Differential expression of transmembrane protein 132C 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 transmembrane protein 132C, TMEM132C, when comparing primary tumors of the breast to the tissue of origin, the normal breast. TMEM132C was also differentially expressed in the brain metastases of patients with metastatic breast cancer. TMEM132C mRNA was present at lower quantities in tumors of the breast as compared to normal breast tissue. Analysis of human survival data revealed that expression of TMEM132C in primary tumors of the breast was correlated with distant metastasis-free survival in patients with luminal B subtype cancer, demonstrating a relationship between primary tumor expression of a differentially expressed gene and patient survival outcomes influenced by PAM50 molecular subtype. TMEM132C may be of relevance to initiation, maintenance or progression of cancers of the female breast.

Keywords: breast cancer, TMEM132C, transmembrane protein 132C, 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. Transmembrane protein 132C emerged as among the most differentially expressed genes in cancer of the female breast.

Methods

We utilized datasets GSE42568² and GSE52604³ 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. GSE52604 was generated using Agilent-014850 Whole Human Genome Microarray 4x44K G4112F with $n=10$ normal breast tissues and $n=35$ brain metastases from patients with breast cancer; analysis was performed using platform GPL6480. 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 TMEM132C 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 TMEM132C mRNA expression levels with distant metastasis-free survival (DMFS) in $n=283$ patients with basal-like subtype cancer, $n=259$ patients with luminal A subtype cancer, $n=183$ patients with luminal B subtype cancer, $n=218$ patients with HER2+ cancer, and $n=15$ 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}.

TMEM132C 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 transmembrane protein 132C, TMEM132C, 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, TMEM132C ranked 16 out of 54675 total transcripts, equating to 99.97% differential expression (Chart 1). Differential expression of TMEM132C in female breast cancer was statistically significant (Chart 1; $p=1.09E-37$).

Analysis of a second microarray dataset³, here studying global gene expression patterns in the brain metastases of patients with metastatic breast cancer, again revealed significant differential expression of TMEM132C in tumor tissue of patients with human breast cancer (Chart 2). When sorting each of the

genes expressed in the brain metastases of patients with breast cancer based on significance of difference as compared to normal breast tissue, TMEM132C ranked 1730 out of 41093 total transcripts, equating to 95.8% differential expression (Chart 2). Differential expression of TMEM132C in the brain metastases of patients with breast cancer was statistically significant (Chart 2; $p=1.92E-06$). These data suggested that differential expression of TMEM132C was not an artifact of a single microarray dataset, nor was it strictly associated with primary tumors of the breast, rather a reflection of the spectrum of transformation in human cancer.

TMEM132C is expressed at lower levels in breast tumors as compared to the breast.

We obtained exact mRNA expression levels for TMEM132C from the breast and from breast tumors to understand the magnitude and direction of TMEM132C expression change. TMEM132C was expressed at lower levels in tumors of the breast as compared to normal breast tissue (Figure 1). Decreased expression of TMEM132C in primary breast tumors approached statistical significance (Figure 1: $p=0.0646$). TMEM132C was expressed at 8.36 ± 1.43 arbitrary units (AU) in normal breast tissue, while it was expressed at 4.56 ± 0.61 AU in tumors of the breast. We calculated a mean fold change of 0.55 in TMEM132C mRNA levels when comparing primary tumors of the breast to normal breast tissues.

TMEM132C expression correlates with survival outcomes in luminal B subtype human breast cancer.

We performed Kaplan-Meier survival analysis to study relationships between tumor TMEM132C mRNA expression levels and survival outcomes in patients with breast cancer. We observed a correlation between TMEM132C expression and distant metastasis-free survival (DMFS) in patients with luminal B subtype breast cancer, in the lower quartile, which was statistically significant (Figure 2; log rank p -value: 0.062 for distant metastasis-free survival, hazard ratio: 0.55 (0.29-1.04) (Fig. 2)). TMEM132C mRNA levels were a positive prognostic indicator in luminal B subtype breast cancer patients. Median DMFS was, on average, markedly shorter for luminal B patients with low tumor expression of TMEM132C as compared to luminal B patients with high tumor expression of TMEM132C (Chart 3). TMEM132C primary tumor expression was not correlated with distant metastasis-free survival in basal-like breast cancer (Figure 2; log rank p -value: 0.13 for DMFS, hazard ratio: 0.68 (0.41-1.12) (Fig. 2)), luminal A subtype breast cancer (Figure 2; log rank p -value: 0.89 for DMFS, hazard ratio: 0.95 (0.45-1.99) (Fig. 2)), in HER2+ breast cancer (Figure 2; log rank p -value: 0.45 for DMFS, hazard ratio: 0.82 (0.48-1.38) (Fig. 2)) or in patients with normal-like breast cancer (Figure 2; log rank p -value: 0.62 for DMFS, hazard ratio: 0.65 (0.12-3.61) (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 TMEM132C was among the most significant transcriptional features in primary tumors from patients with breast cancer. TMEM132C expression in primary tumors of the breast was correlated with distant metastasis-free survival in patients with luminal B subtype disease, with mRNA levels of TMEM132C a positive prognostic indicator for luminal B 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 transmembrane protein 132C, encoded

1 by TMEM132C, is a defining transcriptional feature of human breast cancer: in patients diagnosed after
2 age 50, and in brain metastatic tissues from patients with human breast cancer. TMEM132C was
3 expressed at lower levels in primary tumors from patients with breast cancer as compared to normal breast
4 tissue. Importantly, in patients with luminal B subtype breast cancers, expression of TMEM132C was
5 correlated with distant metastasis-free survival. Transmembrane protein 132C and the molecular
6 processes to which it pertains to may be relevant to the initiation or progression of human breast cancer.
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Rank: 16
Probe ID: 232313_at
p-value: 1.09E-37
t: 1.89E+01
B: 75.1435588
Gene: TMEM132C
Gene name: transmembrane protein 132C

Chart 1: TMEM132C 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: 1730
Probe ID: A_23_P151232
p-value: 1.92E-06
t: 5.4340964
B: 4.760292
Gene: TMEM132C
Gene name: transmembrane protein 132C

Chart 2: TMEM132C is differentially expressed in the brain metastases of patients with metastatic 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 1: TMEM132C is expressed at lower levels in primary breast tumors as compared to normal breast tissue.

The mRNA expression level of TMEM132C in normal breast tissue (left) and in primary tumors of the breast (right) is graphically depicted with the result of a statistical test evaluating significance of difference in TMEM132C expression between normal breast tissue and primary tumors of the breast, a *p*-value, listed above.

Figure 2: TMEM132C expression correlates with distant metastasis-free survival in patients with luminal B breast cancer, but not in patients with basal, luminal A, HER2+ or normal-like cancer.

Depicted in this Kaplan-Meier plot is the probability of distant metastasis-free survival (DMFS) for $n=283$ patients with basal-like breast cancer (upper left), $n=218$ patients with HER2+ breast cancer (upper right), $n=259$ patients with luminal A breast cancer (middle left), $n=183$ patients with luminal B breast cancer (middle right), and $n=15$ patients with normal-like breast cancer (below) stratified into two groups, based on low or high expression of TMEM132C in patient primary tumors, in the lower quartile. The log rank p-value denoting statistical significance of difference in distant metastasis-free 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 TMEM132C 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.