**Claims**

What is claimed is:

1. ~~A method of treating cancer in the lung comprising administering to a patient in need thereof~~Use of a therapeutically effective amount of liposomal annamycin in the manufacture of a medicament for treating cancer in the lung in a patient.

2. The ~~method~~ use of claim 1, wherein said patient has primary or metastatic cancer in the lung.

3. The ~~method~~ use of claim 2, wherein the primary lung cancer is non-small-cell lung carcinoma (NSCLC) or small-cell lung carcinoma (SCLC).

4. The ~~method~~ use of claim 3, wherein the primary lung cancer is a non-small-cell lung cancer selected from adenocarcinoma, squamous-cell carcinoma, and large-cell carcinoma.

5. The ~~method~~ use of claim 2, wherein said patient has metastatic cancer in the lung.

6. The ~~method~~ use of claim 5, wherein the metastatic cancer is a metastasis of a primary cancer selected from bladder cancer, breast cancer, colorectal cancer, head and neck cancer, kidney cancer, melanoma, pancreatic cancer, prostate cancer, and ovarian cancer.

7. The ~~method~~ use of claim 5, wherein the metastatic cancer is a metastasis of a sarcoma.

8. The ~~method~~ use of claim 5, wherein the metastatic cancer is from a cancer of unknown primary origin.

9. The ~~method~~ use of any one of the preceding claims, wherein ~~further comprising administering~~ an effective amount of at least one chemotherapeutic agent is to be further administered to the subject.

10. The ~~method~~ use of claim 9, wherein the at least one chemotherapeutic agent is selected from actinomycin, afatinib, alectinib, asparaginase, azacitidine, azathioprine,

bicalutamide, bleomycin, bortezomib, camptothecin, carboplatin, capecitabine, certinib, cetuximab,cisplatin, chlorambucil, crizotinib, cyclophosphamide, cytarabine, daunorubicin, docetaxel, doxifluridine, doxorubicin, erlotinib, epirubicin, epothilone, etoposide, fludarabine, flutamine, fluorouracil, fostamatinib, gefitinib, gemcitabine, hydroxyurea, ibrutinib, idarubicin, ifosfamide, imatinib, ipilimumab, irinotecan, lapatinib, letrozole, mechlorethamine, mercaptopurine, methotrexate, mitomycin, mitoxantrone, nilotinib, octreotide, oxaliplatin, paclitaxel, palbociclib, panitumumab, pemetrexed, raltitrexed, selumetinib, sorafenib, sunitinib, tamoxifen, temozolomide, teniposide, tioguanine, topotecan, trastuzumab, tremelimumab, valrubicin, vemurafenib, vinblastine, vincristine, vindesine, vinorelbine, and combinations thereof.

11. The ~~method~~ use of claim 9, wherein the at least one chemotherapeutic agent comprises a combination selected from:

    cyclophosphamide, doxorubicin, and vincristine;

    mitomycin, vindesine and cisplatin;

    cisplatin and vinorelbine; and

    cisplatin, etoposide and ifosfamide.

12. ~~.~~The ~~method~~ use of any one of claims 1 to 8, ~~further comprising administering~~wherein an effective amount of at least one immunotherapeutic agent is to be further administered to the subject.

13. The ~~method~~ use of any one of the preceding claims, ~~further comprising one or both of resecting~~ wherein the lung cancer is to be resected and/or ~~administering~~ radiation therapy is to be administered.

14. The ~~method~~ use of any one of the preceding claims, wherein treating cancer in the lung comprises one or more of the following: (a) increasing survival time of the patient; (b) reducing volume of the primary cancer; (c) retarding growth of the primary cancer; (d) reducing number of metastatic tumors; (e) reducing volume of metastatic tumors; and (f) retarding growth of metastatic tumors.

15. The ~~method~~ use of any one of the preceding claims, wherein treating cancer in the lung does not result in annamycin-induced toxicity of the lung of such severity that repeated administration of the annamycin is contraindicated.

16. The ~~method~~ use of any one of the preceding claims, wherein treating cancer in the lung does not result in annamycin-induced systemic toxicity of such severity that repeated administration of the annamycin is contraindicated.

17. The ~~method~~ use of any one of the preceding claims, wherein the therapeutically effective amount of liposomal annamycin ~~said administering~~ is to be administered ~~repeated~~ weekly.

18. The ~~method~~ use of any one of claims 1 to 16, wherein the therapeutically effective amount of liposomal annamycin is to be administered ~~said administering is repeated~~ every two, three, or four weeks.

19. The ~~method~~ use of any one of claims 1 to 18, wherein the liposomal annamycin comprises annamycin, one or more lipids, and one or more non-ionic surfactants.

20. The ~~method~~ use of claim 19, wherein the lipids comprise Dimyristoylphosphatidylcholine (DMPC) and dimyristoylphosphatidylglycerol (DMPG).

21. The ~~method~~ use of claim 19 or 20, wherein the non-ionic surfactant comprises a polysorbate-type surfactant formed from the ethoxylation of sorbitan followed by the addition of a carboxylic acid.

22. The ~~method~~ use of claim 21 wherein the non-ionic surfactant comprises polyoxyethylene sorbitan monolaurate.

23. The ~~method~~ use of any one of the preceding claims, wherein the liposomal annamycin is provided as a preliposomal lyophilizate composition that is reconstituted into an aqueous liposome composition through hydration prior to administration.

24. The ~~method~~ use of claim 23, wherein the preliposomal annamycin lyophilizate comprises: 1.8 – 2.2 wt % annamycin;

3.0 – 3.4 wt. % Polysorbate 20; and

94.4 - 95.2 wt. % of lipids selected from DMPC and DMPG.

25. The ~~method~~ use of claim 24, wherein the DMPC is 65.3 - 67.3 wt. % and the DMPG is 27.1 – 29.9 wt. %.