FEN1 gene variants confer reduced risk of breast cancer in Chinese women: A case-control study

Shuai Lin1*, Meng Wang1*, Xinghan Liu1*, Ye Lu2, Zhuoqing Gong3, Yan Guo4, Pengtao Yang1, Tian Tian1, Cong Dai1, Yi Zheng1, Peng Xu1, Shanli Li1, Yuyao Zhu1, Zhijun Dai1

1Department of Oncology, Second Affiliated Hospital of Xi’an Jiaotong University, Xi’an 710004, China
2Department of Student Affairs, Second Affiliated Hospital of Xi’an Jiaotong University, Xi’an 710004, China
3Department of Health Science Center, Xi’an Jiaotong University, Xi’an 710061, China
4Key Laboratory of Biomedical Information Engineering of Ministry of Education, School of Life Science and Technology, Xi’an Jiaotong University, Xi’an 710049, China

*These authors have contributed equally to this work

Correspondence to: Zhijun Dai, email: dzj0911@126.com

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ABSTRACT

This study aimed to assess the associations of two common Flap endonuclease 1 (FEN1) polymorphisms (rs4246215 and rs174538) with breast cancer risk in northwest Chinese women. We conducted a case-control study with 560 breast cancer patients and 583 age-matched healthy controls from Northwest China. Odds ratios (ORs) and 95% confidence intervals (95% CIs) were used to estimate the associations. We found a significantly reduced risk of breast cancer associated with T allele of rs4246215 (allele model: OR 0.81, 95% CI 0.68–0.96; homozygote model: OR = 0.59, 95% CI = 0.40–0.87; recessive model: OR = 0.61, 95% CI = 0.42–0.89), especially in postmenopausal women (OR = 0.58, 95% CI = 0.35–0.97). Furthermore, the polymorphism showed a decreased association with larger tumor size (heterozygote model: OR = 0.63, 95% CI = 0.44–0.92; dominant model: OR = 0.63, 95% CI = 0.44–0.90). For rs174538, we did not find any difference in all genetic models. However, rs174538 was associated with lymph node metastasis (heterozygote model: OR = 0.57, 95% CI = 0.39–0.81; dominant model: OR = 0.61, 95% CI = 0.43–0.86) and estrogen receptor status (heterozygote model: OR = 1.50, 95% CI = 1.05–2.15; dominant model: OR = 1.42, 95% CI = 1.01–1.98). Haplotype analysis showed that T/rs4246215G/rs174538 haplotype was a protective factor of breast cancer (OR = 0.34, 95% CI = 0.14–0.81). Our results suggest that FEN1 polymorphisms may reduce the risk of breast cancer in Chinese women.

INTRODUCTION

Breast cancer (BC) is the most frequently diagnosed cancer and the leading cause of cancer death among females worldwide [1]. It was estimated that there were 268,600 new cases and 69,500 deaths in Chinese women in 2015 [2]. And, 1,685,210 new cancer cases and 595,690 cancer deaths are projected to occur in the United States in 2016 [1]. Many factors, such as family history, living habits, and emotional expression, are identified as potential risk factors for the development of BC [3, 4]. Recent studies also indicated that the risk of BC may be affected by genetic alterations, including genetic polymorphisms [5].

Genome stability is vital in the transmission of genetic information, and DNA molecule is vulnerable to factors that can induce DNA damage or block replication [6]. Flap structure-specific endonuclease 1 (FEN1) has been well characterized as a key factor in ensuring genomic stability and protecting tissues from tumorigenesis [7]. Human FEN1, located on chromosome 11q12, contains two exons and one intron. FEN1 is
essential for the long-patch pathway, which conducts a repair tract of at least two nucleotides. It plays an important role in the following processes: DNA repair by removing the 5'-flaps generated by Pol δ/e [8], primer removal during lagging-strand DNA synthesis and Okazaki fragment processing [9], and the promotion of apoptotic DNA fragmentation after apoptotic stimuli. It has been reported that FEN1 expression is related to the development and progression of various cancers [10]. Previous studies demonstrated that FEN1 overexpression is common in breast [11, 12], prostate [11], lung, and brain tumors [13], predicting FEN1 might be a marker of tumor progression for many types of tumors [14].

