Differential expression of anti-silencing function 1B histone chaperone in triple negative breast cancer.

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Women diagnosed with triple negative breast cancer can benefit neither from endocrine therapy nor from HER2-targeted therapies¹. 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 anti-silencing function 1B histone chaperone, ASF1B, when comparing the tumor cells of patients with triple negative breast cancer to normal mammary ductal cells². ASF1B was also differentially expressed in bulk tumor in human breast cancer³. ASF1B mRNA was present at significantly increased quantities in TNBC tumor cells relative to normal mammary ductal cells. Analysis of human survival data revealed that expression of ASF1B in primary tumors of the breast was correlated with overall survival in patients with HER2+ type cancer, while within triple negative breast cancer, primary tumor expression of ASF1B was correlated with overall survival in patients with basal-like 2 subtype disease. ASF1B may be of relevance to initiation, maintenance or progression of triple negative breast cancers.

Keywords: breast cancer, ASF1B, anti-silencing function 1B histone chaperone, systems biology of breast cancer, targeted therapeutics in breast cancer.

Patients with hormone receptor positive disease can benefit from endocrine therapy⁴: aromatase inhibition such as that provided by exemestane or letrozole, or estrogen receptor antagonism such as that provided by tamoxifen or fulvestrant. Patients whose tumors overexpress the human epidermal growth factor 2 (HER2) can be treated with HER2-targeted therapies: HER2 inhibitors, immunoglobulin⁵- or small molecule-based⁶. As triple negative breast cancer (TNBC) is defined by its lack of HER2 expression as well as by its lack of expression of the hormone receptor estrogen, patients diagnosed with triple negative breast cancer will not benefit from either of these treatment strategies.

Understanding the transcriptional composition of triple negative breast cancers in humans can facilitate development of novel therapeutics 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 the tumor cells of patients with triple negative breast cancer.

Methods

We utilized datasets GSE38959² and GSE109169³ for this differential gene expression analysis of female breast cancer in conjunction with GEO2R. GSE38959 was generated using Agilent-014850 Whole Human Genome Microarray 4x44K G4112F technology with $n=13$ samples of normal mammary gland ductal cells and $n=30$ samples of tumor cells from patients with triple negative breast cancer; analysis was performed using platform GPL4133. 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 from this dataset are paired tissues (25 tumors matching 25 normal 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 ASF1B expression was significantly different between primary breast tumors and breast tissue using a two-tailed t-test.

For Kaplan-Meier survival analysis, we used the Kaplan-Meier plotter tool⁷ for correlation of ASF1B mRNA expression levels with overall survival (OS) in $n=431$ patients with basal 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. In patients with triple negative breast cancer⁷, we analyzed overall survival (OS) for $n=103$ patients with basal-like 1 subtype TNBC, $n=58$ patients with basal-like 2 subtype TNBC, $n=149$ patients with immunomodulatory subtype TNBC, $n=116$ patients with luminal androgen receptor subtype TNBC, $n=114$ patients with mesenchymal subtype TNBC, and $n=39$ patients with mesenchymal stem-like subtype TNBC.

Results

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

ASF1B is differentially expressed in primary tumors and tumor cells of the breast.

Comparison of the tumor cells of patients with triple negative breast cancer to normal mammary ductal cells² revealed anti-silencing function 1B histone chaperone, encoded by ASF1B, as among the genes whose expression was most significantly different in human TNBC as compared to the normal breast (Chart 1). When sorting each of the genes expressed in the tumor cells of patients with triple negative breast cancer based on significance of difference as compared to normal mammary ductal cells, ASF1B ranked 217 out of 45015 total transcripts, equating to 99.5% differential expression (Chart 1). Differential expression of ASF1B in the tumor cells of patients with triple negative breast cancer was statistically significant (Chart 1; $p=3.58E-11$).

Analysis of a second microarray dataset³, here comparing 25 normal breast tissues to 25 tumors of the breast revealed that ASF1B was also among the genes most differentially expressed in human breast cancer, here in the tumors of women diagnosed with early-onset breast cancer (Chart 2). When sorting each of the genes expressed in tumors of the breast based on significance of difference as compared to normal breast tissue, ASF1B ranked 321 out of 19076 total transcripts, equating to 98.3% differential expression (Chart 2). Differential expression of ASF1B in female breast cancer was statistically significant (Chart 2; $p=5.89E-12$).

