

METHOD FOR MANUFACTURING COMPOSITION COMPRISING LOCAL
ANESTHETIC, HEPARINOID, AND BUFFER

CROSS-REFERENCES

[0001] This application claims priority from United States Provisional Application Serial No. 61/430,468 by C. Lowell Parsons, filed January 6, 2011 and entitled "Method for Manufacturing Composition Comprising Local Anesthetic, Heparinoid, and Buffer," the contents of which are hereby incorporated in their entirety by this reference.

FIELD OF THE INVENTION

[0002] This invention is directed to an improved method of manufacturing a composition suitable for the treatment of urinary bladder symptoms as seen in interstitial cystitis and other lower urinary tract conditions such as, but not limited to, overactive bladder, prostatitis, urethral syndrome in women, and gynecologic chronic pelvic pain. The composition comprises a local anesthetic, typically lidocaine, a glycosaminoglycan, typically heparin, and a buffer, typically sodium bicarbonate which is then instilled directly into the urinary bladder.

BACKGROUND OF THE INVENTION

[0003] Interstitial cystitis (IC) is a chronic progressive disorder of the lower urinary tract that causes urinary urgency and frequency and/or pelvic pain. For many years, urologists regarded IC as a rare disease for which they had no broadly effective treatment. In fact, the condition is quite common. In 1999, prevalence in the United

States was estimated at 750,000 cases (Curhan, et al. J Urol 161(2):549-552 (1999)). However the true prevalence of IC is estimated to be at least 1-2 million patients who are suffering from severe chronic pelvic pain. In addition overactive bladder, urethral syndrome, prostatitis, and gynecologic chronic pelvic pain syndrome comprises millions of patients that also result in bladder symptoms of urgency, frequency, incontinence and or pelvic pain with no effective therapy and all these syndromes share similar symptoms and likely a common pathophysiology with traditionally diagnosed IC (Parsons, CL Int Br J Urol Dec, 2010); there are no broadly effective treatments for these conditions.

[0004] Therefore, treatments that would both benefit a larger portion of the patient population, provide immediate relief of symptoms without causing additional pain, without requiring extensive alterations in diet, and further provide reversal of the disease process over time are necessary.

[0005] Compositions and methods for the treatment of interstitial cystitis are described in United States Patent No. 7,414,039 to Parsons, issued August 19, 2008 and entitled "Interstitial Therapy for Immediate Symptom Relief and Chronic Therapy in Interstitial Cystitis," and in United States Patent Application Publication No. 2008/0300219 by Parsons, published December 4, 2008 and entitled "Novel Interstitial Therapy for Immediate Symptom Relief and Chronic Therapy in Interstitial Cystitis," both of which are incorporated herein in their entirety by this reference, as well as PCT Patent Application Publication No. WO 2006/07663 by Flashner et al., published July 20, 2006 and entitled "Kits and Improved Compositions for Treating Lower Urinary Tract Disorders," and PCT Patent Application Publication No. WO 2007/073397 by Flashner et al., published June 28, 2007 and entitled "Kits and Improved Compositions for Treating Lower Urinary Tract Disorders," both of which are incorporated herein in their entirety by this reference. In general, the compositions disclosed in this issued patent and these published patent applications comprise a local anesthetic, typically lidocaine, a heparinoid, typically heparin, and a buffer, typically sodium bicarbonate that are instilled directly into the urinary bladder.

[0006] Alkalinized lidocaine and heparin can be used to successfully treat bladder symptoms such as, but not limited to, urinary frequency, urgency, incontinence and bladder generated pain. Pain generated by the urinary bladder (a visceral organ) is not always perceived to be arising from the bladder. Pain can be referred anywhere from the navel to the knees and will also refer from the lumbar area down the buttocks to the legs and often has no relation to bladder filling or emptying. Consequently the origin of pelvic pain may not be recognized to be from the bladder. These bladder symptoms can be seen in a variety of "clinical syndromes" which may actually be all from one disease process: a dysfunctional epithelium (Parsons, CL Int Br J Urol, Dec 2010)). Nonetheless all these syndromes that can generate bladder symptoms that can be successfully treated with this solution, including, but not limited to, overactive bladder, interstitial cystitis, urethral syndrome in women, recurrent lower urinary tract infection, prostatitis (male chronic pelvic pain syndrome), radiation cystitis, chemical cystitis, gynecologic chronic pelvic pain syndrome (e.g. endometriosis, vulvodynia, vulvovaginitis, yeast vaginitis).

[0007] However, there is a problem in mixing these three compounds, as the wrong balance will result in the precipitation of lidocaine and loss of efficacy. Lidocaine when exposed to pH's at or above 7.0 will de-ionize and absorb through lipid membranes such as the bladder epithelium. As a result, the absorbed lidocaine will anesthetize the bladder nerves and relieve bladder symptoms noted above. The heparin will "coat" the bladder wall and inhibit the diffusion of potassium that is provoking the bladder symptoms in the first place. So the combination provides prolonged relief of bladder symptoms (Parsons, Urology 2003). However, mixing the heparin, lidocaine, and buffering agent has to be done in an exact way to prevent the precipitation of the lidocaine since lidocaine will precipitate at pH values above 7 depending on the conditions. The precipitation of lidocaine reduces its bioavailability and reduces the efficacy of the composition.

[0008] Accordingly, there is a need for an improved method of preparation for compositions including a heparinoid such as heparin, a local anesthetic such as lidocaine, and a buffer such as sodium bicarbonate, that will avoid precipitation of the local anesthetic such as lidocaine and thus will maintain the bioavailability of the local anesthetic and maintain the efficacy of the composition. In all instances the preparation of these mixtures is for instillation directly into the urinary bladder for the treatment of bladder symptoms. As such the osmolality, specific cation concentrations and pH requirements are not as restrictive as for the preparations of these drugs for intravenous or subcutaneous administration.

SUMMARY OF THE INVENTION

[0009] An improved method of preparation of a composition including a heparinoid such as heparin, a local anesthetic such as lidocaine, and a buffer such as sodium bicarbonate, prevents precipitation of the local anesthetic such as lidocaine and thus maintains the bioavailability of the local anesthetic. This maintains the stability and efficacy of the composition.

[0010] One aspect of the invention is a method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (1) providing a heparinoid in solid form or liquid form;
- (2) providing a local anesthetic in solid form or liquid form;
- (3) adding a liquid buffer to the heparinoid in solid form or liquid form;
- (4) adding the local anesthetic to the mixture of the liquid buffer and the heparinoid; and

(5) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.7 to about 8.3 or even higher is achieved without precipitation of the local anesthetic. An optimal pH is about 7.4.

[0011] Typically, the heparinoid is selected from the group consisting of heparin, chondroitin sulfate, heparan sulfate, hyaluronic acid, keratan sulfate, dermatan sulfate, hyaluronan, and sodium pentosanpolysulfate. Preferably, the heparin is selected from the group consisting of heparin and sodium pentosanpolysulfate. More preferably, the heparinoid is heparin, such as heparin sodium.

[0012] Typically, the local anesthetic is selected from the group consisting of benzocaine, lidocaine, tetracaine, bupivacaine, cocaine, etidocaine, flecainide, mepivacaine, pramoxine, prilocaine, procaine, chloroprocaine, oxyprocaine, proparacaine, ropivacaine, dyclonine, dibucaine, propoxycaine, chloroxylonol, cinchocaine, dexivacaine, diamocaine, hexylcaine, levobupivacaine, propoxycaine, pyrrocaine, risocaine, rodocaine, and pharmaceutically acceptable derivatives and bioisosteres thereof, and a combination thereof. Preferably, the local anesthetic is selected from the group consisting of lidocaine, bupivacaine, benzocaine, tetracaine, etidocaine, flecainide, prilocaine, and dibucaine, and a combination thereof. More preferably, the local anesthetic is lidocaine, such as lidocaine hydrochloride.

