

**IN THE UNITED STATES DISTRICT COURT
FOR THE EASTERN DISTRICT OF MISSOURI
EASTERN DIVISION**

ADVANCED ACCELERATOR
APPLICATIONS USA, INC. and
ADVANCED ACCELERATOR
APPLICATIONS SA,

Plaintiffs,

v.

CURIUM US LLC, CURIUM US
HOLDINGS LLC, CURIUM
NETHERLANDS BV, and CURIUM
INTERNATIONAL TRADING BV,

Defendants.

Case No.

COMPLAINT

Advanced Accelerator Applications USA, Inc. and Advanced Accelerator Applications SA (collectively, “ADACAP” or “Plaintiffs”) by their attorneys hereby allege as follows:

NATURE OF THE ACTION

1. This is an action for patent infringement arising under the patent laws of the United States, Title 35, United States Code and for declaratory judgment pursuant to the Declaratory Judgment Act, 28 U.S.C. §§ 2201, et seq. This action relates to a New Drug Application No. 218525 filed under 21 U.S.C. § 505(b)(2) by the above-named defendants with the U.S. Food and Drug Administration (“FDA”) for approval to engage in the commercial manufacture, use, offer for sale, sale, and/or import of a version of ADACAP’s LUTATHERA® (lutetium Lu 177 dotatate) injection, prior to expiration of U.S. Patent No. 10,596,276 (“the ’276

patent”), U.S. Patent No. 10,596,278 (“the ’278 patent”), and U.S. Patent No. 11,904,027 (“the ’027 patent”).

PARTIES

A. Plaintiffs

2. Advanced Accelerator Applications USA, Inc. is a corporation organized and existing under the laws of the State of Delaware, with its principal place of business at 57 E. Willow Street, Millburn, NJ 07041.

3. Advanced Accelerator Applications SA is a corporation organized and existing under the laws of France, with its principal place of business in France.

4. ADACAP is in the business of creating, developing, and bringing to market new drug therapies to benefit patients against serious diseases, including targeted radioligand therapy for the treatment for cancer. LUTATHERA® is one such treatment. ADACAP markets and sells LUTATHERA® in this judicial district and throughout the United States.

B. Defendants

5. Upon information and belief, Defendant Curium US LLC is a limited liability company organized and existing under the laws of Delaware, having a principal place of business at 111 West Port Plaza Drive Suite 800, St. Louis, MO 63146.

6. Upon information and belief, Defendant Curium US Holdings LLC is a limited liability company organized and existing under the laws of Delaware, having a principal place of business at 111 West Port Plaza Drive Suite 800, St. Louis, MO 63146.

7. Upon information and belief, Defendant Curium Netherlands BV is a Dutch besloten vennootschap organized and existing under the laws of the Netherlands, having a principal place of business at Westerduinweg 3, 1755 LE, Petten, Netherlands.

8. Upon information and belief, Defendant Curium International Trading BV is a Dutch besloten vennootschap organized and existing under the laws of the Netherlands, having a principal place of business at Westerduinweg 3, 1755 LE, Petten, Netherlands.

9. Upon information and belief, Curium US LLC is a wholly owned subsidiary of Curium US Holdings LLC.

10. Upon information and belief, Curium US Holdings LLC is a wholly owned subsidiary of Curium Netherlands BV.

11. Upon information and belief, Curium Netherlands BV is a wholly owned subsidiary of Curium International Trading BV.

12. Curium Netherlands CV and Curium International Trading BV are collectively referred to as “Curium Netherlands” hereinafter, unless otherwise noted.

13. Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV are collectively referred to as “Curium” or “Defendants” hereinafter, unless otherwise noted.

14. Upon information and belief, Curium is in the business of, *inter alia*, developing, manufacturing, marketing, distributing, and directly and/or indirectly selling pharmaceutical products throughout the United States (including in the State of Missouri).

15. By a letter dated September 5, 2024, Curium notified ADACAP that Curium had submitted to the FDA NDA No. 218525 (“Curium Application No. 218525”) for Lutetium Lu-177 Dotatate injection solution, a version of LUTATHERA® (“the 505(b)(2) Product”), seeking approval under the Federal Food, Drug and Cosmetic Act (“FDCA”) to engage in the commercial manufacture, use, offer for sale, and/or sale of Curium’s 505(b)(2) Product prior to the expiration of the ’276, ’278, and ’027 patents.