Previous studies have suggested that single nucleotide polymorphisms (SNPs) in FEN1 may confer susceptibility to cancer. Two SNPs, 4150G>T (rs4246215, in the gene 3'-untranslated region) and −69G>A (rs174538, in the gene promoter region), were identified after thoroughly resequencing the FEN1 locus in 30 Han Chinese healthy volunteers [15]. There have also been several studies that have investigated the associations between these two SNPs and cancer risk [15–18]. And, up till now, there was only one study performed by Lv's et al. referring to BC [18]. It was lack of repeated research to verifying the relationship between these two common SNPs of FEN1 and breast cancer. Therefore, we conducted a case-control study to investigate the associations of the FEN1 rs4246215 and rs174538 polymorphisms with BC risk in the Northwest Chinese population.

RESULTS

Characteristics of the study participants

Basic clinical characteristics of BC patients and the demographic characteristic of both the patients and healthy controls are presented in Table 1. There is no statistically significant difference between the two groups in age (P = 0.612) or the distribution of menopausal status (P = 0.716). However, the BMI was significantly different between BC patients and healthy controls (P = 0.038).

Association between FEN1 polymorphisms and BC risk

The genotype and allele frequencies of the FEN1 rs4246215 and rs174538 polymorphisms are shown in Table 2. The genotype frequencies of the two SNPs in controls both conformed to HWE (P = 0.253, 0.922 for rs4246215 and rs174538, respectively). Compared with the GG genotype, the TT genotype frequency of the rs4246215 polymorphism among patients was significantly different from that of controls (OR = 0.59, 95% CI = 0.42–0.89, corrected p = 0.018). These results suggested that the FEN1 rs4246215 polymorphism had a protective effect on BC. However, we did not observe any significant associations between the FEN1 rs174538 polymorphism and BC risk in any genetic model (shown in Table 2).

Stratified analysis of FEN1 polymorphisms and BC risk

Stratified analysis regarding the effect of rs4246215 and rs174538 polymorphisms on BC by menopausal status was displayed in Table 3. The results indicated that rs4246215 was associated with a decreased BC risk in premenopausal women (OR = 0.58, 95% CI = 0.35–0.97, P = 0.03). However, there was no association between rs174538 and BC risk in either premenopausal patients or postmenopausal patients. Considering the difference of BMI in cases and controls, we infer that BMI may be an important factor of BC. Then, we performed a subgroup analysis by body mass index (BMI), choosing the overweight standard 24 kg/m$^2$ as a cut-point. However, we did not observe any relationship between rs4246215 and breast cancer subjects with BMI $\geq$ 24 kg/m$^2$ (P = 0.10) or BMI $<$ 24 kg/m$^2$ (P = 0.26), as shown in Table 4. Similar results were also obtained between rs174538 and BC patients.

Association between FEN1 polymorphisms and clinical parameters of BC patients

To determine whether the FEN1 polymorphisms had an effect on the different clinical features of BC patients, we then analyzed the associations between the FEN1 polymorphisms and a series of clinic pathological parameters, including tumor size, lymph node metastasis, the status of the estrogen receptor (ER), progesterone receptor (PR), and human epidermal growth factor receptor 2 (Her-2). As shown in Table 5, we found that the mutational genotypes frequencies of the rs4246215 were significantly lower in patients with larger tumor size (> 2 cm) (heterozygote model: OR = 0.63, 95% CI = 0.44–0.92, P = 0.02; dominant model: OR = 0.63, 95% CI = 0.44–0.90, P = 0.01). However, no significant association was detected in other clinical parameters of BC patients.

As shown in Table 6, the same analyses were also performed for the clinical features in relation to the rs174538 polymorphism. We found that the variant genotypes of rs174538 showed a decreased association with lymph node metastasis (GA vs. GG: OR = 0.57, 95% CI = 0.39–0.81, P = 0.002; AA/GA vs. GG: OR = 0.61, 95% CI = 0.43–0.86, P = 0.005). Furthermore, the frequency of the variant genotypes of FEN1 rs174538
polymorphism was significantly higher in ER-positive patients (GA vs. GG: OR = 1.50, 95% CI = 1.05–2.15, \( P = 0.03 \); AA/GA vs. GG: OR = 1.42, 95% CI = 1.01–1.98, \( P = 0.04 \)).

