carlsbad, CA) was used as the template to create syndecan-1 with a TEV cleavage and C-terminal 10xHis tag, and the mutated form of syndecan-1 with the same tag. Mutagenesis was carried out with the Change-It kit (USB/Agilent, Palo Alto, CA), following the protocols therein, as it allows the incorporation of multiple mutations in a single round of PCR. Phosphorylated insertion mutagenic oligonucleotides were synthesized by Integrated DNA Tech., Inc (IDT, Coralville, IA), and the phosphorylated substitution mutagenic oligonucleotides were synthesized by Invitrogen. The phosphorylated mutagenic insertion primer for the tag was 5’P-caccaacagaggaatta tatgccGGTTCTGTTCTGTTCGAATACCT GTATTTTCAGGCGGTTCATCATCATCACCACCA TCACCATCAACCAACCATGAAGCaaagcttaagttt aaacgc-3’, with the complete insertion sequence capitalized. The tag is followed immediately by a termination codon. Potential glycosylation sites at amino acid positions 37, 43, 45, 47, 206, 216 and 231 were replaced with alanine using the following primers in various combinations to incorporate all changes: S37AFor, 5’P- gatcaagatggeGCTggggatgactc-3’; S37ARev, 5’P-gaugatcatccccAGCgccatgtggtc-3’; S206AFor, 5’P-gagcacagagggcGCTgggagtgcgag-3’; S206ARev, 5’P-ctgcccAGCgctctctgcgc-3’; S216AFor, 5’P-cctggagaaccGGCgggaagatagc-3’; S216ARev, 5’P-cgatctctccccGCCgctcttgtgcag-3’; N231AFor, 5’P-ctgacgcggccggGGCagggcagggcag-3’; N231ARev, 5’P-ctggcgaggtgccgGGCagggcagggcag-3’. Sequences were confirmed after each round of PCR by generating DNA with a mini-plasmid kit (Qiagen, Santa Clarita, CA). The plasmids encoding for human wild type syndecan-1 sequence and the sequence for syndecan-1 mutated at glycosylation sites are referred to as WT hSDC1 and ΔGAG hSDC1, respectively. The murine full length syndecan-1 gene in pcDNA3.1(-) vector and the TDM clone have been previously described (24). TDM refers to a triple deletion mutant having all heparan sulfate attachment sites removed. The TDM clone was used as the template for additional alanine substitutions at amino acid residues 43, 207, and 217. A FLAG
tag with a TEV cleavage site was inserted towards the N-terminus, after the signal sequence. Phosphorylated mutagenic oligonucleotides for substitution mutagenesis were synthesized by Invitrogen and the FLAG tag insertion oligonucleotide was synthesized by IDT (Integrated DNA Tech, Inc.) The sequence of the phosphorylated mutagenic insertion primer, FlagNFor, was: 5’P-gctctggegctgggcctgccGATTATAAGGATGACGATGACAAAggtctGA AAACCTGTATTTTCCAGGCGccgctccctcccgca aattttgg -3’, with the FLAG and TEV sites in capital letters. Potential glycosylation sites at amino acid positions 43, 207, and 217 were replaced with alanine (shown in capital letters) using the following primers in various combinations to incorporate all changes: mN43A For, 5’P-ggatgactctgacGCCttcgcgggcgcgg-3’; mN43AREv, 5’P-cgegcegcegcaGCGtcaagacttcc-3’; mS207AFor, 5’P-gcaggagagggcGCTggagaacaacag-3’; mS207AREv, 5’P-tcttgattctcAGCcctctgc-3’; mS217AFor, 5’P-acctttga aacGCTggggagaacacag-3’; and mS217AREv, 5’P-ctgtggatcctcAGCtgtttcaaggt-3’. Sequences were confirmed after each round of PCR by generating DNA with a mini-plasmid kit (Qiagen). DNA was confirmed by the UAB CFAR DNA Sequencing and Analysis Core Facility. The plasmids encoding murine wild type syndecan-1 sequence and the sequence for syndecan-1 mutated at glycosylation sites will be referred as WT mSDC1 and ΔGAG mSDC1 respectively.

Prior to transfection of constructs, ARH-77 (2 \( \times \) 10^6) cells were plated in 5ml of fresh complete medium and grown for 24 h. The cells were then transfected with the murine WT or the ΔGAG SDC1 plasmids using Amaza Nucleofector™ (Amaza, Cologne) as per the manufacturer’s instructions. The transfected cells were then selected with Geneticin (G418; Invitrogen), for 7 days. The selected cells were stained for cell surface syndecan-1 expression and sorted by fluorescence-activated cell sorting (as described below) to sort cells expressing similar levels of either the WT or ΔGAG SDC1 protein. The sorted cells were grown for 7 days and the stable expression and levels of cell surface syndecan-1 was confirmed again by flow cytometry.