16. In its Notice Letter, Curium notified ADACAP that, as a part of Curium Application No. 218525, Curium had filed a certification of the type described in Section 505(b)(2)(A)(iv) of the FDCA (21 U.S.C. § 355(b)(2)(A)(iv)), with respect to the '276, '278, and '027 patents asserting that all three patents are invalid, unenforceable, and/or will not be infringed by the commercial manufacture, use, offer for sale, or sale within the United States, or importation into the United States, of Curium's 505(b)(2) Product.

17. Upon information and belief, Curium Netherlands is in the business of the manufacture and global distribution of radioactive compounds as well as the research and development of radiopharmaceuticals for diagnostics and therapeutic use. Upon information and belief, Curium US LLC, Curium US Holdings LLC, and Curium Netherlands collaborated in the research and development of the drug product that is the subject of Curium Application No. 218525. Upon information and belief, Dr. Maarten de Jong and other scientists at Curium Netherlands have expertise in the formulation and manufacture of radiopharmaceuticals for diagnostics and therapeutic use. Upon information and belief, Dr. de Jong and other scientists at Curium Netherlands were directly involved in the design, research, and development of the formulation and the manufacturing process for producing the drug product that is the subject of Curium Application No. 218525.

18. Upon information and belief, Curium Netherlands recently opened a new Lutetium-177 production facility to supply Lutetium-177 to cancer patients worldwide. *See* <https://www.curiumpharma.com/2024/09/24/netherlands-lutetium-177/> (last accessed October 16, 2024) (Exhibit D). Upon information and belief, following any FDA approval of Curium Application No. 218525, Curium Netherlands will manufacture Lutetium-177 for the production of the drug product that is the subject of Curium Application No. 218525.

19. Upon information and belief, Curium US LLC, Curium US Holdings LLC, and Curium Netherlands acted collaboratively in the preparation and submission of Curium Application No. 218525.

20. Upon information and belief, following any FDA approval of Curium Application No. 218525, Curium US LLC, Curium US Holdings LLC, and Curium Netherlands will work in concert with one another to make, use, offer to sell, and/or sell the drug products that are the subject of Curium Application No. 218525 throughout the United States, and/or import such drug products into the United States, including in this judicial district.

JURISDICTION AND VENUE

21. This action arises under the patent laws of the United States, 35 U.S.C. §§ 100, *et seq.*, and this Court has jurisdiction over the subject matter of this action under 28 U.S.C. §§ 1331 and 1338(a). Venue is proper in this Court under 28 U.S.C. §§ 1391 and 1400(b).

22. Upon information and belief, Curium US LLC and Curium US Holdings LLC are each organized and existing under the laws of the State of Delaware and each have a principal place of business in St. Louis, Missouri in this judicial district.

23. Upon information and belief, Curium Netherlands BV and Curium International Trading BV are foreign corporations not residing in any United States judicial district and may be sued in any judicial district. 28 U.S.C. § 1391(c)(3).

24. Defendants have committed an act of infringement of the '276, '278, and '027 patents in this judicial district by filing Curium Application No. 218525 with the intent to make, use, offer to sell, sell, and/or import the drug products that are the subject of Curium Application No. 218525 in this judicial district, and/or will imminently commit an act of infringement by making, using, offering to sell, selling, and/or importing the same, acts of infringement that will lead to foreseeable harm and injury to ADACAP.

25. Upon information and belief, Defendants have extensive contacts with Missouri, regularly conduct business in Missouri, have purposefully availed themselves of the privilege of doing business in Missouri, and intend to sell in Missouri the drug product described in Curium Application No. 218525 upon approval.

26. For these reasons, and for other reasons that will be presented to the Court if jurisdiction is challenged, the Court has personal jurisdiction over Defendants.

THE PATENTS-IN-SUIT AND LUTATHERA®

27. On March 24, 2020, the U.S. Patent and Trademark Office duly and legally issued the '276 patent, entitled "Stable, Concentrated Radionuclide Complex Solutions." A true and correct copy of the '276 patent is attached hereto as Exhibit A.

