Claims

1. A kynureninase homodimer covalently bound to one or more polyethylene glycol (PEG) molecules, wherein the ratio of PEG molecules to homodimer is from 15:1 to 35:1

2. The homodimer of claim 1, wherein the ratio of PEG molecules to homodimer is from 20:1 to 30:1.

3. The homodimer of claim 1, wherein the ratio of PEG molecules to homodimer is about 25:1.

4. The homodimer of any one of claims 1-3, wherein the PEG molecule has a molecular weight from 1 kDa to 10 kDa.

5. The homodimer of any one of claims 1-4, wherein the homodimer is covalently bound to the PEG molecule by way of one or more lysine or cysteine residues.

6. The homodimer of any one of claims 1-5, wherein the homodimer comprises two polypeptide monomers each having an amino acid sequence that is at least 90% identical to SEQ ID NO: 1, and wherein each polypeptide monomer has one or more amino acid substitutions, relative to SEQ ID NO: 1, selected from the group consisting of L8P, K38E, Y47L, I48F, K50Q, I51M, S60N, K64N, D65G, E66K, N67D, N67P, D67S, A68F, A68T, A68V, F71L, F71M, L72N, K84E, E88N, E89K, E89S, E90Q, D92E, K93N, K93T, A95H, A95Q, K96N, I97H, I97L, I97V, A98G, A99G, A99I, A99R, A99S, A99T, A99V, Y100N, Y100S, Y100T, G101A, H102W, E103F, E103H, E103N, E103Q, E103R, E103V, E103W, V104D, V104E, V104F, V104H, V104K, V104L, V104R, G105A, G105H, G105S, G105T, E106D, K106D, K106E, K106H, K106N, R107P, R107S, P108R, I110A, I110F, I110L, I110M, I110T, T111D, T111H, T111N, T111R, G112A, G112C, G112D, G112K, G112L, G112M, G112Q, G112R, G112S, G112T, G112Y, N127K, I131V, A132V, L133V, A136T, L137T, T138S, N140D, H142Q, Q14R, Y156H, K163T, D168E, H169R, Q175L, I183F, I183L, I183P, I183S, E184A, E184D, E184R, E184T, E184V, E185T, M187L, M189I, K191A, K191G, K191H, K191M, K191N, K191R, K191S, K191T, K191W, E197A, E197D, E197F, E197K, E197M, E197Q, E197S, E197T, E197V, I201C, I201E, I201F, 1201H, I201L, I201S, I201T, I201V, H203K, L219M, L219W, F220L, V223I, F225Y, H230F, H230L, H230Y, N232S, Y246F, F249W, D250E, S274A, S274C, S274G, S274N, S274T, L278M, A280G, A280S, A280T, G281S, A282M, A282P, G284N, I285L, V303L, V303S, F306W, F306Y, S311N, K315E, D317E, D317K, I331C, I331L, I331N, I331S, I331T, I331V, N333T, P334N, P335T, L337T, L338A, L338Q, S341I, K373E, K373N, N375A, N375H, Y376C, Y376F, Y376L, K378G, K378P, K378Q, K378R, K380G, K380S, A382G, A382R, A382T, T383S, K384G, K384N, P386K, P386S, V387L, N389E, I405L, F407Y, S408D, S408N, N411R, D413S, D413V, Q416T, E419A, E419L, K420E, R421N, V424I, K427M, N429E, G432A, and A436T.

7. The homodimer of claim 6, wherein each polypeptide monomer comprises a substitution at A282, F306, F249, A99, G112, E103, V104, and/or S408 relative to SEQ ID NO: 1.

8. The homodimer of claim 6 or 7, wherein each polypeptide monomer comprises (i) the F306W substitution, (ii) the L72N substitution, (iii) the H102W and N333T substitutions, (iv) the I183P substitution, (v) the R107P substitution, and/or (vi) the A436T substitution.

9. The homodimer of any one of claims 6-8, wherein each polypeptide monomer comprises the substitutions L72N, H102W, A282P, F306W, I331S and N333T relative to SEQ ID NO: 1.

10. The homodimer of any one of claims 1-9, wherein the homodimer comprises two polypeptide monomers each having an amino acid sequence that is at least 90% identical to SEQ ID NO: 3.