**Flow cytometry and FACS** - For fluorescence-activated cell sorting (FACS) or flow cytometry, cells were collected by centrifugation and washed twice in phosphate-buffered saline (PBS). Cell pellets were then resuspended in anti-human syndecan-1 (clone BA-38, Cell Sciences, Canton, MA), or anti-mouse syndecan-1 (clone 281.2, (25)) monoclonal antibody diluted in PBS and incubated at 4°C for 1 h. The cells were then washed three times in PBS and incubated with secondary antibody conjugated to Alexa Fluor 647 (Invitrogen) at 4°C for 1 h. Following incubation, the cells were washed three times in PBS and then resuspended in PBS. For flow cytometry, after staining cells were fixed in 4% paraformaldehyde prior to analysis. Cells stained with appropriate isotype-matched IgG antibodies served as a gating control.

**Western blotting** - Cells in culture were pelleted by centrifugation and washed with PBS twice. Cell pellets were then resuspended in lysis buffer (50 mM Tris, pH 7.5, 150 mM NaCl, 1 mM EDTA, 0.5% Triton X-100) containing 10 ug/ml leupeptin and 1X HALT Protease inhibitor cocktail and incubated on ice for 30 m. Lysates were centrifuged at 12,000 \( \times \) g at 4°C for 15 m and the supernatants were removed from the pellets. Proteins isolated from cell lines in the supernatant were quantified by a BCA protein assay reagent kit (Pierce, Rockford, IL). Equal amounts of protein were loaded onto 4–20% gradient SDS-polyacrylamide gels (Bio-Rad Laboratories, Richmond, CA), transferred to a positively charged nylon membrane (Nytran SPC; Schleicher & Schuell, Keene, NH) and probed with goat anti-human syndecan-1 (R&D Systems, Minneapolis, MN) followed by horseradish peroxidase-conjugated secondary anti-mouse antibody (GE Healthcare, Piscataway, NJ). The
levels of actin (loading control) were probed using a beta-actin primary antibody (Sigma-Aldrich). Immunoreactive bands were detected using enhanced chemiluminescence (GE Healthcare).

Preparation of conditioned media - In all experiments, unless mentioned otherwise, cells were seeded and incubated in serum-free RPMI medium for 1 h prior to addition of Hep III and/or chondroitinase ABC. At the end of incubation, cells and cellular debris were separated by centrifuging the conditioned media at 1000 rpm for 5 min and the supernatants were analyzed either by dot-blotting or ELISA. For adherent cells (MDA-MB-231, Panc-1, HeLa, NMuMG), equal numbers were seeded in complete growth medium in a 12 well plate. After overnight incubation, the monolayers were washed with serum-free medium at least twice prior to enzyme treatments.

Dot blotting - Conditioned medium collected from a) ARH-77 cells expressing either human or murine wild-type or mutated syndecan-1 b) NMuMG cells after different treatments, and c) HeLa cells after glycosidase treatments was processed prior to analyses by dot blotting. The conditioned media were treated with Hep III (1.25 mU/ml) and Chase-ABC (12.5 mU/ml) at 37°C for 4 h. This enzymatic digestion ensures that all the glycosaminoglycan chains from the shed syndecan-1 were removed prior to analyses and therefore an increase in shedding was determined only by comparing the differences in the level of syndecan-1 core protein between different samples. After processing, the indicated amounts of the conditioned media were blotted onto nitrocellulose membrane using an immunodot apparatus (Milliblot D, Millipore, Bedford, MA). The membranes were then fixed in Tris-buffered saline containing 0.05% glutaraldehyde for 30 min at room temperature. After washes in distilled water and TBS, the membranes were blocked in TBS containing 3% non-fat dry milk, 0.5% BSA and 0.3% Tween-20 (blocking solution) for 1 h. The membranes were incubated overnight with primary antibodies against murine or human syndecan-1 in the blocking solution. After washes in TBS containing 0.05% Tween-20 (washing solution), membranes were probed with horseradish peroxidase-conjugated secondary anti-mouse antibody (GE Healthcare) and immunoreactive bands were detected by chemiluminescence.

RNA extraction and real-time PCR for human syndecan-1 - 5 X 10^5 CAG cells were seeded in complete growth medium at 37°C and 5% CO₂. Cells were then treated with Hep III (1.25 mU/ml) added every two hours for the next 8 h. Cells treated with water served as a control. At the end of the incubation, cells were washed once with PBS and RNA was extracted using RNeasy columns (Qiagen, Valencia, CA). One microgram of total RNA was reverse transcribed to cDNA using random hexamer primers and Maxima® First Strand cDNA synthesis kit (Fermentas, Hanover, MD) as per manufacturer’s protocol. 25 ng of diluted cDNA was mixed with 2X IQ™ SYBR® green supermix (Bio-Rad, Hercules, CA), along with gene specific primers (200 nM final concentration). The primers used for syndecan-1 were 5’TCTGACAACCTTCTCCGTC3’ (forward); 5’CCACTTTGCGACGACTACA3’ (reverse) and 5’CGCGCGACGACCCATTC GAAC3’ (forward); 5’GAATCGAACCCTG ATTCGGCTC3’ (reverse) for 18S rRNA. The cycle parameters included initial denaturation at 95°C for 3 min followed by 45 cycles of 95°C for 10 s and 55°C for 60 s, and followed by 1 cycle of 95°C for 60 s and 55°C for 60 s. To ensure specific amplification, a melt curve was generated at the end of PCR for each sample. The PCR cycle at which the fluorescence exceeded a set threshold (C_T), for each sample was determined by the iCycler software. Data were analyzed according to the comparative C_T method, as previously described (26), using internal control (18S rRNA) transcript levels to normalize differences in sample loading and preparation.

