Germline PARP4 mutations in patients with primary thyroid and breast cancers

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Abstract

Germline mutations in the PTEN gene, which cause Cowden syndrome, are known to be one of the genetic factors for primary thyroid and breast cancers; however, PTEN mutations are found in only a small subset of research participants with non-syndrome breast and thyroid cancers. In this study, we aimed to identify germline variants that may be related to genetic risk of primary thyroid and breast cancers. Genomic DNAs extracted from peripheral blood of 14 PTEN WT female research participants with primary thyroid and breast cancers were analyzed by whole-exome sequencing. Gene-based case-control association analysis using the information of 406 Europeans obtained from the 1000 Genomes Project database identified 34 genes possibly associated with the phenotype with \( P!1.0!10^{-3} \). Among them, rare variants in the PARP4 gene were detected at significant high frequency (odds ratio \( =5.2; P=1.0!10^{-5} \)). The variants, G496V and T1170I, were found in six of the 14 study participants (43%) while their frequencies were only 0.5% in controls. Functional analysis using HCC1143 cell line showed that knockdown of PARP4 with siRNA significantly enhanced the cell proliferation, compared with the cells transfected with siControl (\( P=0.02 \)). Kaplan–Meier analysis using Gene Expression Omnibus (GEO), European Genome-phenome Archive (EGA) and The Cancer Genome Atlas (TCGA) datasets showed poor relapse-free survival (\( P<0.001 \), Hazard ratio 1.27) and overall survival (\( P=0.006 \), Hazard ratio 1.41) in a PARP4 low-expression group, suggesting that PARP4 may function as a tumor suppressor. In conclusion, we identified PARP4 as a possible susceptibility gene of primary thyroid and breast cancer.

Key Words
thyroid
breast
cancer
PARP4
germline mutation

Introduction

A cancer of a different site and histologic type that develops in a person who already has a cancer diagnosis is considered a secondary primary cancer. Certain primary cancers are associated with a high risk of developing a second primary cancer. Specifically, individuals with a primary thyroid cancer have a high incidence of second
primary cancers even after adjusting for surveillance bias. One study of a cohort of 39,002 research participants suggested that when compared to the general population, thyroid cancer survivors have a 30% increased risk of a second primary breast cancer (Sandeep et al. 2006). The incidence of second primary cancers is also high even in participants with thyroid micro-carcinoma (Kim et al. 2013, Hsu et al. 2014). Because of the high incidence of thyroid cancer in women (Davies & Welch 2014, Siegel et al. 2015), it is important to understand why there is an increased risk of second primary breast cancers (Aschebrook-Kilfoy et al. 2013).

One possible cause for the increased rate of second primary breast cancers is a common genetic pathway. Cowden syndrome (CS) is an autosomal dominant disorder characterized by developing multiple primary cancers in various sites including thyroid and breast (Eng 1993). PTEN is one of the causative genes of CS or Cowden-like syndrome (CLS) (Liaw et al. 1997), and germline mutations are found in 25–85% of CS patients and <5% of CLS patients (Marsh et al. 1998, Tan et al. 2011). Germline mutations in other genes, such as succinate dehydrogenase genes (SDHx) (8%), PIK3CA (8.8%), AKT1 (2.2%), and SEC23B (4%) are also known as CS/CLS susceptibility genes (Ni et al. 2008, Ngeow et al. 2011, Orloff et al. 2013). CS/CLS is one of the plausible causes of primary thyroid and breast cancers, and the standardized incidence rate ratio of second primary cancers in research participants with thyroid cancer is increased to 5.83 per 10,000 person a year (95% CI = 3.01–10.18) by PTEN mutations (Ngeow et al. 2014). However, PTEN germline mutations are responsible for only a small portion of participants who present with non-syndromic primary thyroid and breast cancers (Pal et al. 2001, Ngeow et al. 2014). We hypothesize that germline mutations in genes other than PTEN, SDHx, PIK3CA, and AKT may explain the high rate of primary thyroid and breast cancer. In this study, we sought to identify possible germline mutations associated with thyroid and breast cancers using whole-exome sequencing.

