

Pan-Cancer Analyses Confirmed the Ferroptosis-Related Gene *SLC7A11* as a Prognostic Biomarker for Cancer

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Purpose: Ferroptosis is an iron-dependent and reactive oxygen species (ROS)-reliant form of cell death, exhibiting cellular, molecular, and gene-level characteristics distinct from those of necrosis, autophagy, apoptosis, and pyroptosis. Solute carrier family 7 member 11 (*SLC7A11*), which encodes a cystine/glutamate antiporter transmembrane protein, inhibits ferroptosis by importing cystine and promoting glutathione (GSH) biosynthesis and was found to be overexpressed in multiple human cancers. However, the specific role and underlying mechanism of *SLC7A11* in cancers remains poorly characterized. This research aimed to identify the relationship between *SLC7A11* expression and tumor microenvironment and visualize its prognostic value in pan-cancer.

Patients and Methods: Transcriptomic data for 6313 tumors and normal samples across 20 cancer types were acquired from The Cancer Genome Atlas (TCGA) database. Besides, we presented a novel bioinformatics pipeline that uncovered the impacts of *SLC7A11* on cancer prognosis, tumor mutational burden (TMB), immune cell infiltration in tumor microenvironment, and drug responses. The Genotype-Tissue Expression (GTEx), cBioportal, TCGA and Connectivity Map (CMap) databases were used to explore the expression, genetic alterations, immune microenvironment, and drug responses of *SLC7A11*. A series of deconvolution algorithms, including EPIC, CIBERSORT and GSEA, were utilized for multidimensional analyses of the cancer transcriptomic data.

Results: *SLC7A11* was found to be highly expressed in the 20 types of cancer, especially in solid tumors. Survival analysis uncovered that most cancer patients with up-regulated expression of *SLC7A11* showed poor prognosis, suggesting that *SLC7A11* is a potential oncogene in most cancer types. Furthermore, the expression level of *SLC7A11* was confirmed to be associated with immune cell infiltration in tumor microenvironment, TMB, and drug responses. Gene set enrichment analysis (GSEA) revealed that dysregulation of *SLC7A11* was associated with metabolic and immunity-related signaling pathways in the cancers.

Conclusion: The comprehensive pan-cancer analyses identified *SLC7A11* as an attractive biomarker for immune infiltration and poor prognosis in cancers, shedding new light on the therapeutics of cancers.

Keywords: ferroptosis, *SLC7A11*, pan-cancer, immunotherapy, drug sensitivity

Introduction

The application of immune checkpoint inhibitors (ICIs) has achieved significant success in improving cancer patients' prognosis and is likely to be a crucial part of treatments for patients with a variety of malignancies.¹⁻⁴ However, it should be noted that only a subset of cancer patients are responsive to ICIs.⁵ Several biological indices have been reported as possible biomarkers to identify cancer patients who may benefit from ICIs, such as tumor mutation burden, PD-L1 expression level, and immune cell infiltration in tumor microenvironment.⁶

Ferroptosis is a special form of programmed cell death that shows dependence on active ferrous ions and the accumulation of phospholipid hydroperoxides on the cytoplasmic membrane.⁷⁻⁹ Emerging evidence suggests that ferroptosis exhibits suppressive effects on the development of diverse types of cancer, indicating its potential as a therapeutic target for cancers.¹⁰⁻¹² Similar to apoptosis, increasing evidence suggests that ferroptosis is an essential

mechanism for tumor suppression.^{13,14} However, the clinical application of putative indicator of ferroptosis in cancer therapy is limited, mainly because the specific mechanisms underlying ferroptosis remain unclarified.

SLC7A11 is a cystine/glutamate antiporter transmembrane protein that plays a critical role in mediating cystine uptake in negative modulation of ferroptosis and maintaining cell survival under oxidative stress conditions inhibits and is found to be overexpressed in multiple human cancers.^{15,16} Consistent with this, it was shown that SLC7A11 inactivation has induced potent ferroptosis in many cancer cells; conversely, overexpression of SLC7A11 in cancer cells contributes to the ferroptosis resistance.^{13,17,18} Importantly, recent studies have also demonstrated that immunotherapy and radiotherapy can induce cell ferroptosis partly through increased SLC7A11 expression.^{19–22} Correspondingly, considerable attention has been paid to find out the potential regulatory mechanisms of SLC7A11 in ferroptosis and therapeutically targeting SLC7A11 in cancer therapy. Nevertheless, the role of *SLC7A11* in cancers and the potential molecular mechanisms have not been elucidated, and no studies have been carried out to provide comprehensive pan-cancer analyses for *SLC7A11*.

In the present study, we summarized the possible mechanisms of ferroptosis in cancers and tested the potential of ferroptosis as a putative indicator for cancer immunotherapy via pan-cancer analyses. We further explored the expression patterns and genetic mutations of *SLC7A11* to evaluate their influence on the prognosis of patients with various types of cancer. In addition, we sought to investigate the possible correlation between *SLC7A11* expression and immune cell infiltration in tumor microenvironment. The comprehensive pan-cancer analyses identified *SLC7A11* as an attractive biomarker for tumor microenvironment immune infiltration and poor prognosis in cancers, shedding new light on the therapeutics of cancers.

Materials and Methods

Data Acquisition and Preprocessing

The mRNA expression profiles, genetic data, and related clinical information of pan-cancer cohorts from The Cancer Genome Atlas (TCGA) were downloaded from UCSC xena (<http://xena.ucsc.edu/>), an online database that incorporates gene expression matrix and clinical information from the TCGA and Genotype-Tissue Expression (GTEx) databases. Of the 20 cancer types involved in these two databases, 10 cancer types, including thyroid carcinoma, rectum adenocarcinoma, diffuse large B-cell lymphoma, pheochromocytoma and paraganglioma, thymoma, testicular germ cell tumors, breast invasive carcinoma, lower-grade glioma, chromophobe kidney cancer, and prostate adenocarcinoma, were excluded because the death events were too little for overall survival (OS) analyses.²³ Cases with sarcoma, glioblastoma multiforme, and acute myeloid leukemia were also excluded. Finally, 6313 samples from 20 cancer types, including adrenocortical carcinoma (ACC), cervical squamous cell carcinoma and endocervical adenocarcinoma (CESC), cholangiocarcinoma (CHOL), bladder urothelial carcinoma (BLCA), head and neck squamous cell carcinoma (HNSC), colon adenocarcinoma (COAD), kidney renal papillary cell carcinoma (KIRP), esophageal carcinoma (ESCA), kidney renal clear cell carcinoma (KIRC), lung squamous cell carcinoma (LUSC), liver hepatocellular carcinoma (LIHC), ovarian serous cystadenocarcinoma (OV), lung adenocarcinoma (LUAD), mesothelioma (MESO), stomach adenocarcinoma (STAD), pancreatic adenocarcinoma (PAAD), skin cutaneous melanoma (SKCM), uterine carcinosarcoma (UCS), uterine corpus endometrial carcinoma (UCEC), and uveal melanoma (UVM) with detailed clinical information of the patients from the TCGA and GTEx databases were included in the current study.