Syndecan-1 ELISA - The levels of shed syndecan-1 accumulated in the conditioned media
after different treatments were assessed by enzyme-linked immunosorbent assay (ELISA) using an Eli-pair kit specific for human syndecan-1 core protein (Cell Sciences). The standard curve was linear between 8 and 256 ng/ml, and all samples were diluted to concentrations within that range. All the samples were run in duplicates.

RESULTS

**Syndecan-1 shedding is enhanced by reducing the amount of heparan sulfate present on the core protein** - We have previously demonstrated that syndecan-1 shedding by myeloma cells can be enhanced by increasing endogenous heparanase expression or by treating cells with exogenous heparanase or bacterial heparinase III (Hep III) (21). These results suggest that reducing the amount of heparan sulfate present on syndecan-1 enhances shedding of the proteoglycan. To explore this further, equal numbers of CAG cells were treated with increasing amounts of Hep III enzyme to generate cells having syndecan-1 containing high, intermediate or low levels of heparan sulfate. After treatment with enzyme for 4 h, the cells were harvested and conditioned medium was collected. Western blots of the cell extracts revealed that with an increase in the amount of Hep III enzyme there was a decrease in the amount of heparan sulfate present on syndecan-1. This decrease in amount of heparan sulfate is visualized on the western blots as an increase in syndecan-1 polydispersity and a reduction in the mean molecular size of the proteoglycan (Fig. 1A; note the broadening of the syndecan-1 smear (polydispersity) following treatment with increasing amounts of Hep III, and the clear emergence of a low-molecular weight fraction of syndecan-1 around 80-95 kDa in cells treated with 2.5 mU/ml of Hep III). Although in the western blot there appears to be much higher levels of syndecan-1 present following Hep III digestion (Fig. 1A, compare lanes 1 and 5), quantification of syndecan-1 by ELISA reveals similar levels of syndecan-1 are present in cell extracts before and after Hep III treatment. We speculate that on western blots, the high molecular weight form of syndecan-1 either transfers poorly to the membrane or is not as assessable to the anti-syndecan-1 antibody as is the low molecular weight form, thus giving the false impression that the amount of syndecan-1 in the cell fraction increases following Hep III treatment.

Quantification of soluble syndecan-1 in the medium conditioned by these cells revealed that with the decrease in proteoglycan size there was a corresponding increase in syndecan-1 shedding (Fig. 1B). Inactive Hep III, lacking the ability to degrade heparan sulfate failed to affect syndecan-1 shedding, indicating that the enzymatic activity of Hep III and the consequent decrease in heparan sulfate content are responsible for the increase in shedding (Fig. 1A, B). In separate experiments, it was confirmed that Hep III treatment of syndecan-1 did not alter the ability to accurately quantify syndecan-1 levels by ELISA. In addition, Hep III induced syndecan-1 shedding was confirmed by dot blotting using an antibody different than the one used in ELISA quantification.

Together the results indicated that upon Hep III treatment the amount of syndecan-1 present in the cell fraction remained constant, but there was a dramatic increase in syndecan-1 shedding. This suggested that syndecan-1 synthesis was elevated following Hep III treatment. To confirm this, we tested whether a sustained enzymatic digestion of heparan sulfates over time was accompanied by an enhanced synthesis of syndecan-1. Results revealed there was a significant increase in syndecan-1 synthesis (Fig. 1C) thereby indicating that the amount of heparan sulfate present on the syndecan-1 core protein regulates the rate of core protein synthesis.

We next assessed whether cell lines that express either high molecular weight or low molecular weight forms of syndecan-1 differed in their rate of shedding. When an equal number of
U266, RPMI-8226 and CAG cells were seeded and grown in serum-free medium for 24 h, U266 cells shed a significantly higher level of syndecan-1 than did the other two cell lines (Fig. 2A). Western blots of cell extracts revealed that syndecan-1 from U266 cells ran as a broad smear ranging from approximately 100 kDa to well over 260 kDa, while syndecan-1 from RPMI-8226 and CAG myeloma cells was much less polydisperse and considerably larger in mean molecular size than syndecan-1 from U266 cells (Fig. 2B). Because the polydispersity in syndecan-1 size is due to predominantly to heterogeneity in the size and number of heparan sulfate chains, these results indicate that the overall heparan sulfate content in U266 cell syndecan-1 is less than that in RPMI-8226 and CAG cells. The high rate of syndecan-1 shedding by U266 cells as compared to RPMI-8226 and CAG cells was not due to a higher level of cell surface syndecan-1 on the U266 cells (Fig. 2C), nor was it due to differences in rates of proliferation because the three cell lines grew at similar rates in serum free medium. Consistent with the finding in the myeloma cell lines, we also found when assessing murine cell lines that MPC-11 myeloma cell that have a low molecular weight form of syndecan-1 shed the proteoglycan at a higher rate than do NMuMG cells that have a high molecular weight form of syndecan-1 (see Fig. 5). Together, the results from assessing different cell lines and from treating cells with Hep III (as described in Fig. 1) point to an important role for heparan sulfate in regulating shedding of syndecan-1. Retention of high levels of heparan sulfate on the core protein suppresses syndecan-1 shedding while removal of heparan sulfate enhances syndecan-1 shedding.