11. The homodimer of claim 10, wherein each polypeptide monomer has the amino acid sequence of SEQ ID NO: 3.

12. The homodimer of any one of claims 1-9, wherein the homodimer comprises two polypeptide monomers each having an amino acid sequence that is at least 90% identical to SEQ ID NO: 2.

13. The homodimer of claim 12, wherein each polypeptide monomer has the amino acid sequence of SEQ ID NO: 2.

14. The homodimer of any of claims 1-13, wherein the homodimer has a catalytic activity ($kcat/K_M$) of at least $8000M^{-1}\ s^{-1}$.

15. The homodimer of any one of claims 1-14, wherein the ratio of PEG molecules to homodimer is about 25:1, the PEG molecule has a molecular weight of about 5 kDa and is bound to the homodimer by way of an N-hydroxysuccinimide ester carbonate linking group.

16. A method of producing a PEGylated kynureninase homodimer, wherein the homodimer comprises two polypeptide monomers, the method comprising contacting the homodimer with a PEGylation agent, and wherein the molar input ratio of PEGylation agent to homodimer is from 15:1 to 35:1.

17. The method of claim 16, wherein the molar input ratio of PEGylation agent to homodimer is from 20:1 to 35:1.

18. The method of claim 16, wherein the molar input ratio of PEGylation agent to homodimer is from 25:1 to 35:1.

19. The method of claim 18, wherein the molar input ratio of PEGylation agent to homodimer is about 30:1.

20. The method of any one of claims 16-19, wherein the PEG molecule has a molecular weight from 1 kDa to 10 kDa.

21. The method of any one of claims 16-20, wherein the homodimer is covalently bound to the PEG molecule by way of one or more lysine or cysteine residues.

22. The method of any one of claims 16-21, wherein each polypeptide monomer has an amino acid sequence that is at least 90% identical to SEQ ID NO: 1, and wherein each polypeptide monomer has one or more amino acid substitutions, relative to SEQ ID NO: 1, selected from the group consisting of L8P, K38E, Y47L, I48F, K50Q, I51M, S60N, K64N, D65G, E66K, N67D, N67P, D67S, A68F, A68T, A68V, F71L, F71M, L72N, K84E, E88N, E89K, E89S, E90Q, D92E, K93N, K93T, A95H, A95Q,

K96N, I97H, I97L, I97V, A98G, A99G, A99I, A99R, A99S, A99T, A99V, Y100N, Y100S, Y100T, G101A, H102W, E103F, E103H, E103N, E103Q, E103R, E103V, E103W, V104D, V104E, V104F, V104H, V104K, V104L, V104R, G105A, G105H, G105S, G105T, E106D, K106D, K106E, K106H, K106N, R107P, R107S, P108R, I110A, I110F, I110L, I110M, I110T, T111D, T111H, T111N, T111R, G112A, G112C, G112D, G112K, G112L, G112M, G112Q, G112R, G112S, G112T, G112Y, N127K, I131V, A132V, L133V, A136T, L137T, T138S, N140D, H142Q, Q14R, Y156H, K163T, D168E, H169R, Q175L, I183F, I183L, I183P, I183S, E184A, E184D, E184R, E184T, E184V, E185T, M187L, M189I, K191A, K191G, K191H, K191M, K191N, K191R, K191S, K191T, K191W, E197A, E197D, E197F, E197K, E197M, E197Q, E197S, E197T, E197V, I201C, I201E, I201F, 1201H, I201L, I201S, I201T, I201V, H203K, L219M, L219W, F220L, V223I, F225Y, H230F, H230L, H230Y, N232S, Y246F, F249W, D250E, S274A, S274C, S274G, S274N, S274T, L278M, A280G, A280S, A280T, G281S, A282M, A282P, G284N, I285L, V303L, V303S, F306W, F306Y, S311N, K315E, D317E, D317K, I331C, I331L, I331N, I331S, I331T, I331V, N333T, P334N, P335T, L337T, L338A, L338Q, S341I, K373E, K373N, N375A, N375H, Y376C, Y376F, Y376L, K378G, K378P, K378Q, K378R, K380G, K380S, A382G, A382R, A382T, T383S, K384G, K384N, P386K, P386S, V387L, N389E, I405L, F407Y, S408D, S408N, N411R, D413S, D413V, Q416T, E419A, E419L, K420E, R421N, V424I, K427M, N429E, G432A, and A436T.