OncoPrint, an online cancer microarray data-mining platform (<http://www.oncoPrint.org/>)²⁴ was applied to further compare *SLC7A11* expression in tumor tissues with that in corresponding normal tissues.²⁵ A fold change of 1.5 and a *P* value of 0.05 were applied as the threshold.

Analyses of the Expression and Prognostic Value of SLC7A11

To uncover the correlation between *SLC7A11* expression and OS, the patients were divided into a high-expression group and a low-expression group based on the median level of *SLC7A11* expression in each type of cancer. OS was defined as the duration (days) from the date of diagnosis of cancer to the date of death or the date of the last follow-up for each of the patients. *P* values were calculated using the Log rank test. Survival curves were generated by “survivaal” and

“survminer” R packages, and a p value <0.05 was considered statistically significant. Immunostaining analysis of SLC7A11 protein level was carried out with a rabbit monoclonal anti-SLC7A11 antibody (1:2000, 26864-1-AP, Proteintech). Two experienced pathologists independently assessed the staining of SLC7A11 in one formalin-fixed paraffin-embedding (FFPE) slide. Staining intensity was analyzed and scored from 0 to 3 as follows: 0 = no staining; 1 = weak staining; 2 = moderate staining; and 3 = strong staining. Staining extent was also graded from 0 to 4, according to the percentage of immunoreactive tumor cells (0 = 0%, 1 = 1–25%, 2 = 26–50%, 3 = 51–75%, 4 = 76–100%). The scores of staining intensity were then multiplied by the staining extent scores to yield a total immunohistochemistry (IHC) score (from 0 to 12). A total IHC score of 0 to 3 signified negative staining and a total IHC score of 4 to 12 signified positive staining.

Analyses of SLC7A11 Expression and Clinical Phenotypes

In this study, tumor stages were defined as clinical phenotypes and their relationship with SLC7A11 expression was explored. Statistical analyses (Kruskal–Wallis) for tumor stages were conducted using R software v4.0.3 and plots were generated by the “ggpubr” package in R. $P < 0.05$ was considered statistically significant.

Gene Set Expression Analyses

Gene set enrichment analyses (GSEAs) were performed with the Molecular Signature database (MSigDB) “c2.cp.kegg.v7.2.symbols.gmt” gene set²⁶ to assess enrichment of canonical pathways between the *SLC7A11* high-expression and low-expression groups, using Gene Set Enrichment Analyses software (Java version 4.0).²⁷ Signaling pathways with a normalized $p < 0.05$ and a false discovery rate (FDR) $q < 0.25$ were considered to be significantly enriched pathways.

Analyses of Genetic and Epigenetic Alterations

The open-source cBioPortal database (<http://www.cbioportal.org>) contains cancer genomics data, including information on somatic mutations and copy-number variations (CNVs) for genes, derived from the TCGA and GEO databases.²⁸ Therefore, we utilized this database to analyze single-nucleotide variations in the *SLC7A11* gene in our pan-cancer cohorts. OncoPrints was used to describe the distributions of mutants and copy number alterations (CNAs) in the samples, which were presented in the form of heat map, according to the online instructions at c-BioPortal.

Relationship Between SLC7A11 Expression and Cancer Immunity

Tumor mutational burden (TMB) was defined as the number of somatic, coding, indel mutations and base substitutions per megabases of a genome examined. The TMB in our pan-cancer cohorts was determined based on the TCGA dataset. First, we tested the relationship between TMB and *SLC7A11* gene expression by Spearman correlation analysis. Next, we quantified the proportions of immune cells using the CIBERSORTx²⁹ and EPIC³⁰ algorithms, separately. Spearman correlation analysis was conducted for the relationships among TMB, Immune cell infiltration, and *SLC7A11* expression in multiple tumor tissues.

Drug Responses

Connectivity Map (CMap)³¹ is a public database that contains more than 7000 transcriptional expression profiles from cultured human cells treated with 1309 bioactive small molecules. The expression of *SLC7A11* gene was used to query CMap to predict potential therapeutic drugs for cancer patients. We ranked differentially expressed genes (DEGs) identified between the *SLC7A11* high-expression and low-expression groups according to their p values and the top 500 up-regulated and downregulated genes were deemed eligible for querying against CMap. Drugs with positive connectivity scores are able to induce the input signature in human cell lines. Conversely, drugs with negative connectivity scores are able to induce reverse effects against the input signature in human cell lines.

Results

Expression Patterns of SLC7A11 in Our Pan-Cancer Cohorts

To determine the expression levels of *SLC7A11* in the 20 evaluated cancer types, a total of 6313 samples were enrolled in the study. We found significant differences in the expression pattern of *SLC7A11* between cancer and para-carcinoma tissues in almost all the solid tumors (except for BLCA, SKCM, MESO, and UVM) (Figure 1A), with the expression level of *SLC7A11* being universally higher in almost all the solid tumors (except for BLCA, SKCM, MESO, and UVM) than in the corresponding normal tissues. Moreover, analyses based on the Oncomine database also revealed that, compared with normal tissues, *SLC7A11* expression was up-regulated in solid tumors, such as COAD, ESCA, LUSC, PAAD, BRCA, HNSC, and KIRP (Figure 1B). After comprehensively considering the results of the expression pattern analyses based on different databases, we suggested that *SLC7A11* might be an oncogene for the 20 evaluated types of cancer. To establish if *SLC7A11* is expressed in COAD, we carried out IHC assays on CRC and adjacent normal tissues. As shown in Figure 2, a diffuse pattern of *SLC7A11* expression was observed in both the cytoplasm and the nucleus of colon cancer tissues. Compared with normal adjacent tissues, *SLC7A11* was statistically significantly overexpressed in cancerous tissues ($p < 0.001$).