Heparan sulfates suppress syndecan shedding across multiple cell lineages - Cell lines belonging to different lineages (myeloma - CAG, U266, RPMI-8226; breast cancer - MDA-MB 231; pancreatic cancer - Panc-1; cervical cancer - HeLa) were treated with Hep III and the effect on syndecan-1 shedding was evaluated. In all cell lines, reducing the amount of heparan sulfate chains caused an up-regulation of syndecan-1 shedding by several fold, ranging from ~3 fold in U266 cells to ~10 fold in HeLa cells compared to untreated cells (Fig. 3A, B). Because syndecan-1 also bears chondroitin sulfate chains (27), we treated cells with chondroitin ABC (Chase-ABC) to determine if this would alter shedding. Reducing chondroitin sulfate levels impacted syndecan-1 shedding in only three (U266, CAG, and Panc-1) of the six cell lines tested (Fig. 3A, B). Even in these three cell lines, the maximal effect seen with CAG cells (less than 2 fold increase over untreated controls) was considerably less than levels induced by Hep III treatment in the same cell line (Fig. 3A). Thus, between the two types of glycosaminoglycans present on syndecan-1, heparan sulfates are the major regulators of syndecan-1 shedding.

To test whether reduction in heparan sulfate amount elevated shedding in other members of the syndecan family, the effect of Hep III treatment on syndecan-4 was examined. Among the cell lines, HeLa cells had detectable levels of syndecan-4 and were therefore used to address this question. Similar to syndecan-1, reducing the amount of heparan sulfate, not chondroitin sulfate, elevated the rate of syndecan-4 shedding (Fig. 3C).

A mutant form of human syndecan-1 lacking GAG chains is shed at a high rate - As an independent confirmation of the effect of GAGs on syndecan-1 shedding, we stably expressed either the wild-type human syndecan-1 (WT hSDC1) or mutated form of human syndecan-1 lacking all glycosylation attachment sites including GAG attachment sites (ΔGAG hSDC1; serine to alanine mutations at residues at 37, 45, 47, 206, and 216). These constructs were expressed in ARH-77 cells, an Epstein-Barr virus-transformed human B lymphoblastoid cell line that lacks endogenous expression of syndecan-1 (21). Stable transfectants expressing similar levels of WT hSDC1 and ΔGAG hSDC1 protein on the cell surface were selected and confirmed by flow
cytometric analyses (Fig. 4A). Both transfectant lines proliferated at similar rates. Strikingly, the cells expressing GAGless syndecan-1 shed ~6 fold higher levels of syndecan-1 compared to the cells expressing wild-type syndecan-1 (Fig. 4B), mirroring the effects seen with enzymatic removal of GAGs. As a control, treating ARH-77 cells expressing WT hSDC1 cells with Hep III substantially increased the rate of syndecan-1 shedding while treatment of cells with Chase-ABC had much less impact (Fig. 4C). Together, these findings demonstrate that shedding following Hep III treatment is not due to an induction of shedding by the enzyme; rather it is due to the decrease in the amount of heparan sulfate present on the core protein. Moreover, the results indicate that the default state of the syndecan-1 core protein is to be shed rapidly from the cell surface and that this shedding is suppressed when sufficient heparan sulfate chains are attached to the core protein.

Heparan sulfate regulated shedding of syndecan-1 is not species-specific - In a previously published study, we demonstrated that syndecan-1 from a murine plasmacytoma cell line (MPC-11) had a low level of glycosaminoglycan content compared to syndecan-1 from normal murine mammary epithelial cells (NMuMG) (28). To ascertain whether heparan sulfates regulate murine syndecan-1 shedding, we compared the rate of shed syndecan-1 between these two cell lines using dot blots probed with monoclonal antibody to murine syndecan-1. Results revealed that MPC-11 cells shed syndecan-1 at significantly higher rates compared to NMuMG cells (Fig. 5A). Additionally, removal of heparan sulfates by Hep III treatment dramatically up regulated shedding of murine syndecan-1 from NMuMG cells, whereas chondroitinase treatment failed to affect murine syndecan-1 shedding compared to untreated controls (Fig. 5B). This result demonstrates that heparan sulfate can regulate shedding in murine cells and importantly, also in non-cancerous cells. As a further confirmation of the role of heparan sulfate in murine syndecan-1 shedding, we expressed either the murine wild-type syndecan-1 (WT mSDC1) or a mutated syndecan-1 lacking the GAG chains (ΔGAG mSDC1) in ARH-77 cells. The stable expression of syndecan-1 was confirmed and was found to be similar between the transfectants by flow cytometry (Fig. 5C). Similar to our observations with human syndecan-1, cells expressing ΔGAG mSDC1 shed higher levels of syndecan-1 compared to cells expressing wild-type syndecan-1 (Fig. 5D).