### Haplotype analysis of FEN1 polymorphisms and BC risk

We further conducted haplotype analysis using the Phase 2.1 software to explore whether the interaction of rs4246215 and rs174538 SNPs affected BC risk. Compared with the G\_rs4246215\_G\_rs174538 haplotype, T\_rs4246215\_G\_rs174538 haplotype showed a decreased risk of breast cancer (OR = 0.34, 95% CI = 0.14–0.81, \( P = 0.01 \), shown in Table 7).

### DISCUSSION

FEN1 is a DNA replication/repair protein with pleiotropic functions. As an important tumor suppressor, FEN1 expression is related to the development of cancer and the progression of the disease [9, 10]. In several studies, FEN1 protein expression indicated that altered FEN1 expression might influence the therapeutic response [13]. Van Pel et al. demonstrated that FEN1 and the flap endonuclease inhibitors have potentially broad applicability in the treatment of cancer [19].

Transient transfection and luciferase assays showed that the rs174538 G>A SNP in FEN1 causes increased promoter activity, which is most likely to be due to a higher binding affinity of the G allele with some unknown transcriptional inhibitors. In addition, the rs4246215 G>T SNP is also associated with differential levels of FEN1 RNA expression [16]. Interestingly, the rs174538 polymorphism is significantly associated not only with cancer risk but also with FEN1 mRNA levels in normal tissues. Participants with the rs174538 AA genotype have been found to have significantly higher FEN1 mRNA levels than those with the rs174538 GG and GA genotypes [20].

In our study, we observed that variant genotypes of FEN1 rs4246215, but not rs174538, were associated...
Table 2: Genotype and allele frequencies of *FEN1* polymorphisms among the cases and controls and the associations with BC risk

| Model          | Genotype | Case(560) | Control(583) | OR (95% CI)† | P-value* |
|----------------|----------|-----------|--------------|--------------|----------|
| **rs4246215**  |          |           |              |              |          |
| **Codominant** | G/G      | 260 (46.4%) | 245 (42.0%)  | 1.00 (reference) | 0.026    |
|                | G/T      | 249 (44.5%) | 256 (43.9%)  | 0.92 (0.72-1.17) | 0.49     |
|                | T/T      | 51 (9.1%)   | 82 (14.1%)   | 0.59 (0.40-0.87) | 0.007    |
| **Dominant**   | G/G      | 260 (46.4%) | 265 (43.9%)  | 1.00 (reference) |          |
|                | G/T-T/T  | 300 (53.6%) | 338 (56.1%)  | 0.84 (0.66-1.06) | 0.13     |
| **Recessive**  | G/G-C/T  | 509 (90.9%) | 501 (85.9%)  | 1.00 (reference) |          |
|                | T/T      | 51 (9.1%)   | 82 (14.1%)   | 0.61 (0.42-0.89) | 0.009    |
| **Overdominant** | G/G-C/T | 311 (55.5%) | 327 (56.1%)  | 1.00 (reference) |          |
|                | G/T      | 249 (44.5%) | 256 (43.9%)  | 1.02 (0.81-1.29) | 0.85     |
| **Allele**     | G        | 769 (68.7%) | 746 (64.0%)  | 1.00 (reference) |          |
|                | T        | 351 (31.3%) | 420 (36.0%)  | 0.81 (0.68-0.96) | 0.018    |

* Two-sided χ² test for the distributions of genotype and allele frequencies.
† Adjusted for age and body mass index.

Table 3: Stratified analysis by menopause status on *FEN1* polymorphisms and BC risk

| Genotypes | Case (N= 560) | Control (N= 583) | P * | OR (95% CI)† |
|-----------|---------------|------------------|-----|--------------|
| G/G+G/T   |               |                  |     |              |
| N (%)     | N (%)         |                  |     |              |
| Pre-menopause |               |                  |     |              |
| G/G+G/T   | 187 (89.9%)   | 271 (87.2%)      | 1.00 (reference) |          |
| TT        | 21 (10.1%)    | 45 (12.8%)       | 0.16 | 0.68 (0.39-1.17) |
| Post-menopause |            |                  |     |              |
| G/G+G/T   | 322 (91.5%)   | 230 (84.7%)      | 1.00 (reference) |          |
| TT        | 30 (8.5%)     | 37 (15.3%)       | 0.03 | 0.58 (0.35-0.97) |

