

trials to treat NMIBC by instillation in the bladder with various weekly schedules. Gemcitabine is typically instilled over 1 to 2 hours once or twice a week for several weeks at doses typically ranging from 500 to 2000 mg in up to 100 mL of saline.

[0009] It is known that such liquid formulations are voided from the bladder after short dwell times of 1 to 2 hours thus limiting their therapeutic benefit. In addition, high concentrations (40 mg/mL) and high doses (up to 2 grams per instillation) are used in an attempt to achieve therapeutic tissue levels with limited dwell time. However, in addition to local tolerability issues, intravesical delivery of high doses of gemcitabine can lead to significant systemic absorption and cause gastrointestinal, bladder and bone marrow toxicity further limiting its utility.

[0010] Systemic gemcitabine is often used in combination with systemic cisplatin both neoadjuvantly and adjuvantly to radical cystectomy to treat muscle invasive bladder cancer and to treat metastatic disease. However, many patients are ineligible or refuse treatment with these agents due to their side effects which include bone marrow suppression.

[0011] Accordingly, there remains a need for methods for treating tumors at distant sites in an individual with a tumor in the bladder.

[0012] The disclosures of all publications, patents, patent applications and published patent applications referred to herein are hereby incorporated herein by reference in their entirety.

SUMMARY

[0013] The present disclosure in various examples provides methods of treating tumor metastasis, reducing (such as eradicating) preexisting tumor metastasis, reducing incidence or burden of preexisting tumor metastasis, suppressing or delaying tumor metastasis, inhibiting tumor cells at a second site distinct from a first tumor site in the bladder of an individual comprising locally delivering to the bladder an effective amount of a chemotherapeutic agent, wherein the chemotherapeutic agent is delivered continuously to the bladder for period of time.

[0014] In some examples, there is provided a method of inhibiting tumor cell growth at a second tumor site distinct from a first tumor site in the bladder of an individual, comprising locally delivering to the bladder an effective amount of a chemotherapeutic agent,

wherein the chemotherapeutic agent is delivered continuously to the bladder for at least about 24 hours. In some examples, the tumor cells at the second tumor site are located at a pelvic node. In some examples, the tumor cells at the second tumor site are located at a distant node. In some examples, the tumor cells at the second tumor site are circulating tumor cells. In some examples, the tumor cells at the first tumor site result from metastasis of a primary tumor at the second tumor site. In some examples, the tumor cells at the second site are selected from the group consisting of liver, lung, bone, brain, lymph node, pelvic node, peritoneum, skin, prostate, breast, colon, rectum, and cervix. In some examples, the individual has a urothelial carcinoma of lower tract.

[0014a] In one aspect, the present disclosure refers to use of an effective amount of a chemotherapeutic agent in the manufacture of a medicament for inhibiting tumor cell growth at a second tumor site distinct from a first tumor site in the bladder of an individual, wherein the chemotherapeutic agent is to be delivered locally and continuously to the bladder for at least about 24 hours.

[0015] In some examples, also provided herein is a method of treating or suppressing tumor metastasis at a site distinct from the bladder in an individual having a urothelial carcinoma of lower tract, comprising locally delivering to the bladder an effective amount of a chemotherapeutic agent, wherein the chemotherapeutic agent is delivered continuously to the bladder for at least about 24 hours. In some examples, the tumor metastatic site is at one or more of: liver, lung, bone, brain, lymph node, pelvic node, peritoneum, skin, prostate, breast, colon, rectum, and cervix. In some examples, the tumor metastasis is at two or more different sites.

[0015a] In another aspect, the present disclosure refers to use of an effective amount of a chemotherapeutic agent in the manufacture of a medicament for treating or suppressing tumor metastasis at a site distinct from the bladder in an individual having a urothelial carcinoma of lower tract, wherein the chemotherapeutic agent is to be delivered locally and continuously to the bladder for at least about 24 hours.

[0016] In some examples according to any of the methods described above, the individual has bladder cancer. In some examples, the individual has muscle invasive bladder cancer or carcinoma in situ (CIS). In some examples, the individual is unfit for or refuses cystectomy.

[0017] In some examples according to any of the methods described above, the individual has not undergone transurethral resection of bladder tumors (TURBT).

[0018] In other examples, the individual has undergone transurethral resection of bladder tumors (TURBT). In some of these examples, the individual has undergone transurethral resection of bladder tumors (TURBT) and has residual tumor at the site of resection.

[0019] In some examples according to any of the methods described above, the chemotherapeutic agent is delivered at a dose of from about 1 mg/day to about 300 mg/day. In some examples, the concentration of the chemotherapeutic agent in the urine is from about 0.1 µg/mL to about 200 µg/mL during the delivery period.

[0020] In some examples according to any of the methods described above, the chemotherapeutic agent is delivered continuously to the bladder of the individual for a period of about 7 days to about three weeks. In some examples, the method comprises an induction delivery period followed by a maintenance delivery period. In some examples, the induction delivery period and the maintenance delivery period are separated by a rest period of about 7 to about 14 days. In some examples, the chemotherapeutic agent is delivered at a first release rate during the induction delivery period followed and a second release rate during the maintenance delivery period.

