Differential expression of solute carrier family 7 member 10 in cancers of the breast.

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Breast cancer affects women at relatively high frequency¹. We mined published microarray datasets^{2,3} to determine in an unbiased fashion and at the systems level genes most differentially expressed in the primary tumors of patients with breast cancer. We report here significant differential expression of the gene encoding solute carrier family 7 member 10, SLC7A10, when comparing primary tumors of the breast to the tissue of origin, the normal breast. SLC7A10 mRNA was present at significantly lower quantities in tumors of the breast as compared to normal breast tissue. Analysis of human survival data revealed that expression of SLC7A10 in primary tumors of the breast was correlated with overall survival in patients with basal-like, HER2+ and normal-like subtype cancer, demonstrating a relationship between primary tumor expression of a differentially expressed gene and patient survival outcomes influenced by PAM50 molecular subtype. SLC7A10 may be of relevance to initiation, maintenance or progression of cancers of the female breast.

Keywords: breast cancer, SLC7A10, solute carrier family 7 member 10, systems biology of breast cancer, targeted therapeutics in breast cancer.

Invasive breast cancer is diagnosed in over a quarter of a million women in the United States each year¹ and in 2018, breast cancer was the leading cause of cancer death in women worldwide⁴. While patients with localized breast cancer are provided a 99% 5-year survival rate, patients with regional breast cancer, cancer that has spread to lymph nodes or nearby structures, are provided an 86% 5-year survival rate^{5,6}. Patients with metastasis to distant sites, like the brain, are provided a 27% 5-year survival rate^{5,6}. Understanding how primary tumors are most transcriptionally different from the tissue from which they originate, the breast, can facilitate development of novel diagnostic and therapeutics to promote early detection and enhanced treatment, and contribute to efforts to prevent progression to metastatic stages. We mined published microarray data^{2,3} to understand at the transcriptome level and in an unbiased fashion genes most differentially expressed in primary tumors of the breast as compared to normal breast tissue. Solute carrier family 7 member 10 emerged as among the most differentially expressed genes in cancer of the female breast.

Methods

We utilized datasets GSE42568² and GSE109169³ for this global differential gene expression analysis of female breast cancer. GSE42568 was generated using Affymetrix Human Genome U133 Plus 2.0 array technology with $n=17$ normal breast tissue biopsies and $n=104$ primary breast tumor biopsies from patients with breast cancer; analysis was performed using platform GPL570; the majority of patients whose tumors were analyzed were age 50 or older. GSE109169 was generated using Affymetrix Human Exon 1.0 ST Array technology with $n=25$ normal breast tissue and $n=25$ tumors of the breast; analysis was performed using platform GPL5175. The tissues whose expression was profiled in this dataset are paired tissues (25 tumors matching 25 breast tissues from 25 patients). The Benjamini and Hochberg method of p -value adjustment was used for ranking of differential expression but raw p -values were used to assess statistical significance of global differential expression. Log-transformation of data was auto-detected, and the NCBI generated category of platform annotation was used. A statistical test was performed to evaluate whether SLC7A10 expression was significantly different between primary breast tumors and normal breast tissue using a two-tailed t-test. For Kaplan-Meier survival analysis, we used the Kaplan-Meier plotter online tool⁷ for correlation of SLC7A10 mRNA expression levels with overall survival (OS) in $n=431$ patients with basal-like subtype cancer, $n=596$ patients with luminal A subtype cancer, $n=439$ patients with luminal B subtype cancer, $n=362$ patients with HER2+ cancer, and $n=51$ patients with normal-like subtype cancer.

Results

We performed discovery of genes associated with breast cancer in females by mining two independently published microarray datasets^{2,3}.

SLC7A10 is differentially expressed in primary tumors of the breast.

