

# Combined effect of polymorphisms in *Rad51* and *XRCC3* on breast cancer risk and chromosomal radiosensitivity

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**Abstract.** Enhanced *in vitro* chromosomal radiosensitivity (CRS) has been proposed as a marker for low-penetrance gene mutations predisposing to breast cancer (BC). Since the double strand break (DSB) is the most detrimental form of DNA damage induced by ionizing radiation, it is possible that mutations in genes encoding proteins involved in DSB repair affect breast cancer risk. The purpose of the present study was to examine whether five single nucleotide polymorphisms (SNPs) in *Rad51* and *Xrcc3* (rs1801320, rs1801321, rs1799796, rs861539 and rs1799794) exhibited an association with breast cancer susceptibility in a Belgian population of BC patients with a known or putative genetic predisposition. We also ascertained whether a relationship exists between the occurrence of the 'variant' alleles of these variations and *in vitro* CRS. Blood samples were obtained from BC patients and from healthy female individuals. Variations in the 5' UTR of *Rad51* and *Xrcc3* were genotyped, and statistical analysis was performed. The results showed that low-penetrant variations in *Rad51* and *Xrcc3*, two proteins belonging to the homologous recombination DSB repair pathway, may modify BC risk in patients already carrying a pathological mutation in the highly penetrant BC genes *BRCA1* and *BRCA2*. Combined risk genotype analysis revealed that *Rad51* SNPs enhance BC risk in *BRCA2* patients, whereas *Xrcc3* SNPs significantly enhance BC risk in carriers of *BRCA1* mutations and in patients with hereditary BC. When four putative risk genotypes of *Rad51* and *Xrcc3* were combined, positive significant odds ratios were obtained in the entire patient population and in patients with a hereditary history of disease. Although obtained from a limited number of patients, our data are supportive of a polygenic model whereby combinations of weak variations

are responsible for an enhanced BC risk by acting jointly with high-penetrant mutations in *BRCA1* or *BRCA2*.

## Introduction

Breast cancer (BC) is the most frequently occurring cancer in women worldwide, with an estimated 1.15 million new cases in 2002 (23% of all cancers). More than half of the cases occurred in the industrialized countries of Europe and North America (1). In Europe, BC accounted for 28.9% of all cancer cases in women in 2006 and was the leading cause of cancer death in the European Union (16.7%) (2).

In a number of independent studies, enhanced *in vitro* chromosomal radiosensitivity (CRS) has been demonstrated in a significant number of breast cancer patients (3-8). *In vitro* CRS is therefore considered to be a hallmark for BC and a marker for low-penetrant gene mutations predisposing to BC (3,4,6,7,9). The most detrimental form of ionizing radiation-induced DNA damage is the double strand break (DSB) due to its ability to generate chromosomal aberrations when misrepaired or left unrepaired. The fact that DSB-initiated chromosomal instability may eventually trigger carcinogenesis in the breast epithelium is supported by evidence that breast cancer susceptibility genes such as *BRCA1* and *BRCA2*, and ATM and TP53 play crucial roles in DNA DSB repair and chromosome stability (10,11).

DSB repair in mammalian cells involves the error-free homologous recombination (HR) and the error-prone non-homologous end joining (NHEJ) pathways (reviewed in ref. 12). The cell cycle determines which pathway is activated after double strand breakage in a eukaryotic cell. The more accurate HR response mechanism is activated in the S/G<sub>2</sub> phase, when the sister chromatid provides a correct template of the damaged sequence for alignment. On the other hand, while the more error-prone mechanism of NHEJ is the favoured pathway in cells in the G<sub>0</sub>/G<sub>1</sub> phase. The inactivity of either pathway was shown to lead to potentially oncogenic translocations and other karyotypic changes in animal models (13). In humans, inherited genetic defects in these pathways often manifest as an increased CRS (14). For example, downregulation or mutation of DNA DSB repair proteins involved in the NHEJ pathway were shown to be associated with both BC risk and enhanced CRS (15-18).