[0013] Typically, the buffer is selected from the group consisting of bicarbonate buffer, Tris (Tris(hydroxymethyl)aminomethane) buffer, MOPS buffer (3-(N-morpholino)propanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N-(2-ethanesulfonic acid) buffer, ACES (2-[(2-amino-2-oxoethyl)amino]ethanesulfonic acid) buffer, ADA (N-(2-acetamido)2-iminodiacetic acid) buffer, AMPSO (3-[(1,1-dimethyl-2-hydroxyethyl)amino]-2-propanesulfonic acid) buffer, BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid buffer, Bicine (N,N-bis(2-hydroxyethyl)glycine) buffer, Bis-Tris (bis-(2-hydroxyethyl)imino-tris(hydroxymethyl)methane buffer, CAPS (3-

(cyclohexylamino)-1-propanesulfonic acid) buffer, CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) buffer, CHES (2-(N-cyclohexylamino)ethanesulfonic acid) buffer, DIPSO (3-[N,N-bis(2-hydroxyethyl)amino]-2-hydroxy-propanesulfonic acid) buffer, HEPPS (N-(2-hydroxyethyl)piperazine)-N'-(3-propanesulfonic acid) buffer, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid) buffer, MES (2-(N-morpholino)ethanesulfonic acid) buffer, triethanolamine buffer, imidazole buffer, glycine buffer, ethanolamine buffer, phosphate buffer, MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid) buffer, PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid) buffer, POPSO (piperazine-N,N'-bis(2-hydroxypropaneulfonic acid) buffer, TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid) buffer; TAPSO (3-[N-tris(hydroxymethyl)methylamino]-2-hydroxy-propanesulfonic acid) buffer, TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid) buffer, tricine (N-tris(hydroxymethyl)methylglycine buffer), 2-amino-2-methyl-1,3-propanediol buffer, 2-amino-2-methyl-1-propanol buffer, and a combination thereof. Preferably, the buffer is selected from the group consisting of bicarbonate buffer, Tris buffer, and a combination thereof. More preferably, the buffer is bicarbonate buffer, such as sodium bicarbonate.

[0014] The method can further comprise the step of adding one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative, subsequent to preparation of a buffered composition including the heparinoid, the local anesthetic, and the buffer.

[0015] In one alternative, both the heparinoid and the local anesthetic are provided in solid form; the solid form can be a powdered form. In another alternative, both the heparinoid and the local anesthetic are provided in liquid form. In yet another alternative, the heparinoid is provided in solid form and the local anesthetic is provided in liquid form; the solid form for the heparinoid can be a powdered form. In still another alternative, the heparinoid is provided in liquid form and the local anesthetic is provided in solid form; the solid form for the local anesthetic can be a powdered form. In

preparation, powders of all three components: (i) a heparinoid such as heparin; (ii) a local anesthetic such as lidocaine; and (iii) a buffer can be hydrated simultaneously if the lidocaine does not precipitate.

[0016] Preferably, the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is bicarbonate buffer. More preferably, the heparin is heparin sodium, the local anesthetic is lidocaine hydrochloride, and the bicarbonate buffer is sodium bicarbonate.

[0017] Another aspect of the invention is an alternative method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (1) providing a heparinoid in solid form or liquid form;
- (2) providing a local anesthetic in solid form or liquid form;
- (3) mixing the heparinoid in solid form or liquid form and the local anesthetic in solid form or liquid form;
- (4) adding a liquid buffer to the mixture of the heparinoid and the local anesthetic to form a mixture of liquid buffer, the heparinoid, and the local anesthetic; and
- (5) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.7 to about 8.3 without precipitation of the local anesthetic. An optimal pH is about 7.4.

[0018] For this alternative method, the heparinoid, local anesthetic, and buffer are as described above.

[0019] This alternative method can further comprise the step of adding one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an

antifungal agent; (v) a vasoconstrictor; or (vi) a preservative, subsequent to preparation of a buffered composition including the heparinoid, the local anesthetic, and the buffer.

[0020] In one alternative, both the heparinoid and the local anesthetic are provided in solid form; the solid form can be a powdered form. In another alternative, both the heparinoid and the local anesthetic are provided in liquid form. In yet another alternative, the heparinoid is provided in solid form and the local anesthetic is provided in liquid form; the solid form for the heparinoid can be a powdered form. In still another alternative, the heparinoid is provided in liquid form and the local anesthetic is provided in solid form; the solid form for the local anesthetic can be a powdered form.

[0021] In yet another alternative, liquid solutions of the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffer can be simultaneously mixed provided that the lidocaine does not precipitate. For example, this can be done with lower levels of buffer when the desired pH is closer to pH 7.0. In general, this alternative comprises the steps of:

- (1) providing a heparinoid in liquid form;
- (2) providing a local anesthetic in liquid form;
- (3) providing a buffer in liquid form;
- (4) simultaneously mixing the heparinoid, the local anesthetic, and the buffer, avoiding precipitation of the local anesthetic; and
- (5) if necessary, adjusting the pH of the mixture of the liquid buffer, the liquid local anesthetic, and the liquid heparinoid so that a pH of from about 6.7 to about 8.3 is achieved without precipitation of the local anesthetic.

[0022] Yet another aspect of the invention is a premixed liquid composition comprising a heparinoid, a local anesthetic, and a buffer that is stable for at least three months without precipitation of the local anesthetic. The premixed liquid composition can be prepared by the methods described above. The heparinoid, local anesthetic, and buffer are as described above. Preferably, the heparinoid is heparin, the local

anesthetic is lidocaine, and the buffer is bicarbonate buffer. More preferably, the heparin is heparin sodium, the local anesthetic is lidocaine hydrochloride, and the bicarbonate buffer is sodium bicarbonate. Typically, the pH of this composition is from about 6.7 to about 8.3. Preferably, the pH of this composition is from about 7.0 to about 7.8. More preferably, the pH of this composition is about 7.4.

[0023] The composition can further comprise one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative.

[0024] The composition can be suitable for treating, ameliorating, or preventing a lower urinary tract disorder including, but not limited to, a lower urinary tract disorder selected from the group consisting of bacterial cystitis, fungal/yeast cystitis, vulvar vestibulitis, vulvodynia, dyspareunia, urethral syndrome, and endometriosis in women; prostatitis and chronic pelvic pain syndrome in men; and radiation-induced cystitis, chemotherapy-induced cystitis, interstitial cystitis, and overactive bladder in men or women. Typically, the lower urinary tract disorder is interstitial cystitis.

DETAILED DESCRIPTION OF THE INVENTION

[0025] An improved method of preparation of a composition including a heparinoid such as heparin, a local anesthetic such as lidocaine, and a buffer such as sodium bicarbonate comprises the following steps:

[0026] (1) The first step is obtaining a heparinoid in solid form, such as in a powdered form, such as heparin or sodium pentosanpolysulfate containing no other agents such as an appropriate cation (i.e., sodium) and a local anesthetic such as lidocaine hydrochloride in solid form, such as a powdered form. Buffer (powdered or

liquid) should not be added to the local anesthetic in solid form at this stage as it will result in precipitation of the lidocaine when the mixture is hydrated.

[0027] (2) In one alternative, the buffer can be added to the heparinoid in solid form, such as a powdered form, and then the local anesthetic such as lidocaine can then be added. In another alternative, the heparinoid in solid form, such as a powdered form, and the local anesthetic in solid form, such as a powdered form, can be mixed and then buffer can be added.

[0028] (3) Once a composition including both heparinoid and local anesthetic is prepared, the composition, which is a solution at this point, is then buffered by gradually adding the buffering agent, such as sodium bicarbonate, until a pH of from about 6.7 to about 8.3 or possibly higher is achieved without precipitation of the local anesthetic. Preferably, the pH of the resulting composition is from about 7.0 to about 7.8. More preferably, the pH of the resulting composition is about 7.4. The amount of the buffering agent required at this stage may vary with each preparation, so the pH should be continually monitored to ensure that the quantity of buffer added is appropriate.

[0029] (4) Lastly, a preservative can be added as necessary; this does not result in precipitation.

[0030] (5) The mixing of heparin in a solid form and lidocaine hydrochloride in a solid form may be done first with the buffering agent and water added later. In another alternative, the various liquid components could all be mixed (in proper sequence) when ready for production of the final product. In yet another alternative, powdered heparin and buffer could be mixed with liquid lidocaine to obtain the final product. In yet another alternative, powders of all three components, namely the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffering agent, can be hydrated simultaneously if the lidocaine does not precipitate. In another alternative, liquid solutions of all three components, namely the heparinoid such as heparin, the local

anesthetic such as lidocaine, and the buffering agent, can be mixed simultaneously provided that the lidocaine does not precipitate; for example, this can be done with lower levels of buffer when the desired pH is closer to a pH value of 7.0. In essence, to produce the final product, any combination of liquid or powdered agents could be employed as long as the proper sequence of mixing the compounds is followed. The key is to avoid contact of the local anesthetic with buffer unless the heparinoid is present.