Prognostic Value of SLC7A11 in Our Pan-Cancer Cohorts

The correlation between *SLC7A11* expression and prognosis (OS) in our pan-cancer cohorts was examined by survival analysis of the TCGA datasets. As shown in Figure 3, we found that *SLC7A11* expression significantly affected patients' OS in seven cancer types, including ACC (HR=4.88, $P < 0.001$), KIRP (HR=3.44, $P < 0.001$), LIHC (HR=1.85, $P < 0.001$), LUAD (HR=1.53, $P = 0.004$), MESO (HR=1.95, $P = 0.003$), OV (HR=0.74, $P = 0.019$), UVM (HR=7.10, $P < 0.001$). For most of these

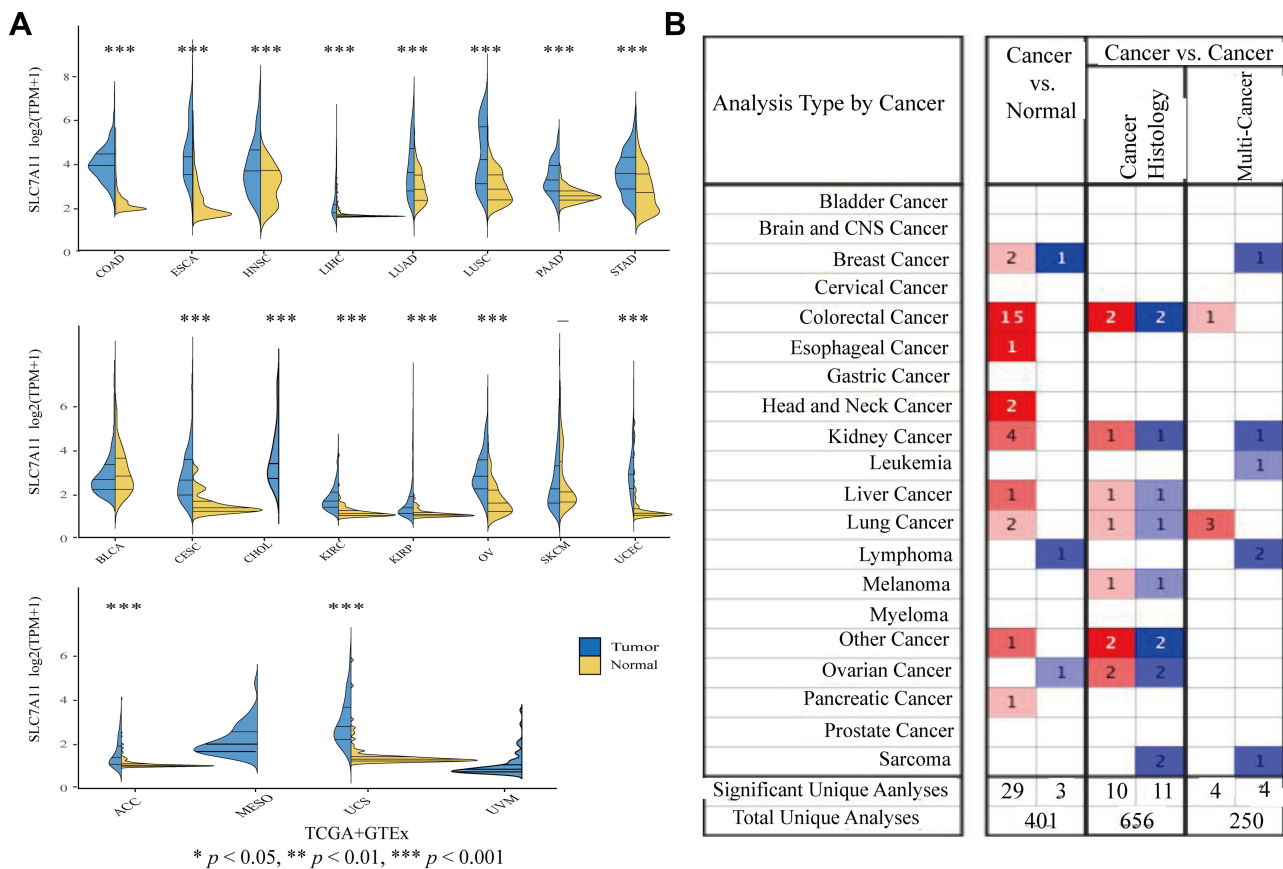


Figure 1 Pan-cancer analysis of *SLC7A11* expression. **(A)** Expression levels of *SLC7A11* in tumor and normal tissues were assessed based on the TCGA and GTEx databases (Wilcoxon test was used for statistical analyses, * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$). **(B)** Expression levels of *SLC7A11* in tumor and normal tissues were assessed based on Oncomine. A fold change of 2 and a P value of 0.01 were designed as the cutoff values.

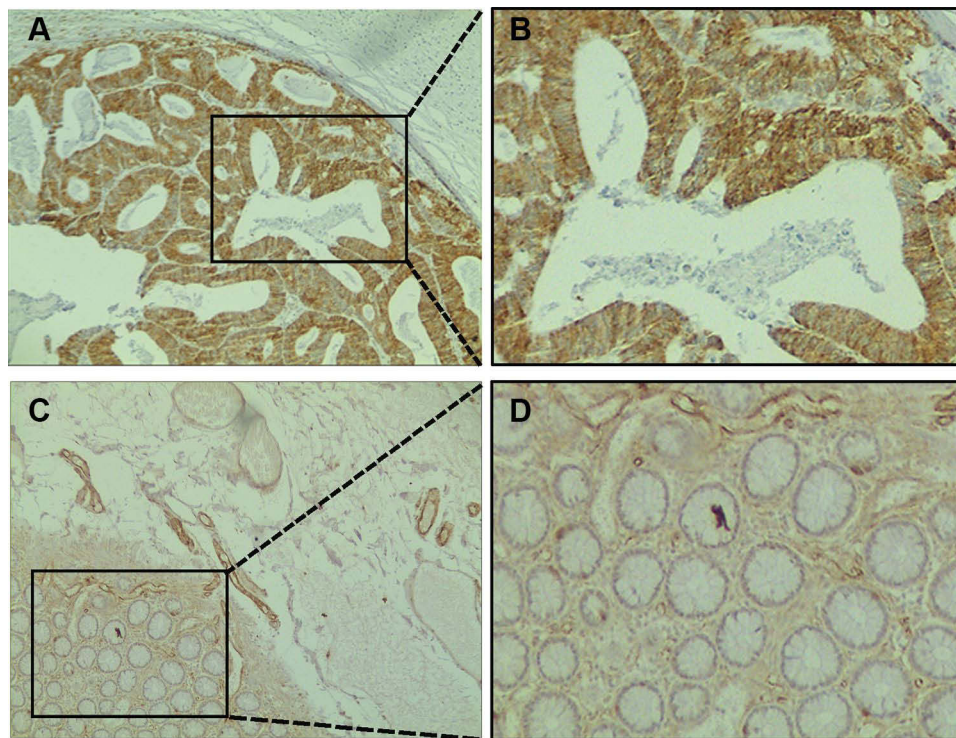


Figure 2 Immunohistochemical staining of SLC7A11 in colon cancer tissues and normal adjacent tissues. (A) Primary cancer tissues magnification 100×. (B) Magnification 400×. (C) Normal adjacent tissues magnification 100×. (D) Magnification 400×.

cancer types, high *SLC7A11* expression predicted a poor prognosis. However, in OV, high *SLC7A11* expression was associated with better OS (Figure 3B). Overall, these results indicated that *SLC7A11* was an important prognostic factor in various cancers and might be a potential pan-cancer prognostic marker.

