Differential expression of potassium voltage-gated channel interacting protein 2 in cancers of the breast.

Shahan Mamoor, MS¹

¹shahanmamoor@gmail.com

East Islip, NY 11730

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 potassium voltage-gated channel interacting protein 2, KCNIP2, when comparing primary tumors of the breast to the tissue of origin, the normal breast. KCNIP2 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 KCNIP2 in primary tumors of the breast was correlated with recurrence-free survival in patients with basal-like and 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. KCNIP2 may be of relevance to initiation, maintenance or progression of cancers of the female breast.

Keywords: breast cancer, KCNIP2, potassium voltage-gated channel interacting protein 2, 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. Potassium voltage-gated channel interacting protein 2 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 KCNIP2 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 KCNIP2 mRNA expression levels with recurrence-free survival (RFS) in $n=953$ patients with basal-like subtype cancer, $n=1809$ patients with luminal A subtype cancer, $n=1353$ patients with luminal B subtype cancer, $n=695$ patients with HER2+ cancer, and $n=119$ 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}.

KCNIP2 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 potassium voltage-gated channel interacting protein 2, KCNIP2, 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, KCNIP2 ranked 15 out of 54675 total transcripts, equating to 99.97% differential expression (Chart 1). Differential expression of KCNIP2 in female breast cancer was statistically significant (Chart 1; $p=9.36E-38$).

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

KCNIP2 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, KCNIP2 ranked 1666 out of 19076 total transcripts, equating to 91.3% differential expression (Chart 2). Differential expression of KCNIP2 in the tumors of patients with breast cancer was statistically significant (Chart 2; $p=1.94E-07$). These data suggested that differential expression of KCNIP2 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.

KCNIP2 is expressed at significantly lower levels in breast tumors as compared to the breast.

We obtained exact mRNA expression levels for KCNIP2 from the breast and from breast tumors to understand the magnitude and direction of KCNIP2 expression change. KCNIP2 was expressed at lower levels in tumors of the breast as compared to normal breast tissue (Figure 1). Decreased expression of KCNIP2 in primary breast tumors was statistically significant (Figure 1: $p=0.0033$). KCNIP2 was expressed at 7.93 ± 2.11 arbitrary units (AU) in normal breast tissue, while it was expressed at 3.50 ± 0.50 AU in tumors of the breast. We calculated a mean fold change of 0.44 in KCNIP2 mRNA levels when comparing primary tumors of the breast to normal breast tissues.

KCNIP2 expression correlates with survival outcomes in basal-like and luminal B subtype human breast cancer.

We performed Kaplan-Meier survival analysis to study relationships between tumor KCNIP2 mRNA expression levels and survival outcomes in patients with breast cancer. We observed a correlation between KCNIP2 expression and recurrence-free survival (RFS) in patients with luminal B subtype breast cancer which was statistically significant (Figure 2; log rank p -value: 0.0091 for recurrence-free survival, hazard ratio: 0.79 (0.67-0.94) (Fig. 2)). KCNIP2 mRNA levels were a positive prognostic indicator in luminal B subtype breast cancer patients. Median RFS was 127.4 months for luminal B patients with low tumor expression of KCNIP2 while median RFS was 164.3 months for luminal B patients with high tumor expression of KCNIP2 (Chart 3).

We observed a correlation between KCNIP2 expression and recurrence-free survival (RFS) in patients with basal-like subtype breast cancer which approached statistical significance (Figure 2; log rank p -value: 0.058 for recurrence-free survival, hazard ratio: 0.81 (0.66-1.01) (Fig. 2)). KCNIP2 mRNA levels were a positive prognostic indicator in basal-like subtype breast cancer patients. Median RFS was 20 months for basal-like patients with low tumor expression of KCNIP2 while median RFS was 30 months for basal-like patients with high tumor expression of KCNIP2 (Chart 3).

KCNIP2 primary tumor expression was not correlated with recurrence-free survival in luminal A subtype breast cancer (Figure 2; log rank p -value: 0.19 for RFS, hazard ratio: 0.87 (0.71-1.07) (Fig. 2)), in HER2+ breast cancer (Figure 2; log rank p -value: 0.25 for RFS, hazard ratio: 0.87 (0.68-1.11) (Fig. 2)) or in patients with normal-like breast cancer (Figure 2; log rank p -value: 0.52 for RFS, hazard ratio: 0.8 (0.41-1.58) (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 KCNIP2 was among the most significant transcriptional features in primary tumors from patients with breast cancer. KCNIP2 expression in primary tumors of the breast was correlated with recurrence-free survival in patients with luminal B and basal-like subtype disease, with mRNA levels of KCNIP2 a positive prognostic indicator for luminal B and basal-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 potassium voltage-gated channel interacting protein 2, encoded by KCNIP2, is a defining transcriptional feature of human breast cancer: in patients diagnosed after age 50, and in early onset-breast cancer. KCNIP2 was expressed at significantly lower levels in primary tumors from patients with breast cancer as compared to normal breast tissue. Importantly, in patients with luminal B and basal-like breast cancers, expression of KCNIP2 was correlated with recurrence-free survival. Potassium voltage-gated channel interacting protein 2 and the molecular processes to which it pertains to may be relevant to the initiation or progression of human breast cancer.