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The main event of HR is the homology search and DNA strand invasion by the Rad51-ssDNA presynaptic filament, positioning the invading 3'-end on a template duplex DNA to initiate repair synthesis. Additionally, cofactors that promote assembly or stabilization of the Rad51 ssDNA filament as well as the disassembly of the Rad51-DNA complexes are significant. These cofactors include five Rad51 paralogs (Rad51B, Rad51C, Rad51D, Xrcc2 and Xrcc3) that are required for the *in vivo* Rad51 filament formation (19,20). In addition, the Xrcc3/Rad51C heterodimer appears to be required for the resolution of Holliday Junctions generated during HR (21). In this process, BRCA2 supports Rad51 filament formation, nucleation and filament stabilisation and is required for the ionizing radiation-induced formation of Rad51 foci *in vivo*. In addition, the formation of radiation damage-induced Rad51 foci requires functional Xrcc3 (22) and BRCA1 (10).

The relationship between mutations in *BRCA1/BRCA2* and BC risk is well known. Since HR is a key pathway in the maintenance of genomic stability, mutations in additional genes encoding proteins involved in HR are likely to affect cancer risk. To prove this hypothesis, different polymorphisms in *Rad51* (23-33) and *Xrcc3* (22,25,26,33-43) were previously investigated, with conflicting results. In this study, we examined whether two single nucleotide polymorphisms (SNPs) in the 5' untranslated region (UTR) of the *Rad51* gene and three SNPs in *Xrcc3* exhibited an association with breast cancer susceptibility in a Belgian population of BC patients with a known or putative genetic predisposition. We also investigated whether a relationship existed between the occurrence of the variant alleles of these variations and *in vitro* CRS.

## Materials and methods

**Study population.** The control population for this study included 172 healthy women (mainly staff members of Ghent University and Ghent University Hospital). Blood samples were obtained during the annual occupational medical examination. This control group did not include aged individuals. Thus, for age-matching purposes, blood samples from elderly healthy women were obtained during local senior club meetings. Written informed consent was provided by all members of the control group.

Samples of breast cancer patients were obtained in co-operation with the Centre for Medical Genetics of the Ghent University Hospital. The patients met at least one of the following criteria: i) patients were required to have at least 3 first-degree relatives with breast and/or ovarian cancer; or ii) belong to families, including at least two first- and/or second-degree relatives diagnosed with breast and/or ovarian cancer before the average age of 50; or iii) be affected by bilateral breast cancer with both tumors diagnosed before the age of 50; or iv) were required to have been diagnosed with BC prior to the age of 40, but not have a family history of the disease. Since a positive family history and/or diagnosis at a young age are regarded as significant risk factors for the development of hereditary breast cancer, the patients were screened for mutations in the *BRCA1* and *BRCA2* genes at the Centre for Medical Genetics, as previously described (44).

Patients were provided with genetic counselling and signed informed consent was provided by all of the patients.

The present study was approved by the Review Board of the first author's institution.

For statistical analysis, the breast cancer population was divided into groups of patients showing: i) hereditary BC; ii) familial BC and iii) sporadic cases at young age, but without a family history of BC. In 'hereditary' BC families, an autosomal dominant Mendelian pattern indicated that the disease was inherited, whereas BC was considered to be 'familial' when the frequency of the disease was clearly higher within the family when compared to the general population, but the criteria for hereditary BC were not met.

To evaluate whether the presence of a known mutation in *BRCA1* and/or *BRCA2* affected the association between the SNPs under study and BC susceptibility, the patient population was also divided into groups of carriers of *BRCA1* and/or *BRCA2* mutations.

Depending on the availability of genetic material, different numbers of patients were included for analysis of the different SNPs. The number of controls and patients evaluated for each SNP and the mean age of the population are shown in Table I.