Analyses of SLC7A11 Expression and Clinical Phenotypes

Next, we performed subgroup analyses of the differential expression of *SLC7A11* according to tumor stage and found that *SLC7A11* expression significantly correlated with tumor stage in KIRP, LIHC, STAD, and SKCM (Figure 4). Notably, the majority of significant differences in *SLC7A11* expression were observed between tumor stages II, III and IV. Interestingly, *SLC7A11* expression increased significantly in stages III and IV compared with stages I and II in KIRP and LIHC, but not in SKCM and STAD.

Mechanisms Underlying the Prognostic Value of SLC7A11 in Our Pan-Cancer Cohorts

Given the important role of SLC7A11 in ferroptosis and the interesting results we found, we determined to explore the role of SLC7A11 dysregulation in the pathogenesis of the 20 investigated types of cancer. We examined the enriched gene sets in samples with high- and low-SLC7A11 mRNA expression, according to the SLC7A11 median expression level, and the first two most significant pathways are shown in Figure 5. The GSEA analysis demonstrated that several cancer-associated pathways and immune-related pathways were hyperactivated in the high SLC7A11 expression group in most tumors. INTERFERON SIGNALING and IL_12 STAT4 PATHWAY were found in ACC, KIRP, and UVM. GPCR LIGAND BINDING was found in COAD, PAAD, LUSC, and KIRC. CELL CYCLE MITOTIC, and CYTOKINE_CYTOKINE RECEPTOR INTERACTION PATHWAY were found in HNSC, and CHOL. The specific function of SLC7A11 in the pathogenesis of pan-cancer is needed to be experimentally verified.

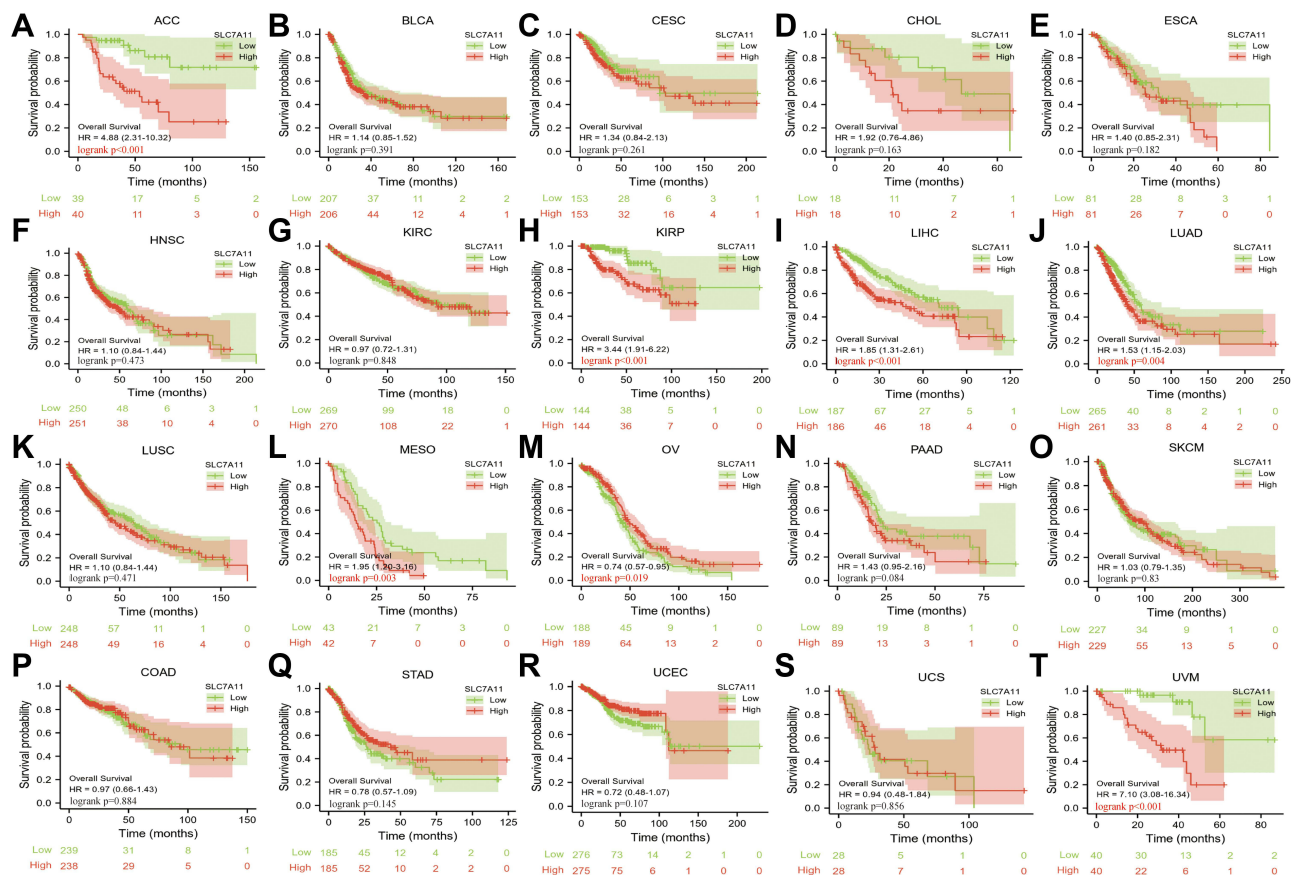


Figure 3 Kaplan-Meier analysis of OS of *SLC7A11* high expression versus low expression patients among the 20 cancer types. (A) ACC; (B) BLCA; (C) CESC; (D) CHOL; (E) ESCA; (F) HNSC; (G) KIRC; (H) KIRP; (I) LIHC; (J) LUAD; (K) LUSC; (L) MESO; (M) OV; (N) PAAD; (O) SKCM; (P) COAD; (Q) STAD; (R) UCEC; (S) UCS; (T) UVM.

Analyses of Genetic Alterations in *SLC7A11* in Our Pan-Cancer Cohorts

Given that genetic changes play a critical role in regulating cancer oncogenesis and development, genetic alterations in *SLC7A11* were further explored based on cBioPortal. A total of 1602 cases retrieved from three tissue studies (10,945 samples from MSKCC, Nat Med 2017; 500 cases from Umich, Nature 2017; 249 cases from Broad/Dana-Farber, Nat Genet 2018) and three cell-line studies (1739 samples from Broad 2019; 1020 samples from Novartis/Broad, Nature 2012; 67 samples NCI, Cancer Res 2012) were analyzed. We found that *SLC7A11* mutations were one of the most important single factors for its deregulated expression. The overall mutation rate of *SLC7A11* in cancers was relatively low (less than 8%). A total of 43 mutations (40 missense and 3 truncation mutations) were found in *SLC7A11*. These mutations affect the protein region of amino acids 0 to 501 (Figure 6B). Among the 20 types of cancer, endometrial carcinoma demonstrated the highest rate of *SLC7A11* mutation followed by Melanoma (Figure 6A). These results implied that genetic alterations in *SLC7A11* played vital roles in tumorigenesis of specific types of cancer, such as endometrial cancer, lymphoma, and colorectal cancer.

