WHAT IS CLAIMED IS:

1. A method for preparing an enriched and expanded cell population of tumour reactive T-cells for use in cancer therapy comprising identifying and/or obtaining a cell population expressing CD150/SLAM/SLAMF1 and expanding the cell population.
2. The method according to claim 1, wherein the T-cells expressing CD150/SLAM/SLAMF1 are T-cells selected from cells originating from a subject.
3. The method according to claim 1 or 2, wherein selecting the T-cells expressing CD150/SLAM/SLAMF1 comprises one or more of (i) flow cytometry, (ii) antibody panning, (iii) magnetic selection, (iv) biomarker targeted cell enrichment.
4. The method according to claim 1 or 2 wherein selecting the T-cells comprises contacting the cell population with an anti-CD150/SLAM/SLAMF1 antibody.
5. The method according to any one of claims 1-4, wherein the expanding comprises:
    a. irradiating feeder cells and co-stimulating with an anti-CD150/SLAM/SLAMF1 antibody and optionally one or more cytokines; or
    b. stimulating or activating CD150/SLAM/SLAMF1 or
    c. rapid expansion protocol (REP) wherein cells are mixed with irradiated feeder cells and one or more cytokines until at least $1x10^9$ or at least $5x10^9$ or at least $1x10^{10}$ cells are obtained.
6. The method according to any one of claims 1-5 wherein the cell population is:
    a. a population of tumour infiltrating lymphocytes (TILs) from a tumour biopsy, lymphnode or ascites; and/or
    b. a population of T-cells engineered to express a CAR and/or a TCR
    c. a population of T-cells isolated from blood.
7. The method according to any one of claims 1-6 wherein the cancer is a melanoma, lung cancer, squamous cell cancer, cancer of the peritoneum, hepatocellular cancer, gastric or stomach cancer including gastrointestinal cancer, pancreatic cancer, glioblastoma, cervical cancer, ovarian cancer, liver cancer, bladder cancer, hepatoma, breast cancer, colon cancer, rectal cancer, colorectal cancer, endometrial cancer or uterine carcinoma, salivary gland carcinoma, kidney or renal cancer, prostate cancer, vulval cancer, thyroid cancer, hepatic carcinoma, anal carcinoma, penile carcinoma, head cancer or neck cancer.

8. The method according to any one of claims 1-7 wherein the cancer therapy is adoptive cell therapy.
9. The method of any one of claims 1-8, wherein the method further comprises exposing the T-cells to a modulator that increases SLAM production.
10. The method of claim 9, wherein the modulator comprises a cytokine.
11. The method of claim 10, wherein the cytokine is IL-2, IL-15, IL-18, IL-12, IL-23, IL-27, IL-35, IL-39, IL-18, IL-36, IL-37, IL-38, IFN-alpha, IFN-beta, IFN-gamma or a combination thereof.
12. The method of claim 9 or 10, wherein the modulator is a cytokine combination of IL-7+ IL-15, IL-2 + IL-7 + IL-15, IL-2 + IL-12, IL-2 + IL-18, IL-2 + IL-12 + IL-7 + IL-15, IL-2 + IL-12 + IL-7 + IL-15 + IL-6, IL-2 + IL-12 + IL-7 + IL-15 + IL-21, IL-2 + IL-12 + IL-7 + IL-15 + IL-6 + IL-21, IL-7 + IL-15 + IL-6, IL-7 + IL-15 + IL-21, IL-7 + IL-15 + IL-6 + IL-21, IL-2 + IL-12 + IL-6, IL-2 + IL-12 + IL-21, or IL-2 + IL-12 + IL-6 + IL-21.
13. The method of any one of claims 9-12 wherein the modulator further comprises a Th2 blocking agent.
14. The method of claim 13, wherein the Th2 blocking agent is an antibody.
15. The method of claim 14 wherein the antibody is selected from anti-IL-4 (αIL4), anti IL-4R (αIL4R), anti IL-5R (αIL5R), anti IL-5 (αIL5), anti-IL13R (αIL13R) or anti-IL13 (αIL13).
16. The method of claim 15 wherein the antibody is selected from Mepolizumab, Resilizumab, Benralizumab, Tralokinumab, Lebrikizumab or Dupilumab.
17. The method of claim 13 or 14, wherein the modulator comprises IL-2 + αIL4.
18. The method of claim 13 or 14, wherein the modulator comprises IL-2 + IL-12 + αIL4.
19. The method of claim 13 or 14, wherein the modulator comprises IL-2+ IL-12 + IL-7 + IL-15 + αIL4.
20. The method of any one of claims 11-19, wherein IL-12 is briefly added for 1-96 h at 1-150 ng/ml added at the expanding step and subsequently replaced with another cytokine.
21. The method according to any of claims 10-20 wherein the cytokine is produced by an artificial antigen presenting cell and the tumor reactive T cells are co-cultured with the antigen presenting cell.