**Collection and preparation of blood samples.** *In vitro* CRS was examined in a total of 62 patients. Samples (20 ml) of heparinized blood were obtained from the 62 patients and from matched control female individuals. Within 6 h after venepuncture, blood cultures (from 5 ml whole blood) were started to assess *in vitro* CRS using the G<sub>2</sub> chromatid break and the G<sub>0</sub> micronucleus (MN) assays (3,4). The two cytogenetic assays quantify the extent of DNA damage induced by irradiation during the G<sub>0</sub> or G<sub>2</sub> phases of the cell cycle. Lymphocyte separation was performed on part of the blood samples within 24 h after venepuncture, using lymphoprep (Axis-Shield, Lucron, Dieren, The Netherlands). Isolated lymphocytes were stored in liquid nitrogen until DNA extraction was performed (QIAamp DNA Blood Mini Kit, Qiagen, Hilden, Germany).

**Genotyping.** The c.-98 G>C (rs1801320) and c.-61 G>T (rs1801321) variations in the 5' UTR of *Rad51*, and the c.562-14 A>G (rs1799796), c.722 C>T (rs861539) and c.-316 A>G (rs1799794) variations in *Xrcc3* were genotyped using a polymerase chain reaction (PCR), combined with restriction fragment length polymorphism analysis or a SnapShot technique. The SNPs were named using the current nomenclature of the Human Genome Variation Society (45). The reference sequences of the corresponding genes are shown in Table II.

PCR products of control individuals were amplified using 100-200 ng DNA in a 25 µl reaction containing 0.5 mM dNTPs (Amersham Bioscience, Gent, Belgium), 1X PCR buffer (Invitrogen, Merelbeke, Belgium), 1.5 mM MgCl<sub>2</sub> (Invitrogen), 1 mM forward and reverse primers (Invitrogen) and 0.6 units Platinum Taq polymerase (Invitrogen). The PCR reaction of patient DNA contained 60 ng DNA in a 16 µl reaction, including 0.3 mM dNTPs, 0.6X PCR buffer, 0.9 mM MgCl<sub>2</sub>, 0.9 mM forward and reverse primers and 0.36 units Platinum Taq polymerase. Table I shows the sequences of primers used in this study.

DNA fragments containing *Rad51* SNPs were amplified by an initial denaturation step of 94°C, followed by 12 cycles including steps of denaturation at 94°C for 20 sec, annealing at 58°C and elongation at 72°C for 1 min. At each cycle, the

Table I. The number (n) of controls and patients evaluated for each single nucleotide polymorphism (SNP) and the mean, the range and standard deviation (SD) of the age of the populations and different patient groups are shown, together with the PCR primers and genotyping conditions.

	Control	All cases	Her <sup>a</sup> cases	Fam <sup>b</sup> cases	Spor <sup>c</sup> cases	No BRCA <sup>d</sup>	BRCA1 <sup>e</sup>	BRCA2 <sup>f</sup>	Genotyping
<i>Rad51</i> c.-98 G>C	n Mean age (range) SD	345 44.61 (23-80) 11.80	118 47.27 (23-75) 11.24	179 46.73 (26-80) 11.36	48 32.02 (24-38) 3.90	254 45.80 (24-80) 12.13	57 38.67 (23-65) 9.03	34 44.41 (25-71) 10.52	PCR primers Digest conditions 3 h, 60°C F-gcaggatcaagctctcgag R-tcgacttctgctcgaac <i>Bst</i> NI
<i>Rad51</i> c.-61 G>T	n Mean age (range) SD	254 44.79 (25-75) 11.60	86 47.88 (25-75) 10.73	133 46.78 (26-72) 11.29	35 32.32 (25-38) 3.46	188 45.77 (25-75) 11.89	39 40.10 (27-65) 9.59	27 44.08 (25-71) 10.86	PCR primers Snapshot primer F-gcaggatcaagctctcgag R-tcgacttctgctcgaac cgtcgggtcggcgctgcac
<i>Xrcc3</i> c.562-14 A>G	n Mean age (range) SD	350 44.80 (23-80) 11.78	122 47.55 (23-75) 11.31	179 46.92 (26-80) 11.22	49 32.11 (24-38) 3.91	256 45.91 (24-80) 12.09	58 38.98 (23-65) 8.78	36 45.53 (25-71) 11.30	PCR primers Digest conditions 3 h, 37°C F-gacacctctacagaggacg R-ttctcgatggttaggcacag <i>Pvu</i> II
<i>Xrcc3</i> Thr241 Met	n Mean age (range) SD	323 44.75 (23-80) 11.74	109 47.05 (23-75) 10.98	170 47.10 (26-80) 11.33	44 31.95 (24-38) 3.96	239 45.86 (24-80) 12.06	52 39.28 (23-65) 9.23	32 44.53 (25-71) 10.73	PCR primers Digest conditions 3 h, 37°C F-gacacctctacagaggacg R-ttctcgatggttaggcacag <i>Nla</i> III
<i>Xrcc3</i> c.-316 A>G	n Mean age (range) SD	343 44.64 (23-80) 11.80	117 47.28 (23-75) 11.18	179 46.70 (26-80) 11.38	47 31.96 (24-38) 3.92	253 45.86 (24-80) 12.13	56 38.73 (23-65) 9.12	34 44.41 (25-71) 10.52	PCR primers Digest Conditions 2 h, 37°C F-tgaggcgcctaatcagctg R-tggactgtgtaagcagcg <i>Fok</i> I

