Genome-wide identification of *OTP* gene as a novel methylation marker of breast cancer

MYUNG SOON KIM^{1,4}, JINSUN LEE², TAEJEONG OH¹, YOUNGHO MOON¹, EILSUNG CHANG², KWANG SUN SEO³, BENJAMIN DOUGLAS HOEHN¹, SUNGWHAN AN¹ and JEUNG-HOON LEE⁴

¹Genomictree Inc., Daejeon 301-721; Departments of ²Surgery and ³Pathology, Chungnam National University College of Medicine, Daejeon 301-721; ⁴Department of Dermatology, School of Medicine, Chungnam National University, Daejeon 301-721, Republic of Korea

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Abstract. Aberrant DNA methylation occurs early and frequently in tumorigenesis. Identification of DNA methylation biomarkers is a field that provides potential for improving the clinical process of breast cancer diagnosis. We utilized a genome-wide technique, methylated DNA isolation assay (MeDIA), in combination with high-resolution CpG microarray analysis to identify hypermethylated genes in breast cancer. Among differentially methylated genes between tumor and adjacent normal tissues, 3 candidate genes (*LHX2*, *WT1* and *OTP*) were finally selected through a step-wise filtering process and examined for methylation status in normal tissues, primary tumor, and paired adjacent normal-appearing tissues from 39 breast cancer patients. Based on the calculated cut-off values, all genes showed significantly higher frequencies of aberrant hypermethylation in primary tumors (43.6% for *LHX2*, 89.7% for *WT1* and 100% for *OTP*, $P < 0.05$) while frequencies were intermediate in paired adjacent normal tissues and absent in normal tissues. On further analysis, the methylation level in primary tumors was not significantly correlated with clinicopathological features. Interestingly, DNA methylation of a novel gene *OTP* was detected in adjacent normal tissues even 6 cm away from primary tumors, suggesting that *OTP* methylation may qualify as a biomarker for the early detection of breast cancer. In conclusion, we successfully identified a novel gene *OTP* frequently methylated in breast cancer by genome-wide screening. Our results suggest that the *OTP* gene may

play a crucial role in breast carcinogenesis, although further clinical validation will be needed to evaluate the potential application of *OTP* in the early detection of breast cancer.

Introduction

Breast cancer is the most frequently diagnosed cancer in women with 1.15 million cases reported worldwide every year (1,2). Higher quality mammography and magnetic resonance imaging have increased the detection of breast cancer leading to a overall decline in mortality. These screening procedures, however, may also lead to over diagnosis and result in unnecessary treatments and procedures (<http://www.breastcancer.org/>). Therefore, development of new screening tools with high accuracy would be of great value.

It is now widely recognized that aberrant epigenetic modifications play a crucial role in altering gene expression and inducing tumor formation (3,4). In recent years much attention has been paid to epigenetic gene silencing. This loss of gene transcription is due to hypermethylation of CpG islands in the promoter region and is known to frequently occur in the early stages of cancer development. These findings have suggested that such aberrant events could be utilized as molecular diagnostic biomarkers for early detection or risk identification for cancers (5).

There have been several studies profiling gene methylation at a genomic level or using panels of known genes (6-11). These studies have attempted to define methylation sites in genes and determine their frequency of methylation in clinical samples, which may lead to their use in molecular diagnostic testing for early detection of breast cancer. However, these established cancer-associated genes do not yet have a sufficient usability to be considered efficient clinical diagnostic markers.

In the present study, we performed a CpG microarray analysis in conjunction with a proprietary method for enrichment of methylated DNA we developed, using optimally truncated methyl DNA binding domain (MBD2bt), named methylated DNA isolation assay (MeDIA) (12). We used this system to characterize frequently methylated genes in early stage primary tumors and matched normal appearing tissues from breast cancer patients. Through step-wise filtering and subsequent verification of methylation-positivity in clinical

Correspondence to: Dr Sungwhan An, Genomictree Inc., 829 Tamnip-dong Yuseong, Daejeon 305-501, Republic of Korea
E-mail: genomictreel@korea.com

Professor Jeung-Hoon Lee, Department of Dermatology, School of Medicine, Chungnam National University, 33 Munhwa-ro, Jung-gu, Daejeon 301-721, Republic of Korea
E-mail: jhoon@cnu.ac.kr

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samples using a pyrosequencing-methylation assay, we identified new candidate genes which may provide additive value to other potential methylation biomarkers for the early detection of breast cancer.