Studying the global gene expression profiles of 104 breast cancers from patients aged 31 to 89 revealed that the gene encoding solute carrier family 7 member 10, SLC7A10, was among the genes most differentially expressed in tumors of the breast in human breast cancer (Chart 1). When sorting each of the genes expressed in tumors of the breast based on significance of difference as compared to normal breast tissue, SLC7A10 ranked 68 out of 54675 total transcripts, equating to 99.9% differential expression (Chart 1). Differential expression of SLC7A10 in female breast cancer was statistically significant (Chart 1; $p=9.66E-33$).

Analysis of a second microarray dataset³, here studying global gene expression patterns in the tumors of 25 patients with early-onset breast cancer, again revealed significant differential expression of

SLC7A10 in human breast cancer (Chart 2). When sorting each of the genes expressed in the tumors of patients with breast cancer based on significance of difference as compared to normal breast tissue, SLC7A10 ranked 3315 out of 19076 total transcripts, equating to 82.6% differential expression (Chart 2). Differential expression of SLC7A10 in the tumors of patients with breast cancer was statistically significant (Chart 2; $p=2.49E-05$). These data suggested that differential expression of SLC7A10 was not an artifact of a single microarray dataset, nor was it strictly associated with early-onset breast cancer, rather a general feature of cancers of the breast.

SLC7A10 is expressed at significantly lower levels in breast tumors as compared to the breast.

We obtained exact mRNA expression levels for SLC7A10 from the breast and from breast tumors to understand the magnitude and direction of SLC7A10 expression change. SLC7A10 was expressed at lower levels in tumors of the breast as compared to normal breast tissue (Figure 1). Decreased expression of SLC7A10 in primary breast tumors was statistically significant (Figure 1: $p<0.0001$). SLC7A10 was expressed at 5.04 ± 1.61 arbitrary units (AU) in normal breast tissue, while it was expressed at 2.37 ± 0.21 AU in tumors of the breast. We calculated a mean fold change of 0.47 in SLC7A10 mRNA levels when comparing primary tumors of the breast to normal breast tissues.

SLC7A10 expression correlates with survival outcomes in basal-like, HER2+ and normal-like subtype human breast cancer.

We performed Kaplan-Meier survival analysis to study relationships between tumor SLC7A10 mRNA expression levels and survival outcomes in patients with breast cancer. We observed a correlation between SLC7A10 expression and overall survival (OS) in patients with basal-like subtype breast cancer which was statistically significant (Figure 2; log rank p -value: 0.04 for overall survival, hazard ratio: 1.5 (1.02-2.2) (Fig. 2)). SLC7A10 mRNA levels were a negative prognostic indicator in basal-like subtype breast cancer patients. Median OS was 95.08 months for basal-like patients with low tumor expression of SLC7A10 while median OS was 59.88 months for basal-like patients with high tumor expression of SLC7A10 (Chart 3).

We also observed a correlation between SLC7A10 expression and overall survival (OS) in patients with HER2+ subtype breast cancer which approached statistical significance (Figure 2; log rank p -value: 0.064 for overall survival, hazard ratio: 1.44 (0.98-2.13) (Fig. 2)). SLC7A10 mRNA levels were a negative prognostic indicator in HER2+ subtype breast cancer patients. Median OS was 69.99 months for HER2+ patients with low tumor expression of SLC7A10 while median OS was 41.39 months for HER2+ patients with high tumor expression of SLC7A10 (Chart 3).

We also observed a correlation between SLC7A10 expression and overall survival (OS) in patients with normal-like subtype breast cancer which approached statistical significance (Figure 2; log rank p -value: 0.051 for overall survival, hazard ratio: 2.74 (0.95-7.91) (Fig. 2)). SLC7A10 mRNA levels were a negative prognostic indicator in normal-like subtype breast cancer patients. OS was, on average, markedly greater for normal-like patients with low tumor expression as compared to normal-like patients with high tumor expression of SLC7A10 (Chart 3).

SLC7A10 primary tumor expression was not correlated with overall survival in luminal A breast cancer (Figure 2; log rank p -value: 0.79 for OS, hazard ratio: 0.94 (0.61-1.46) (Fig. 2)) or in luminal B subtype breast cancer (Figure 2; log rank p -value: 0.99 for OS, hazard ratio: 1 (0.7-1.42) (Fig. 2)).