[0031] (6) A final liquid product that has been produced by a method such as those described above will be stable and can be stored for three months or more with no precipitation. Accordingly, a premixed liquid composition prepared according to the present invention that is stable for at least three months with no precipitation of the lidocaine is within the scope of the present invention.

[0032] It is necessary to begin the preparation of the composition with powdered heparinoid (heparin) and powdered local anesthetic, as commercially available heparin and lidocaine solutions that are sold in the United States are intended for systemic injection into the body. These are the only products available should someone want to compound them for use into the bladder. These available heparin and lidocaine solutions are not compatible when the buffer is added; the lidocaine precipitates regardless of subsequent attempts to avoid precipitation and maintain the lidocaine in solution.

[0033] The composition can further comprise an osmolar component as described further below.

[0034] As used herein, "heparinoid" refers to any molecule comprising a glycosaminoglycan which refers to a molecule comprising a network of long, branched chains of sugars (e.g., chondroitin sulfate, heparan sulfate, hyaluronic acid, keratan sulfate, dermatan sulfate, hyaluronan, sodium pentosanpolysulfate, and the like) and

optimally further comprising smaller, nitrogen-containing molecules (e.g. low molecular weight molecules). It is not meant to limit the present invention to any one glycosaminoglycan (GAG) or source of GAG. GAG molecules include but are not limited to low molecular weight (LMW) GAGs, naturally derived GAGs, biotechnologically prepared GAGs, chemically modified GAGs, synthetic GAGs, and the like. Heparinoids can also be comprised of pentoses instead of hexoses (GAGs are comprised of hexoses) such as pentosanpolysulfate. It is not meant to limit the present invention to any one heparinoid molecule or source of heparinoid molecule. As used herein, "heparin" refers to a heterogeneous group of straight-chain anionic glycosaminoglycans, as described above, having anticoagulant properties with a molecular weight ranging from 2,000 to 40,000 Da. Heparin is measured by its specific anticoagulation activity in units. In some embodiments, heparin is a higher molecular weight species ranging from 8,000-40,000 daltons. As used herein, "low-molecular-weight heparins" refers to a lower molecular weight (LMW) species ranging from 2,000-8,000 daltons. Sodium pentosanpolysulfate can range from 4,000-6,000 daltons. Also included within the scope of the invention are polymers such as dalteparin or enoxaparin. LMW heparins are made by enzymatic or chemical controlled hydrolysis of unfractionated heparin and have very similar chemical structure as unfractionated heparin except for some changes that may have been introduced due to the enzymatic or chemical treatment. While not intending to limit the mechanism of action of the invention's compositions, the mechanism of action of these drugs may be similar to that of full-length heparin. LMW heparins are usually isolated from bulk heparin. In one embodiment, heparin or another heparinoid is a heparin salt. As used herein, the phrases "pharmaceutically acceptable salts", "a pharmaceutically acceptable salt thereof" or "pharmaceutically accepted complex" for the purposes of this application are equivalent and refer to derivatives prepared from pharmaceutically acceptable non-toxic acids or bases including inorganic acids and bases and organic acids and bases.

[0035] Because of the negative charges of these polysaccharides due to the occurrence of sulfate groups and/or carboxylic acid groups in them, they are

administered in the form of salts, with an appropriate cation to neutralize the negative charges on the acid groups. Typically, the cation is sodium. However, other physiologically tolerable counterions that do not induce urinary tract dysfunctions, such as magnesium, aluminum, calcium, ammonium, or salts made from physiologically acceptable organic bases such as, but not limited to, trimethylamine, triethylamine, morpholine, pyridine, piperidine, picoline, dicyclohexylamine, N,N'-dibenzylethylenediamine, 2-hydroxyethylamine, bis-(2-hydroxyethyl)amine, tri-(2-hydroxyethyl)amine, dibenzylpiperidine, N-benzyl-*p*-phenethylamine, dehydroabietylamine, N,N'-bisdehydroabietylamine, glucamine, N-methylglucamine, collidine, quinine, quinoline, and basic amino acids such as lysine and arginine, can be used. These cationic counterions can alternatively be used as the counterions with anionic buffers such as bicarbonate, as well. Sodium is typically employed as the positively-charged counterion as indicated above. These salts may be prepared by methods known to those skilled in the art. However, it is generally undesirable to use potassium as a counterion due to its role in the etiology of the conditions and syndromes being treated. Other polysaccharides that have the required activity include, but are not limited, to dextran sulfate and carrageenan. Other glycosaminoglycans can be used in methods according to the invention, including low molecular weight (LMW) glycosaminoglycans, naturally derived glycosaminoglycans, biotechnologically prepared glycosaminoglycans, chemically modified glycosaminoglycans, and synthetic glycosaminoglycans and linear anionic polysaccharides comprised of pentoses.

[0036] In some embodiments, a heparinoid comprises a heparin-like molecule (e.g. heparan sulfate). For example, a heparin-like molecule such as heparan sulfate is a glycosaminoglycan with a structure similar to heparin with the difference being that heparan sulfate has undergone less polymerization than heparin and so has more glucuronic acid and N-acetyl glucosamine than heparin. Heparan sulfate contains fewer sulfate groups, so is not as effective as an anticoagulant as heparin. Heparin exists in a variety of forms characterized by different degrees of sulfation. Typically, heparin has a molecular weight of from about 2 kDa to about 40 kDa. Heparin and heparan sulfate

are both characterized by repeating units of disaccharides containing a uronic acid (glucuronic or iduronic acid) and glucosamine, which is either N-sulfated or N-acetylated. The sugar residues may be further O-sulfated at the C-6 and C-3 positions of the glucosamine and the C-2 position of the uronic acid. There are at least 32 potential unique disaccharide units in this class of compounds. Five examples of sugars occurring in heparin are: (1) α -L-iduronic acid 2-sulfate; (2) 2-deoxy-2-sulfamino- α -D-glucose 6-sulfate; (3) β -D-glucuronic acid; (4) 2-acetamido-2-deoxy- α -D-glucose; and (5) α -L-iduronic acid.

[0037] In one embodiment, heparin contains at least 130 USP units per mg. Heparin is measured by its specific anticoagulation activity in units; either USP units or international units (IU) are specified in stating the activity of heparin. As used herein, "USP unit" refers to the quantity of heparin that prevents 1.0 ml of citrated sheep plasma from clotting for 1 hour after the addition of 0.2 ml of 1% CaCl_2 at 20° C when compared to a USP reference standard (defined as units/ml). As used herein, "IU" refers to the quantity of heparin that is active in assays as established by the Fifth International standard for Unfractionated Heparin (WHO-5) (defined as International Units/ml) (Linhardt, R. J. & Gunay, N. S. (1999) *Semin Thromb Hemost* 25, 5-16.).

[0038] Particularly preferred heparinoids for use in methods according to the present invention and compositions prepared by those methods include heparin and sodium pentosanpolysulfate. A most particularly preferred heparinoid for use in methods according to the present invention and compositions prepared by those methods is heparin. A preferred form of heparin is heparin sodium, although, as described above, other counterions can be used. The quantity of heparin in compositions prepared according to methods of the present invention can range up to as high as from about 200,000 units to about 250,000 units per unit dose of the composition; any intermediate quantity of heparin, such as, but not limited to, 5,000 units, 10,000 units, 15,000 units, 20,000 units, 25,000 units, 30,000 units, 35,000 units, 40,000 units, 45,000 units, 50,000 units, 55,000 units, 60,000 units, 65,000 units,

70,000 units, 75,000 units, 80,000 units, 85,000 units, 90,000 units, 95,000 units, 100,000 units, 110,000 units, 120,000 units, 130,000 units, 140,000 units, 150,000 units, 160,000 units, 170,000 units, 180,000 units, 190,000 units, 210,000 units, 220,000 units, 230,000 units, or 240,000 units per unit dose of the composition can be used.

[0039] The quantity of heparinoid in the composition will vary depending on the subject, the severity and course of the disease, the subject's health, the response to treatment, pharmacokinetic considerations such as liver and kidney function, and the judgment of the treating physician. Accordingly, a number of compositions including differing quantities of heparin per unit dose can be prepared by methods according to the present invention.