Materials and methods

Reagents. All chemical reagents used were purchased from Sigma-Aldrich (MA, USA) unless otherwise noted. Oligonucleotides were synthesized by Bioneer Co. (Daejeon, Republic of Korea).

Cells and clinical specimens. The human breast cancer cell lines, MCF-7 (HTB-22) and MDA-MB-231 (HTB-26), were obtained from the American Type Culture Collection and cultured in RPMI-1640 supplemented with 10% fetal bovine serum.

Fresh-frozen tissue specimens were obtained from the School of Medicine, ChungNam National University, Daejeon, Republic of Korea. All specimens and pertinent patient information were treated in accordance with the Institutional Review Board of the School of Medicine, ChungNam National University, Daejeon, Republic of Korea. Each tumor specimen was histologically verified by a board-certified pathologist and archived for further DNA study. Genomic DNA of normal tissues without any history of malignancy was purchased from BioChain, Inc. (CA, USA). Patient characteristics used in study are described in Table I.

Genome-wide methylation microarray analysis. CpG microarray analysis in conjunction with the enrichment of methylated DNA by methylated DNA isolation assay (MeDIA) using optimally truncated methyl DNA binding domain (MBD2bt) was performed as previously described with slight modification (12). The genomic DNA was isolated from samples taken from breast tumor blocks and their adjacent normal tissues of 3 patients (stage I) and the equal amount of DNA was mixed. Briefly, 0.5 μ g of sonic fragmented genomic DNA from primary tumor and their match-paired normal-appearing adjacent tissues was incubated with 2 μ g of recombinant MBD2bt protein for 4 h at 4°C with rocking platform, respectively. The enriched methylated DNA was amplified using whole genome amplification kit (GenomePlex®, Sigma, USA) as recommended by the manufacturer. The common reference DNA (CRD) was prepared by amplifying genomic DNA from normal placenta control without enrichment as an internal control. Enriched methylated DNA and CRD were labeled with Cy5 and Cy3, respectively. The labeled DNA samples were purified by PCR purification kit (Qiagen, USA) and then co-hybridized to human CpG island microarrays containing 237,000 oligonucleotide probes covering 27,800 CpG islands (Agilent, Santa Clara Technologies, Inc., CA, USA) according to the manufacturer's instructions.

Microarray data analysis. The hybridized images were analyzed by Agilent DNA microarray Scanner (Agilent Technologies, Inc.) and the data quantification was performed using Feature Extraction software version 9.3.2.1 (Agilent Technologies, Inc.). Preprocessing of raw data and normalization steps were performed using GeneSpring 7.3.1 (Agilent

Table I. Clinicopathological features of the 39 breast cancer patients.

Features	Patients (n=39)
Age	
≤50	21
>50	18
Stage	
I	17
II	19
III	3
Tumor size	
T1	17
T2	21
T3	1
Histological grade	
I	8
II	16
III	15
LN metastasis	
pN0	25
pN1	12
pN2	1
pN3	1
ER	
Negative	10
Positive	29
PR	
Negative	12
Positive	27
HR status	
Positive (ER ⁺ and/or PR ⁺)	8
Negative (ER ⁻ and PR ⁻)	31
HER-2/neu status	
Positive	6
Negative	33
Histological type	
Ductal	32
Lobular	1
Mixed ductal and lobular	1
Other	5

ER, estrogen receptor; PR, progesterone receptor; HR, hormone receptor.

Technologies, Inc.). The background-corrected intensity data were normalized using the intensity dependent LOWESS method to remove the dye bias within each array as recommended by the manufacturer (Agilent Technologies, Inc.). The relative methylation level of each probe between tumor and their adjacent normal tissues was calculated by comparison of ratio in each of enriched DNA/CRD. The multiple-probe

Table II. Primers for bisulfite PCR and pyrosequencing.