Heparan sulfates suppress MMP-mediated shedding of syndecan-1 - Various agonists such as PMA (phorbol ester 12-myristate 13-acetate), insulin, growth factors and chemokines can stimulate syndecan-1 shedding through activation of diverse molecular pathways involving ERK (extracellular signal-regulated kinase) (29), MAP kinases (19), Protein kinase C (PKC) (29,30), Protein-tyrosine kinases (PTKs) (23,31), or Rab5 signaling (20). To determine if removal of heparan sulfate by Hep III enhances shedding by any of these pathways, CAG cells were incubated in the presence of specific inhibitors for individual pathways prior to Hep III treatment (Fig. 6A). The concentrations of inhibitors used in the current study have been reported in previously published studies (29-31). Cells treated with Hep III in the presence of DMSO served as a control. Blocking PKC activation or MEK activity had no significant effect on the level of syndecan-1 shedding (Fig. 6A). Inhibition of receptor tyrosine kinase signaling by genistein caused a modest (~30%) but significant decrease in shedding (Fig. 6A). Blocking actin polymerization or protein synthesis failed to block the shedding induced by removal of heparan sulfate by Hep III in this short term assay.

Matrix metalloproteinases (MMPs) like MMP-7, MMP-9, and MMP-14 are known to regulate syndecan-1 shedding by proteolytic cleavage of the core protein (32-34). To test the possibility that heparan sulfates regulate MMP-mediated proteolysis, we utilized BB-94, a broad-based MMP inhibitor. Addition of BB-94 dramatically inhibited Hep III-induced syndecan-1shedding.
shedding (~3 fold decrease) in CAG cells (Fig. 6A). Confirming these results in another model system, addition of BB-94 significantly reduced shed syndecan-1 from ARH-77 cells expressing GAGless syndecan-1 (ΔGAG hSDC1) (Fig. 6B). This data provides strong evidence that heparan sulfate present on syndecan-1 suppresses MMP-mediated shedding. Importantly, addition of either purified heparan sulfate or heparin (100 ug/ml) failed to inhibit syndecan-1 shedding from these cells (Fig. 6B), demonstrating that not merely the presence of heparan sulfates but their localization on the syndecan-1 core protein is critical to suppress syndecan-1 shedding.

We next used a different model to verify that heparan sulfate chains of syndecan-1 inhibit core protein shedding. Panc-1 cells were treated with PMA, an agonist that has been shown to dramatically increase MMP-mediated syndecan-1 shedding (30). We hypothesized that in this model, removal of heparan sulfates by Hep III would dramatically enhance the effect of PMA on syndecan-1 shedding. As predicted, addition of PMA to Hep III treated cells significantly increased levels of shed syndecan-1 compared to cells treated with either Hep III alone or PMA alone (Fig. 6C). Further, this elevated shedding by combination treatment with Hep III and PMA was blocked upon addition of BB-94 (Fig. 6C).

**DISCUSSION**

The importance of shed syndecan-1 in human disease has been well documented by studies showing its role in cancer progression, wound healing, inflammation and infection (4,5). Although the mechanisms regulating shedding are not fully understood, it is known that agonist-induced signaling upregulates syndecan shedding via the action of MMPs. In the present work we provide evidence for a mechanism that suppresses syndecan shedding. This occurs via the heparan sulfate chains present on syndecan-1 which act to prevent MMP cleavage of the core protein. Specifically, we found that i) the rate of shedding from the cell surface is significantly enhanced when heparan sulfate chains of cell surface syndecan-1 are trimmed by enzymes, or when the syndecan-1 core protein lacks heparan sulfate chains, ii) heparan sulfate-mediated suppression of shedding occurs in multiple cell types, for both syndecan-1 and syndecan-4 and in both human and murine cells, iii) to suppress shedding, the heparan sulfate chains must be attached to the core protein of syndecan-1 because exogenous heparan sulfate or heparin does not suppress shedding, iv) when heparan sulfate chains are removed from syndecan-1, shedding is not increased if MMP inhibitors are present indicating that heparan sulfate chains protect the core protein from MMP-mediated proteolysis, and v) the heparan sulfate chains on syndecan-1, by way of its regulation of shedding, also regulate syndecan-1 core protein synthesis. These findings are important because the negative regulation of syndecan-1 shedding ensures that syndecan-1 is maintained on the cell surface and that excess syndecan-1 is not released into the extracellular compartment. During disease progression, enhanced degradation of heparan sulfate chains likely shifts the balance between stimulation and suppression of syndecan-1 shedding leading to excessive shedding and further dysregulation of tissue homeostasis.