[0040] In accordance with the practice of the invention, merely by way of example, when the heparinoid is sodium pentosanpolysulfate, the amount of heparinoid in the composition may be about 1 mg to about 1500 mg of sodium pentosanpolysulfate per unit dose (for example about 100 mg to about 600 mg per unit dose of sodium pentosanpolysulfate). In accordance with the practice of the invention, merely by way of example, when the heparinoid is heparan sulfate, the amount of heparinoid in the composition may be about 0.5 mg to about 10,000 mg of heparan sulfate per unit dose (for example about 100 mg to about 300 mg per unit dose of heparan sulfate). In accordance with the practice of the invention, merely by way of example, when the heparinoid is hyaluronic acid, the amount of heparinoid in the composition may be about 5 mg to about 1500 mg of hyaluronic acid per unit dose (for example about 10 mg to about 100 mg per unit dose of hyaluronic acid). In accordance with the practice of the invention, merely by way of example, when the heparinoid is chondroitin sulfate, the amount of heparinoid in the composition may be about 1 mg to about 10,000 mg of chondroitin sulfate per unit dose (for example about 100 mg to about 300 mg per unit dose of chondroitin sulfate). In accordance with the practice of the invention, merely by

way of example, when the heparinoid is heparin sodium, the amount of heparinoid in the composition may be about 10 mg to about 1000 mg of heparin sodium per unit dose.

[0041] The local anesthetic is typically a sodium channel blocker, such as, but not limited to, the drugs referred to commonly as the "caine" drugs, as well as other sodium channel blockers. The local anesthetic in a composition prepared according to the methods of the present invention can be, but is not limited to, any of benzocaine, lidocaine, tetracaine, bupivacaine, cocaine, etidocaine, flecainide, mepivacaine, pramoxine, prilocaine, procaine, chloroprocaine, oxyprocaine, proparacaine, ropivacaine, dyclonine, dibucaine, propoxycaine, chloroxylenol, cinchocaine, dexivacaine, diamocaine, hexylcaine, levobupivacaine, propoxycaine, pyrrocaine, risocaine, rodocaine, and pharmaceutically acceptable derivatives and bioisosteres thereof, or a combination thereof. Preferably, the anesthetic (e.g., local anesthetic) is selected from the group consisting of lidocaine, bupivacaine, benzocaine, tetracaine, etidocaine, flecainide, prilocaine, and dibucaine, or a combination thereof. A particularly preferred local anesthetic is lidocaine, such as lidocaine hydrochloride. As used herein, the recitation of a local anesthetic includes all salts of that local anesthetic that are compatible with the desired pH, the buffer used, and any counterions present; the recitation of a local anesthetic is not intended to limit the salt form or counterion used beyond these criteria.

[0042] The quantity of local anesthetic in the composition will vary depending on the subject, the severity and course of the disease, the subject's health, the response to treatment, pharmacokinetic considerations such as liver and kidney function, and the judgment of the treating physician. Accordingly, a number of compositions including differing quantities of local anesthetic per unit dose can be prepared by methods according to the present invention. For example, the amount of anesthetic agent in the compositions may be in the range of about 10 mg to about 400 mg per unit dose. For example, the amount of lidocaine can be 10 mL of 1% lidocaine per unit dose or 16 mL of 2% lidocaine per unit dose.

[0043] The buffer in a composition prepared according to the methods of the present invention can be, but is not limited to, bicarbonate buffer, Tris (Tris(hydroxymethyl)aminomethane) buffer, MOPS buffer (3-(N-morpholino)propanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N-(2-ethanesulfonic acid) buffer, ACES (2-[(2-amino-2-oxoethyl)amino]ethanesulfonic acid) buffer, ADA (N-(2-acetamido)2-iminodiacetic acid) buffer, AMPSO (3-[(1,1-dimethyl-2-hydroxyethyl)amino]-2-propanesulfonic acid) buffer, BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid buffer, Bicine (N,N-bis(2-hydroxyethyl)glycine) buffer, Bis-Tris (bis-(2-hydroxyethyl)imino-tris(hydroxymethyl)methane buffer, CAPS (3-(cyclohexylamino)-1-propanesulfonic acid) buffer, CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) buffer, CHES (2-(N-cyclohexylamino)ethanesulfonic acid) buffer, DIPSO (3-[N,N-bis(2-hydroxyethyl)amino]-2-hydroxy-propanesulfonic acid) buffer, HEPPS (N-(2-hydroxyethyl)piperazine)-N'-(3-propanesulfonic acid) buffer, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid) buffer, MES (2-(N-morpholino)ethanesulfonic acid) buffer, triethanolamine buffer, imidazole buffer, glycine buffer, ethanolamine buffer, phosphate buffer, MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid) buffer, PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid) buffer, POPSO (piperazine-N,N'-bis(2-hydroxypropaneulfonic acid) buffer, TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid) buffer; TAPSO (3-[N-tris(hydroxymethyl)methylamino]-2-hydroxy-propanesulfonic acid) buffer, TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid) buffer, tricine (N-tris(hydroxymethyl)methylglycine buffer), 2-amino-2-methyl-1,3-propanediol buffer, and 2-amino-2-methyl-1-propanol buffer, or a combination thereof. Particularly preferred buffers are bicarbonate buffer and Tris buffer or a combination thereof. A particularly preferred bicarbonate buffer is sodium bicarbonate.

[0044] The quantity of buffer in the composition will vary depending on the subject, the severity and course of the disease, the subject's health, the response to treatment, pharmacokinetic considerations such as liver and kidney function, and the

judgment of the treating physician. Accordingly, a number of compositions including differing quantities of buffer per unit dose can be prepared by methods according to the present invention. For example, when the buffer is sodium bicarbonate, the amount of sodium bicarbonate may be about 3 mL of 8.4% sodium bicarbonate per unit dose.

[0045] A particularly preferred composition prepared by methods according to the present invention can comprise heparin sodium as the heparinoid, lidocaine hydrochloride as the local anesthetic, and sodium bicarbonate as the buffer.

[0046] Compositions prepared by methods according to the present invention can include one or more additional optional components. Such additional optional components can include:

- (1) an osmolar component that provides an isotonic or nearly isotonic solution compatible with human cells and blood;
- (2) a compound that enables persistence of the composition to the surface of the bladder epithelium in a quantity sufficient to treat, ameliorate, or prevent a lower urinary tract disorder;
- (3) an antibacterial agent in a quantity sufficient to treat, ameliorate, or prevent a lower urinary tract disorder;
- (4) an antifungal agent in a quantity sufficient to treat, ameliorate, or prevent a lower urinary tract disorder;
- (5) a vasoconstrictor in a quantity sufficient to treat, ameliorate, or prevent a lower urinary tract disorder; and
- (6) a preservative.

[0047] When present, the optional osmolar component is a salt, such as sodium chloride, or a sugar or a combination of two or more of these components. The sugar may be a monosaccharide such as dextrose, a disaccharide such as sucrose or lactose, a polysaccharide such as dextran 40, dextran 60, or starch, or a sugar alcohol such as mannitol. It should be obvious to those skilled in the art that all components of the

solution contribute to the osmolality of the solution but to achieve an isotonic or near-isotonic solution, the contributions of those components should be taken into account to ensure that the proper proportion of osmolar component is added and an excess of osmolar component is not added which would result in a hypertonic solution. In fact, when the composition described above including heparin sodium as the heparinoid, lidocaine hydrochloride as the anesthetic, and sodium bicarbonate as the buffer, the osmolar contributions of the sodium ion from the heparin sodium and sodium bicarbonate, the chloride ion from the lidocaine hydrochloride, and the carbonate/bicarbonate ion from the sodium bicarbonate are sufficient not to require an additional osmolar component. However, the solution does not need to be isotonic; it can be hypotonic, isotonic, or hypertonic since the bladder epithelium can tolerate a large range of osmolality.