Genes	Primer (5'-3')	Assayed CpG sites ^a	Annealing T _m (°C)	Amplicon size (bp)
<i>OTP</i>	Forward: TATTYGGAAATTGGTTTTTA Reverse: Biotin-TCRATTTTAAAAAAGTTCTT Sequencing (F): TTATYGTATTTTTTAATGG	+2064, +2073, +2079, +2091, +2099	52	98
<i>LHX2</i>	Forward: TYGTTTAGAAATGGAAATGGGTT Reverse: Biotin-CCACAACCTCCCTCTATAACCTTAAC Sequencing (F): TTGGGGTGGGGGGAA	+2212, +2214, +2217, +2239	58	106
<i>WT1</i>	Forward: YGGTTTGATTYGGTAATTTAGTA Reverse: Biotin-AAAAACAAAACCTTCTC Sequencing (F): TTTTTTTTGGAGTAAATTAT	+1991, +2006, +2024	54	175

Y, C or T, R, A or G. ^aDistance (nt) from transcription start site (+1).

enriched genes were selected as methylation target genes when their probes showed positive call ≥ 2.0 -fold in methylation of tumor compared with paired adjacent normal tissues in at least two adjacent probes allowing a one-probe gap within the CpG islands.

The raw DNA microarray data were submitted for public access to the Gene Expression Omnibus (<http://www.ncbi.nlm.nih.gov/geo/>) and can be obtained using the accession number: GSE27374.

Methylation assessment by quantitative bisulfite-pyrosequencing. Bisulfite PCR and pyrosequencing primers were designed to amplify 3 to 5 CpG dinucleotide sites in the target regions of genes using PSQ Assay Design software (Biotage, Sweden). Sequences of primers used and PCR condition are given in Table II. Genomic DNA (200 ng) was modified by sodium bisulfate using the EZ DNA Methylation kit (Zymo Research, CA, USA) according to the manufacturer's instructions. Bisulfite-modified DNA was amplified in a 50 μ l reaction with the primer set and Taq polymerase (Enzymomics, Daejeon, South Korea). The PCR amplification was run for 45 cycles with an optimal annealing temperature. The PCR products were visualized on 2% agarose gel electrophoresis by ethidium bromide staining.

Pyrosequencing was performed using the Pyro Gold kit and PSQ 96MA instrument (Qiagen, Hilden, Germany) as instructed by the manufacturer. The methylation index (MtI) of each gene in each sample was calculated as the average value of $^mC/(^mC+C)$ for all examined CpGs in target regions. All experiments included samples without templates as negative control.

Statistical analysis. Statistical analysis was completed using the MedCalc software, version 9.3.2.0 (MedCalc, Mariakerke, Belgium). A P-value of <0.05 was considered as statistically significant. Wilcoxon Rank sum test was used to determine whether there were differences in methylation levels in tumors and adjacent normal tissues. The relationship between methylation level and clinicopathological features was analyzed by ANOVA test.

Results

Comprehensive methylation pattern analysis comparing primary breast tumors and matched-adjacent tissue. To comprehensively investigate CpG sites of genes which are aberrantly methylated early in breast cancer, we compared the DNA methylation status of pooled genomic DNA from three primary tumors with match-paired normal-appearing adjacent tissue (hereby, termed adjacent normal) from stage I breast cancer patients by MEDIA-CpG microarray analysis (12). Using this approach we observed that several hundred genes, including polycomb-target as a major group, demonstrated differential methylation between tumor and adjacent normal tissues. In order to reduce the number of genes and further verify methylation we adopted the following sequential filtering criteria: i) initial gene lists were chosen through an arbitrary selection process from probes in microarrays with signal intensities $\geq 2,000$ in tumor tissue; ii) methylation levels were compared in tumor tissues and paired adjacent normal tissue, selected genes had higher (≥ 2.0 -fold) levels of methylation in at least two adjacent probes allowing a one-probe gap within the CpG islands; and iii) genes not harboring CpG islands in the regulatory region were excluded (Fig. 1). From this, we obtained 6 candidate genes, *AXIN2*, *LHX2*, *NFIX*, *OTP*, *PTGER4* and *WT1* for methylation assessment (Table III).

Examination of methylation status of 6 candidates in cell lines and normal tissue. We confirmed the methylation status of the 6 candidate genes derived by microarray analysis using a pyrosequencing assay in breast cancer cell lines MCF7 and MDA-MB-231 (Fig. 2A). In this assay, sodium bisulfite-treated genomic DNA is amplified by PCR using gene specific primer sets with methylation-independent sequences. We demonstrated that all genes appear to have high level of methylation in both cell lines. We also examined the DNA methylation status in normal breast tissues from 6 healthy individuals. Three genes, *AXIN2*, *NFIX*, *PTGER4* showed a relatively higher range of methylation level (MtI $>33\%$) while the level of methylation of the remaining 3 genes, *LHX2*, *OTP* and *WT1* were either lower range (MtI range from 11% to 20.6%)

