Supplementary methods

- 1. This T cell-inflamed score model is shown as follows: T cell-inflamed score = $0.008346 \times EXP_{CCL5} + 0.072293 \times EXP_{CD27} + 0.042853 \times EXP_{CD274} 0.0239 \times EXP_{CD276} + 0.031021 \times EXP_{CD8A} + 0.151253 \times EXP_{CMKLR1} + 0.074135 \times EXP_{CXCL9} + 0.004313 \times EXP_{CXCR6} + 0.020091 \times EXP_{HLA-DQA1} + 0.058806 \times EXP_{HLA-DRB1} + 0.07175 \times EXP_{HLA-E} + 0.060679 \times EXP_{ID01} + 0.123895 \times EXP_{LAG3} + 0.075524 \times EXP_{NKG7} + 0.003734 \times EXP_{PDCD1LG2} + 0.032999 \times EXP_{PSMB10} + 0.250229 \times EXP_{STAT1} + 0.084767 \times EXP_{IIGIT}.$
- 2. Cancer-Immunity Cycle: the status of anti-cancer immunity across seven-step Cancer-Immunity Cycle included: Step 1, release of cancer cell antigens; Step 2, cancer antigen presentation; Step 3, priming and activation; Step 4, trafficking of immune cells to tumors; Step 5, infiltration of immune cells into tumors; Step 6, recognition of cancer cells by T cells; Step 7, killing of cancer cell.

Supplementary Figures

Figure S1



Figure S1: CTLA4 shapes a non-inflamed TME in CGGA GBM cohort. (A) Differences in the expression of 108 immunomodulators (chemokines,

receptors, MHC, and immunostimulators) between high- and low-CTLA4 groups in GBM. (B) Differences in the various steps of the cancer immunity cycle between high- and low-CTLA4 groups. Step1: release of cancer cell antigens; Step 2: cancer antigen presentation; Step 3: priming and activation; Step 4_1- Step 4_17: B cell, Basophil CD4+ T cell, CD8+ T cell, Dendritic cell, Eosinophil, Macrophage, MDSC, Monocyte, Neutrophil, NK cell, T cell, Th1 cell, Th17 cell, Th2 cell, Th22 cell, Treg cell recruiting, respectively; Step 5: infiltration of immune cells into tumors; Step 6: recognition of cancer cells by T cells; Step 7: killing of cancer cells. (C) Differences in the expression of 17 inhibitory immune checkpoints between high- and low-CTLA4 groups. (D) Differences in the effector genes of the above tumor-associated immune cells between high- and low-CTLA4 groups. **P < 0.01, ***P < 0.001, ns nonsignificant. MDSC = myeloid derived suppress cell.



Figure S2: (A-S) Kaplan–Meier survival analysis of hub genes calculated by MEGENA algorithm in three GBM cohorts.



Figure S3: Immune landscape between high- and low-CTLA4 groups in CGGA GBM cohort. (A) Relative proportion of TIICs based on CIBERSORT algorithm in CGGA GBM cohort. (B-H) The association between CTLA4 and macrophage using seven independent algorithms. **P < 0.01, ***P < 0.001, ns non-significant.





Figure S4: (A-G) Correlation between CTLA4 and classical chemokines and surface markers of macrophage in CGGA GBM cohort.