[0048] If an antibacterial agent is present, the antibacterial agent can be selected from the group consisting of a sulfonamide, a penicillin, a combination of trimethoprim plus sulfamethoxazole, a quinolone, methenamine, nitrofurantoin, a cephalosporin, a carbapenem, an aminoglycoside, a tetracycline, and a macrolide. Suitable sulfonamides include, but are not limited to, sulfanilamide, sulfadiazine, sulfamethoxazole, sulfisoxazole, sulfamethizole, sulfadoxine, and sulfacetamide. Suitable penicillins include, but are not limited to, methicillin, nafcillin, oxacillin, cloxacillin, dicloxacillin, ampicillin, amoxicillin, bacampicillin, carbenicillin, ticarcillin, mezlocillin, and piperacillin. Suitable quinolones include, but are not limited to, nalidixic acid, cinoxacin, norfloxacin, ciprofloxacin, ofloxacin, sparfloxacin, lomefloxacin, fleroxacin, pefloxacin, and amifloxacin. Suitable cephalosporins include, but are not limited to, cephalothin, cephazolin, cephalexin, cefadroxil, cefamandole, cefoxatin, cefaclor, cefuroxime, loracarbef, cefonicid, cefotetan, ceforanide, cefotaxime, cefpodoxime proxetil, ceftizoxime, ceftriaxone, cefoperazone, ceftazidime, and cefepime. Suitable carbapenems include, but are not limited to, imipenem, meropenem, and aztreonam. Suitable aminoglycosides include, but are not limited to, netilmycin and gentamicin. Suitable tetracyclines include, but are not limited to, tetracycline,

oxytetracycline, demeclocycline, minocycline, doxycycline, and chlortetracycline. Suitable macrolides include, but are not limited to, erythromycin, clarithromycin, and azithromycin.

[0049] If an antifungal agent is present, the antifungal agent can be selected from the group consisting of amphotericin B, itraconazole, ketoconazole, fluconazole, miconazole, and flucytosine.

[0050] If a vasoconstrictor is present, the vasoconstrictor can be epinephrine.

[0051] If a compound that enables persistence of the composition to the surface of the bladder epithelium is present, the compound is typically an activatable gelling agent. The activatable gelling agent is typically a thermoreversible gelling agent. The thermoreversible gelling agent can be selected from the group consisting of Pluronic F127 gel, Lutrol gel, N-isopropylacrylamide, ethylmethacrylate, N-acryloxysuccinimide, xyloglucan sols of 1-2%, graft copolymers of pluronic and poly(acrylic acid), pluronic-chitosan hydrogels, and a [poly(ethylene glycol)-poly(lactic acid-co-glycolic acid)]-poly(ethylene glycol)] (PEG-PLGA-PEG) copolymer.

[0052] If a preservative is present, the preservative can be selected from the group consisting of parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and benzoic acid; benzoic acid is now being used clinically with good success. However, typically, compositions prepared by methods according to the present invention do not require a preservative component, as the compositions are prepared and dispensed in sealed single-unit-dose vials.

[0053] If any of these optional components, i.e., the osmolar component, the compound that enables persistence of the composition to the surface of the bladder epithelium, the antibacterial component, the antifungal compound, the vasoconstrictor, or the preservative, are present, they are typically added after a stable solution including

the heparinoid, the local anesthetic, and the buffer has been prepared. The quantities of these additional optional components, if used, are chosen such that the solution of the heparinoid, the local anesthetic, and the buffer remains stable and precipitation of the local anesthetic is avoided and the final pH of the solution is achieved; the final pH is typically from about 6.7 to about 8.3 as described above. An optimum pH is about 7.4.

[0054] If sterilization of the composition is required, it is typically performed by filtration. Other sterilization methods are known in the art.

[0055] Accordingly, one aspect of the present invention is a method for preparing a composition useful for treatment of a lower urinary tract condition comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (1) providing a heparinoid in solid form or liquid form;
- (2) providing a local anesthetic in solid form or liquid form;
- (3) adding a liquid buffer to the heparinoid in solid form or liquid form;
- (4) adding the local anesthetic to the mixture of the liquid buffer and the heparinoid; and
- (5) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved that is from about 6.7 to about 8.3 without precipitation of the local anesthetic. An optimum pH is about 7.4.

[0056] As detailed above, the heparinoid or the local anesthetic can be provided in powdered form at the initiation of the mixing process when the mixing process starts with the heparinoid or local anesthetic in solid form. All possible combinations of solid form and liquid form are possible for this method; in other words, it is possible to use: (i) both the heparinoid and the local anesthetic in solid form; (ii) both the heparinoid and the local anesthetic in liquid form; (iii) the heparinoid in solid form, with the local anesthetic in liquid form; or (iv) the heparinoid in liquid form with the local anesthetic in solid form. Typically, as used herein, "liquid form" refers to a solution, such as an

aqueous solution. In another alternative, as described above, powders of all three components, namely the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffering agent, can be hydrated simultaneously if the lidocaine does not precipitate. In yet another alternative, liquid solutions of all three components, namely the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffering agent, can be mixed simultaneously provided that the lidocaine does not precipitate; for example, this can be done with lower levels of buffer when the desired pH is closer to a pH value of 7.0.

[0057] Another aspect of the present invention is a method for preparing a composition useful for treatment of a lower urinary tract condition comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (1) providing a heparinoid in solid form or liquid form;
- (2) providing a local anesthetic in solid form or liquid form;
- (3) mixing the heparinoid in solid form or liquid form and the local anesthetic in solid form or liquid form;
- (4) adding a liquid buffer to the mixture of the heparinoid and the local anesthetic to form a mixture of liquid buffer, the heparinoid, and the local anesthetic; and
- (5) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.7 to about 8.3 without precipitation of the local anesthetic. An optimum pH is about 7.4.

[0058] As detailed above, the heparinoid or the local anesthetic can be provided in powdered form at the initiation of the mixing process when the mixing process starts with the heparinoid or local anesthetic in solid form. All possible combinations of solid form and liquid form are possible for this method; in other words, it is possible to use: (i) both the heparinoid and the local anesthetic in solid form; (ii) both the heparinoid and the local anesthetic in liquid form; (iii) the heparinoid in solid form, with the local anesthetic in liquid form; or (iv) the heparinoid in liquid form with the local anesthetic in

solid form. In another alternative, as described above, powders of all three components, namely the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffering agent, can be hydrated simultaneously if the lidocaine does not precipitate. In yet another alternative, liquid solutions of all three components, namely the heparinoid such as heparin, the local anesthetic such as lidocaine, and the buffering agent, can be mixed simultaneously provided that the lidocaine does not precipitate; for example, this can be done with lower levels of buffer when the desired pH is closer to a pH value of 7.0.

[0059] In these methods, the heparinoid, the local anesthetic, and the buffer are as described above. In particular, a preferred heparinoid is heparin, a preferred local anesthetic is lidocaine, in the form of lidocaine hydrochloride, and a preferred buffer is sodium bicarbonate.

[0060] These methods can further comprise an additional step of adding one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative, subsequent to preparation of a buffered composition including the heparinoid, the local anesthetic, and the buffer.

[0061] Yet another aspect of the present invention is a method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (1) providing a heparinoid in liquid form;
- (2) providing a local anesthetic in liquid form;
- (3) providing a buffer in liquid form;
- (4) simultaneously mixing the heparinoid, the local anesthetic, and the buffer, avoiding precipitation of the local anesthetic; and

(5) if necessary, adjusting the pH of the mixture of the liquid buffer, the liquid local anesthetic, and the liquid heparinoid so that a pH of from about 6.7 to about 8.3 is achieved without precipitation of the local anesthetic.

[0062] Preferred components for the heparinoid, the local anesthetic, and the buffer are as described above.

[0063] Yet another aspect of the present invention is a premixed liquid composition comprising a heparinoid, a local anesthetic, and a buffer that is stable for at least three months without precipitation of the local anesthetic. Typically, the composition is prepared by a method according to the present invention as described above. Typically, in this composition, the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is sodium bicarbonate. Typically, the pH of this composition is from about 6.7 to about 8.3. Preferably, the pH of this composition is from about 7.0 to about 7.8. More preferably, the pH of this composition is about 7.4.

[0064] Compositions according to the present invention are suitable for treating, ameliorating, or preventing a lower urinary tract disorder selected from the group consisting of bacterial cystitis, fungal/yeast cystitis, vulvar vestibulitis, vulvodynia, dyspareunia, urethral syndrome, and endometriosis in women; prostatitis and chronic pelvic pain syndrome in men; and radiation-induced cystitis, chemotherapy-induced cystitis, interstitial cystitis, and overactive bladder in men or women. Compositions according to the present invention are particularly useful in treating interstitial cystitis.

