DUSP26 is differentially expressed in the lymph node and brain metastases of patients with metastatic breast cancer.

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Metastasis to the brain is a clinical problem in patients with breast cancer¹⁻³. We mined published microarray data^{4,5} to compare primary and metastatic tumor transcriptomes to discover genes associated with brain metastasis in patients with metastatic breast cancer. We found that the dual specificity phosphatase 26, encoded by DUSP26, was among the genes whose expression was most different in the brain and lymph node metastases of patients with metastatic breast cancer as compared to primary tumors of the breast and normal breast tissues, respectively. DUSP26 mRNA was present at higher quantities in brain metastatic tissues as compared to primary tumors of the breast. These data combined suggest some level of common origin for metastases that reside in the lymph nodes and colonize the brain.

Keywords: breast cancer, metastasis, brain metastases, central nervous system metastases, lymph node metastasis, DUSP26, dual specificity phosphatase 26, systems biology of breast cancer, targeted therapeutics in breast cancer.

One report described a 34% incidence of central nervous system metastases in patients treated with trastuzumab for breast cancer². More recently, the NEfERT-T clinical trial⁶ which compared administration of either neratinib or trastuzumab in conjunction with paclitaxel demonstrated that in a randomized, controlled setting, in breast cancer patients treated with neratinib, not only was the incidence of central nervous system recurrence significantly lower, the time to central nervous system metastasis was significantly delayed as compared to patients administered trastuzumab. The alarmingly high rate of central nervous system metastasis described, as well as data, both anecdotal² and from a randomized, controlled setting⁶ illustrating that treatment with trastuzumab may be associated with these events demands an enhanced understanding of the transcriptional makeup of brain metastatic tissues to support identification of therapeutic targets, whether they are treatment related or not. We performed a global comparative analysis of primary and metastatic tumors in patients with brain metastatic breast cancer^{4,5}. We discovered significant differential and increased expression of DUSP26, in brain metastatic tissues of patients with metastatic breast cancer.

Methods

We used datasets GSE10893⁴ and GSE124648⁵ and for this global differential gene expression analysis of brain metastatic breast cancer in conjunction with GEO2R. GSE10893 was generated using Agilent-011521 Human 1A Microarray G4110A technology with $n=11$ primary breast tumors and $n=3$ brain metastases from patients with breast cancer; analysis performed using platform GPL885. GSE124648, a microarray dataset, was generated using Affymetrix Human Genome U133A Array technology with $n=10$ normal breast tissues and $n=44$ lymph node metastases from patients with breast cancer; platform GPL96 was used for this analysis. 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 DUSP26 gene expression was significantly between normal breast tissues and brain metastases in humans with breast cancer using a two-tailed, unpaired t-test with Welch's correction. We used PRISM for all statistical analyses of differential gene expression in human breast cancer (Version 8.4.0)(455).

Results

We performed global comparative transcriptome analysis of metastatic tumor tissues of patients with metastatic breast cancer using published microarray data^{4,5} to describe the transcriptional landscape of brain and lymph node metastasis in human breast cancer in an unbiased fashion and for the discovery of novel therapeutic targets.

DUSP26 is differentially expressed in the brain metastases of patients with brain metastatic breast cancer.

Through analysis of published microarray data⁴, we identified the dual specificity phosphatase 26, encoded by DUSP26, as a differentially expressed gene in the breast metastatic tissues of humans with breast cancer (Table 1). When sorting each of the genes expressed in brain metastases based on significance of difference as compared to primary tumors of the breast in patients with breast cancer, DUSP26 ranked 201 out of 22575 total transcripts (Table 1), equating to 99.1% differential expression. Differential expression of GRM8 in the brain metastases of patients with metastatic breast cancer was statistically significant (Table 1; $p=6.58E-04$).

DUSP26 is differentially expressed in the lymph node metastases of patients with metastatic breast cancer.

1 Next, we queried a second microarray dataset⁵ generated using lymph node metastases and
2 normal breast tissues from patients with breast cancer to determine whether we could validate differential
3 expression of DUSP26 in metastatic tissues of humans with breast cancer, and whether differential
4 expression of DUSP26 was specific to brain metastases or could be observed in metastases to other sites
5 in patients with metastatic breast cancer. We observed significant differential expression of DUSP26 in
6 lymph node metastases of patients with metastatic breast cancer. When sorting each of the genes
7 expressed in lymph node metastases based on significance of difference as compared to normal breast
8 tissues in patients with breast cancer, DUSP26 ranked 7445 out of 22283 total transcripts (Table 2),
9 equating to 66.6% differential expression. Differential expression of DUSP26 in the lymph node
10 metastases of patients with metastatic breast cancer was statistically significant (Table 2; $p=1.79E-04$).
11 Thus, differential expression of DUSP26 was conserved in the brain and lymph node metastases of
12 patients with metastatic breast cancer in two independent microarray datasets^{4,5}.

13 **DUSP26 is expressed at higher levels in the brain metastases of patients with brain metastatic** 14 **breast cancer.**

15 We obtained exact mRNA expression levels for DUSP26, in normal breast tissues and in brain
16 metastasis of patients with brain metastatic breast cancer to determine direction and statistical
17 significance of change in DUSP26 expression in brain metastatic tissues. DUSP26 was expressed at
18 higher levels in the brain metastases of patients with breast cancer as compared to primary tumors of the
19 breast, but this difference was not deemed statistically significant (Figure 1; $p=0.1617$).

20 Thus, by mining published microarray data^{4,5} in an unbiased and systematic fashion, we identified
21 the interleukin DUSP26 as among the genes whose expression was most different, transcriptome-wide, in
22 the brain and lymph node metastases of patients with breast cancer when compared to tissues of the
23 breast, transformed and benign; DUSP26 expression, as compared to the transformed breast, was higher
24 in metastasis to the brain.

25 **Discussion**

26 We provided evidence here that DUSP26 is among the genes whose expression is most different
27 in the brain and lymph node metastases of patients with brain metastatic brain cancer. Evaluation of the
28 effects of depletion of DUSP26 in mouse models of metastatic breast cancer on metastasis to the central
nervous system are merited. Differential expression of DUSP26 in metastasis to the lymph nodes and to
the brain suggests some level of common origin for metastases that reside in the lymph nodes,
successfully evade immune clearance, penetrate the central nervous system by unknown means and
subsequently colonize the brain.

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Rank	ID	p-value	t	B	Gene	Gene name
201	1598	6.58E-04	4.30	-0.18711	DUSP26	dual specificity phosphatase 26 (putative)

Table 1: DUSP26 is differentially expressed in brain metastases in patients with metastatic breast cancer when compared to primary tumors of the breast.

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

Rank	ID	p-value	t	B	Gene	Gene name
7445	219144_at	1.79E-04	-3.790624	-0.5014	DUSP26	dual specificity phosphatase 26 (putative)

Table 2: DUSP26 is differentially expressed in lymph node metastases of patients with metastatic breast cancer when compared to normal breast tissue.

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

Figure 1: DUSP26 is expressed at higher levels in the brain metastases of patients with brain metastatic breast cancer as compared to primary tumors of the breast.

The mRNA expression level of DUSP26 in primary tumors of the breast (left) and in the brain metastases of brain metastatic breast cancer patients (right) is graphically represented here with mean DUSP26 levels marked and the result of a statistical test evaluating difference in mRNA expression level between brain metastases and primary tumors of the breast, a *p*-value, listed above.