22. The method of any one of claims 1-21, wherein the cell population is isolated, cultured with a modulator that increases CD150/SLAM/SLAMF1 production and wherein the cultured cells are separated from unselected cells.
23. The method of any of claims 1-22 wherein the T cells are engineered to express a cytokine receptor which provides cytokine signaling upon engagement of another drug or cytokine.
24. The method of any one of claims 1-23, wherein the T cell is a CD4+ or a CD8+ T cell.
25. A population of cells obtained according to the method of any preceding claim.
26. The population of cells of claim 25 for use in the treatment of cancer.
27. An enriched and expanded cell population of tumour reactive T-cells expressing CD150/SLAM/SLAMF1 for use in the treatment of cancer.
28. The population of cells according to any of claims 25-27, wherein the population comprises T-cells and wherein >25%, >30% or >40% of the T-cells express CD150/SLAM/SLAMF1.
29. The population of cells according to any one of claims 25-28, wherein the cells are TILs.
30. A method of adoptive cell therapy, comprising administering the population of cells according to any of claims 25-29, to a subject, wherein the T-cells are autologous or allogenic.
31. A method for treating a cancer in a subject, which comprises the step of administering a cell population according to any of claims 25-29, to the subject.
32. A pharmaceutical composition comprising a population of cells according to any of claims 25-29.
33. The pharmaceutical composition according to claim 32 for use in the treatment of cancer.
34. A method for assessing the tumour reactivity of a cell population which method comprises quantifying the proportion of T-cells expressing SLAM/SLAMF1/CD150 in the cell population.
35. A method according to claim 34 wherein the cell population is i) TILs from a patient and the T-cells expressing SLAM/SLAMF1/CD150 are quantified pre-REP and/or post-REP; or ii) a population of T-cells engineered to express an exogenous CAR and/or a TCR.

36. A method according to claim 34 or 35 wherein the proportion of T-cells expressing SLAM/SLAMF1/CD150 being at least 25% of the cell population indicates that the cell population is tumour reactive.
37. A method for identifying an agent for increasing SLAM/SLAMF1/CD150 expression in isolated ex vivo expansion of T cells for use in adoptive cell therapy, comprising
    a. contacting tumor-infiltrating lymphocytes with a candidate modulator that upregulates the expression of SLAM in T cells.
    b. screening the effect of the modulator on the expression of SLAM in the cells; and
    c. identifying a modulator that increases the expression of SLAM in the cells.
38. The method of claim 37, wherein the T cell is a CD4+ or a CD8+ T cell.
39. An in vitro method for the prognosis of patients who may receive Tumour Infiltrating Lymphocyte (TIL) therapy for cancer comprising:
    a. quantifying in a sample of tumour digest, TIL material during the manufacturing process or TIL product, from said patient, at least one biological marker on the TIL; and
    b. comparing the value obtained at step a) for said at least one biological marker with a predetermined reference value for the same biological marker; which predetermined reference value is correlated with a specific prognosis of progression of said cancer.
40. The method of claim 39, wherein the at least one biological marker is SLAM/SLAMF1/CD150.
41. A cancer therapy method comprising preparing an enriched and expanded cell population of tumour reactive T-cells comprising identifying and/or obtaining a cell population expressing CD150/SLAM/SLAMF1, expanding the cell population, and administering to a cancer patient in need thereof cells from the cell population.
42. The method according to claim 41, wherein the T-cells expressing CD150/SLAM/SLAMF1 are T-cells selected from cells originating from a subject.
43. The method according to claim 41 or 42, wherein selecting the T-cells expressing CD150/SLAM/SLAMF1 comprises one or more of (i) flow cytometry, (ii) antibody panning, (iii) magnetic selection, (iv) biomarker targeted cell enrichment.