SLC7A11 Expression was Correlated with TMB and Immune Cell Infiltration in Tumor Microenvironment in Our Pan-Cancer Cohorts

Emerging evidence has highlighted that targeting ferroptosis-associated metabolisms could improve the efficacy of cancer immunotherapy.³² Therefore, we explored the relationship between *SLC7A11* expression and immune cell infiltration in tumor microenvironment. According to a previous study, an immunosuppressive tumor microenvironment lacking T lymphocyte infiltration is associated with low tumor mutational burden (TMB).³³ Therefore, we explored whether *SLC7A11* expression influenced TMB in our pan-cancer cohorts and found that *SLC7A11* expression was positively correlated with TMB in most of the studied cancer types, especially in ACC, PAAD, STAD, COAD, and

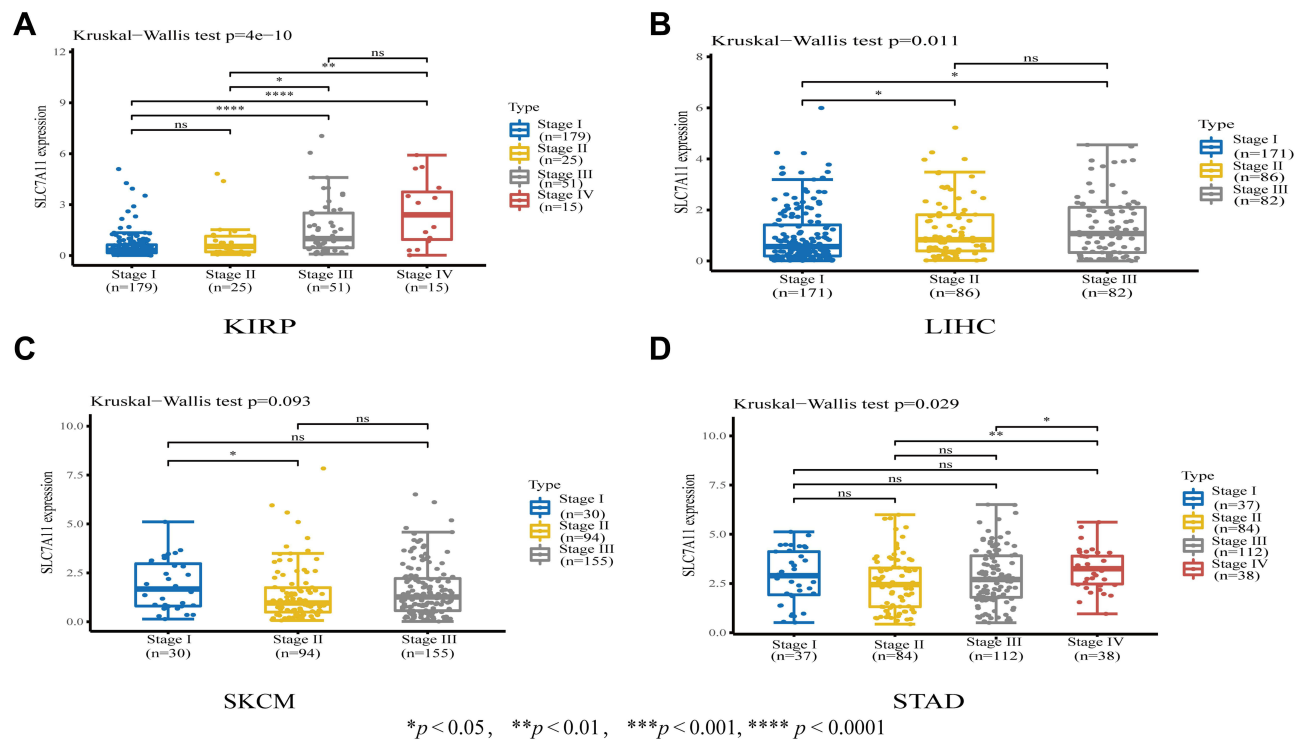


Figure 4 Associations between *SLC7A11* expression and tumor stages in (A) KIRP, (B) LIHC, (C) SKCM, and (D) STAD. * $P < 0.05$, ** $P < 0.01$, **** $P < 0.0001$.

HNSC, but was negatively correlated with the TMB of UVM. Furthermore, by using the EPIC and TIMER algorithms, we found that *SLC7A11* expression correlated with the level of immune cell infiltration in tumor microenvironment in different types of cancer, especially in LUSC, LUAD, and LIHC (Figure 7B). Consistent with our findings, immunotherapy was found to exhibit almost identical significant curative effects on hepatocellular carcinoma (HCC) and LUSC clinically.^{34–36} These results revealed that *SLC7A11* expression was strongly correlated with TMB and immune cell infiltration in tumor microenvironment, suggesting that *SLC7A11* may be a predictor of responses of cancers to immunotherapy.

Drug Responses

Based on the results of CMap database inquiry, a total of 8 small molecular agents were obtained (Figure 8). The eight small molecules showed negative scores in the analyses, meaning that they have the potential to treat corresponding cancers. Among the small molecules, MS-275 had the highest absolute value of score, meaning that the drug has the potential to treat the 20 types of cancer. These results suggested that MS-275 might be a potential broad-spectrum anti-cancer drug. The therapeutic effects of MS-275 on cancers warrant further investigation.

Discussion

In this research, we revealed the roles of *SLC7A11*, a ferroptosis-related gene, as a predictor of survival and a potential regulator of responses to ICIs in 20 types of cancer. Our findings suggested that *SLC7A11* expression was up-regulated in a variety of cancers, especially in solid tumors, and that its high expression could be used as a prognostic indicator. We also found that genetic changes and expression level of *SLC7A11* had potential effects on prognosis, immune cell infiltration in tumor microenvironment, TMB, and responses to putative small-molecule drugs. To the best of our knowledge, this is the first pan-cancer analysis of *SLC7A11* to comprehensively investigate its potential role in underlying tumor prognosis and treatment in cancers.