* Two-sided χ² test for the distributions of genotype frequencies.
Table 4: Stratified analysis by body mass index (BMI) on FEN1 polymorphisms and BC risk

| Genotypes | BMI<24 | | | P* | OR (95%CI) | | Genotypes | BMI<24 | | | P* | OR (95%CI) |
|-----------|--------|--------|--------|------|-------------|--------|-----------|--------|--------|--------|-------------|--------|-----------|
| G/G+G/T   | 479    | 466    | 1.00   | reference | 1.51 | (0.95-2.41) | G/G+G/A | 471    | 465    | 1.00   | reference | 1.11  | (0.71-1.73) |
| TT        | 32     | 47     | 0.10   | | 1.51 | (0.95-2.41) | AA      | 41     | 45     | 0.64   | | 1.11  | (0.71-1.73) |
| BMI≥24    |        |        |        |        |        |        |        |        |        |        |        |        |        |
| G/G+G/T   | 30     | 35     | 1.00   | reference | 1.51 | (0.95-2.41) | G/G+G/A | 28     | 45     | 1.00   | reference | 1.11  | (0.71-1.73) |
| TT        | 19     | 35     | 0.26   | | 1.58 | (0.75-3.31) | AA      | 20     | 33     | 0.94   | | 1.03  | (0.50-2.13) |

* Two-sided χ² test for the distributions of genotype frequencies.
BMI ≥24 is the judgement standard for overweight.

Table 5: The associations between the FEN1 rs4246215 polymorphism and clinical characteristics of BC patients

| Variables   | GG (%) | TT (%) | P* | OR (95%CI) | GT (%) | P* | OR (95%CI) | GT+TT (%) | P* | OR (95%CI) |
|-------------|--------|--------|----|-------------|--------|----|-------------|-----------|----|-------------|
| Tumor size  |        |        |    |             |        |    |             |           |    |             |
| <2 cm       | 73 (38.8) | 20 (10.6) | 1.00 | (reference) | 95 (50.5) | 1.00 | (reference) | 115 (61.2) | 1.00 | (reference) |
| ≥2 cm       | 187 (50.3) | 31 (8.3) | 0.11 | 0.61 (0.32-1.13) | 154 (41.4) | **0.02** | 0.63 (0.44-0.92) | 185 (49.7) | **0.01** | 0.63 (0.44-0.90) |
| LN metastasis |        |        |    |             |        |    |             |           |    |             |
| Negative    | 110 (46.6) | 23 (9.7) | 1.00 | (reference) | 103 (43.6) | 1.00 | (reference) | 126 (53.4) | 1.00 | (reference) |
| Positive    | 150 (46.3) | 28 (8.6) | 0.71 | 0.89 (0.49-1.63) | 146 (45.1) | 0.83 | 1.04 (0.73-1.48) | 174 (53.7) | 0.94 | 1.01 (0.72-1.42) |
| ER          |        |        |    |             |        |    |             |           |    |             |
| Negative    | 111 (44.9) | 26 (10.5) | 1.00 | (reference) | 110 (44.5) | 1.00 | (reference) | 136 (55.1) | 1.00 | (reference) |
| Positive    | 149 (47.6) | 25 (8.0) | 0.28 | 0.72 (0.39-1.31) | 139 (44.4) | 0.74 | 0.94 (0.66-1.34) | 164 (52.4) | 0.53 | 0.90 (0.64-1.26) |
| PR          |        |        |    |             |        |    |             |           |    |             |
| Negative    | 124 (48.6) | 25 (9.8) | 1.00 | (reference) | 106 (41.6) | 1.00 | (reference) | 131 (51.4) | 1.00 | (reference) |
| Positive    | 136 (44.6) | 26 (8.5) | 0.86 | 0.95 (0.52-1.73) | 143 (46.9) | 0.98 | 1.00 (0.70-1.41) | 169 (55.4) | 0.34 | 1.18 (0.84-1.64) |
| HER-2       |        |        |    |             |        |    |             |           |    |             |
| Negative    | 185 (47.5) | 33 (8.5) | 1.00 | (reference) | 171 (44.0) | 1.00 | (reference) | 204 (52.4) | 1.00 | (reference) |
| Positive    | 75 (43.9) | 18 (10.5) | 0.36 | 1.35 (0.71-2.54) | 78 (45.6) | 0.54 | 1.13 (0.77-1.64) | 96 (56.1) | 0.42 | 1.16 (0.81-1.67) |