28. The claims of the '276 patent are valid and enforceable. The '276 patent is wholly owned by Advanced Accelerator Applications SA, which therefore has the right to sue for and obtain equitable relief and damages for infringement of the '276 patent.

29. On March 24, 2020, the U.S. Patent and Trademark Office duly and legally issued the '278 patent, entitled "Stable, Concentrated Radionuclide Complex Solutions." A true and correct copy of the '278 patent is attached hereto as Exhibit B.

30. The claims of the '278 patent are valid and enforceable. The '278 patent is wholly owned by Advanced Accelerator Applications SA, which therefore has the right to sue for and obtain equitable relief and damages for infringement of the '278 patent.

31. On February 20, 2024, the U.S. Patent and Trademark Office duly and legally issued the '027 patent, entitled "Stable, Concentrated Radionuclide Complex Solutions." A true and correct copy of the '027 patent is attached hereto as Exhibit C.

32. The claims of the '027 patent are valid and enforceable. The '027 patent is wholly owned by Advanced Accelerator Applications SA, which therefore has the right to sue for and obtain equitable relief and damages for infringement of the '027 patent.

33. Advanced Accelerator Applications USA, Inc. is the holder of New Drug Application ("NDA") No. 208700 by which the FDA granted approval for the commercial manufacturing, marketing, sale, and use of LUTATHERA®. LUTATHERA® is a radiolabeled somatostatin analog indicated for the treatment of somatostatin receptor-positive gastroenteropancreatic neuroendocrine tumors (GEP-NETs), including foregut, midgut, and hindgut neuroendocrine tumors in adults.

34. LUTATHERA® and the use of LUTATHERA® is covered by one or more claims of each of the '276, '278, and '027 patents.

35. The FDA's official publication of approved drugs (the "Orange Book") lists the '276, '278, and '027 patents in connection with LUTATHERA®.

**COUNT I: INFRINGEMENT BY EACH DEFENDANT OF THE '276 PATENT UNDER
35 U.S.C. § 271(e)**

36. ADACAP incorporates each of the proceeding paragraphs 1–35 as if fully set forth herein.

37. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

38. Defendants' submission of Curium Application No. 218525 to obtain approval to engage in the commercial manufacture, use, offer to sell, or sale of Defendants'

505(b)(2) Product, prior to the expiration of the '276 patent constitutes infringement of one or more of the claims of the '276 patent under 35 U.S.C. § 271(e)(2)(A).

39. Upon information and belief, Defendants had actual and constructive knowledge of the '276 patent at least as of September 5, 2024, and have since been aware that filing of the aforementioned Curium Application No. 218525 with the FDA constituted an act of infringement of the '276 patent.

40. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product with its proposed labeling immediately and imminently upon approval of Curium Application No. 218525.

41. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '276 patent.

42. Upon information and belief, Defendants know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '276 patent.

43. Upon information and belief, Defendants plan and intend to, and will, actively induce infringement of the '276 patent when Curium Application No. 218525 is approved, and plan and intend to, and will, do so immediately and imminently upon approval.

44. Upon information and belief, Defendants know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '276 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use.

45. The foregoing acts by Defendants constitute and/or will constitute direct and indirect infringement of the '276 patent.

46. If Defendants' infringement of the '276 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

COUNT II: DECLARATORY JUDGMENT ON INFRINGEMENT BY EACH DEFENDANT OF THE '276 PATENT UNDER 35 U.S.C. §§ 271(a), (b), AND (c)

47. ADACAP incorporates each of the proceeding paragraphs 1–46 as if fully set forth herein.

48. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product immediately and imminently upon final approval of Curium Application No. 218525. Therefore, a case or controversy exists between each Defendant or group of Defendants and ADACAP as to infringement of the '276 patent.

49. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

50. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '276 patent.

51. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '276 patent.

52. Upon information and belief, Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will each have actual and constructive knowledge of the '276 patent, and each will actively induce the other to infringe the '276 patent when Curium Application No. 218525 is approved, and will each do so immediately and imminently upon final approval.

53. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '276 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will contribute to the infringement of the '276 patent immediately and imminently upon approval of Curium Application No. 218525.

54. The foregoing acts by Defendants constitute and/or will constitute direct infringement of the '276 patent, active inducement of infringement of the '276 patent, and/or contribution to the infringement by others of the '276 patent under 35 U.S.C. §§ 271(a), (b), and (c).

55. If Defendants' infringement of the '276 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

COUNT III: INFRINGEMENT BY EACH DEFENDANT OF THE '278 PATENT
UNDER 35 U.S.C. § 271(e)

56. ADACAP incorporates each of the proceeding paragraphs 1–55 as if fully set forth herein.

57. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same

active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

58. Defendants' submission of Curium Application No. 218525 to obtain approval to engage in the commercial manufacture, use, offer to sell, or sale of Defendants' 505(b)(2) Product, prior to the expiration of the '278 patent constitutes infringement of one or more of the claims of the '278 patent under 35 U.S.C. § 271(e)(2)(A).

59. Upon information and belief, Defendants had actual and constructive knowledge of the '278 patent at least as of September 5, 2024, and have since been aware that filing of the aforementioned Curium Application No. 218525 with the FDA constituted an act of infringement of the '278 patent.

60. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product with its proposed labeling immediately and imminently upon approval of Curium Application No. 218525.

61. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '278 patent.

62. Upon information and belief, Defendants know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '278 patent.

63. Upon information and belief, Defendants plan and intend to, and will, actively induce infringement of the '278 patent when Curium Application No. 218525 is approved, and plan and intend to, and will, do so immediately and imminently upon approval.

64. Upon information and belief, Defendants know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '278 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use.

65. The foregoing acts by Defendants constitute and/or will constitute direct and indirect infringement of the '278 patent.

66. If Defendants' infringement of the '278 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

COUNT IV: DECLARATORY JUDGMENT ON INFRINGEMENT BY EACH DEFENDANT OF THE '278 PATENT UNDER 35 U.S.C. §§ 271(a), (b), AND (c)

67. ADACAP incorporates each of the proceeding paragraphs 1–66 as if fully set forth herein.

68. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product immediately and imminently upon final approval of Curium Application No. 218525. Therefore, a case or controversy exists between each Defendant or group of Defendants and ADACAP as to infringement of the '278 patent.

69. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

70. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '278 patent.

71. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '278 patent.

72. Upon information and belief, Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will each have actual and constructive knowledge of the '276 patent, and each will actively induce the other to infringe the '278 patent when Curium Application No. 218525 is approved, and will each do so immediately and imminently upon final approval.

73. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '278 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will contribute to the infringement of the '278 patent immediately and imminently upon approval of Curium Application No. 218525.

74. The foregoing acts by Defendants constitute and/or will constitute direct infringement of the '278 patent, active inducement of infringement of the '276 patent, and/or contribution to the infringement by others of the '278 patent under 35 U.S.C. §§ 271(a), (b), and (c).

75. If Defendants' infringement of the '278 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

**COUNT V: INFRINGEMENT BY EACH DEFENDANT OF THE '027 PATENT UNDER
35 U.S.C. § 271(e)**

76. ADACAP incorporates each of the proceeding paragraphs 1–75 as if fully set forth herein.

77. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

78. Defendants' submission of Curium Application No. 218525 to obtain approval to engage in the commercial manufacture, use, offer to sell, or sale of Defendants' 505(b)(2) Product, prior to the expiration of the '027 patent constitutes infringement of one or more of the claims of the '027 patent under 35 U.S.C. § 271(e)(2)(A).

79. Upon information and belief, Defendants had actual and constructive knowledge of the '027 patent at least as of September 5, 2024, and have since been aware that filing of the aforementioned Curium Application No. 218525 with the FDA constituted an act of infringement of the '027 patent.

80. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product with its proposed labeling immediately and imminently upon approval of Curium Application No. 218525.

81. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '027 patent.

82. Upon information and belief, Defendants know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '027 patent.

83. Upon information and belief, Defendants plan and intend to, and will, actively induce infringement of the '027 patent when Curium Application No. 218525 is approved, and plan and intend to, and will, do so immediately and imminently upon approval.