These data indicated that differential expression of ASF1B was not an artifact of a single microarray dataset, was not an artifact of laser microdissection, and suggested that perturbed expression of ASF1B was likely a transcriptional feature of triple negative breast cancers and early-onset human breast cancers.

ASF1B is expressed at significantly higher levels in triple negative tumor cells as compared to normal cells of the breast.

We obtained exact mRNA expression levels for ASF1B - from normal mammary ductal cells of the breast and from tumor cells of patients with triple negative breast cancer - to understand the magnitude and direction of ASF1B expression change in TNBC. ASF1B mRNA transcript was present at higher levels in triple negative tumor cells of the breast as compared to normal mammary ductal cells (Figure 1). Increased expression of ASF1B in primary tumor-isolated triple negative breast cancer cells was statistically significant (Figure 1: $p<0.0001$). ASF1B was expressed at 29.9 ± 10.0 arbitrary units (A.U.) in normal mammary ductal cells, while it was expressed at 90.1 ± 29.6 A.U. in tumor cells of patients with triple negative breast cancer. We calculated a mean fold change of 3.01 when comparing triple negative breast cancer cells to normal mammary ductal cells.

ASF1B expression correlates with survival outcomes in HER2+ human breast cancer.

We performed Kaplan-Meier survival analysis to evaluate potential relationships between primary tumor ASF1B mRNA expression levels and survival outcomes in patients with breast cancer, when analyzing outcomes based on the molecular subtype of the patient (PAM50 classification): luminal A, luminal B, HER2+, normal-like, and basal-like. We observed a statistically significant correlation between ASF1B expression and overall survival (OS) in patients with HER2+ subtype breast cancer (Figure 2; log rank p -value: 0.072 for overall survival, hazard ratio: 1.43 (0.97-2.1) (Fig. 2)).

ASF1B mRNA levels were a negative prognostic indicator in HER2+ breast cancer patients. Median OS was 55.2 months for HER2+ patients with low tumor expression of ASF1B while median OS was 48 months for HER2+ patients with high tumor expression of ASF1B (Chart 3).

ASF1B primary tumor expression was not correlated with overall survival in basal subtype (Figure 2; log rank p -value: 0.94 for OS, hazard ratio: 0.98 (0.67-1.44) (Fig. 2), luminal A subtype (Figure 2; log rank p -value: 0.73 for OS, hazard ratio: 0.93 (0.61-1.42) (Fig. 2)), in luminal B breast cancer (Figure 2; log rank p -value: 0.57 for OS, hazard ratio: 1.11 (0.78-1.57) (Fig. 2)), or in the normal-like subtype (Figure 2; log rank p -value: 0.9 for OS, hazard ratio: 1.06 (0.4-2.83) (Fig. 2)).

Influence of primary tumor ASF1B expression on survival outcomes based on TNBC molecular subtype.

We also performed Kaplan-Meier survival analysis to examine relationships between ASF1B mRNA expression levels with survival strictly in patients with triple negative breast cancer, analyzing

outcomes based on the TNBC molecular subtype⁸ of the patient: basal-like 1, basal-like 2, immunomodulatory, mesenchymal, mesenchymal stem-like, and luminal androgen receptor. We observed a correlation between ASF1B expression and overall survival (OS) in TNBC patients with basal-like 2 subtype disease which trended towards statistical significance (Figure 3; log rank *p*-value: 0.091 for overall survival, hazard ratio: 2.42 (0.84-6.96) (Fig. 3)). ASF1B mRNA levels were a negative prognostic indicator in patients with basal-like 2 TNBC. Median OS was 90 months for basal-like 2 TNBC patients with low tumor expression of ASF1B while median OS was 30 months for basal-like 2 TNBC patients with high tumor expression of ASF1B (Chart 4).