* Two-sided χ² test for the distributions of genotype frequencies.
LN: Axillary lymph node; ER: Estrogen receptor; PR: Progesterone receptor; HER-2: human epidermal growth factor receptor 2.
Table 6: The associations between the FEN1 rs174538 polymorphism and clinical characteristics of BC patients

| Variables     | GG (%) | TT (%) | P* | OR (95%CI) | GT (%) | P* | OR (95%CI) | GT+TT (%) | P* | OR (95%CI) |
|---------------|--------|--------|----|------------|--------|----|------------|-----------|----|------------|
| **Tumor size**|        |        |    |            |        |    |            |           |    |            |
| <2 cm         | 72 (38.3) | 20 (10.6) | 1.00 (reference) | 96 (51.1) | 1.00 (reference) | 116 (61.7) | 1.00 (reference) |
| ≥2 cm         | 171 (46.0) | 41 (11.0) | 0.63 | 0.86 (0.47-1.57) | 160 (43.0) | 0.22 | 0.79 (0.55-1.15) | 201 (54.0) | 0.08 | 0.73 (0.51-1.04) |
| **LN metastasis**|      |        |    |            |        |    |            |           |    |            |
| Negative      | 86 (36.4) | 24 (10.2) | 1.00 (reference) | 126 (53.4) | 1.00 (reference) | 150 (63.6) | 1.00 (reference) |
| Positive      | 157 (48.5) | 37 (11.4) | 0.57 | 0.84 (0.47-1.50) | 130 (40.1) | 0.002 | 0.57 (0.39-0.81) | 167 (51.5) | 0.005 | 0.61 (0.43-0.86) |
| **ER**        |        |        |    |            |        |    |            |           |    |            |
| Negative      | 119 (48.2) | 23 (9.3) | 1.00 (reference) | 105 (42.5) | 1.00 (reference) | 128 (51.8) | 1.00 (reference) |
| Positive      | 114 (37.6) | 38 (12.5) | 0.06 | 1.72 (0.97-3.07) | 151 (49.8) | 0.03 | 1.50 (1.05-2.15) | 189 (62.3) | 0.04 | 1.42 (1.01-1.98) |
| **PR**        |        |        |    |            |        |    |            |           |    |            |
| Negative      | 120 (47.1) | 22 (8.6) | 1.00 (reference) | 113 (44.3) | 1.00 (reference) | 135 (52.9) | 1.00 (reference) |
| Positive      | 123 (40.3) | 39 (12.8) | 0.06 | 1.73 (0.97-3.09) | 143 (46.9) | 0.24 | 1.23 (0.87-1.76) | 182 (59.7) | 0.11 | 1.32 (0.94-1.84) |
| **HER-2**     |        |        |    |            |        |    |            |           |    |            |
| Negative      | 167 (42.9) | 43 (11.1) | 1.00 (reference) | 179 (46.0) | 1.00 (reference) | 222 (57.1) | 1.00 (reference) |
| Positive      | 76 (44.5) | 18 (10.5) | 0.79 | 0.92 (0.50-1.70) | 77 (45.0) | 0.77 | 0.95 (0.65-1.38) | 95 (55.5) | 0.74 | 0.94 (0.65-1.35) |

* Two-sided χ² test for the distributions of genotype frequencies.

LN: Axillary lymph node; ER: Estrogen receptor; PR: Progesterone receptor; HER-2: human epidermal growth factor receptor 2.