References

1. DeSantis, C.E., Ma, J., Goding Sauer, A., Newman, L.A. and Jemal, A., 2017. Breast cancer statistics, 2017, racial disparity in mortality by state. *CA: a cancer journal for clinicians*, 67(6), pp.439-448.
2. Clarke, C., Madden, S.F., Doolan, P., Aherne, S.T., Joyce, H., O'driscoll, L., Gallagher, W.M., Hennessy, B.T., Moriarty, M., Crown, J. and Kennedy, S., 2013. Correlating transcriptional networks to breast cancer survival: a large-scale coexpression analysis. *Carcinogenesis*, 34(10), pp.2300-2308.
3. Chang, J.W., Kuo, W.H., Lin, C.M., Chen, W.L., Chan, S.H., Chiu, M.F., Chang, I.S., Jiang, S.S., Tsai, F.Y., Chen, C.H. and Huang, P.H., 2018. Wild-type p53 upregulates an early onset breast cancer-associated gene GAS7 to suppress metastasis via GAS7–CYFIP1- mediated signaling pathway. *Oncogene*, 37(30), pp.4137-4150.
4. Bray, F., Ferlay, J., Soerjomataram, I., Siegel, R.L., Torre, L.A. and Jemal, A., 2018. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA: a cancer journal for clinicians*, 68(6), pp.394-424.
5. ACS Cancer Facts & Figures 2019.
<https://www.cancer.net/cancer-types/breast-cancer/metastatic/statistics>.
6. Survival Rates for Breast Cancer. <https://www.cancer.org/cancer/breast-cancer/understanding-a-breast-cancer-diagnosis/breast-cancer-survival-rates.html>
7. Györfy, B., Lanczky, A., Eklund, A.C., Denkert, C., Budczies, J., Li, Q. and Szallasi, Z., 2010. An online survival analysis tool to rapidly assess the effect of 22,277 genes on breast cancer prognosis using microarray data of 1,809 patients. *Breast cancer research and treatment*, 123(3), pp.725-731.

Rank: 15
Probe ID: 223727_at
p-value: 9.36E-38
t: 1.89E+01
B: 75.296953
Gene: KCNIP2
Gene name: potassium voltage-gated channel interacting protein 2

Chart 1: KCNIP2 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: 1666
Probe ID: 3304073
p-value: 1.94E-07
t: 6.01
B: 6.798586
Gene: KCNIP2
Gene name: potassium voltage-gated channel interacting protein 2

Chart 2: KCNIP2 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 1: KCNIP2 is expressed at significantly lower levels in primary breast tumors as compared to normal breast tissue.

The mRNA expression level of KCNIP2 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 KCNIP2 expression between normal breast tissue and primary tumors of the breast, a *p*-value, listed above.

Figure 2: KCNIP2 expression correlates with recurrence-free survival in patients with basal-like and luminal B breast cancer, but not in patients with luminal A, HER2+ or normal-like disease.

Depicted in this Kaplan-Meier plot is the probability of recurrence-free survival (RFS) for $n=953$ patients with basal-like breast cancer (upper left), $n=695$ patients with HER2+ breast cancer (upper right), $n=1809$ patients with luminal A breast cancer (middle left), $n=1353$ patients with luminal B breast cancer (middle right), and $n=119$ patients with normal-like breast cancer (below) stratified into two groups, based on low or high expression of KCNIP2 in patient primary tumors. The log rank p-value denoting statistical significance of difference in recurrence-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 KCNIP2 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 KCNIP2 expression: 20 months

High KCNIP2 expression: 30 months

Luminal B patients

Low KCNIP2 expression: 127.4 months

High KCNIP2 expression: 164.3 months

Chart 3: In basal-like and luminal B breast cancer, median recurrence-free survival is superior in patients with high primary tumor expression of KCNIP2.

The recurrence-free survival of $n=953$ basal-like breast cancer and $n=1353$ luminal B subtype breast cancer patients based on stratification into low or high tumor expression of KCNIP2 is listed in this chart.