Figure 1. Schematic diagram of MeDIA-assisted high-resolution CpG microarray analysis in breast tissues. The methylated DNA was separately enriched from 3 tumors and paired adjacent normal tissues by MeDIA technique. The enriched methylated DNAs were amplified and subjected to CpG microarray analysis. Amplified methylated DNAs (Cy5) were individually compared with amplified-CRD (Cy3) without enrichment as an internal control. The relative methylation level of each probe across samples was calculated and 6 candidate genes were identified by step-wise filtering processes.

or unmethylated (Mtl <10%) (Fig. 2B). We then decide not to include *AXIN2*, *NFIX*, *PTGER4* genes which had Mtl >33% in normal breast tissue, from further clinical validation studies assuming that they would not be able to meet necessary specificity standards in future testing.

Clinical validation of *LHX2*, *OTP* and *WT1* genes in primary tumor and adjacent normal breast tissue. To verify whether *LHX2*, *OTP* and *WT1* genes are hypermethylated in primary breast tumors, we examined their methylation status in 39 primary breast tumors and matched-adjacent normal tissue from breast cancer patients representing stages I, II, and III as well as other clinicopathological parameters (Table I). Most of the samples from cancer patients showed a higher methylation level of each gene in primary tumors when compared to matched normal tissue (61.5% for *LHX2*, 92.3% for *WT1* and 100% for *OTP*).

We then used the highest methylation levels of each gene in normal samples as a cut-off value for methylation-positive calls. All genes showed a high frequency of methylation positivity in tumors (43.6% for *LHX2*, 89.7% for *WT1* and 100% for *OTP*). We also observed that aberrant methylation of each gene existed in tumor-adjacent normal tissue, although the levels and frequencies were intermediate between those of tumor and normal tissues (Fig. 3 and Table IV). In addition, we investigated the potential relationship between methylation levels of *LHX2*, *OTP* and *WT1* and clinicopathological parameters including patients' age, histological type, tumor size, histological grade, clinical stage, presence of lymph node

Figure 2. The quantitative pyrosequencing methylation assay of 6 candidate genes in 2 cell lines and 6 healthy normal tissues. Bisulfite-treated genomic DNA was amplified by specific primers. The Mtl is presented as the mean value with standard deviation (SD) for all examined CpGs in target region. (A) Methylation status of 6 candidate genes in 2 two breast cancer cell lines, MCF-7 and MDA-MB-231. Gene names are indicated at the bottom and left axis of the methylation level, respectively. (B) The methylation levels of 6 genes in healthy normal tissues. The left axis indicates methylation level, and gene names are indicated at the bottom.

metastases, steroid receptor (ER and PR) status and HER2 amplification. There were no significant correlations between methylation levels and any of the analyzed clinicopathological parameters (Table V).

Discussion

The number of genes showing aberrant methylation in breast cancer is increasing, although only a few genes have demonstrated promise as methylation markers for early detection and risk assessment (13,14). To expand these findings to novel genes in breast cancer, both gene-specific and genomic-level methylation profiling approaches have been performed (6-11).

From among those approaches, Pfeifer and colleagues focused on finding new genes which are frequently found to be methylated in early on-set breast cancer. To accomplish this they used CpG microarray analysis in conjunction with the methylated CpG island recovery assay (MIRA), which is based on the affinity of a two protein complex (MBD2/MBD3L1) attached to methylated DNA. Their data highlighted homeobox genes such as *THX1*, *HOXB13* and *HNF1B*

Table III. Candidate genes identified by MeDIA-assisted CpG microarray analysis.

Gene name	Description	Chr. location	Function
<i>AXIN2</i>	Axin 2 (conductin, axil)	17q23-q24	Multicellular organismal development
<i>LHX2</i>	LIM homeobox 2	9q33.3	Regulation of transcription, DNA-dependent
<i>NFIX</i>	Nuclear factor I/X (CCAAT-binding transcription factor)	19p13.3	DNA replication; regulation of transcription, DNA-dependent
<i>PTGER4</i>	Prostaglandin E receptor 4 (subtype EP4)	5p13.1	G-protein signaling
<i>OTP</i>	Orthopedia homeobox	5q13.3	Multicellular organismal development; regulation of transcription, DNA-dependent; induction of apoptosis
<i>WT1</i>	Wilms tumor 1c	11p13	Negative regulation of cell proliferation; epithelial cell differentiation; germ cell development

Table IV. Methylation levels and frequencies of hypermethylation of 3 genes in 39 pairs of breast tissues.