If the amount of heparan sulfate present on syndecan-1 regulates its shedding, then it is important to understand what controls degradation of heparan sulfate chains at the cell surface. First, we know that heparanase, an enzyme that cleaves heparan sulfate chains, reduces the amount of heparan sulfate present on syndecan-1 and increases syndecan-1 shedding (21). Enhanced syndecan-1 shedding is also detected in mice expressing a heparanase transgene, indicating that the link between heparanase expression and enhanced shedding occurs in vivo (21). Heparanase expression also leads to upregulation of MMP-9, a known sheddase of syndecan-1 (22). Thus, heparanase likely plays a dual role in
promoting shedding by directly reducing the amount of heparan sulfate on syndecan-1 (thereby making it more susceptible to MMPs), and indirectly by upregulating expression of MMP-9. It is also interesting that heparanase is known to be upregulated in cancer, inflammation and wound healing, all pathological states that are associated with increased syndecan-1 shedding. A second mechanism for degradation of heparan sulfate is through the action of reactive oxygen species which can chemically fragment heparan sulfate (35). This has not been studied extensively, but there is evidence that oxidative damage leads to syndecan-1 shedding which contributes to lung fibrosis (9). Because enhanced levels of reactive oxygen species are often found in cancer and inflammatory diseases, it is possible that oxidative damage to heparan sulfates contributes significantly to the increased levels of shed syndecan-1 that help drive these diseases.

In addition to degradation of heparan sulfate, the rate of syndecan shedding may also be regulated by the length and number of heparan sulfate chains assembled onto a core protein during proteoglycan synthesis. Although the mechanisms controlling chain length and number remain unclear, the length of chains can be influenced by levels of EXT enzymes, the glycosyltransferases responsible for heparan sulfate chain elongation. For example, the length of heparan sulfate chains was reduced when EXT1 and EXT2 genes were knocked down in HEK 293 cells. In contrast, knock down of EXTL3 resulted in longer heparan sulfate chains (36). Thus, dysregulation of these enzymes during disease could indirectly lead to enhanced syndecan shedding.

Elevated shedding of syndecan in response to a reduction in heparan sulfate content was common for both human and murine syndecan-1 and for syndecan-4. This also held for the murine/human chimeric proteoglycan composed of a murine syndecan-1 core protein with attached human heparan sulfate chains that was generated when human ARH-77 cells, which are negative for syndecan-1 expression, were transfected with the cDNA for murine syndecan-1. Moreover, this shows that the intrinsic cellular mechanisms that control shedding can be present in cells that do not normally express the proteoglycan. Together these results indicate that regulation of syndecan shedding by heparan sulfate is a highly conserved molecular phenomenon, and supports the importance of this mechanism for control of shedding by heparan sulfate.

The precise molecular mechanism of how the heparan sulfates on syndecan-1 protect the core protein from metalloproteinases is not clear. Cell surface heparan sulfate can bind to and inhibit the activity of A Disintegrin And Metalloprotease 12 (ADAM12), a protease that mediates ectodomain shedding of pro-EGF. This inhibition by heparan sulfate occurs via its binding to the ADAM12 catalytic domain (37). However, in contrast to what we found regarding heparan sulfate inhibition of syndecan shedding, ADAM12 activity was also inhibited by exogenous heparin. We do not know if the heparan sulfates of syndecan bind to MMPs and inhibit their activity directly. It has been shown in some models that MMP-7 activity is enhanced by exogenous heparin, although it is not enhanced by heparan sulfate (38,39). However, in our model we found that exogenous heparin had no effect on shedding, suggesting that it is unlikely that syndecans are binding to MMPs and directly inhibiting their activity. Another possible mechanism could be related to a change in protein folding and stabilization which can be altered by changes in glycosylation (40,41). Removal of heparan sulfate could alter syndecan-1 core protein conformation and expose cryptic sites that are susceptible to protease cleavage. Moreover, heparan sulfate chains could simply provide a physical barrier that blocks accessibility of the protease to the core protein. Another important consideration regarding mechanism is that syndecan-1 shedding can be regulated by intracellular mechanisms.
When the cytoplasmic domain of syndecan-1 associates with the protein transport regulator Rab5, shedding of syndecan-1 is inhibited (20). Removal of heparan sulfates from the core protein might disrupt formation or maintenance of the syndecan/Rab5 complex and lead to elevation of shedding.