<sup>a</sup>Hereditary patient group; <sup>b</sup>familial patient group; <sup>c</sup>sporadic patient group; <sup>d</sup>patient group without *BRCA1* or *BRCA2* mutation; <sup>e</sup>patient groups with *BRCA1* or *BRCA2* mutation, respectively.

Table II. Chromosomal location of *Rad51* and *XRCC3*, and reference sequences used to name the polymorphisms under investigation.<sup>a</sup>

Gene, chromosomal location and reference sequence	SNP (alternative name, rs no. on NCBI)	Variant allele frequency	Variant allele frequency (published in NCBI)	HWE (p>0.05)
<b>Rad51</b>				
15q15.1	c.-98 G>C (5' UTR 135 G>C, rs1801320)	Controls: 4% Cases: 6%	European pop: 6.7%	Controls: 0.834 Cases: 0.543
Ref. sequence: NM_002875.3	c.-61 G>T (5' UTR 172 G>T, rs1801321)	Controls: 41% Cases: 42%	European pop: 46.7% Caucasian pop: 43.3%	Controls: 0.571 Cases: 0.582
<b>XRCC3</b>				
14q32.3	c.562-14A>G (IVS5-14 A>G, A17893G, rs1799796)	Controls: 29% Cases: 31%	European pop: 31.7% Caucasian pop: 33.3%	Controls: 0.631 Cases: 0.615
Ref. sequence: NM_001100119.1	c.722C>T (Thr241Met, C18067T, rs861539)	Controls: 43% Cases: 38%	European pop: 41.7% Caucasian pop: 45%	Controls: 0.964 Cases: 0.858
	c.-316A>G (5' UTR 4541 A>G, rs1799794)	Controls: 17% Cases: 21%	European pop: 18.4% Caucasian pop: 15.5%	Controls: 0.590 Cases: 0.743

<sup>a</sup>The five single nucleotide polymorphisms with variant allele frequencies in the control and patient populations are compared to the variant allele frequencies shown in NCBI (<http://www.ncbi.nlm.nih.gov/>). The genotype distributions were compared to those expected from the Hardy-Weinberg equilibrium (HWE) using a  $\chi^2$  test.

annealing temperature was decreased by 1°C. This was followed by 24 cycles at 94°C (40 sec), at 46°C (40 sec) and at 72°C (30 sec), and by a final extension step at 72°C (10 min). The DNA fragments containing c.562-14 A>G and c.722C>T in *Xrcc3* and the fragment surrounding the *Xrcc3* c.-316 A>G SNP were amplified using a 35-cycle PCR program consisting of an initial denaturation step at 95°C for 5 min, followed by 35 cycles of 3 min, including a denaturation step at 95°C (1 min), an annealing step at 58°C (1 min) and an elongation step at 72°C (1 min), followed by a final extension step of 10 min at 72°C. The efficiency of the PCR reaction was confirmed by electrophoresis on a 1.5% agarose gel. A negative control containing water was included during each PCR reaction to exclude contamination.