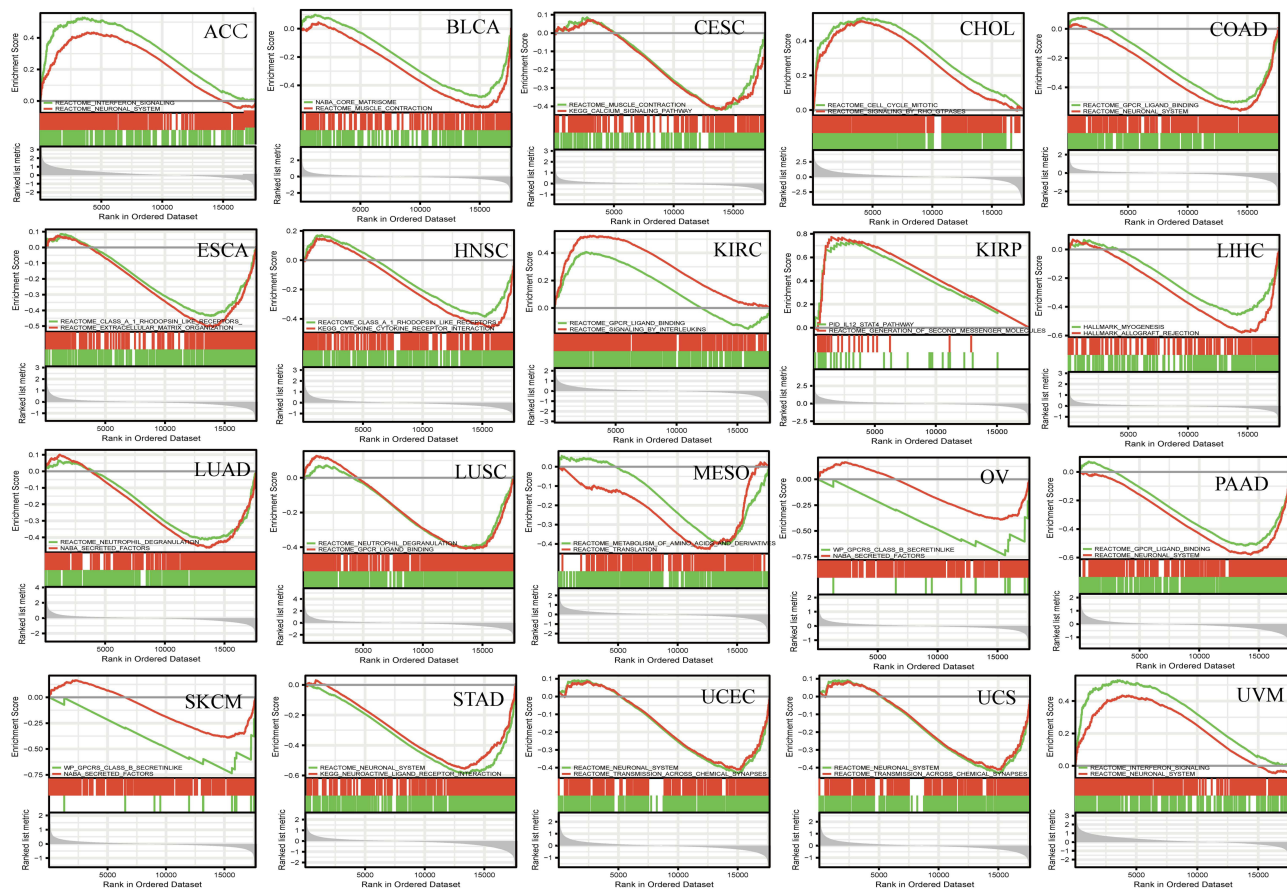


Figure 5 GSEA of *SLC7A11* RNA-seq data in 20 cancer types. GSEA results and GSEA plots of representative data are shown. Normalized enrichment scores (NESs), nominal *P* values (NOMS), and false discovery rates (FDRs) are indicated. Peaks on the upward curve indicate positive regulation and peaks on the downward curve indicate negative regulation.

Ferroptosis is a newly identified form of regulated cell death, which was found to be triggered by the extensive accumulation of iron-dependent reactive oxygen species and lipid peroxides in cancers.^{7,37,38} In addition, researchers have found that ferroptosis-related genes play increasingly important roles in predicting prognosis and shaping tumor microenvironment in cancers.^{39–41} Specifically, induction of ferroptosis is a new therapeutic strategy for advanced cancers. *SLC7A11* is a critical ferroptosis regulator. Previous studies have identified *SLC7A11* as a new UFMylation substrate and revealed that targeting the UFM1/*SLC7A11* pathway is a promising cancer treatment strategy.⁴² Besides, Hagiwara et al found that *SLC7A11* was regulated by the PBAF (SWI/SNF) complex, a subunit of which (PBRM1) formed a complex with MUC1-C and NRF2 to increase chromatin accessibility and induce *SLC7A11* expression.⁴³ *SLC7A11* has also been found to be associated with drug resistance. Lei et al reported that p53 deficiency promoted radiotherapy resistance in cancer cells or tumor tissues at least partly through inhibition of *SLC7A11*-mediated ferroptosis. Collectively, these data motivated us to focus on *SLC7A11*-related mechanisms and carry out in-depth explorations of a variety of cancers.

SLC7A11 has been suggested to play an oncogenic role in various tumors. In order to understand the potential influence of *SLC7A11* on the prognosis of various types of cancer, we performed pan-cancer analyses of the expression patterns of *SLC7A11* across 6313 tumor samples from 20 cancer types deposited by different research groups in TCGA. We observed that *SLC7A11* was highly expressed in various tumors, which is consistent with the observations from previous studies.^{43–46} Notably, by looking at the genetic alterations of *SLC7A11*, we found that the overall mutation rate of *SLC7A11* in cancers was less than 8% and that the main mutation type was missense mutation. We speculate that the mutations in *SLC7A11* are responsible for its aberrant expression in cancers. In addition, by exploring the pathways enriched in cancers with high *SLC7A11* expression, we found that most of the enriched pathways were metabolic, tumor