In summary, this work defines a new role for heparan sulfate and dramatically broadens the scope of importance of this glycosaminoglycan. Although previous studies have clearly established the role of heparan sulfate in binding various ligands, our findings assign two new functions to heparan sulfate. First, heparan sulfate chains attached to the core protein retain syndecan-1 at the cell surface by inhibiting its shedding. Thus, heparan sulfate controls the location of syndecan-1 thereby impacting its function as either a cell surface molecule or as a soluble effector that can regulate distal cells. Second, heparan sulfate regulates syndecan-1 core protein synthesis. When heparan sulfate is diminished, shedding is enhanced and the rate of core protein synthesis dramatically increases. This indicates the existence of a feedback loop in which the cell attempts to maintain a critical level of syndecan-1 on the cell surface. Lastly, our results indicate that therapies designed to preserve the heparan sulfate chains of syndecans, or to increase heparan sulfate chain length or number could diminish syndecan shedding and help to control the progression of aggressive cancers or inflammatory diseases.

REFERENCES


**FOOTNOTES**

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1The abbreviations used are: ADAM, a disintegrin and metalloproteinase; ADAMTS, a disintegrin and metalloproteinase with thrombospondin motifs; BSA, bovine serum albumin; ELISA, enzyme-linked immunosorbent assay; ERK, extracellular signal-regulated kinase; EXT, exostosin; EXTL, exostosin-like; FBS, fetal bovine serum; GAG, glycosaminoglycan; Hep III, heparinase III; HGF, hepatocyte growth factor; MMP, matrix metalloproteinase; MT1-MMP, membrane type 1 matrix metalloproteinase; PKC, protein kinase C; PMA, phorbol ester 12-myristate 13-acetate; PTK, protein tyrosine kinase; SDC1, syndecan-1; uPA, urokinase-type plasminogen activator; uPAR, urokinase-type plasminogen activator receptor; VEGF, vascular endothelial growth factor; WT, wild-type.

2Unpublished observation, V. C. Ramani

FIGURE LEGENDS

FIGURE 1. Reducing the amount of heparan sulfate by enzymatic digestion enhances syndecan-1 shedding from cells. CAG cells were plated at a density of 10⁶ cells/ml in serum free medium for 2 h and then the individual wells were treated with different amounts of Hep III enzyme for 4 h or with heat inactivated Hep III enzyme as a control. At the end of incubation the cells and conditioned media were collected and analyzed separately. (A) Cell lysates were analyzed by immunoblotting for syndecan-1. β-actin was probed in the same blots to demonstrate equal loading of the total protein. (B) Conditioned media were analyzed for their level of shed syndecan-1 (SDC1). Data are mean ± S.E. of three independent experiments. *, p < 0.005 versus untreated controls. (C) CAG cells were treated with Hep III for 8 h and the level of syndecan-1 was determined by real-time PCR. Control included CAG cells treated with distilled water. Transcript levels were normalized against 18S rRNA levels. Data are mean ± S.E. of three independent experiments. *, p < 0.005 versus control.

FIGURE 2. U266 myeloma cells shed more syndecan-1 than RPMI-8226 or CAG myeloma cells. (A) 10⁶ cells/ml of U266, RPMI-8226 or CAG myeloma cells were seeded and grown in serum-free media for 24 h. The level of shed syndecan-1 in the conditioned medium from each cell line was determined by ELISA. Data are mean ± S.E. of three independent experiments. *, p < 0.01 versus U266 cells. (B) Equal protein from total cell extracts of the three myeloma cell lines were probed for syndecan-1 and β-actin by immunoblotting. (C) The level of cell surface expression of syndecan-1 on each cell line was determined by staining with BA38 antibody and analyzed by flow cytometry: IgG control antibody (shaded), U266 cells (dotted line), RPMI-8226 cells (solid line), CAG cells (dashed line).

FIGURE 3. Heparan sulfate suppression of syndecan-1 shedding is not restricted to myeloma cells. Shedding of syndecan-1 in response to reduction of glycosaminoglycans in tumor cell lines of different origins, (A) 10⁶ cells/ml of myeloma cell lines (U266, RPMI-8226, and CAG) were seeded in serum-free medium for 6 h either in the presence of Hep III (1.25 mU/ml) or Chase-ABC (12.5 mU/ml) separately. The level of shed syndecan-1 in the conditioned medium was determined by ELISA. Untreated cells for each cell line tested served as the control. Data are mean ± S.E. of three independent experiments. *, p < 0.01 versus untreated control for each individual cell line tested **, p < 0.05 versus untreated control for each individual cell line tested. (B) 0.5 x 10⁶ cells of tumor cells of different origins: MDA-MB-231 (breast cancer), Panc-1 (pancreatic cancer), HeLa (cervical cancer) were seeded separately in serum free medium for 24 h either in the presence of Hep III (1.25 mU/ml) or Chase-ABC (12.5 mU/ml). Untreated cells of each cell line were included as control. At the end of incubation, the level of shed syndecan-1 in
the conditioned medium was determined by ELISA. Data are mean ± S.E. of three independent experiments. *, \( p < 0.005 \) versus untreated controls. (C) Shedding of syndecan-4 is elevated upon reduction of the amount of heparan sulfate. 0.5 \( \times 10^6 \) HeLa cells were seeded in serum free medium in the presence of either Hep III (1.25 mU/ml) or Chase-ABC (12.5 mU/ml) for 16 h. Untreated cells served as a control. The designated amounts of treated conditioned medium were probed by dot blotting for shed syndecan-4. The experiment was repeated at least three times and a representative blot is shown. The blot reflects a greater than two-fold increase in shed syndecan-4 upon treatment with Hep III compared to untreated controls.