23. The method of any one of claims 16-22, wherein each polypeptide monomer comprises a substitution at A282, F306, F249, A99, G112, E103, V104, and/or S408 relative to SEQ ID NO: 1.

24. The method of claim 22 or 23, wherein each polypeptide monomer comprises the (i) F306W substitution, (ii) the L72N substitution, (iii) the H102W and N333T substitutions, (iv) the I183P substitution, (v) the R107P substitution, and/or (vi) the A436T substitution.

25. The method of any one of claims 22-24, wherein each polypeptide monomer comprises the substitutions L72N, H102W, A282P, F306W, I331S and N333T relative to SEQ ID NO: 1.

26. The method of any one of claims 16-25, wherein each polypeptide monomer has an amino acid sequence that is at least 90% identical to SEQ ID NO: 3.

27. The method of claim 26, wherein each polypeptide monomer has the amino acid sequence of SEQ ID NO: 3.

28. The method of any one of claims 16-25, wherein each polypeptide monomer has an amino acid sequence that is at least 90% identical to SEQ ID NO: 2.

29. The method of claim 28, wherein each polypeptide monomer has the amino acid sequence of SEQ ID NO: 2.

30. The method of any of claims 16-29, wherein the homodimer has a catalytic activity (k_{cat}/K_M) of at least $8000 M^{-1} s^{-1}$.

31. The method of any one of claims 16-30, wherein the ratio of PEG molecules to homodimer is about 25:1, the PEG molecule has a molecular weight of about 5 kDa and is bound to the homodimer by way of an N-hydroxysuccinimide ester carbonate linking group.

32. The method of any of claims 16-31, wherein the homodimer is contacted with the PEGylation agent in an aqueous buffer.

33. The method of claim 32, wherein the aqueous buffer comprises disodium phosphate and/or sodium acetate.

34. The method of claim 32 or 33, wherein the aqueous buffer ranges in pH from 7.5 to 9.5.

35. The method of any one claims 32-34, wherein the aqueous buffer ranges in ionic strength from 250 mM to 350 mM.

36. The method of any one of claims 16-35, wherein size exclusion chromatography is used to isolate the homodimer.

37. A pharmaceutical formulation comprising the kynureninase homodimer of any one of claims 1-15 in a pharmaceutically acceptable carrier.

38. The formulation of claim 38 or the kynureninase homodimer of any one of claims 1-15 for use in treating a subject having an IDO1, IDO2, or TDO expressing tumor.

39. The formulation or kynureninase homodimer for use in accordance with claim 38, wherein the tumor is a solid tumor or a hematological tumor.

40. The formulation or kynureninase homodimer for use in accordance with claim 38 or 39, wherein the subject is a human patient.

41. The formulation or kynureninase homodimer for use in accordance with any one of claims 38-40, wherein the formulation is to be administered intratumorally, intravenously, intradermally, intraarterially, intraperitoneally, intralesionally, intracranially, intraarticularly, intraprostaticaly, intrapleurally, intratracheally, intraocularly, intranasally, intravitreally, intravaginally, intrarectally, intramuscularly, subcutaneously, subconjunctival, intravesicularlly, mucosally, intrapericardially, intraumbilically, orally, by inhalation, by injection, by infusion, by continuous infusion, by localized perfusion bathing target cells directly, via a catheter, or via a lavage.

42. The formulation or kynureninase homodimer for use in accordance with any one of claims 38-41, wherein the formulation or kynureninase homodimer is to be administered to the subject with a second anti-cancer therapy.

43. The formulation or kynureninase homodimer for use in accordance with claim 42, wherein the second anticancer therapy is radiation therapy, surgical therapy, an immunotherapy or a second anticancer compound.

44. The formulation or kynureninase homodimer for use in accordance with claim 43, wherein the second anticancer compound is an immune checkpoint inhibitor.

45. The formulation or kynureninase homodimer for use in accordance with claim 43, wherein the second anticancer compound comprises an anti- PDI, anti-CTLA-4, or anti-PD-LI antibody.

46. The formulation of claim 38 or the kynureninase homodimer of any one of claims 1-15 for use in a method of providing a T-cell response in a human subject having an IDO1, IDO2, or TDO expressing tumor.