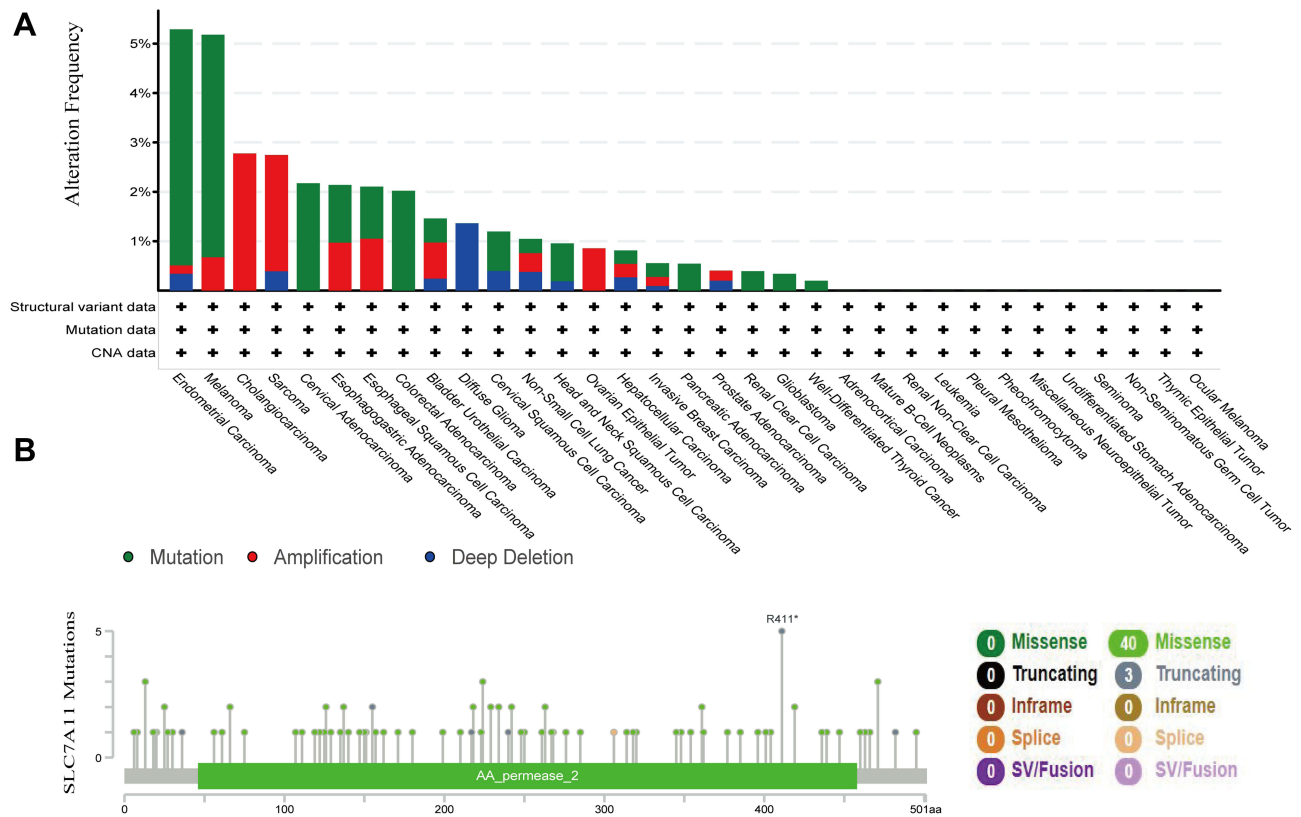


Figure 6 Aberrant expression and mutations of *SLC7A11* in the investigated types of cancer. **(A)** A total of 43 mutation sites were detected, which are located between amino acids 0 to 501 of the *SLC7A11* protein. **(B)** The alteration frequency of *SLC7A11* was determined using cBioportal. The alterations included deletions (blue), amplifications (red), and mutations (green); according to the HGVS (Human Genome Variation Society) gene mutation naming rule R411* indicates the arginine at position 411 becomes a termination code (nonsense mutation); an asterisk (*) indicates the termination code.

microenvironment-related and tumorigenesis-related pathways. These findings suggested that aberrant expression of *SLC7A11* induced diverse biological effects of tumorigenesis and cancer development. Subsequently, survival analysis revealed that the overexpression of *SLC7A11* was associated significantly with reduced OS, indicating that the expression level of *SLC7A11* may be a valuable biomarker for the prognosis of cancer patients.

Accumulating evidence suggests that ferroptosis and lipid metabolism play critical roles in modulating tumor immunity.^{17,47} The mobilization of immune cells to (or within) tumor microenvironment is controlled by various chemokines, cytokines, and pathways, such as ferroptosis.⁴⁸ To clarify the association between *SLC7A11* expression and tumor immunity, we performed Pearson's correlation analysis and observed a strong positive correlation between *SLC7A11* expression and the level of immune cell infiltration in tumor microenvironment in various cancers. Our findings suggest that *SLC7A11* expression may be a valuable biomarker for predicting which cancer patients will benefit from immunotherapy. Consistent with our findings, Dingshan Li and Yongsheng Li pointed out that a combination of inducing ferroptosis and blocking tumor immune escape may become the basis for effective antitumor therapies in clinics, and that the expression of *SLC7A11* may be a valuable predictive biomarker for the prognosis and responses to immunotherapy in patients with breast carcinoma.⁴⁷ These findings need to be further confirmed to examine if *SLC7A11* could be a potential therapeutic target for cancers.

In addition, we utilized the CMap database to identify several small-molecule drugs with potential therapeutic efficacy against cancers with abnormal *SLC7A11* expression. Some of the drugs were shown to have potential anti-cancer effects, such as MS-275/entinostat (an HDAC inhibitor)⁴⁹ and puromycin,⁵⁰ which also exhibited inhibitory effects on most types of cancer investigated in our study. Entinostat, combined with exemestane, has shown encouraging

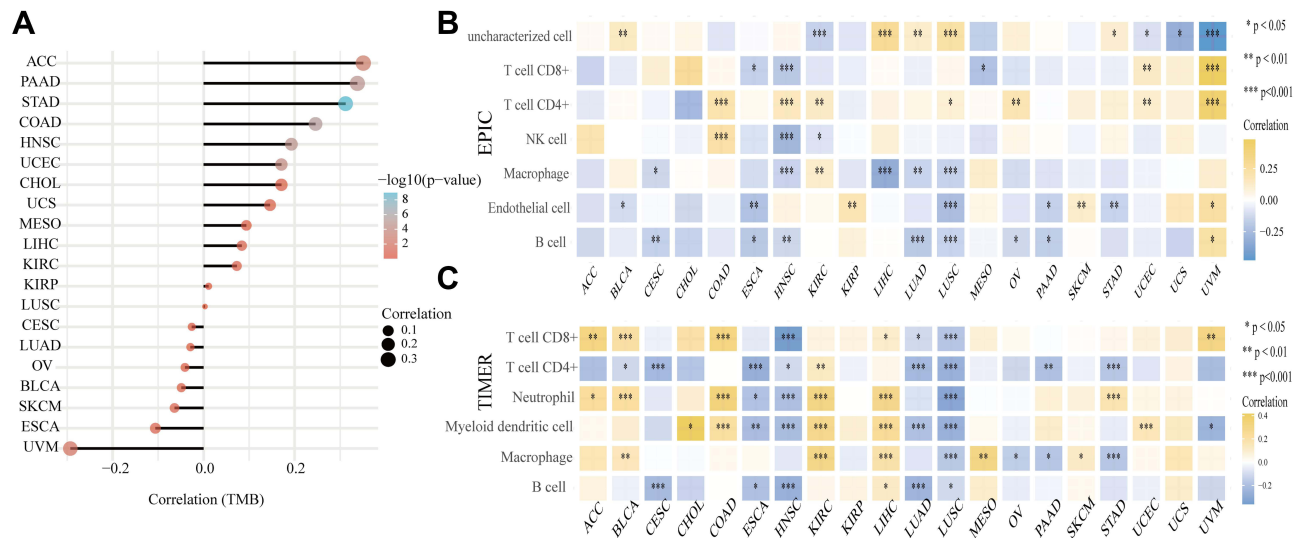


Figure 7 (A) Correlation coefficients between *SLC7A11* expression and TMB in different cancers. The size of the dots in the figure represents the value of the correlation coefficients, and the different colors represent different levels of significance (reflected by *P* values). The vertical axis represents tumor types, while the horizontal axis represents correlation coefficient between *SLC7A11* expression and TMB; a positive relationship is represented by a positive value, and the stronger the correlation, the darker the color (**P* < 0.05, ***P* < 0.01, ****P* < 0.001). (B and C) Spearman correlation analysis heat maps showing the correlations between immune scores of microenvironment infiltrating immune cells (CD8+ T cells, CD4+ T cells, NK cells, macrophages, endothelial cells, B cells, and myeloid dendritic cells) and *SLC7A11* gene expression in 20 tumor tissues. Two latest algorithms, including TIMER (B) and EPIC (C) were used to evaluate immune scores. The horizontal axis represents different tumor tissues, the vertical axis represents different immune scores, and different colors represent correlation coefficients. A positive value represents a positive correlation and a negative value represents a negative correlation. The stronger the correlation, the darker the color, and the asterisk represents the importance (**p*) (**p* < 0.05, ***p* < 0.01, ****p* < 0.001). The significance of the two groups of samples passed the Wilcoxon test.