FIGURE 4. A mutated form of human syndecan-1 lacking glycosaminoglycan chains gets shed at a much higher rate than wild-type syndecan-1. (A) ARH-77 cells were transfected with a cDNA encoding either wild-type human syndecan-1 or mutated syndecan-1 lacking glycosaminoglycan attachment sites. Stable expression of cell surface syndecan-1 was confirmed by flow cytometry following staining with BA38 antibody: IgG control (black line), WT hSDC1 (grey line), and ΔGAG hSDC1 (dotted line). (B) Equal numbers of WT hSDC1 or ΔGAG hSDC1 cells were seeded in serum free medium for 24 h. The conditioned media were harvested and levels of syndecan-1 were assessed by ELISA. Data are mean ± S.E. of three independent experiments. *, \( p < 0.0005 \) versus WT hSDC1 cells. (C) WT hSDC1 cells were treated with either Hep III (1.25 mU/ml) or Chase-ABC (12.5 mU/ml) for 24 h. After incubation, the conditioned media were harvested and the level of shed syndecan-1 was assessed by ELISA. Data are mean ± S.E. of three independent experiments. *, \( p < 0.01 \) versus untreated control.

FIGURE 5. Shedding of murine syndecan-1 is enhanced by heparan sulfate removal. (A) Equal numbers of murine MPC-11 myeloma cells and murine NMuMG epithelial cells were seeded in serum-free medium for 24 h and the level of shed syndecan-1 determined by dot blotting with monoclonal antibody 281.2. (B) Equal numbers of NMuMG cells were seeded in serum-free medium and treated with either Hep III (1.25 mU/ml) or Chase-ABC (12.5 mU/ml) for 24 h. Untreated monolayers served as a control. After incubation, the conditioned medium was harvested and the level of shed syndecan-1 determined by dot blotting. The experiments were repeated at least three times and a representative dot blot is shown. (C) ARH-77 cells were transfected with a cDNA encoding either wild-type murine syndecan-1 or mutated syndecan-1 lacking heparan sulfate chain attachment sites. Stable expression of cell surface syndecan-1 was confirmed by flow cytometry by staining with 281.2 antibody: Isotype IgG control (black line), WT mSDC1 (dotted line), and ΔGAG mSDC1 (dashed line). (D) Equal numbers of WT mSDC1 or ΔGAG mSDC1 cells were seeded in serum free medium for 24 h, the conditioned medium was harvested and the level of shed syndecan-1 determined by dot blotting. The experiments were repeated at least three times and a representative dot blot is shown.

FIGURE 6. Heparan sulfate chains on syndecan-1 limit metalloproteinase-mediated shedding of syndecan-1. (A) CAG cells were incubated in the presence of different inhibitors for 2 h in serum free medium then treated with Hep III (1.25 mU/ml) for 4 h and the levels of shed syndecan-1 were determined by ELISA. Data are mean ± S.E. of three independent experiments. *, \( p < 0.05 \) versus cells treated with Hep III in the presence of DMSO. The specific inhibitors and their concentrations used- RTK signaling (genistein, 37 µM), MEK activity (U0126, 1µM), PKC (bisindolylmaleimide, 1µM), metalloproteinase activity (BB-94, 5µM), actin polymerization (cytochalasin D, 10µM), protein synthesis (cycloheximide, 0.3mM). (B) Equal numbers of WT hSDC1 or ΔGAG hSDC1 cells were seeded in serum
free medium for 24 h. Experimental groups also included ∆GAG hSDC1 cells incubated in the presence of either purified heparan sulfate, heparin or BB-94 for 24 h. The level of shed syndecan-1 after 24 h was assessed by ELISA. Data are mean ± S.E. of three independent experiments. *, \( p < 0.005 \) versus untreated ∆GAG hSDC1 cells. (C) Panc-1 cells were seeded at equal density in serum-free medium and were treated with either Hep III (1.25 mU/ml) or with Hep III and DMSO for 4 h. Treatment groups also included Hep III treated cells incubated in the presence of PMA (0.5µM) or with PMA and BB-94 (5µM) for the next 2 h. Cells treated with either PMA or with BB-94 alone served as additional controls. Data are mean ± S.E. of three independent experiments. *, \( p < 0.05 \) versus groups treated with PMA alone. #, \( p < 0.01 \) versus groups treated with Hep III alone.
Figure 2
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A

Shed SDC1 (ng/ml)

<table>
<thead>
<tr>
<th></th>
<th>U266</th>
<th>RPMI-8226</th>
<th>CAG</th>
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B

- U266
- RPMI-8226
- CAG

Actin

C

Count

FL4-H::ALEXA 647

![Graph](image4)
Figure 3

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Figure 6
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A

B

C
Heparan sulfate chains of syndecan-1 regulate ectodomain shedding
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