To genotype the 5' UTR c.-61 G>T SNP, the SnapShot multiplex system was utilized (Applied Biosystems, Halle, Belgium). SnapShot primers are shown in Table I. Analysis of 2 µl of the SnapShot products [with 10 µl Amresco Capillary Electrophoresis buffer (Lucron Bioproducts, Dieren, The Netherlands) and 0.3 µl LIZ-120 Size Standard (Applied Biosystems)] was performed on the capillary system of the ABI Prism 3730 Genetic Analyser (Applied Biosystems). Results were visualised on an Applied Biosystems Peak scanner, version 1.0. To genotype the c.-98 G>C SNP in *Rad51* and the three SNPs in *Xrcc3*, PCR products were digested with sequence-specific restriction endonucleases (Table I). Digested products were then analysed by electrophoresis on

a 2% agarose gel and visualised under ultraviolet light after ethidium bromide staining.

**Statistical analysis.** Statistical analysis of the data was performed using Microsoft office Excel 2007 or the Statistical Package for Social Sciences (SPSS), version 15.0.

The association of the five SNPs with BC risk was evaluated by calculating age-corrected odds ratios (ORs) by means of logistic regression. To improve the statistical power of the analytical investigation, particularly for the rare homozygous variant (HV) genotypes, the OR was calculated for the heterozygous (He) or HV genotypes using the homozygous normal (HN) genotype as reference, and for the He and HV genotypes combined in a single group. The complete control population was always used to compare genotype frequencies with those found in the different patient groups (divided by family history or presence of a *BRCA1/BRCA2* mutation).

The observed genotype distributions were compared with those expected from the Hardy-Weinberg equilibrium (HWE) using a standard  $\chi^2$  test. The linkage disequilibrium (LD) coefficient  $r^2$  between the analyzed SNPs was calculated using Haploview (version 4.0) (46).

## Results

**Comparison of variant allele and standard frequencies.** The observed genotype distributions of the five analyzed SNPs



Table III. Genotype frequencies in control subjects and in the total patient population (all cases).

		Controls		All Cases		
		% (No.)	% (No.)	Corr OR	(95% CI)	p ( $\alpha=0.05$ )
<i>Rad51</i>	GG	91.18 (155)	87.54 (302)	Ref.		
c.-98	GC	8.82 (15)	12.46 (43)	1.42	(0.76-2.67)	0.275
G>C	CC	0.00 (0)	0.00 (0)			
	GC+CC	8.82 (15)	12.46 (43)	1.42	(0.76-2.67)	0.275
<i>Rad51</i>	GG	32.47 (50)	33.86 (86)	Ref.		
c.-61	GT	52.60 (81)	45.28 (115)	0.93	(0.58-1.47)	0.742
G>T	TT	14.94 (23)	20.87 (53)	1.40	(0.76-2.59)	0.287
	GT+TT	67.53 (104)	66.14 (168)	1.03	(0.67-1.60)	0.882
<i>XRCC3</i>	AA	48.21 (81)	48.29 (169)	Ref.		
c.562-14	AG	44.64 (75)	43.14 (151)	0.93	(0.63-1.38)	0.714
A>G	GG	7.14 (12)	8.57 (30)	1.22	(0.58-2.55)	0.600
	AG+GG	51.79 (87)	51.71 (181)	0.97	(0.66-1.42)	0.869
	CC	32.14 (54)	36.84 (119)	Ref.		
<i>XRCC3</i>	CT	50.00 (84)	49.23 (159)	0.86	(0.56-1.31)	0.479
Thr241Met	TT	17.86 (30)	13.93 (45)	0.78	(0.44-1.41)	0.414
	CT+TT	67.86 (114)	63.16 (204)	0.84	(0.56-1.26)	0.396
<i>XRCC3</i>	AA	68.02 (117)	64.14 (220)	Ref.		
c.-316	AG	30.23 (52)	31.49 (108)	1.03	(0.68-1.56)	0.892
A>G	GG	1.74 (3)	4.37 (15)	2.47	(0.68-8.94)	0.168
	AG+GG	31.98 (55)	35.86 (123)	1.11	(0.74-1.65)	0.619