Materials and methods

Samples

The study cohort consisted of 14 women known to have both primary breast and thyroid cancer malignancies. They were identified from a cohort of study participants who consented to participate in Cleveland Clinic IRB No.8458. These 14 research participants were selected under informed consent due to characteristics making them high risk for a possible genetic association. Particularly, they all developed both primary cancers prior to 50 years of age; none had radiation exposure to the head and neck or chest prior to developing the cancers, and there was a high frequency of familial breast and thyroid cancers in the cohort. In addition, all participants had already been ruled out for a PTEN germline mutation (Ngeow et al. 2014). Genomic DNA was extracted from peripheral blood samples, and the quality and quantity of each DNA sample were examined by Qubit 2.0 Fluorometer (LifeTechnologies, Carlsbad, CA, USA) and 2200 Tape Station (Agilent Technologies, Santa Clara, CA, USA). This study is approved by the Institutional Review Board at the University of Chicago (No. 8962) and Cleveland Clinic (No. 8458).

Whole-exome sequencing

DNA libraries for whole-exome sequencing were prepared from 200–1000 ng genomic DNA using SureSelectXT Human All Exon V5 (Agilent Technologies). The prepared whole-exome libraries were quantified on the 2200 Tape Station (Agilent Technologies), and then sequenced by 150 bp paired-end reads on NextSeq 500 Desktop Sequencer (Illumina, San Diego, CA, USA). All procedures were performed according to the manufacturers’ protocol.

Read mapping

After the exclusion of low-quality reads (base quality of <20 for more than 80% of bases) using FASTX toolkit, sequence reads were mapped to the human reference genome GRCh37/hg19 using Burrows–Wheeler Aligner (BWA) (v0.7.10) (Li 2014). After read pairs with a mapping quality of <30 and with mismatches more than 5% of read length were excluded, BAM files were generated using SAMTools (v1.1), and possible PCR duplicated reads were removed using Picard v1.91 (http://broadinstitute.github.io/picard/). As a population control, BAM files of 406 European (including Caucasian (CEU), British from England and Scotland (GBR), Iberian populations in Spain (IBS), and Tuscan in Italy (TSI), except for Finnish in Finland (FIN)) subjects, which were indicated to be healthy volunteers, were obtained from the 1000 Genomes Project database (www.1000genomes.org).

Variant calling

BCFtools (v1.1) was used to call variants (single nucleotide variants and indels) with the following criteria of i) sequence depth of ≥10, ii) variant depth of ≥4, and iii) base quality of
Variants were annotated using the hg19 database in single nucleotide polymorphism (SNP) effect prediction tools (SnpEff) (Cingolani et al. 2012). The primary SnpEff genomic effects include splice site acceptor, splice site donor, indel frameshift, indel non-frameshift, non-sense, non-synonymous, and synonymous variants. For variants that have multiple different annotations, the highest impact effect was selected. The predicted impact of amino acid substitutions was annotated using five algorithms of latest release time (LRT), MutationTaster, PolyPhen-2 HumDiv, PolyPhen-2 HumVar, and scale invariant feature transform (SIFT). Variants predicted as ‘deleterious’ in LRT, ‘disease causing automatic’ and ‘disease causing’ in MutationTaster, ‘probably damaging’ and ‘possibly damaging’ in PolyPhen-2 HumDiv and PolyPhen-2 HumVar, and ‘damaging’ in SIFT were considered as ‘deleterious’.

Sanger sequencing

Target regions were amplified by PCR from genomic DNA (Supplementary Table 1, see section on supplementary data given at the end of this article). The amplified products were sequenced using BigDye Terminator v3.1 Cycle Sequencing Kit and 3500 Series Genetic Analyzer (Life Technologies) according to the manufacturer’s protocols. Variants not validated by Sanger sequencing were excluded from the following analysis.

Association analysis

Gene-based association analysis was performed using a burden test with 100 000-time permutation in PLINK/SEQ v0.10. In case of multiple results due to alternative transcripts, we selected the result with the smaller \( P \) value. \( P \) values of \(<0.05\) were considered as statistically significant. To adjust multiple testing by the strict Bonferroni correction, we used significance level of \( P<3.65\times10^{-6}\) (0.05/13 705) in the gene-based association analysis.