Table 7: The haplotype frequencies of FEN1 polymorphisms and breast cancer risk

| Haplotypes | Cases (N=1120) n, % | Controls (N=1164) n, % | OR (95% CI) | p |
|------------|---------------------|------------------------|-------------|---|
| rs4246215 G | G | 735 (65.6%) | 718 (61.7%) | 1.00 (reference) |
| rs174538 G | A | 34 (3.0%) | 28 (2.4%) | 1.19 (0.71-1.98) | 0.51 |
| T | G | 7 (0.6%) | 20 (1.7%) | 0.34 (0.14-0.81) | 0.01 |
| T | A | 344 (30.7%) | 400 (34.4%) | 0.84 (0.70-1.00) | 0.05 |
with decreased BC risk. Our results are partly consistent with those of some other studies [15–18]. We searched PubMed, the ISI Web of Knowledge, Embase and China National Knowledge Infrastructure about the genome-wide association study of FEN1, only find one study in east China (Huaian, Jiangsu Province and Jinan, Shandong Province) about the association between the FEN1 and breast cancer risk. The authors indicated that rs174538GG and rs4246215GG genotypes were significantly correlated to increased risk for developing breast cancer compared with the mutation homozygote [18], which were in accordance with ours. Though the samples of our study were smaller than theirs, we conducted more details subgroup analyses, which may offer evidences for clinical treatment and prognosis evaluation of breast cancer.

We also observed that the variant rs4246215 genotypes in the FEN1 gene demonstrated a decreased association with tumor size. Furthermore, the variant genotypes of rs174538 were associated with negative lymph node metastasis and ER-positive status. These results suggested that the two polymorphisms in the FEN1 gene are related to the development and progression of BC, and may help to accurately predict the clinical course of BC. In addition, the rs4246215 polymorphism showed an increased association with postmenopausal status. We observed that patients with cancer have lower BMI in this study, it may be inferred that patients with BC lost weight after the onset of the cancer. We conducted the stratified analysis by BMI, but we did not observe any association between BMI and the two polymorphism.

Moreover, we did haplotype anaysis and find the T
rs4246215G
rs174538
haplotype may decrease the breast cancer risk. But, in Lv’s et al. [18] study, they showed that G
rs4246215A
rs174538
, T
rs4246215A
rs174538
, and G
rs4246215G
rs174538
haplotypes were associated with increased risks of developing breast cancer compared with the T
rs4246215G
rs174538
haplotype. This was broadly consistent with our study. But, we did not find G
rs4246215A
rs174538
and T
rs4246215A
rs174538
haplotypes had any associations with breast cancer. The differences may be attributed to the geographical and life style differences in northwest and east Chinese women. Thus, more large well-designed repeated studies are needed to evaluate the results.

Our study had some limitations. First, all people recruited in this study were from one hospital in the northwest China. Larger sample size and multi center experiment are needed for further verification. Second, more predisposing factors should be investigated, such as high-dose radiation exposure, and postmenopausal obesity. Third, we hypothesized that the two polymorphisms may enhance the expression of FEN1 gene via up-regulating the mRNA, thus strengthen the DNA repair ability of the injured breast cells. But, further experiments on cell or animal level are needed to explain the specific mechanisms.

In summary, our case-control study indicates that FEN1 polymorphisms have effects in reducing the BC risk in northwest Chinese women. Further functional studies and large population-based prospective studies are still required to elucidate the influence of FEN1 polymorphisms on BC.

MATERIALS AND METHODS

Ethics statement

The Institutional Review Board of the Xi’an Jiaotong University (Xi’an, China) approved the study. At the time of recruitment, written informed consent was obtained from all participants involved in the study.

Study population

A total of 560 sporadic BC patients were recruited between January 2013 and October 2014 at the Second Affiliated Hospital of Xi’an Jiaotong University, China. The patients were recruited without age restrictions. All of the patients had pathologically confirmed. Patients were excluded if they had received chemotherapy or radiotherapy before surgery (guaranteeing the accuracy of tumor information we collected) or had other types of cancer. For comparison, 583 cancer-free controls were randomly selected from participants who were seeking health care in the outpatient departments at the hospital and were frequency-matched to the patients with age (± 5 years). All participants were interviewed with a self-administered questionnaire after obtaining the written informed consent.