## Risk genotype analyses.

Risk genotypes		Controls		All cases		
(No.)		% (No.)	% (No.)	Corr OR	(95% CI)	p ( $\alpha=0.05$ )
<i>Rad51</i> <sup>a</sup>	0	75.32 (116)	68.11 (173)	Ref.		
	>1	24.68 (38)	31.11 (89)	1.39	(0.87-2.20)	0.387
<i>XRCC3</i> <sup>b</sup>	0	91.67 (154)	86.18 (293)	Ref.		
	>1	8.33 (14)	13.82 (47)	<b>1.79</b>	<b>(0.94-3.41)</b>	<b>0.076</b>
<i>Rad51</i> and <i>XRCC3</i> <sup>c</sup>	0	70.67 (106)	60.32 (152)	Ref.		
	>1	29.33 (44)	39.68 (100)	<b>1.59</b>	<b>(1.02-2.47)</b>	<b>0.041</b>

<sup>a</sup>c.-98 G>T and c.-61 G>T; <sup>b</sup>c.562-14 and c.-316 A>G; <sup>c</sup>c.-98 G>T and c.-61 G>T (*Rad51*), and c.562-14 and c.-316 A>G (*Xrcc3*). Age-corrected ORs (Corr OR) for each individual single nucleotide polymorphism (SNP) and for grouped 'risk genotypes' are shown, together with 95% confidence intervals (95% CI). Significant and borderline-significant findings are highlighted.

showed no systematic deviation from the HWE ( $p>0.05$ ). Comparison of the variant allele frequency of our control population with standard frequencies published on NCBI (47) resulted in no significant deviations (Table II).

**Variations in *Rad51* and *Xrcc3* and breast cancer risk.** Age-corrected ORs indicative of an association between individual SNPs in *Rad51* or *Xrcc3* and BC risk in the total patient population are recorded in Table III. Table IV shows the ORs for patient groups classified on the basis of familial history, whereas in Table V patient groups are subdivided depending on the presence or absence of a mutation in *BRCA1* or *BRCA2*. No significant associations were found between individual SNPs and breast cancer risk. On the other hand, the

c.-316 A>G SNP in *Xrcc3* exhibited high, borderline-significant ORs in *BRCA1* mutation carriers (OR<sub>He+HV</sub>=1.92,  $p=0.053$ , Table V). The ORs calculated for the c.722 C>T (Thr241Met) variation in *Xrcc3* were found to be  $\leq 1$  in most cases (Tables III-V), indicating that the variant 'T' allele is not a risk allele for BC. Concerning the c.-98 G>C SNP in *Rad51*, a non-significant enhancement of BC risk was observed for the heterozygous genotype (OR>1) in the total patient population (Table III) as well as in patient groups divided on the basis of familial history or BRCA mutations (Tables IV and V), with the exception of sporadic cases diagnosed at a young age (Table IV). As regards the c.-61 G>T variation in *Rad51*, and the c.562-14 A>G and c.-316 A>G SNPs in *Xrcc3*, the 'homozygous variant' genotypes mostly yielded ORs >1.

Table IV. Genotype frequencies in control subjects and in breast cancer patient groups divided by family history.