ASF1B primary tumor expression was not correlated with overall survival in basal-like 1 subtype (Figure 3; log rank *p*-value: 0.97 for OS, hazard ratio: 1.01 (0.47-2.19) (Fig. 3), immunomodulatory subtype (Figure 3; log rank *p*-value: 0.68 for OS, hazard ratio: 1.18 (0.53-2.61) (Fig. 3)), in the mesenchymal subtype (Figure 3; log rank *p*-value: 0.52 for OS, hazard ratio: 0.81 (0.42-1.56) (Fig. 3)), in the mesenchymal stem-like subtype (Figure 3; log rank *p*-value: 0.66 for OS, hazard ratio: 0.8 (0.29-2.23) (Fig. 3)), or in the luminal androgen receptor subtype (Figure 3; log rank *p*-value: 0.47 for OS, hazard ratio: 0.8 (0.44-1.46) (Fig. 3)).

Discussion

We performed comparative transcriptome analysis using two independently published microarray datasets^{2,3} to define and catalog transcriptional differences in human triple negative breast cancer. We provided evidence here that differential expression of anti-silencing function 1B histone chaperone, encoded by the ASF1B gene, is a defining transcriptional feature of triple negative breast cancer. ASF1B was also differentially expressed in the bulk tumors of patients with early-onset breast cancer. Expression of ASF1B was correlated with overall survival in patients with HER2+ subtype breast cancer, and within triple negative breast cancer patients, primary tumor expression of ASF1B was correlated with overall survival in patients with basal-like 2 subtype disease. Anti-silencing function 1B histone chaperone and the molecular processes to which it pertains to may be relevant to the initiation or progression of triple negative breast cancer in humans.

References

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Rank: 217
Probe ID: 18700
p-value: 3.58E-11
t: -8.672945
B: 15.3677
Gene: ASF1B
Gene name: anti-silencing function 1B histone chaperone

Chart 1: ASF1B is differentially expressed in triple negative 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: 321
Probe ID: 3852565
p-value: 5.89E-12
t: -8.9
B: 17.042783
Gene: ASF1B
Gene name: anti-silencing function 1B histone chaperone

Chart 2: ASF1B is differentially expressed in 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: ASF1B is expressed at significantly higher levels in primary TNBC tumor cells as compared to normal cells of the breast.

The mRNA expression level of ASF1B in normal mammary ductal cells (left) and in the isolated primary tumor cells of patients with triple negative breast cancer (right) is graphically depicted with the result of a statistical test evaluating significance of difference in ASF1B expression between normal cells of the breast and TNBC tumor cells, a *p*-value, listed above.

Figure 2: ASF1B expression correlates with overall survival in patients with HER2+ breast cancer, but not in patients with basal, luminal A, luminal B, or normal-like subtype cancer.

Depicted in this Kaplan-Meier plot is the probability of overall survival (OS) for $n=431$ patients with basal subtype 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 ASF1B 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 ASF1B 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.

1 Low ASF1B expression: 55.2 months

2 High ASF1B expression: 48 months

3 **Chart 3: Median overall survival is superior in HER2+ subtype breast cancer patients with low**
4 **primary tumor expression of ASF1B.**

5 The overall survival of $n=362$ HER2+ subtype breast cancer patients based on stratification into low or
6 high tumor expression of ASF1B is listed in this chart.

Figure 3: ASF1B expression correlates with overall survival in triple negative breast cancer patients whose tumors are of the basal-like 2 molecular subtype.

Depicted in this Kaplan-Meier plot is the probability of overall survival (OS) for $n=103$ patients with basal-like 1 subtype TNBC (upper left), $n=58$ patients with basal-like 2 subtype TNBC (upper right), $n=149$ patients with immunomodulatory subtype TNBC (middle left), $n=116$ patients with luminal androgen receptor subtype TNBC (middle right), $n=114$ patients with mesenchymal subtype TNBC (lower left), and $n=39$ patients with mesenchymal stem-like subtype TNBC (lower right), stratified into two groups, based on low or high expression of ASF1B in the primary tumors of patients with TNBC. 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 ASF1B 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.

1 Low ASF1B expression: 90 months

2 High ASF1B expression: 30 months

3 **Chart 4: In patients with basal-like 2 subtype disease, median overall survival is superior in triple**
4 **negative breast cancer patients with low primary tumor expression of ASF1B.**

5 The overall survival of $n=58$ basal-like 2 subtype TNBC patients based on stratification into low or high
6 tumor expression of ASF1B is listed in this chart.