Genotyping assay

Peripheral venous blood sample of about 2 mL was collected into tubes containing ethylene diamine tetraacetic acid from each participant and then stored at −80 °C for later use. The DNA concentration was measured by spectrophotometry (DU 530 UV/VIS spectrophotometer, Beckman Coulter, Inc., Fullerton, CA, USA), as described in our previous studies [21–24]. Two tag SNPs (rs4246215 and rs174538), which captured the majority of the known common variation in FEN1 according to the data on the Chinese population from HapMap (http://www.hapmap.org), were selected in our study. Sequenom MassARRAY Assay Design 3.0 Software (Agena Bioscience, San Diego, CA, USA) was used to design a multiplexed SNP MassEXTEND assay [25]. SNP genotyping was performed using the Sequenom MassARRAY RS1000 according to the standard protocol recommended by the manufacturer. The corresponding primers used for each SNP in our study are listed in Table 8. Sequenom Typer 3.0 Software was used for data analyses [25, 26].
Statistical analyses

The Student t-test or the $\chi^2$ test was used to examine the differences in the distributions of demographic characteristics, selected variables, and genotypes distributions of the two SNPs between the patients and controls. Hardy-Weinberg equilibrium (HWE) was tested by chi-square test for each SNP before the analysis. We conducted a case-control study for all of the subjects with adjustment for age and body mass index (BMI), and then performed the stratified analyses by menopausal status and BMI. The associations between FEN1 rs4246215 and rs174538 polymorphisms and breast cancer susceptibility were estimated by odds ratios (ORs) and 95% confidence intervals (CIs). Corresponding primers used for each SNP in our study are listed in Table 8. Five genetic models were used in our study, namely allele model, co-dominant model (contains of homozygote model and heterozygote model), recessive model, dominant model, and over-dominant model. All statistical analyses were performed with SPSS 18.0 for Windows software (PASW Statistics, SPSS Inc., Chicago, IL, USA). A $P$-value < 0.05 was considered as the criterion of statistical significance, and all statistical tests were two-sided. Phase2.1 software (download from http://stephenslab.uchicago.edu/phase/download.html) was used to conduct all common haplotypes and $\chi^2$ test was used to estimate the ORs and 95% CIs for each haplotype.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest.

REFERENCES

1. Siegel RL, Miller KD, Jemal A. Cancer statistics, 2016. CA Cancer J Clin. 2016; 66:7-30.
2. Chen W, Zheng R, Baade PD, Zhang S, Zeng H, Bray F, Jemal A, Yu XQ, He J. Cancer statistics in China, 2015. CA Cancer J Clin. 2016; 66:115-132.
3. Wolff MS, Weston A. Breast cancer risk and environmental exposures. Environ Health Perspect. 1997; 105:891-896.
4. Goldblatt H, Cohen M, Azaiza F. Expression of emotions related to the experience of cancer in younger and older Arab breast cancer survivors. Etnh Health. 2016:1-14.
5. Lichtenstein P, Holm NV, Verkasalo PK, Iliadou A, Kaprio J, Koskenvuo M, Pukkala E, Skytthe A, Hemminki K. Environmental and heritable factors in the causation of cancer--analyses of cohorts of twins from Sweden, Denmark, and Finland. N Engl J Med. 2000; 343:78-85.
6. Dai ZJ, Liu XH, Ma YF, Kang HF, Jin TB, Dai ZM, Guan HT, Wang M, Liu K, Dai C, Yang WX, Wang XJ. Association Between Single Nucleotide Polymorphisms in DNA Polymerase Kappa Gene and Breast Cancer Risk in Chinese Han Population: A STROBE-Compliant Observational Study. Medicine. 2016; 95:e2466.
7. Lieber MR. The FEN-1 family of structure-specific nucleases in eukaryotic DNA replication, recombination and repair. Bioessays. 1997; 19:233-240.
8. Klungland A, Lindahl T. Second pathway for completion of human DNA base excision-repair: reconstitution with purified proteins and requirement for DNase IV (FEN1). EMBO J. 1997; 16:3341-3348.
9. Hennke G, Friedrich-Heineken E, Hubscher U. Flap endonuclease 1: a novel tumour suppressor protein. Trends Biochem Sci. 2003; 28:384-390.
10. Zheng L, Dai H, Zhou M, Li M, Singh P, Qiu J, Tsark W, Huang Q, Kernstine K, Zhang X, Lin D, Shen B. Fen1 mutations result in autoimmunity, chronic inflammation and cancers. Nature medicine. 2007; 13:812-819.
11. Lam JS, Seligson DB, Yu H, Li A, Eeva M, Pantuck AJ, Zeng G, Horvath S, Belldegrun AS. Flap endonuclease 1 is overexpressed in prostate cancer and is associated with a high Gleason score. BJU Int. 2006; 98:445-451.
of flap endonuclease 1 gene in breast and other cancers. Mol Cancer Res. 2008; 6:1710-1717.