Quantitative real-time PCR

Total RNAs from 18 cell lines previously extracted (Park et al. 2010) were reverse transcribed to cDNA using SuperScript III (LifeTechnologies). Real-time PCR using TaqMan gene expression assay for \( PARP4 \) (Hs00173105_m1) and \( GAPDH \) (Hs02758991_g1) was conducted on ViiA7 instrument (Life Technologies). mRNA levels of \( PARP4 \) were normalized to \( GAPDH \) expression.

siRNA transfection and cell proliferation assay

HCC1143 cell line authenticated by short tandem repeat analysis was cultured under the recommendations of their respective depositors. siRNA oligonucleotides for targeting \( PARP4 \) transcripts were purchased from Sigma–Aldrich. siNegative control (Cosmo Bio, Tokyo, Japan) was used as a control siRNA. siRNAs were transfected into cells with Lipofectamine RNAi max (Life Technologies). After 48-h incubation, cell viability assay was performed with the Cell Counting Kit-8 (Dojindo, Kumamoto, Japan).

Statistical analysis

Expression analysis and cell viability experiment were repeated three times at triplicates and evaluated using the Student’s \( t \) test. The prognostic value of \( PARP4 \) was analyzed by Kaplan–Meier plotter (http://kmplot.com/), which includes gene expression data of 4142 breast cancer cases from GEO, EGA, and TCGA using Affymetrix HG-U133A, HG-U133 Plus 2.0, and HG-U133A 2.0 (Gyorffy et al. 2010, 2013).

Results

The median age at diagnosis for thyroid and breast cancer was 38 (range: 23–49) and 46 (range: 36–49) years old respectively (Table 1). Five participants were diagnosed with both thyroid and breast cancer at the same age. Six cases were diagnosed to have thyroid cancer before breast cancer diagnosis. In total, 13 thyroid cancers were either follicular or papillary subtype, and 11 breast cancers were a ductal subtype (Table 1). Seven (50%) and five (36%) participants had a family history of thyroid or breast cancer within three generations of the proband respectively, although no additional family members had primary thyroid and breast cancers.

In exome sequencing of the 14 cases, we obtained a total sequencing output with an average read depth of 138\( \times \) per base. We also analyzed 406 European white controls obtained from the 1000 Genomes Project database using the same algorithm to minimize the calling discordance due to different algorithms. We identified a...
total of 105,064 and 345,793 variants in our cases and the 1000 Genomes controls respectively (Table 2). Of these variants, 4.8 and 18.8% were rare (defined as having a minor allele frequency (MAF) of <1% in the controls in both cases and controls). After excluding common SNPs with MAF of ≥1% in the controls, 5073 variants in 2909 genes in cases and 65,170 variants in 14,232 genes in controls were identified within the coding regions (Table 2). In the 14 participants, we inspected rare variants known to be responsible for hereditary cancer syndromes such as SDHx and KLLN (for CS), and BRCA1/2 (for hereditary breast and ovarian cancer syndrome) (Supplementary Figure 1); however, no participant had more than one of these rare variants identified. In addition, no statistically significant enrichment of variants was observed in the recurrently somatically mutated genes (known to occur in more than 10% of cancers) that have been previously reported in epithelial thyroid cancer (such as BRAF, RET, and TERT) and breast cancers (such as PIK3CA, TP53, and CDH1) (Supplementary Figure 2).

Using five different prediction tools, variants predicted as deleterious substitutions in at least one algorithm were identified in 2316 and 13,554 genes in cases and controls respectively (Table 2). Gene-based association analysis of these deleterious variants identified 34 genes, in which rare variants were enriched with a P value of <3.65×10⁻⁶. However, none of these genes showed a genome-wide significant level of association when we considered the Bonferroni correction for multiple testing (P<3.65×10⁻⁶) (Supplementary Figure 3). Since several