[0065] As used herein, the terms "treat, ameliorate, or prevent" refer to any detectable improvement, whether subjective or objective, in the lower urinary tract disorder of the subject to whom the composition is administered. For example, the terms "treat, ameliorate, or prevent" can refer to an improvement as determined by the PORIS scale, the PUF scale, or any component of those scales; reduction of pain; reduction of urinary frequency; reduction of urinary urgency; reduction of requirement

for narcotic administration; reduction of incontinence; reduction of abnormal permeability of the urothelium to potassium; or improvement in more than one of these parameters. The terms "treat, ameliorate, or prevent" do not state or imply a cure for the underlying lower urinary tract disorder.

[0066] The invention is illustrated by the following Example. This Example is included for illustrative purposes only, and is not intended to limit the invention.

EXAMPLE

[0067] The following experiment was performed to test the efficacy of compositions prepared according to the present invention and compare them with compositions prepared by previous methods.

[0068] Solution 1 was prepared as follows: Commercially available heparin, lidocaine (approved for human use by the FDA) were mixed as follows: 5 ml heparin solution containing 50,000 units of heparin, 5 ml of 4% lidocaine (200 mg) and 2 ml of 8.4% sodium bicarbonate (168 mg) were mixed by adding the heparin and bicarbonate together and then lastly adding lidocaine with a final pH of 7.4. The solution became turbid after 2-3 minutes reflecting loss of lidocaine from solution.

[0069] Solution 2 was prepared as follows: Powdered sodium heparin (50,000 units) and powdered lidocaine hydrochloride (200 mg) were dissolved in a total of 13 ml water and 2 ml of 8.4% sodium bicarbonate (168 mg) were then added, yielding a final pH of 7.4. The solution was sterile filtered and put into a stoppered vial and remained clear and stable. However if the first step was to mix the liquid lidocaine with buffer without the presence of the heparin, the solution became turbid reflecting the loss of lidocaine from solution.

[0070] To test clinically the two different solutions (Solutions 1 and 2, above), they were each given to 5 different patients with diagnosed interstitial cystitis. The patients received in a blinded fashion each solution on different days and were then asked to compare the solutions. All five found Solution 2 to be at least 50% more effective than Solution 1.

[0071] These results indicate improved efficacy of a composition prepared by a method according to the present invention, with improved bioavailability of the local anesthetic, in this case, lidocaine.

[0072] It proved possible to compose the solution and avoid precipitation of the lidocaine at a pH value greater than pH 8.3.

[0073] Clinically patients have done well with three pH's tested; one was 7.06, another was 7.45 and yet another was 7.72. In addition at all pH's the lidocaine was in solution when starting with powdered heparin and lidocaine being mixed and then buffered accordingly with either tris or sodium bicarbonate.

ADVANTAGES OF THE INVENTION

[0074] The present invention provides an improved method of preparing compositions for treatment of a lower urinary tract disorder that include a heparinoid, a local anesthetic, and a buffer, typically heparin, lidocaine, and sodium bicarbonate. Compositions prepared by the method of the present invention are stable for three months or more and do not undergo precipitation of the local anesthetic. Accordingly, they retain efficacy and the local anesthetic retains bioavailability, an improvement over previously available compositions.

[0075] Compositions according to the present invention possess industrial applicability as compositions intended for medical use, specifically to treat lower urinary tract diseases and conditions. Methods according to the present invention possess industrial applicability for the preparation of a medicament to treat lower urinary tract diseases and conditions.

[0076] With respect to ranges of values, the invention encompasses each intervening value between the upper and lower limits of the range to at least a tenth of the lower limit's unit, unless the context clearly indicates otherwise. Moreover, the invention encompasses any other stated intervening values and ranges including either or both of the upper and lower limits of the range, unless specifically excluded from the stated range.

[0077] Unless defined otherwise, the meanings of all technical and scientific terms used herein are those commonly understood by one of ordinary skill in the art to which this invention belongs. One of ordinary skill in the art will also appreciate that any methods and materials similar or equivalent to those described herein can also be used to practice or test this invention.

[0078] The publications and patents discussed herein are provided solely for their disclosure prior to the filing date of the present application. Nothing herein is to be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention. Further the dates of publication provided may be different from the actual publication dates which may need to be independently confirmed.

[0079] All the publications cited are incorporated herein by reference in their entireties, including all published patents, patent applications, and literature references, as well as those publications that have been incorporated in those published documents. However, to the extent that any publication incorporated herein by

reference refers to information to be published, applicants do not admit that any such information published after the filing date of this application to be prior art.

[0080] As used in this specification and in the appended claims, the singular forms include the plural forms. For example the terms "a," "an," and "the" include plural references unless the content clearly dictates otherwise. Additionally, the term "at least" preceding a series of elements is to be understood as referring to every element in the series. The inventions illustratively described herein can suitably be practiced in the absence of any element or elements, limitation or limitations, not specifically disclosed herein. Thus, for example, the terms "comprising," "including," "containing," etc. shall be read expansively and without limitation. Additionally, the terms and expressions employed herein have been used as terms of description and not of limitation, and there is no intention in the use of such terms and expressions of excluding any equivalents of the future shown and described or any portion thereof, and it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the inventions herein disclosed can be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of the inventions disclosed herein. The inventions have been described broadly and generically herein. Each of the narrower species and subgeneric groupings falling within the scope of the generic disclosure also form part of these inventions. This includes the generic description of each invention with a proviso or negative limitation removing any subject matter from the genus, regardless of whether or not the excised materials specifically resided therein. In addition, where features or aspects of an invention are described in terms of the Markush group, those schooled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group. It is also to be understood that the above description is intended to be illustrative and not restrictive. Many embodiments will be apparent to those of in the art upon reviewing the above description. The scope of the invention should therefore,

be determined not with reference to the above description, but should instead be determined with reference to the appended claims, along with the full scope of equivalents to which such claims are entitled. Those skilled in the art will recognize, or will be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described. Such equivalents are intended to be encompassed by the following claims.

I claim:

1. A method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:
 - (a) providing a heparinoid in solid form or liquid form;
 - (b) providing a local anesthetic in solid form or liquid form;
 - (c) adding a liquid buffer to the heparinoid in solid form or liquid form;
 - (d) adding the local anesthetic to the mixture of the liquid buffer and the heparinoid; and
 - (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.7 to about 8.3 without precipitation of the local anesthetic.
2. The method of claim 1 wherein the heparinoid is selected from the group consisting of heparin, chondroitin sulfate, heparan sulfate, hyaluronic acid, keratan sulfate, dermatan sulfate, hyaluronan, and sodium pentosanpolysulfate.
3. The method of claim 2 wherein the heparinoid is selected from the group consisting of heparin and sodium pentosanpolysulfate.
4. The method of claim 3 wherein the heparinoid is heparin.
5. The method of claim 4 wherein the heparin is heparin sodium.
6. The method of claim 4 wherein the heparin has a molecular weight of from about 8,000 daltons to about 40,000 daltons.

7. The method of claim 4 wherein the heparin has a molecular weight of from 2,000 daltons to about 8,000 daltons.

8. The method of claim 3 wherein the heparinoid is sodium pentosanpolysulfate.

9. The method of claim 1 wherein the local anesthetic is selected from the group consisting of benzocaine, lidocaine, tetracaine, bupivacaine, cocaine, etidocaine, flecainide, mepivacaine, pramoxine, prilocaine, procaine, chlorprocaine, oxyprocaine, proparacaine, ropivacaine, dyclonine, dibucaine, propoxycaine, chloroxylonol, cinchocaine, dexivacaine, diamocaine, hexylcaine, levobupivacaine, propoxycaine, pyrrocaine, risocaine, rodocaine, and pharmaceutically acceptable derivatives and bioisosteres thereof, and a combination thereof.

10. The method of claim 9 wherein the local anesthetic is selected from the group consisting of lidocaine, bupivacaine, benzocaine, tetracaine, etidocaine, flecainide, prilocaine, and dibucaine, and a combination thereof.