84. Upon information and belief, Defendants know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '027 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use.

85. The foregoing acts by Defendants constitute and/or will constitute direct and indirect infringement of the '027 patent.

86. If Defendants' infringement of the '027 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

COUNT VI: DECLARATORY JUDGMENT ON INFRINGEMENT BY EACH DEFENDANT OF THE '027 PATENT UNDER 35 U.S.C. §§ 271(a), (b), AND (c)

87. ADACAP incorporates each of the proceeding paragraphs 1–86 as if fully set forth herein.

88. Upon information and belief, Defendants intend to engage in the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product immediately and imminently upon final approval of Curium Application No. 218525. Therefore, a case or controversy exists between each Defendant or group of Defendants and ADACAP as to infringement of the '027 patent.

89. By filing Curium Application No. 218525, Defendants have necessarily represented to the FDA that, upon approval, Defendants' 505(b)(2) Product will have the same

active ingredient, method of administration, dosage form, and strength as LUTATHERA®, and will be bioequivalent to LUTATHERA®.

90. The commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '027 patent.

91. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that the commercial manufacture, use, offer for sale, sale, marketing, distributing, and/or importation of Defendants' 505(b)(2) Product would infringe one or more claims of the '027 patent.

92. Upon information and belief, Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will each have actual and constructive knowledge of the '027 patent, and each will actively induce the other to infringe the '027 patent when Curium Application No. 218525 is approved, and will each do so immediately and imminently upon final approval.

93. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will know that Defendants' 505(b)(2) Product is especially made or adapted for use in infringing the '027 patent, and that Defendants' 505(b)(2) Product is not suitable for any substantial non-infringing use. Upon information and belief, each of Curium US LLC, Curium US Holdings LLC, Curium Netherlands BV, and Curium International Trading BV will contribute to the infringement of the '027 patent immediately and imminently upon approval of Curium Application No. 218525.

94. The foregoing acts by Defendants constitute and/or will constitute direct infringement of the '027 patent, active inducement of infringement of the '027 patent, and/or

contribution to the infringement by others of the '027 patent under 35 U.S.C. §§ 271(a), (b), and (c).

95. If Defendants' infringement of the '027 patent is not permanently enjoined, ADACAP will suffer substantial and irreparable harm for which there is no remedy at law.

PRAYER FOR RELIEF

WHEREFORE, ADACAP prays that this Court grant the following relief:

1. A declaration and judgment that Defendants' imminent making, using, offering to sell, or selling in the United States, or importing into the United States, or inducing or contributing to the same, of Defendants' 505(b)(2) Product will infringe the '276, '278, and '027 patents.

2. A judgment that one or more claims of the '276, '278, and '027 patents is not invalid, is enforceable, and is infringed by Defendants' submission of Curium Application No. 218525, and that Defendants' making, using, offering to sell, or selling in the United States, or importing into the United States of Defendants' 505(b)(2) Product, will infringe the '276, '278, and '027 patents.

3. An order pursuant to 35 U.S.C. § 271(e)(4)(A) providing that the effective date of any approval of Curium Application No. 218525 shall be a date which is not earlier than the expiration of the '276, '278, and '027 patents, including any extensions and/or additional periods of exclusivity to which ADACAP is or becomes entitled.

4. An order permanently enjoining Defendants, their affiliates, subsidiaries, and each of their officers, agents, servants, and employees and those acting in privity or concert with them, from making, using, offering to sell, or selling in the United States, or importing into the United States Defendants' 505(b)(2) Product, until after the expiration of the '276, '278, and

'027 patents, including any extensions and/or additional periods of exclusivity to which ADACAP is or becomes entitled.

5. Damages or other monetary relief to ADACAP if Defendants engage in commercial manufacture, use, offers to sell, sale, or importation in or into the United States of Defendants' 505(b)(2) Product, prior to the expiration date(s) of the '276, '278, and '027 patents, including any extensions and/or additional periods of exclusivity to which ADACAP is or becomes entitled.

6. Such further and other relief as this Court deems proper and just, including any appropriate relief under 35 U.S.C. § 285.

DATED: October 18, 2024

Respectfully submitted,

/s/ James F. Bennett

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