[0021] In some examples according to any of the methods described above, the method comprises a) an induction delivery period, wherein the concentration of chemotherapeutic agent in the urine of the individual is at least about 0.1 $\mu\text{g/mL}$; b) a rest period; and c) a maintenance delivery period, wherein the concentration of the chemotherapeutic agent in the urine of the individual is greater than about 0.1 $\mu\text{g/mL}$.

[0022] In some examples according to any of the methods described above, the individual does not receive a radiation therapy. In other examples, the method further comprises a radiation therapy.

[0023] In some examples according to any of the methods described above, the chemotherapeutic agent is delivered by an intravesical delivery device. In some examples, the intravesical device contains 100 mg to 500 mg of the chemotherapeutic agent. In some examples, the intravesical device comprises a housing configured for intravesical insertion; and a dosage form comprising chemotherapeutic agent, wherein the housing holds the dosage form and is configured to release chemotherapeutic agent. In some examples, the intravesical drug delivery device comprises: a housing defining a reservoir; a first unit contained within the reservoir, the first unit comprising an chemotherapeutic agent; and a second unit contained within the reservoir in a position distinct from the first unit, wherein the second unit comprises a functional agent that facilitates in vivo release of the chemotherapeutic agent from the housing. In some examples, the intravesical drug delivery device comprises a housing which contains and controllably releases the chemotherapeutic agent and is elastically deformable between a retention shape configured to retain the device in the individual's bladder and a deployment shape for passage of the device through the individual's urethra. In some examples, the device comprises a drug reservoir lumen bounded by a first wall and a second wall, wherein the first wall is impermeable to the

drug and the second wall is permeable to the chemotherapeutic agent. In some example, the chemotherapeutic agent is released from the device by osmotic pressure. In some examples, the chemotherapeutic agent is released from the device by diffusion. In some example, the chemotherapeutic agent contained in the housing is in a non-liquid form. In some examples, the non-liquid form is selected from the group consisting of tablets, granules, powders, semisolids, capsules, and combinations thereof.

[0024] In some examples according to any of the methods described above, the chemotherapeutic agent is selected from the group consisting of a nucleoside analog, a taxane, a platinum-based agent, and an anthracycline analogue. In some examples, the chemotherapeutic agent is a nucleoside analog. In some of these examples, the nucleoside analog is gemcitabine.

[0025] In some examples according to any of the methods described above, the individual is human. In some examples, the individual is unsuitable for systemic chemotherapy. In some examples, the individual has a compromised immune system. In some examples, the individual has a high tumor burden.

BRIEF DESCRIPTION OF THE DRAWINGS

[0026] FIG. 1 shows a modified bladder perfusion system to introduce tumors into bladder and subcutaneous tissue.

[0027] FIG. 2 shows the treatment and dosing schedule for four groups of rats: 1) untreated rats inoculated with subcutaneous tumor only; 2) untreated rats inoculated with bladder tumor and then subcutaneous tumor; 3) rats inoculated with bladder tumor, perfused with 180 µg/mL gemcitabine, and then inoculated with subcutaneous tumor and again perfused with 180µg/mL gemcitabine; and 4) rats inoculated with bladder tumor and subcutaneous tumor simultaneously and perfused with 180 µg/mL gemcitabine twice.

[0028] FIG. 3 shows the average volume of subcutaneous tumors in four groups of rats after introducing tumors into bladder and/or subcutaneous tissue and during the course of the treatment.

[0029] FIG. 4 shows the volume of subcutaneous tumors in each individual rat in the four groups after introducing tumors into bladder and/or subcutaneous tissue and during the course of the treatment.

CLAIMS

1. Use of an effective amount of a chemotherapeutic agent in the manufacture of a medicament for treating or suppressing tumor metastasis at a site distinct from the bladder in an individual having a urothelial carcinoma of lower tract, wherein the chemotherapeutic agent is to be delivered locally and continuously to the bladder for at least about 24 hours.
2. The use of claim 1, wherein the tumor metastatic site is at one or more of: liver, lung, bone, brain, lymph node, pelvic node, peritoneum, skin, prostate, breast, colon, rectum, and cervix.
3. The use of any one of claims 1-2, wherein the tumor metastasis is at two or more different sites.
4. Use of an effective amount of a chemotherapeutic agent in the manufacture of a medicament for inhibiting tumor cell growth at a second tumor site distinct from a first tumor site in the bladder of an individual, wherein the chemotherapeutic agent is to be delivered locally and continuously to the bladder for at least about 24 hours.
5. The use of claim 4, wherein the tumor cells at the second tumor site are located at a pelvic node.
6. The use of claim 4, wherein the tumor cells at the second tumor site are located at a distant node.
7. The use of claim 4, wherein the tumor cells at the second tumor site are circulating tumor cells.
8. The use claim 4, wherein the tumor cells at the first tumor site result from metastasis of a primary tumor at the second tumor site.