11. The method of claim 10 wherein the local anesthetic is lidocaine.

12. The method of claim 11 wherein the lidocaine is lidocaine hydrochloride.

13. The method of claim 1 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris (Tris(hydroxymethyl)aminomethane) buffer, MOPS buffer (3-(N-morpholino)propanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N-(2-ethanesulfonic acid) buffer, ACES (2-[(2-amino-2-oxoethyl)amino]ethanesulfonic acid) buffer, ADA (N-(2-acetamido)2-iminodiacetic acid) buffer, AMPSO (3-[(1,1-dimethyl-2-hydroxyethyl)amino]-2-propanesulfonic acid) buffer, BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid buffer, Bicine (N,N-bis(2-hydroxyethylglycine)

buffer, Bis-Tris (bis-(2-hydroxyethyl)imino-tris(hydroxymethyl)methane buffer, CAPS (3-(cyclohexylamino)-1-propanesulfonic acid) buffer, CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) buffer, CHES (2-(N-cyclohexylamino)ethanesulfonic acid) buffer, DIPSO (3-[N,N-bis(2-hydroxyethyl)amino]-2-hydroxy-propanesulfonic acid) buffer, HEPPS (N-(2-hydroxyethyl)piperazine)-N'-(3-propanesulfonic acid) buffer, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid) buffer, MES (2-(N-morpholino)ethanesulfonic acid) buffer, triethanolamine buffer, imidazole buffer, glycine buffer, ethanolamine buffer, phosphate buffer, MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid) buffer, PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid) buffer, POPSO (piperazine-N,N'-bis(2-hydroxypropaneulfonic acid) buffer, TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid) buffer; TAPSO (3-[N-tris(hydroxymethyl)methylamino]-2-hydroxy-propanesulfonic acid) buffer, TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid) buffer, tricine (N-tris(hydroxymethyl)methylglycine buffer), 2-amino-2-methyl-1,3-propanediol buffer, 2-amino-2-methyl-1-propanol buffer, and a combination thereof.

14. The method of claim 13 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris buffer, and a combination thereof.

15. The method of claim 14 wherein the buffer is bicarbonate buffer.

16. The method of claim 15 wherein the buffer is sodium bicarbonate.

17. The method of claim 13 wherein the buffer is Tris buffer.

18. The method of claim 1 wherein the method further comprises the step of adding one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative,

subsequent to preparation of a buffered composition including the heparinoid, the local anesthetic, and the buffer.

19. The method of claim 1 wherein both the heparinoid and the local anesthetic are provided in solid form.

20. The method of claim 19 wherein both the heparinoid and the local anesthetic are provided in powdered form.

21. The method of claim 1 wherein both the heparinoid and the local anesthetic are provided in liquid form.

22. The method of claim 1 wherein the heparinoid is provided in solid form and the local anesthetic is provided in liquid form.

23. The method of claim 22 wherein the heparinoid is provided in powdered form.

24. The method of claim 1 wherein the heparinoid is provided in liquid form and the local anesthetic is provided in solid form.

25. The method of claim 24 wherein the local anesthetic is provided in powdered form.

26. The method of claim 1 wherein the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is bicarbonate buffer.

27. The method of claim 26 wherein the heparin is heparin sodium, the local anesthetic is lidocaine hydrochloride, and the bicarbonate buffer is sodium bicarbonate.

28. The method of claim 1 wherein the pH achieved is from about 7.0 to about 7.8.
29. The method of claim 1 wherein the pH achieved is about 7.4.
30. The method of claim 1 wherein the heparinoid, the local anesthetic, and the buffer are originally provided in solid form and hydrated simultaneously.
31. A method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:
- (a) providing a heparinoid in solid form or liquid form;
 - (b) providing a local anesthetic in solid form or liquid form;
 - (c) mixing the heparinoid in solid form or liquid form and the local anesthetic in solid form or liquid form;
 - (d) adding a liquid buffer to the mixture of the heparinoid and the local anesthetic to form a mixture of liquid buffer, the heparinoid, and the local anesthetic; and
 - (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH of from about 6.7 to about 8.3 is achieved without precipitation of the local anesthetic.
32. The method of claim 31 wherein the heparinoid is selected from the group consisting of heparin, chondroitin sulfate, heparan sulfate, hyaluronic acid, keratan sulfate, dermatan sulfate, hyaluronan, and sodium pentosanpolysulfate.
33. The method of claim 32 wherein the heparinoid is selected from the group consisting of heparin and sodium pentosanpolysulfate.

34. The method of claim 33 wherein the heparinoid is heparin.
35. The method of claim 31 wherein the heparin is heparin sodium.
36. The method of claim 34 wherein the heparin has a molecular weight of from about 8,000 daltons to about 40,000 daltons.
37. The method of claim 34 wherein the heparin has a molecular weight of from 2,000 daltons to about 8,000 daltons.
38. The method of claim 32 wherein the heparinoid is sodium pentosanpolysulfate.
39. The method of claim 31 wherein the local anesthetic is selected from the group consisting of benzocaine, lidocaine, tetracaine, bupivacaine, cocaine, etidocaine, flecainide, mepivacaine, pramoxine, prilocaine, procaine, chlorprocaine, oxyprocaine, proparacaine, ropivacaine, dyclonine, dibucaine, propoxycaine, chloroxylenol, cinchocaine, dexivacaine, diamocaine, hexylcaine, levobupivacaine, propoxycaine, pyrrocaine, risocaine, rodocaine, and pharmaceutically acceptable derivatives and bioisosteres thereof, and a combination thereof.
40. The method of claim 39 wherein the local anesthetic is selected from the group consisting of lidocaine, bupivacaine, benzocaine, tetracaine, etidocaine, flecainide, prilocaine, and dibucaine, and a combination thereof.
41. The method of claim 40 wherein the local anesthetic is lidocaine.
42. The method of claim 41 wherein the lidocaine is lidocaine hydrochloride.

43. The method of claim 31 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris (Tris(hydroxymethyl)aminomethane) buffer, MOPS buffer (3-(N-morpholino)propanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N-(2-ethanesulfonic acid) buffer, ACES (2-[(2-amino-2-oxoethyl)amino]ethanesulfonic acid) buffer, ADA (N-(2-acetamido)2-iminodiacetic acid) buffer, AMPSO (3-[(1,1-dimethyl-2-hydroxyethyl)amino]-2-propanesulfonic acid) buffer, BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid buffer, Bicine (N,N-bis(2-hydroxyethylglycine) buffer, Bis-Tris (bis-(2-hydroxyethyl)imino-tris(hydroxymethyl)methane buffer, CAPS (3-(cyclohexylamino)-1-propanesulfonic acid) buffer, CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) buffer, CHES (2-(N-cyclohexylamino)ethanesulfonic acid) buffer, DIPSO (3-[N,N-bis(2-hydroxyethyl)amino]-2-hydroxy-propanesulfonic acid) buffer, HEPPS (N-(2-hydroxyethyl)piperazine)-N'-(3-propanesulfonic acid) buffer, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid) buffer, MES (2-(N-morpholino)ethanesulfonic acid) buffer, triethanolamine buffer, imidazole buffer, glycine buffer, ethanolamine buffer, phosphate buffer, MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid) buffer, PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid) buffer, POPSO (piperazine-N,N'-bis(2-hydroxypropaneulfonic acid) buffer, TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid) buffer; TAPSO (3-[N-tris(hydroxymethyl)methylamino]-2-hydroxy-propanesulfonic acid) buffer, TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid) buffer, tricine (N-tris(hydroxymethyl)methylglycine buffer), 2-amino-2-methyl-1,3-propanediol buffer, 2-amino-2-methyl-1-propanol buffer, and a combination thereof.

44. The method of claim 43 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris buffer, and a combination thereof.

45. The method of claim 44 wherein the buffer is bicarbonate buffer.

46. The method of claim 44 wherein the buffer is sodium bicarbonate.
47. The method of claim 44 wherein the buffer is Tris buffer.
48. The method of claim 31 wherein the method further comprises the step of adding one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative, subsequent to preparation of a buffered composition including the heparinoid, the local anesthetic, and the buffer.
49. The method of claim 31 wherein both the heparinoid and the local anesthetic are provided in solid form.
50. The method of claim 49 wherein both the heparinoid and the local anesthetic are provided in powdered form.
51. The method of claim 31 wherein both the heparinoid and the local anesthetic are provided in liquid form.
52. The method of claim 31 wherein the heparinoid is provided in solid form and the local anesthetic is provided in liquid form.
53. The method of claim 52 wherein the heparinoid is provided in powdered form.
54. The method of claim 31 wherein the heparinoid is provided in liquid form and the local anesthetic is provided in solid form.