Thus, through comparative transcriptome analysis of primary tumors of the breast and normal breast tissue, we found that differential expression and down-regulation of SLC7A10 was among the most significant transcriptional features in primary tumors from patients with breast cancer. SLC7A10 expression in primary tumors of the breast was correlated with overall survival in patients with basal-like, HER2+ and normal-like subtype disease, with mRNA levels of SLC7A10 a negative prognostic indicator

for basal-like, HER2+ and normal-like breast cancer patients.

Discussion

Invasive breast cancer is a medical problem with a 27% 5-year survival rate for women whose disease has spread to distant sites^{5,6}. To facilitate understanding of the basic transcriptional differences between primary tumors of the breast and the tissues from which these tumors originate, normal breast tissues, we performed comparative transcriptome analysis using two independently published microarray datasets^{2,3}, providing evidence here that differential expression of solute carrier family 7 member 10, encoded by SLC7A10, is a defining transcriptional feature of human breast cancer: in patients diagnosed after age 50, and in early onset-breast cancer. SLC7A10 was expressed at significantly lower levels in primary tumors from patients with breast cancer as compared to normal breast tissue. Importantly, in patients with basal-like, HER2+ and normal-like subtype breast cancers, expression of SLC7A10 was correlated with overall survival. Solute carrier family 7 member 10 and the molecular processes to which it pertains to may be relevant to the initiation or progression of human breast cancer.

References

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Rank: 68
Probe ID: 220868_s_at
p-value: 9.66E-33
t: 1.65E+01
B: 63.9181008
Gene: SLC7A10
Gene name: solute carrier family 7 member 10

Chart 1: SLC7A10 is differentially expressed in the primary tumors of patients with breast cancer.

Rank of differential expression, probe ID, p-value of global differential expression, t, a moderated t-statistic, B, the log-odds of differential expression between the groups compared, gene and gene name are listed in this chart.

Rank: 3315
Probe ID: 3858907
p-value: 2.49E-05
t: 4.64
B: 2.068927
Gene: SLC7A10
Gene name: solute carrier family 7 member 10

Chart 2: SLC7A10 is differentially expressed in the primary tumors of patients with early-onset human breast cancer.

Rank of differential expression, probe ID, p-value of global differential expression, t, a moderated t-statistic, B, the log-odds of differential expression between the groups compared, gene and gene name are listed in this chart.

Figure 2: SLC7A10 expression correlates with overall survival in patients with basal-like, HER2+ and normal-like subtype breast cancer, but not in patients with luminal A or in luminal B subtype cancer.

Depicted in this Kaplan-Meier plot is the probability of overall survival (OS) for $n=431$ patients with basal-like breast cancer (upper left), $n=362$ patients with HER2+ breast cancer (upper right), $n=596$ patients with luminal A breast cancer (middle left), $n=439$ patients with luminal B breast cancer (middle right), and $n=51$ patients with normal-like breast cancer (below) stratified into two groups, based on low or high expression of SLC7A10 in patient primary tumors. The log rank p-value denoting statistical significance of difference in overall survival when comparing the two groups, as well as hazard ratio for this comparison is listed above. Listed below is the number of patients at risk (number of patients alive) per interval, after stratification based on SLC7A10 expression; in the first interval, number at risk is number of patients alive; in each subsequent interval, number at risk is the number at risk less those who have expired or are censored.

Basal-like patients

Low SLC7A10 expression: 95.08 months

High SLC7A10 expression: 59.88 months

HER2+ patients

Low SLC7A10 expression: 69.99 months

High SLC7A10 expression: 41.39 months

Chart 3: In basal-like and HER2+ subtype breast cancer, median overall survival is superior in patients with low primary tumor expression of SLC7A10.

The overall survival of $n=431$ basal-like subtype breast cancer patients and $n=362$ HER2+ subtype breast cancer patients based on stratification into low or high tumor expression of SLC7A10 is listed in this chart.