<table>
<thead>
<tr>
<th>ID no.</th>
<th>Race</th>
<th>Cancer type</th>
<th>Age of diagnosis</th>
<th>Family history of thyroid and breast cancer</th>
<th>Histology</th>
<th>Metastasis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>White</td>
<td>Thyroid</td>
<td>36</td>
<td>Papillary</td>
<td>Unknown</td>
<td>No</td>
</tr>
<tr>
<td>2</td>
<td>White</td>
<td>Breast</td>
<td>36</td>
<td>Medullary</td>
<td>Unknown</td>
<td>No</td>
</tr>
<tr>
<td>3</td>
<td>Hispanic</td>
<td>Thyroid</td>
<td>48</td>
<td>Papillary</td>
<td>Unknown</td>
<td>No</td>
</tr>
<tr>
<td>4</td>
<td>Hispanic</td>
<td>Breast</td>
<td>48</td>
<td>Follicular</td>
<td>Unknown</td>
<td>No</td>
</tr>
<tr>
<td>5</td>
<td>White</td>
<td>Thyroid</td>
<td>47</td>
<td>Follicular/papillary</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>6</td>
<td>White</td>
<td>Breast</td>
<td>47</td>
<td>Ductal</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>7</td>
<td>White</td>
<td>Thyroid</td>
<td>25</td>
<td>Ductal</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>8</td>
<td>White</td>
<td>Breast</td>
<td>46</td>
<td>Ductal/papillary</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>9</td>
<td>Unknown</td>
<td>Thyroid</td>
<td>38</td>
<td>Uncal (maternal, 51 years)</td>
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<td>No</td>
</tr>
<tr>
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<td>Breast</td>
<td>35</td>
<td>Ductal</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>11</td>
<td>Black</td>
<td>Thyroid</td>
<td>46</td>
<td>Papillary</td>
<td>Unknown</td>
<td>No</td>
</tr>
<tr>
<td>12</td>
<td>White</td>
<td>Breast</td>
<td>23</td>
<td>Papillary</td>
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</tr>
<tr>
<td>13</td>
<td>American</td>
<td>Thyroid</td>
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</tr>
<tr>
<td>14</td>
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<td>Breast</td>
<td>49</td>
<td>Ductal</td>
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</tr>
<tr>
<td></td>
<td></td>
<td>Thyroid</td>
<td>48</td>
<td>Ductal</td>
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<td>No</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Breast</td>
<td>44</td>
<td>Ductal</td>
<td>No</td>
<td>No</td>
</tr>
</tbody>
</table>

NA, not available.
previous reports indicated that the mutations in the genes involved in DNA repair pathways are genetic risk factors for several types of familial cancer, we focused on the DNA repair pathway and found that 13 of the 14 participants had rare variants in at least one DNA repair-related gene (Fig. 1 and Supplementary Table 2). Among them, PARP4 showed the most significant association with the risk for primary thyroid and breast cancers (odds ratio $Z 5.2; P = 1.0 \times 10^{-5}$; Fig. 1). In PARP4, two different variants, G496V and T1170I, were found in six of the 14 (43%) participants, and both of these were validated by Sanger sequencing (Fig. 2a). The majority of these were T1170I substitutions found in five of the 14 (36%) cases including three White, one American Indian, and one unknown race case. In contrast, this variant was found in only two of the 406 (0.5%) control individuals. Interestingly, among the 17 mutations found in 960 breast cancer cases in TCGA database, seven cases (41%) showed the two-hit mutation: a somatic mutation and loss of one PARP4 allele. Furthermore, one of these 17 somatic mutations, p.T1170I, is identical to one of germline mutations we detected (Supplementary Table 3). G496V substitution was uniquely found in the remaining one American Indian case and not found in any of the 406 controls (Fig. 2b).

Next, we examined whether PARP4 shows tumor-suppressive function (Fig. 3). mRNA expression of PARP4 in 18 breast cancer cell lines was examined by qPCR. Since HCC1143 cells showed the highest expression of PARP4...

### Table 2 Summary of number of cases, variants, and genes with rare variants

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Variants</th>
<th>Rare variants</th>
<th>Genes with rare variants</th>
<th>Genes with variants predicted as deleterious</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cases</td>
<td>14</td>
<td>105 064</td>
<td>5073</td>
<td>2909</td>
</tr>
<tr>
<td>Controls</td>
<td>406</td>
<td>345 793</td>
<td>65 170</td>
<td>14 232</td>
</tr>
<tr>
<td>aControls are 406 European individuals in the 1000 Genomes Project database.</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Figure 1
Association of the burden of rare variants in DNA repair-related genes with risk for primary thyroid and breast cancer.
among the 18 cell lines tested, we used HCC1143 cells for the further experiments. When PARP4 expression was knocked down by siRNA, the proliferation of HCC1143 cells was significantly enhanced threefold compared to the cells transfected with siControl ($P < 0.02$; Fig. 3c), suggesting that PARP4 works as a tumor suppressor.