13. Nikolova T, Christmann M, Kaina B. FEN1 is overexpressed in testis, lung and brain tumors. Anticancer Res. 2009; 29:2453-2459.

14. Zheng L, Jia J, Finger LD, Guo Z, Zer C, Shen B. Functional regulation of FEN1 nuclease and its link to cancer. Nucleic Acids Res. 2011; 39:781-794.

15. Yang M, Guo H, Wu C, He Y, Yu D, Zhou L, Wang F, Xu J, Tan W, Wang G, Shen B, Yuan J, Wu T, Lin D. Functional FEN1 polymorphisms are associated with DNA damage levels and lung cancer risk. Human mutation. 2009; 30:1320-1328.

16. Liu L, Zhou C, Zhou L, Peng L, Li D, Zhang X, Zhou M, Kuang P, Yuan Q, Song X, Yang M. Functional FEN1 genetic variants contribute to risk of hepatocellular carcinoma, esophageal cancer, gastric cancer and colorectal cancer. Carcinogenesis. 2012; 33:119-123.

17. Chen YD, Zhang X, Qiu XG, Li J, Yuan Q, Jiang T, Yang M. Functional FEN1 genetic variants and haplotypes are associated with glioma risk. J Neurooncol. 2013; 111:145-151.

18. Lv Z, Liu W, Li D, Liu L, Wei J, Zhang J, Ge Y, Wang Z, Chen H, Zhou C, Yuan Q, Zhou L, Yang M. Association of functional FEN1 genetic variants and haplotypes and breast cancer risk. Gene. 2014; 538:42-45.

19. van Pel DM, Barrett IJ, Shimizu Y, Sajesh BV, Guppy BJ, Pfeifer T, McManus KJ, Hieter P. An evolutionarily conserved synthetic lethal interaction network identifies FEN1 as a broad-spectrum target for anticancer therapeutic development. PLoS genetics. 2013; 9:e1003254.

20. Gao XR, Zhang SL, Yang YF, Han GR. FEN1 -69G>A and 4150G>T polymorphisms and cancer risk in Chinese population. Sci Rep. 2014; 4:6183.

21. Wang Z, Liu X, Wang X, Chong T, Lin S, Wang M, Ma X, Liu K, Xu P, Feng Y, Dai Z. Polymorphisms in TIM-3 and breast cancer susceptibility in Chinese women: A case-control study. Oncotarget. 2016; 7:43703-43712. doi: 10.18632/oncotarget.9665.

22. Ren HT, Li YM, Wang XJ, Kang HF, Jin TB, Ma XB, Liu XH, Wang M, Liu K, Xu P, Yao QL, Dai ZJ. PD-1 rs2227982 Polymorphism Is Associated With the Decreased Risk of Breast Cancer in Northwest Chinese Women: A Hospital-Based Observational Study. Medicine. 2016; 95:e3760.

23. Liu X, Wang X, Fu SW, Wang M, Kang H, Guan H, Zhang S, Ma X, Lin S, Liu K, Feng Y, Dai C, Dai Z. Genetic association of deleted in colorectal carcinoma variants with breast cancer risk: A case-control study. Oncotarget. 2016; 7:32765-32773. doi: 10.18632/oncotarget.9024.

24. Wang M, Wang X, Fu SW, Liu X, Jin T, Kang H, Ma X, Lin S, Guan H, Zhang S, Liu K, Dai C, Zhu Y, Dai Z. Single-nucleotide polymorphisms in PSCA and the risk of breast cancer in a Chinese population. Oncotarget. 2016; 7:27665-27675. doi: 10.18632/oncotarget.8491.

25. Jin TB, Li XL, Yang H, Jiri M, Shi XG, Yuan DY, Kang LL, Li SQ. Association of polymorphisms in FLT3, EGFR, ALOX5, and NEIL3 with glioblastoma in the Han Chinese population. Medical oncology (Northwood, London, England). 2013; 30:718.

26. Thomas RK, Baker AC, Debiasi RM, Winckler W, Laframboise T, Lin WM, Wang M, Feng W, Zander T, MacConaill L, Lee JC, Nicoletti R, Hatton C, et al. High-throughput oncogene mutation profiling in human cancer. Nature genetics. 2007; 39:347-351.