44. The method according to claim 41 or 42 wherein selecting the T-cells comprises contacting the cell population with an anti-CD150/SLAM/SLAMF1 antibody.
45. The method according to any one of claims 41-44, wherein the expanding comprises:
    a. irradiating feeder cells and co-stimulating with an anti- CD150/SLAM/SLAMF1 antibody and optionally one or more cytokines; or
    b. stimulating or activating CD150/SLAM/SLAMF1 or
    c. rapid expansion protocol (REP) wherein cells are mixed with irradiated feeder cells and one or more cytokines until at least $1x10^9$ or at least $5x10^9$ or at least $1x10^{10}$ cells are obtained.
46. The method according to any one of claims 41-45 wherein the cell population is:
    a. a population of tumour infiltrating lymphocytes (TILs) from a tumour biopsy, lymphnode or ascites; and/or
    b. a population of T-cells engineered to express a CAR and/or a TCR
    c. a population of T-cells isolated from blood.
47. The method according to any one of claims 41-46 wherein the cancer is a melanoma, lung cancer or ovarian cancer.
48. The method according to any one of claims 41-47 wherein the cancer therapy is adoptive cell therapy.
49. The method of any one of claims 41-48, wherein the method further comprises exposing the T-cells to a modulator that increases SLAM production.
50. The method of claim 49, wherein the modulator comprises a cytokine.
51. The method of claim 50, wherein the cytokine is IL-2, IL-15, IL-18, IL-12, IL-23, IL-27, IL-35, IL-39, IL-18, IL-36, IL-37, IL-38, IFN-alpha, IFN-beta, IFN-gamma or a combination thereof.
52. The method of claim 50 or 51, wherein the modulator is a cytokine combination of IL-7+ IL-15, IL-2 + IL-7 + IL-15, IL-2 + IL-12, IL-2 + IL-18, IL-2 + IL-12 + IL-7 + IL-15, IL-2 + IL-12 + IL-7 + IL-15 + IL-6, IL-2 + IL-12 + IL-7 + IL-15 + IL-21, IL-2 + IL-12 + IL-7 + IL-15 + IL-6 + IL-21, IL-7 + IL-15 + IL-6, IL-7 + IL-15 + IL-21, IL-7 + IL-15 + IL-6 + IL-21, IL-2 + IL-12 + IL-6, IL-2 + IL-12 + IL-21, or IL-2 + IL-12 + IL-6 + IL-21.
53. The method of any one of claims 49-52 wherein the modulator further comprises a Th2 blocking agent.

54. The method of claim 53, wherein the Th2 blocking agent is an antibody.
55. The method of claim 54 wherein the antibody is selected from anti-IL-4 (αIL4), anti IL-4R (αIL4R), anti IL-5R (αIL5R), anti IL-5 (αIL5), anti-IL13R (αIL13R) or anti-IL13 (αIL13).
56. The method of claim 55 wherein the antibody is selected from Mepolizumab, Resilizumab, Benralizumab, Tralokinumab, Lebrikizumab or Dupilumab.
57. The method of claim 54 or 55, wherein the modulator comprises IL-2 + αIL4.
58. The method of claim 54 or 55, wherein the modulator comprises IL-2 + IL-12 + αIL4.
59. The method of claim 54 or 55, wherein the modulator comprises IL-2+ IL-12 + IL-7 + IL-15 + αIL4.
60. The method of any one of claims 51-59, wherein IL-12 is briefly added for 1-96 h at 1-150 ng/ml at the expanding step and subsequently replaced with another cytokine.
61. The method according to any of claims 49-60 wherein the modulator is produced by an artificial antigen presenting cell and the tumor reactive T cells are co-cultured with the antigen presenting cell.
62. The method of any one of claims 41-61, wherein the cell population is isolated, cultured with a modulator that increases CD150/SLAM/SLAMF1 production and separating the cultured cells from unselected cells.
63. The method of any of claims 41-62 wherein the T cells are engineered to express a cytokine receptor which provides cytokine signaling upon engagement of another drug or cytokine.
64. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer, comprising administering to a cancer patient in need thereof cells from a cell population, wherein the cell population is prepared by a method comprising identifying and/or obtaining a cell population expressing CD150/SLAM/SLAMF1 and expanding the cell population.
65. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to claim 64, wherein the T-cells expressing CD150/SLAM/SLAMF1 are T-cells selected from cells originating from a subject.
66. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to claim 64 or 65, wherein selecting the T-cells expressing

CD150/SLAM/SLAMF1 comprises one or more of (i) flow cytometry, (ii) antibody panning, (iii) magnetic selection, (iv) biomarker targeted cell enrichment.

67. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to claim 64 or 65 wherein selecting the T-cells comprises contacting the cell population with an anti-CD150/SLAM/SLAMF1 antibody.
68. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to any one of claims 64-67, wherein the expanding comprises:
    a. irradiating feeder cells and co-stimulating with an anti- CD150/SLAM/SLAMF1 antibody and optionally one or more cytokines; or
    b. stimulating or activating CD150/SLAM/SLAMF1 or
    c. rapid expansion protocol (REP) wherein cells are mixed with irradiated feeder cells and one or more cytokines until at least $1x10^9$ or at least $5x10^9$ or at least $1x10^{10}$ cells are obtained.
69. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to any one of claims 64-68 wherein the cell population is:
    a. a population of tumour infiltrating lymphocytes (TILs) from a tumour biopsy, lymphnode or ascites; and/or
    b. a population of T-cells engineered to express a CAR and/or a TCR
    c. a population of T-cells isolated from blood.
70. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to any one of claims 64-69 wherein the cancer is a melanoma, lung cancer, squamous cell cancer, cancer of the peritoneum, hepatocellular cancer, gastric or stomach cancer including gastrointestinal cancer, pancreatic cancer, glioblastoma, cervical cancer, ovarian cancer, liver cancer, bladder cancer, hepatoma, breast cancer, colon cancer, rectal cancer, colorectal cancer, endometrial cancer or uterine carcinoma, salivary gland carcinoma, kidney or renal cancer, prostate cancer, vulval cancer, thyroid cancer, hepatic carcinoma, anal carcinoma, penile carcinoma, head cancer or neck cancer.

71. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer according to any one of claims 64-70 wherein the cancer treatment is adoptive cell therapy.
72. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of any one of claims 64-71, wherein the method further comprises exposing the T-cells to a modulator that increases SLAM production.
73. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 72, wherein the modulator comprises a cytokine.
74. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 73, wherein the cytokine is IL-2, IL-15, IL-18, IL-12, IL-23, IL-27, IL-35, IL-39, IL-18, IL-36, IL-37, IL-38, IFN-alpha, IFN-beta, IFN-gamma or a combination thereof.
75. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 73 or 74, wherein the modulator is a cytokine combination of IL-7+ IL-15, IL-2 + IL-7 + IL-15, IL-2 + IL-12, IL-2 + IL-18, IL-2 + IL-12 + IL-7 + IL-15, IL-2 + IL-12 + IL-7 + IL-15 + IL-6, IL-2 + IL-12 + IL-7 + IL-15 + IL-21, IL-2 + IL-12 + IL-7 + IL-15 + IL-6 + IL-21, IL-7 + IL-15 + IL-6, IL-7 + IL-15 + IL-21, IL-7 + IL-15 + IL-6 + IL-21, IL-2 + IL-12 + IL-6, IL-2 + IL-12 + IL-21, or IL-2 + IL-12 + IL-6 + IL-21.
76. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of any one of claims 72-75 wherein the modulator further comprises a Th2 blocking agent.
77. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 76, wherein the Th2 blocking agent is an antibody.
78. The method of claim 77 wherein the antibody is selected from anti-IL-4 (αIL4), anti IL-4R (αIL4R), anti IL-5R (αIL5R), anti IL-5 (αIL5), anti-IL13R (αIL13R) or anti-IL13 (αIL13).
79. The method of claim 78 wherein the antibody is selected from Mepolizumab, Resilizumab, Benralizumab, Tralokinumab, Lebrikizumab or Dupilumab.
80. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 77 or 78, wherein the modulator comprises IL-2 + αIL4.

81. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 77 or 78, wherein the modulator comprises IL-2 + IL-12 + αIL4.
82. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of claim 77 or 78, wherein the modulator comprises IL-2+ IL-12 + IL-7 + IL-15 + αIL4.
83. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of any one of claims 74-82, wherein IL-12 is briefly added for 1-96 h at 1-150 ng/ml at the expanding step and subsequently replaced with another cytokine.
84. An enriched and expanded cell population according to any of claims 64-83 wherein the modulator is produced by an artificial antigen presenting cell and the tumor reactive T cells are co-cultured with the antigen presenting cell.
85. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of any one of claims 64-83, wherein the cell population is isolated, cultured with a modulator that increases CD150/SLAM/SLAMF1 production and wherein the cultured cells are separated from unselected cells.
86. An enriched and expanded cell population of any of claims 64-85 wherein the T cells are engineered to express a cytokine receptor which provides cytokine signaling upon engagement of another drug or cytokine
87. An enriched and expanded cell population of tumour reactive T-cells for use in the treatment of cancer of any one of claims 64-87, wherein the T cell is a CD4+ or a CD8+ T cell.
88. The use of a cytokine in increasing expression of CD150/SLAM/SLAMF1 in a method for preparing an enriched and expanded cell population of tumour reactive T-cells for use in cancer therapy comprising exposing the T cells to a cytokine.
89. The use of a cytokine of claim 88 wherein the cytokine is used in combination with a Th2 blocking agent.
90. A method for increasing expression of CD150/SLAM/SLAMF1 in tumour reactive T-cells for use in cancer therapy comprising identifying and/or obtaining a cell population expressing CD150/SLAM/SLAMF1 and expanding the cell population in the presence of a cytokine that increases expression of CD150/SLAM/SLAMF1.

91. The method of claim 90 wherein the cytokine is used in combination with a Th2 blocking agent.