**Discussion**

In this study, we comprehensively analyzed exome variants in 14 participants with primary thyroid and breast cancer, and identified PARP4 as a possible susceptibility gene candidate for primary thyroid and breast cancer. PARP4, also known as VPARP, is a family member of the poly (ADP-ribose) polymerase (PARP) (Kickhoefer et al. 1999). The PARP protein superfamily has 17 members and controls a wide array of cellular processes such as DNA repair, transcriptional regulation, and RNA interference (Gibson & Kraus 2012). So far, the function of a few PARPs including PARP1, PARP2, and PARP5 have been well-characterized (Gibson & Kraus 2012). PARP1 and PARP4 contain BRCA1 carboxy-terminal (BRCT) domain repeats, which are thought to bind phosphorylated DNA damage-sensing proteins recruiting PARPs to sites of DNA damage (Manke et al. 2003). Although little is known about the biological function of PARP4, it is suspected to be involved in the DNA repair pathway due to its BRCT domain. Many hereditary diseases responsible for synchronous cancers such as Lynch syndrome ($MLH1$, $MSH2$, $MSH6$, and $PMS2$ are responsible genes) and hereditary breast and ovarian cancer ($BRCA1$ and $BRCA2$) are known to be caused by germline mutations in genes involved in the DNA repair pathway (Lynch et al. 2009). Moreover, several DNA repair genes are reported to be mutated to a significant degree in both aggressive papillary thyroid carcinoma and breast cancers (Cancer Genome Atlas Network 2012, Cancer Genome Atlas Research Network 2014). As shown in

**Figure 2**

Mutations in PARP4 gene in 14 participants with synchronous thyroid and breast cancers and 406 control individuals. (a) Validation of two variants, G496V and T1170I, by Sanger sequencing. (b) Location of each mutation in PARP4. Closed and open circles represent the location of PARP4 mutations found in cases and controls respectively.
Fig. 3, we suggest that PARP4 might have a tumor suppressor function. In addition, Kaplan–Meier analysis using gene expression data from 4142 cases of breast cancer which was available on Kaplan–Meier plotter (http://kmplot.com/) showed worse relapse-free survival \( (P = 0.001, \text{Hazard ratio } 1.27) \) and overall survival \( (P = 0.006, \text{Hazard ratio } 1.41) \) in the \( \text{PARP4} \) low-expression group. These lines of evidence support the possibility that PARP4 plays a critical role in thyroid and breast tumorigenesis.

Radiation therapy is known to increase the risk of second primary cancers (Tucker et al. 1991). Generally, a radiation-induced second primary cancer arising at the radiation-associated site typically arises 10 years after the initial radiation exposure (Grantzau & Overgaard 2015). However, since both thyroid and breast cancers were diagnosed prior to any therapeutic radiation exposure, including radioactive iodine (RAI) our cohort does not reflect a link between radiation-induced secondary primary cancer risk and \( \text{PARP4} \) germline mutations.

In conclusion, through the whole-exome sequencing approach, we have implicated \( \text{PARP4} \) germline mutations as possible susceptibility factors for the risk of synchronous thyroid and breast cancers. It is obvious that our study has a limitation in its sample size; thus, further analysis is warranted with a larger number of samples to verify the results obtained in this study. Although we show evidence of the tumor suppressor function of PARP4 in breast cancer cells, further analysis is warranted to clarify the development of primary thyroid and breast cancer. Regardless, the high rate of the rare variant of \( \text{PARP4} \) found in our study when compared to the very low rate in the controls is compelling evidence of a possible new genetic syndrome that is responsible for both primary thyroid and breast cancer.
Supplementary data
This is linked to the online version of the paper at http://dx.doi.org/10.1530/ERC-15-0359.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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