## **SUPPLEMENTARY FIGURES**

Study ID	r	HR (95% CI)	Weight %
TCGA LGG dataset		2.77 (1.92, 3.99)	20.35
CGGA dataset 1	_	• 7.32 (5.42, 9.89)	20.85
CGGA dataset 2		4.47 (2.82, 7.08)	19.50
CGGA dataset 3 -		1.40 (0.89, 2.22)	19.53
Rembrandt dataset		2.06 (1.34, 3.18)	19.77
<b>Overall</b> (I-squared = 91.5%, p = 0.000)	$\langle \rangle$	3.07 (1.68, 5.63)	100.00
NOTE: Weights are from random effects analysis			

Supplementary Figure 1. Meta-analysis of the overall prognostic effect of *SLC10A3* among five LGG datasets.



Supplementary Figure 2. Survival analysis of immune cells and correlation with *SLC10A3* expression in LGG. Low expression of immune cells. ((A) B cell, (B) CD8+T cell, (C) CD4+T cell, (D) Macrophage, (E) Neutrophil, (F) Dentritic cell) is correlated with better overall survival in LGG. *SLC10A3* expression is positively associated with the abundance of immune cells ((G) B cell, (H) CD8+T cell, (I) CD4+T cell, (J) Macrophage, (K) Neutrophil, (L) Dentritic cell).



Supplementary Figure 3. Correlation of *SLC10A3* expression with immune sub-types and immune check points in LGG, which was obtained from TISIDB database. (A) Expression of *SLC10A3* is remarkably different among the four immune sub-types. (B) Expression of *SLC10A3* is remarkably different among the six molecular sub-types. Expression of *SLC10A3* is positively linked with immune check points ((C) PDCD1. (D) CD274. (E) PDCD1LG2. (F) HAVCR2. (G) IDO1. (H) LAG3).



Supplementary Figure 4. Correlation analysis of immune check points and *SLC10A3* expression in LGG, which was obtained from TIMER database. (A) PDCD1. (B) CD274. (C) PDCD1LG2. (D) HAVCR2. (E) IDO1. (F) LAG3.



**Supplementary Figure 5. Predictive ability of** *SLC10A3* **to the response of immunotherapy in pan-cancer ICIs cohort.** (A) There is no significant difference of *SLC10A3* between non-responders and responders in ICIs cohort. (B) There is no significant difference of *SLC10A3* between non-responders with primary tumor. (C) There is significant difference of *SLC10A3* between non-responders and responders with primary tumor. (C) There is significant difference of *SLC10A3* between non-responders and responders with primary tumor. (C) There is significant difference of *SLC10A3* between non-responders and responders with recurrent tumor. The predictive ability of *SLC10A3* for immunotherapy response is 0.505 for all the cancer population (D) 0.507 for the primary tumor individuals (E) and 0.674 for the recurrent tumor individuals (F).



**Supplementary Figure 6. Enrichment analysis of** *SLC10A3* **co-expressed genes in LGG. (A)** The typical pathways of GO and KEGG analysis of *SLC10A3* in LGG. (B) The detailed genes of the most significant GO and KEGG pathways.



Supplementary Figure 7. Multiplex immunohistochemistry profiling of *SLC10A3* and immune markers in normal brain tissues. (A) PD1(pink), (B) CD4(yellow), (C) PD-L1(white), (D) *SLC10A3* (blue). (E) CD20(red), (F) CD68(green). (G) The merged image of seven markers. (H) Each marker stands for one special color. (I) Cell phenotype image constructed by the seven markers in the multiplex staining.



Supplementary Figure 8. Comparison of *SLC10A3* expression between LGG and normal tissues in total area, tumor area and stromal area. (A) total *SLC10A3* positive cells, (B) total *SLC10A3* cytoplasm intensity, (C) total *SLC10A3* cell intensity. (D) tumor *SLC10A3* positive cells, (E) tumor *SLC10A3* cytoplasm intensity, (F) tumor *SLC10A3* cell intensity. (G) stromal *SLC10A3* positive cells, (H) stromal *SLC10A3* cell intensity. (I) stromal *SLC10A3* cell intensity.