Gene	Methylation index (% , mean $\pm$ SD) ^a		Number and percentage of hypermethylation ^b		
	Paired tumor-adjacent tissues	Tumor tissues	Normal (%)	Paired tumor-adjacent tissues (%)	Tumor tissues (%)
<i>OTP</i>	27.0 $\pm$ 11.8	56.2 $\pm$ 16.3	0/6 (0.0)	34/39 (87.2)	39/39 (100)
<i>LHX2</i>	16.0 $\pm$ 8.8	22.7 $\pm$ 17.2	0/6 (0.0)	8/39 (20.5)	17/39 (43.6)
<i>WT1</i>	6.8 $\pm$ 3.3	17.3 $\pm$ 12.2	0/6 (0.0)	23/39 (59.0)	35/39 (89.7)

^aAverage methylation level with standard deviation of 39 tissue samples in each assay. ^bPercentage of hypermethylation: the percentage of methylation-positive samples that showed higher methylation level than cut-off value. Each cut-off value was determined as the highest Mtls of each gene in 6 normal samples (*OTP*, 16.6; *LHX2*, 20.6 and *WT1*, 5.9). P-values were calculated with the use of the Wilcoxon Rank sum test ($P \leq 0.05$).

Figure 3. Methylation status of *LHX2*, *OTP* and *WT1* in healthy normal and breast tissues from breast cancer patients. Multiple line/scatter plots of Mtls for *OTP*, *LHX2* and *WT1* were plotted from pyrosequencing analysis of 39 clinical tumor tissue samples (T) and 39 paired tumor-adjacent normal tissue samples (NT) and 6 healthy normal tissues (N). The samples from the same patient are linked with a straight line.

as new potential methylation biomarkers and demonstrated a high frequency of methylation in significant portion of breast cancers (10).

In the present study, we employed a similar approach to screen differentially methylated CpG sites in primary tumor

tissues and tumor-adjacent normal tissue of early stage breast cancer patients. However, we used an optimally truncated methylated DNA binding domain for the enrichment of methylated DNA (MBD2bt) instead of the MBD2/MBD3L1 complex (12). We found that hundreds of genes are aberrantly

Table V. The relationship between clinicopathological parameters and Mtls of 3 genes in breast cancer samples.

Features	Mtls (% , mean $\pm$ SD) ^a						Case
	<i>OTP</i>	P-value ^a	<i>LHX2</i>	P-value ^a	<i>WT1</i>	P-value ^a	
Age (mean, range)							
≤50	55.9±15.8	0.92	22.6±16.9	0.95	14.4±10.8	0.22	21
>50	56.4±17.4		22.9±18.0		20.7±13.2		18
Stage							
I	54.2±13.3	0.51	22.2±15.1	0.85	15.2±8.8	0.35	17
II-III	57.7±18.5		23.2±19.0		18.9±14.3		22
Tumor size							
T1	56.6±13.3	0.51	27.8±19.0	0.11	15.2±8.7		17
T2-3	57.7±18.5		18.9±14.9		18.9±14.4	0.35	22
Histological grade							
I	53.8±9.2	0.26	28.3±21.4	0.57	14.1±7.5	0.23	8
II	61.3±16.7		22.4±12.9		21.3±12.6		16
III	52.0±18.2		20.2±19.2		14.7±13.2		15
LN metastasis							
pN0	55.8±13.8	0.86	21.0±13.3	0.41	17.3±12.3	0.99	25
pN1-3	56.8±20.7		25.8±22.8		17.3±12.5		14
ER							
Negative	51.8±18.9	0.34	22.5±22.7	0.97	16.0±14.8	0.70	10
Positive	57.7±15.5		22.8±15.3		17.7±11.5		29
PR							
Negative	52.4±21.5	0.34	25.7±22.0	0.48	14.7±13.9	0.38	12
Positive	57.8±13.6		21.4±14.8		18.5±11.5		27
HR status							
Positive (ER ⁺ and/or PR ⁺)	52.3±20.4	0.47	27.1±23.3	0.43	17.1±16.6	0.96	8
Negative (ER ⁻ and PR ⁻)	57.1±15.4		21.6±15.5		17.3±11.2		31
HER-2/neu status							
Negative	56.1±16.6	0.94	23.2±17.6	0.73	16.9±10.7	0.67	33
Positive	56.6±16.4		20.5±15.6		19.3±19.8		6

^aP-values were calculated with the use of ANOVA test (P<0.05). ER, estrogen receptor; PR, progesterone receptor; HR, hormone receptor.

methyated in early primary breast tumors. These genes were composed of previously reported and novel genes in breast cancer. From the methylated target genes of Tommasi and colleagues (10), we found that 60/81 genes matched our data in breast cancer. This observation suggested that our strategy for selection of putative methylation biomarkers was sound. For clinical validation, we narrowed down the number of genes to three, *LHX2*, *WT1* and *OTP*, through step-wise selection procedure (described in the results section in this study) which demonstrated a higher level and frequency of methylation in these 3 genes in breast tumor tissue when compared to adjacent normal tissue.