efficacy in patients with HR+/HER2-metastatic breast cancer.⁵¹ Notably, epigenetic modulators were demonstrated to increase the number of CD8+ effector T cells and improve the efficacy of ICIs. A Phase I clinical study (ETCTN-9844) exhibited that the combination of entinostat, nivolumab and ipilimumab was safe and tolerable with expected rates of immune-related adverse events.⁵² Thus, these identified small-molecule drugs could be potential therapeutic drugs for cancers with abnormal *SLC7A11* expression.

Limitations

Our study has some limitations. First, this is a retrospective study, the data of which come entirely from open databases and the results have not been verified experimentally. Second, the mechanisms underlying *SLC7A11*'s prognostic value in cancers have not been explored. Third, although *SLC7A11* expression exhibited correlations with TMB, the level of immune cell infiltration in tumor microenvironment, and drug responses in cancers, the underlying molecular mechanisms must be investigated by further in vivo and in vitro experiments before these results can be generalized to clinical practice.

Conclusions

Through comprehensive pan-cancer analyses using various databases and algorithms, we found that *SLC7A11* was highly expressed in various cancers, and this high expression was associated with poor overall survival in most of the cancers investigated. In addition, *SLC7A11* expression exhibited significant correlations with TMB, the level of immune cell infiltration in tumor microenvironments, and drug responses in cancers. In summary, these results confirmed the significance of *SLC7A11* expression in cancer prognosis and as a potential predictor for the responses to ICIs.

Data Sharing Statement

The datasets analyzed during the current study are available from the corresponding author upon reasonable request.

specific inhibitors

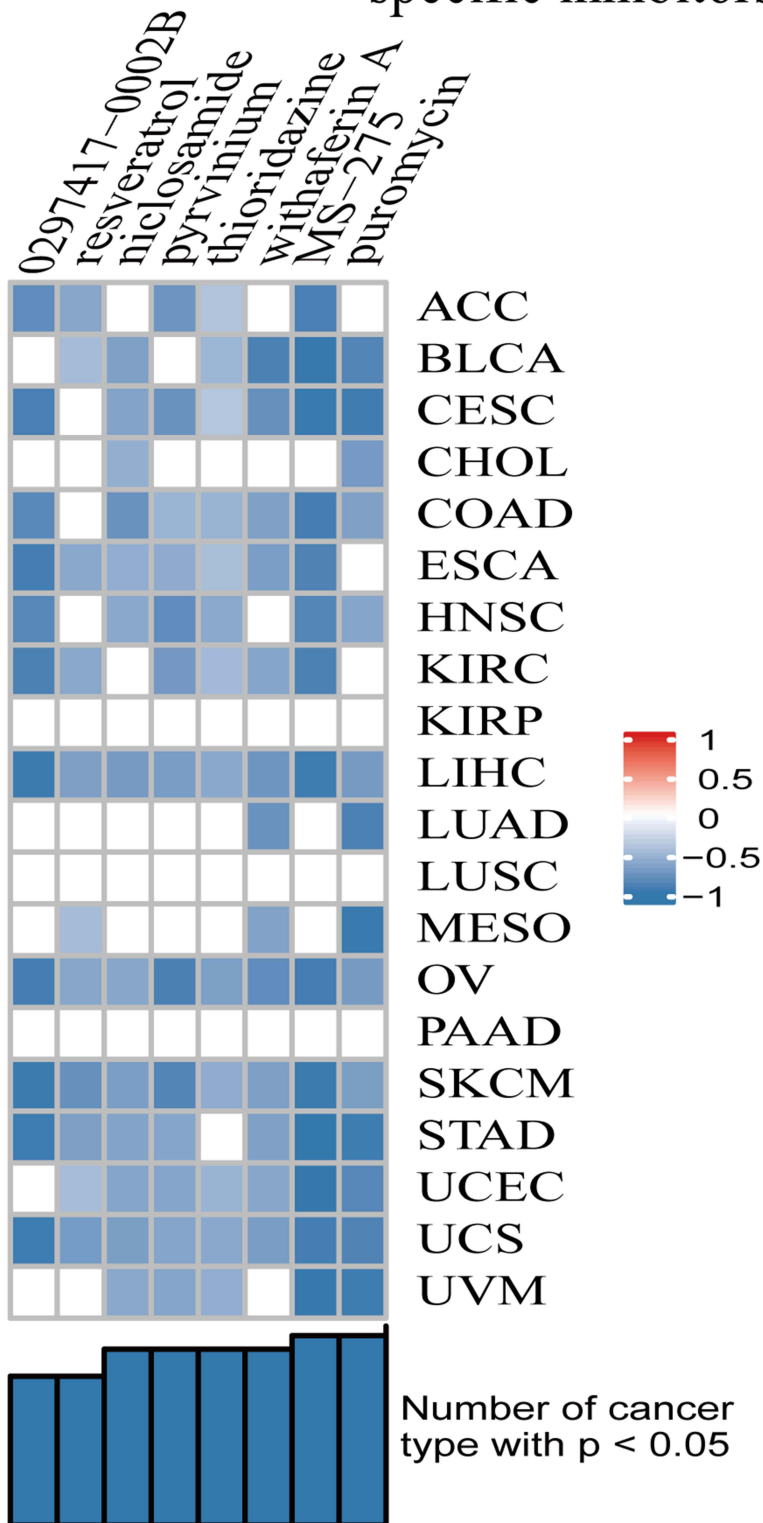


Figure 8 Small molecule drugs with significant correlations as revealed by the results of CMap analysis.

Ethics Approval

The study protocol was approved by the Ethics Committee of the First affiliated Hospital of Dalian Medical University (YJ-KY-FB-2021-21).

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Author Contributions

All authors made a significant contribution to the conception, study design, execution, acquisition of data, analysis and interpretation of data of this work; they also took part in drafting, revising or critically reviewing the manuscript; gave approval of the final version to be published; had agreed on the journal to which the article had been submitted; and agreed to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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