55. The method of claim 54 wherein the local anesthetic is provided in powdered form.

56. The method of claim 31 wherein the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is bicarbonate buffer.

57. The method of claim 56 wherein the heparin is heparin sodium, the local anesthetic is lidocaine hydrochloride, and the bicarbonate buffer is sodium bicarbonate.

58. The method of claim 29 wherein the pH achieved is about 7.4.

59. A method for preparing a composition useful for treatment of a lower urinary tract condition by instilling the composition into the urinary bladder, the composition comprising a mixture comprising a heparinoid, a local anesthetic, and a buffer, the method comprising the steps of:

- (a) providing a heparinoid in liquid form;
- (b) providing a local anesthetic in liquid form;
- (c) providing a buffer in liquid form;
- (d) simultaneously mixing the heparinoid, the local anesthetic, and the buffer, avoiding precipitation of the local anesthetic; and
- (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the liquid local anesthetic, and the liquid heparinoid so that a pH of from about 6.7 to about 8.3 is achieved without precipitation of the local anesthetic.

60. The method of claim 59 wherein the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is sodium bicarbonate.

61. The method of claim 59 wherein the pH achieved is about 6.8 to about 7.2 and the quantity of the buffer required is reduced.

62. A premixed liquid composition comprising a heparinoid, a local anesthetic, and a buffer that is stable for at least three months without precipitation of the local anesthetic.

63. The composition of claim 62 wherein the composition is prepared by a method comprising the steps of:

- (a) providing a heparinoid in solid form or liquid form;
- (b) providing a local anesthetic in solid form or liquid form;
- (c) adding a liquid buffer to the heparinoid in solid form or liquid form;
- (d) adding the local anesthetic to the mixture of the liquid buffer and the heparinoid; and
- (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.8 to about 8.3 is achieved without precipitation of the local anesthetic.

64. The composition of claim 62 wherein the composition is prepared by a method comprising the steps of:

- (a) providing a heparinoid in solid form or liquid form;
- (b) providing a local anesthetic in solid form or liquid form;
- (c) mixing the heparinoid in solid form or liquid form and the local anesthetic in solid form or liquid form;
- (d) adding a liquid buffer to the mixture of the heparinoid and the local anesthetic to form a mixture of liquid buffer, the heparinoid, and the local anesthetic; and
- (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the local anesthetic, and the heparinoid so that a pH is achieved of from about 6.8 to about 8.3 is achieved without precipitation of the local anesthetic.

65. The composition of claim 62 wherein the composition is prepared by a method comprising the steps of:

- (a) providing a heparinoid in liquid form;
- (b) providing a local anesthetic in liquid form;
- (c) providing a buffer in liquid form;
- (d) simultaneously mixing the heparinoid, the local anesthetic, and the buffer, avoiding precipitation of the local anesthetic; and
- (e) if necessary, adjusting the pH of the mixture of the liquid buffer, the liquid local anesthetic, and the liquid heparinoid so that a pH of from about 6.7 to about 8.3 is achieved without precipitation of the local anesthetic.

66. The composition of claim 62 wherein the heparinoid is selected from the group consisting of heparin, chondroitin sulfate, heparan sulfate, hyaluronic acid, keratan sulfate, dermatan sulfate, hyaluronan, and sodium pentosanpolysulfate.

67. The composition of claim 66 wherein the heparinoid is selected from the group consisting of heparin and sodium pentosanpolysulfate.

68. The composition of claim 67 wherein the heparinoid is heparin.

69. The composition of claim 68 wherein the heparin is heparin sodium.

70. The composition of claim 68 wherein the heparin has a molecular weight of from about 8,000 daltons to about 40,000 daltons.

71. The composition of claim 68 wherein the heparin has a molecular weight of from 2,000 daltons to about 8,000 daltons.

72. The composition of claim 67 wherein the heparinoid is sodium pentosanpolysulfate.

73. The composition of claim 62 wherein the local anesthetic is selected from the group consisting of benzocaine, lidocaine, tetracaine, bupivacaine, cocaine, etidocaine, flecainide, mepivacaine, pramoxine, prilocaine, procaine, chloroprocaine, oxyprocaine, proparacaine, ropivacaine, dyclonine, dibucaine, propoxycaine, chloroxylenol, cinchocaine, dexivacaine, diamocaine, hexylcaine, levobupivacaine, propoxycaine, pyrrocaine, risocaine, rodocaine, and pharmaceutically acceptable derivatives and bioisosteres thereof, and a combination thereof.

74. The composition of claim 73 wherein the local anesthetic is selected from the group consisting of lidocaine, bupivacaine, benzocaine, tetracaine, etidocaine, flecainide, prilocaine, and dibucaine, and a combination thereof.

75. The composition of claim 74 wherein the local anesthetic is lidocaine.

76. The composition of claim 75 wherein the lidocaine is lidocaine hydrochloride.

77. The composition of claim 63 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris (Tris(hydroxymethyl)aminomethane) buffer, MOPS buffer (3-(N-morpholino)propanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N-(2-ethanesulfonic acid) buffer, ACES (2-[(2-amino-2-oxoethyl)amino]ethanesulfonic acid) buffer, ADA (N-(2-acetamido)2-iminodiacetic acid) buffer, AMPSO (3-[(1,1-dimethyl-2-hydroxyethyl)amino]-2-propanesulfonic acid) buffer, BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid buffer, Bicine (N,N-bis(2-hydroxyethylglycine) buffer, Bis-Tris (bis-(2-hydroxyethyl)imino-tris(hydroxymethyl)methane buffer, CAPS (3-(cyclohexylamino)-1-propanesulfonic acid) buffer, CAPSO (3-(cyclohexylamino)-2-hydroxy-1-propanesulfonic acid) buffer, CHES (2-(N-cyclohexylamino)ethanesulfonic acid) buffer, DIPSO (3-[N,N-bis(2-

hydroxyethyl)amino]-2-hydroxy-propanesulfonic acid) buffer, HEPPS (N-(2-hydroxyethylpiperazine)-N'-(3-propanesulfonic acid) buffer, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid) buffer, MES (2-(N-morpholino)ethanesulfonic acid) buffer, triethanolamine buffer, imidazole buffer, glycine buffer, ethanolamine buffer, phosphate buffer, MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid) buffer, PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid) buffer, POPSO (piperazine-N,N'-bis(2-hydroxypropaneulfonic acid) buffer, TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid) buffer; TAPSO (3-[N-tris(hydroxymethyl)methylamino]-2-hydroxy-propanesulfonic acid) buffer, TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid) buffer, tricine (N-tris(hydroxymethyl)methylglycine buffer), 2-amino-2-methyl-1,3-propanediol buffer, 2-amino-2-methyl-1-propanol buffer, and a combination thereof.

78. The composition of claim 77 wherein the buffer is selected from the group consisting of bicarbonate buffer, Tris buffer, and a combination thereof.

79. The composition of claim 78 wherein the buffer is bicarbonate buffer.

80. The composition of claim 79 wherein the buffer is sodium bicarbonate.

81. The composition of claim 78 wherein the buffer is Tris buffer.

82. The composition of claim 63 wherein the composition further comprises one or more of: (i) an osmolar component; (ii) a compound that enables persistence of the composition to the surface of the bladder epithelium; (iii) an antibacterial agent; (iv) an antifungal agent; (v) a vasoconstrictor; or (vi) a preservative.

83. The composition of claim 63 wherein the heparinoid is heparin, the local anesthetic is lidocaine, and the buffer is bicarbonate buffer.

84. The composition of claim 83 wherein the heparin is heparin sodium, the local anesthetic is lidocaine hydrochloride, and the bicarbonate buffer is sodium bicarbonate.

85. The composition of claim 63 wherein the composition is suitable for treating, ameliorating, or preventing a lower urinary tract disorder selected from the group consisting of bacterial cystitis, fungal/yeast cystitis, vulvar vestibulitis, vulvodynia, dyspareunia, urethral syndrome, and endometriosis in women; prostatitis and chronic pelvic pain syndrome in men; and radiation-induced cystitis, chemotherapy-induced cystitis, interstitial cystitis, and overactive bladder in men or women.

86. The composition of claim 85 wherein the lower urinary tract disorder is interstitial cystitis.

87. The composition of claim 63 wherein the pH of the composition is from about 7.0 to about 7.8.

88. The composition of claim 87 wherein the pH of the composition is about 7.4.