9. The use of claim 8, wherein the tumor cells at the second site are selected from the group consisting of liver, lung, bone, brain, lymph node, pelvic node, peritoneum, skin, prostate, breast, colon, rectum, and cervix.
10. The use of any one of claims 4-9, wherein the individual has a urothelial carcinoma of lower tract.
11. The use of any one of claims 1-10, wherein the individual has bladder cancer.
12. The use of claim 11, wherein the individual has muscle invasive bladder cancer or carcinoma in situ (CIS).
13. The use of claim 11 or 12, wherein the individual is unfit for or refuses cystectomy.
14. The use of any one of claims 11-13, wherein the individual has not undergone transurethral resection of bladder tumors (TURBT).
15. The use of any one of claims 11-13, wherein the individual has undergone transurethral resection of bladder tumors (TURBT).
16. The use of claim 15, wherein the individual has residual tumor at the site of resection.
17. The use of any one of claims 1-16, wherein the chemotherapeutic agent is to be delivered continuously to the bladder of the individual for a period of about 7 days to about three weeks.
18. The use of any one of claims 1-17, wherein the use comprises an induction delivery period followed by a maintenance delivery period.
19. The use of claim 18, wherein the induction delivery period and the maintenance delivery period are separated by a rest period of about 7 to about 14 days.
20. The use of claim 18, wherein the chemotherapeutic agent is to be delivered at a first release rate during the induction delivery period followed and a second release rate during the maintenance delivery period.

21. The use of any of claims 1-20, wherein the chemotherapeutic agent is to be delivered at a dose of from about 1 mg/day to about 300 mg/day.
22. The use of any of claims 1-21, wherein the concentration of the chemotherapeutic agent in the urine is from about 0.1 µg/mL to about 200 µg/mL during the delivery period.
23. The use of any one of claims 1-22, comprising
 - a) an induction delivery period, wherein the concentration of chemotherapeutic agent in the urine of the individual is at least about 0.1 µg/mL;
 - b) a rest period; and
 - c) a maintenance delivery period, wherein the concentration of the chemotherapeutic agent in the urine of the individual is greater than about 0.1 µg/mL.
24. The use of any of claims 1-23, wherein the individual does not receive a radiation therapy.
25. The use of any one of claims 1-24, wherein the medicament is to be delivered with a radiation therapy.
26. The use of any one of claims 1-25, wherein the chemotherapeutic agent is to be delivered by an intravesical delivery device.
27. The use of claim 26, wherein the intravesical device contains 100 mg to 500 mg of the chemotherapeutic agent.
28. The use of any of claims 26-27, wherein the intravesical device comprises a housing configured for intravesical insertion; and a dosage form comprising chemotherapeutic agent, wherein the housing holds the dosage form and is configured to release chemotherapeutic agent.

29. The use of any one of claims 26-28, wherein the intravesical drug delivery device comprises:
- a housing defining a reservoir;
 - a first unit contained within the reservoir, the first unit comprising an chemotherapeutic agent; and
 - a second unit contained within the reservoir in a position distinct from the first unit, wherein the second unit comprises a functional agent that facilitates in vivo release of the chemotherapeutic agent from the housing.
30. The use of any one of claims 26-29, wherein the intravesical drug delivery device comprises a housing which contains and controllably releases the chemotherapeutic agent and is elastically deformable between a retention shape configured to retain the device in the individual's bladder and a deployment shape for passage of the device through the individual's urethra.
31. The use of any one of claims 26-30, wherein the device comprises a drug reservoir lumen bounded by a first wall and a second wall, wherein the first wall is impermeable to the drug and the second wall is permeable to the chemotherapeutic agent.
32. The use of any one of claims 26-31 wherein the chemotherapeutic agent is released from the device by osmotic pressure.
33. The use of any one of claims 26-31, wherein the chemotherapeutic agent is released from the device by diffusion.
34. The use of any of claims 22-29, wherein the chemotherapeutic agent contained in the housing is in a non-liquid form.
35. The use of claim 34, wherein the non-liquid form is selected from the group consisting of tablets, granules, powders, semisolids, capsules, and combinations thereof.

36. The use of any one of claims 1-35, wherein the chemotherapeutic agent is selected from the group consisting of a nucleoside analog, a taxane, a platinum-based agent, and an anthracycline analogue.
37. The use of any one of claims 1-36, wherein the chemotherapeutic agent is a nucleoside analog.
38. The use of claim 37, wherein the nucleoside analog is gemcitabine.
39. The use of any one of claims 1-38, wherein the individual is human.
40. The use of any one of claims 1-39, wherein the individual is unsuitable for systemic chemotherapy.
41. The use of any one of claims 1-40, wherein the individual has a compromised immune system.
42. The use of any one of claims 1-41, wherein the individual has a high tumor burden.