Notably our finding of *LHX2* and *WT1*, are also present in the previous gene list (10), indicating that an affinity-based CpG methylation microarray with methylated DNA binding domain of MBD2t or MBD2/MBD3L1 complex is a powerful approach for global methylation mapping.

Considering that one of the ultimate goals of tumor specific molecular markers is to detect tumors non-invasively in an early stage of development, tumor-specific DNA methylation might be the appropriate candidate for this purpose since aberrant methylation at CpG islands occurs frequently and is an early epigenetic event in tumorigenesis (5,15,16). Conceptually a promising methylation marker with the purpose of early cancer detection is thought to be retaining the tendency of methylation positivity across all tumor types, and from early to advanced states regardless of clinicopathological status including stage. Some portion of cell-free DNA in serum or plasma is thought to have originated from tumor cells undergoing either apoptosis or necrosis in cancer patients (17-19). Accordingly, identification of tumor-specifically methylated sites of genes is a critical step in the sequential process of molecular marker development for early cancer detection in body fluids including blood derivatives from breast cancer

Figure 4. Methylation status of *OTP* gene in breast tumors and adjacent histologically normal tissues. Breast tissue was obtained from tumor and its surrounding zones 2, 4 and 6 cm from the grossly visible tumor boundary from patients who underwent mastectomy. The Mtl of each sample is depicted by gradient black scale (right). Each row and column represents individual patients (n=20) and Mtl from tumor (T) and adjacent normal tissue (2, 4 and 6 cm), respectively.

patients. Therefore, we postulate that the *OTP* gene would be a strong candidate marker for early detection considering its methylation frequency (sensitivity/specificity) and the early time of occurrence.

First, this gene showed methylation in 100% of our 39 primary tumor samples. Second, although the level of methylation is lower than that of matched-primary tumor tissue, there is a high frequency of methylation positivity that clearly appears in 87% of tumor-adjacent normal tissue when given the cut-off values are used (Fig. 3 and Table IV).

To further assess the extent of aberrant methylation of the *OTP* gene in adjacent normal tissues, we examined the methylation status of the *OTP* gene quantitatively in 20 cases of available adjacent normal tissues at various distances away from the tumor lesion boarder. Aberrant methylation of the *OTP* gene was extended up to 6 cm away from the primary tumor (Fig. 4). In addition, we could not find a significant correlation between *OTP* gene methylation status and various clinicopathological characteristics (Table V). This suggests that this gene may gain aberrant methylation at a very early stage and be an indicator of predisposition in precancerous regions due to a field methylation effect.

Most gene methylation is barely detectable in the serum or plasma of healthy individuals. Recently, several groups, using a panel of multiple genes which are well known to be tumor-associated, have demonstrated the prevalence of methylation positive genes in sera or plasma from breast cancer patients (20-27). Nevertheless, current multiple gene panel assays still lack the necessary sensitivity for application in the clinical setting.

To our knowledge, the *OTP* gene has never been reported to be linked to DNA methylation modification in breast cancer.

Taken together, our findings strengthen the possibility of *OTP* gene methylation as a biomarker candidate for early detection of breast cancer. The *OTP* gene has been described to be important for brain development (28,29). In addition, Wu and colleagues have recently reported that *OTP* is frequently methylated in astrocytomas (30). However, little is known about the biological function of this novel gene in cancer.

In summary, we demonstrated that microarray analysis using a minimum size of methylated DNA binding domain for methylated DNA enrichment successfully works for genomic level methylation profiling. With this approach we have identified the *OTP* gene which can provide an additive sensitivity to current set of candidate genes for early detection of breast cancer.

More sample validation and defining a specific region or regions of *OTP* that are methylated will be necessary to determine if the methylation status of this gene can be utilized as a biomarker for large scale, early detection of breast cancer.

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