	Controls			Hereditary cases			Familial cases			Sporadic cases				
	RG (No.)	% (No.)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)
<i>Rad51</i> c.-98 G>C	GG	91.18 (155)	87.29 (103)	Ref.	-	-	85.47 (153)	Ref.	-	-	95.83 (46)	Ref.	-	-
	GC	8.82 (15)	12.71 (15)	1.39	(0.65-3.00)	0.396	14.53 (26)	1.69	(0.86-3.34)	0.130	4.17 (2)	0.33	(0.06-1.71)	0.185
	CC	0.00 (0)	0.00 (0)	-	-	-	0.00 (0)	-	-	-	0.00 (0)	-	-	-
	GC+CC	8.82 (15)	12.71 (15)	1.39	(0.65-3.00)	0.396	14.53 (26)	1.69	(0.86-3.34)	0.130	4.17 (2)	0.33	(0.06-1.71)	0.185
<i>Rad51</i> c.-61 G>T	GG	32.47 (50)	32.56 (28)	Ref.	-	-	31.58 (42)	Ref.	-	-	45.71 (16)	Ref.	-	-
	GT	52.60 (81)	50.00 (43)	1.01	(0.55-1.84)	0.977	45.86 (61)	0.95	(0.55-1.62)	0.840	31.43 (11)	0.63	(0.24-1.65)	0.345
	TT	14.94 (23)	17.44 (15)	1.23	(0.55-2.75)	0.619	22.56 (30)	1.58	(0.79-3.14)	0.196	22.86 (8)	1.48	(0.46-4.79)	0.510
	GT+TT	67.53 (104)	67.44 (58)	1.06	(0.60-1.87)	0.848	68.42 (91)	1.09	(0.66-1.81)	0.736	54.29 (19)	0.83	(0.35-1.96)	0.670
<i>XRCC3</i> c.562-14 A>G	AA	48.21 (81)	46.72 (57)	Ref.	-	-	50.28 (90)	Ref.	-	-	44.90 (22)	Ref.	-	-
	AG	44.64 (75)	40.98 (50)	0.92	(0.56-1.52)	0.745	41.90 (75)	0.86	(0.55-1.35)	0.504	53.06 (26)	0.99	(0.45-2.15)	0.976
	GG	7.14 (12)	12.30 (15)	1.66	(0.71-3.87)	0.243	7.82 (14)	1.07	(0.46-2.48)	0.876	2.04 (1)	0.48	(0.05-4.88)	0.531
	AG+GG	51.79 (87)	53.28 (65)	1.02	(0.64-1.64)	0.927	49.72 (89)	0.89	(0.58-1.36)	0.585	55.10 (27)	0.94	(0.44-2.03)	0.877
<i>XRCC3</i> Thr241 Met	CC	32.14 (54)	42.20 (46)	Ref.	-	-	35.29 (60)	Ref.	-	-	29.55 (13)	Ref.	-	-
	CT	50.00 (84)	45.87 (50)	0.71	(0.41-1.21)	0.204	51.18 (87)	0.92	(0.57-1.50)	0.746	50.00 (22)	0.81	(0.33-2.00)	0.644
	TT	17.86 (30)	11.93 (13)	0.59	(0.27-1.27)	0.175	13.53 (23)	0.78	(0.40-1.54)	0.480	20.45 (9)	1.93	(0.59-6.36)	0.279
	CT+TT	67.86 (114)	57.80 (63)	0.68	(0.41-1.12)	0.131	64.71 (110)	0.89	(0.56-1.41)	0.617	70.45 (31)	1.00	(0.43-2.35)	0.995
<i>XRCC3</i> c.-316 A>G	AA	68.02 (117)	61.54 (72)	Ref.	-	-	63.13 (113)	Ref.	-	-	74.47 (35)	Ref.	-	-
	AG	30.23 (52)	34.19 (40)	1.20	(0.72-2.00)	0.492	32.40 (58)	1.06	(0.66-1.70)	0.817	21.28 (10)	0.42	(0.17-1.04)	0.062
	GG	1.74 (3)	4.27 (5)	2.64	(0.60-11.56)	0.197	4.47 (8)	2.73	(0.69-10.71)	0.151	4.26 (2)	2.65	(0.19-36.86)	0.467
	AG+GG	31.98 (55)	38.46 (45)	1.28	(0.78-2.10)	0.338	36.87 (66)	1.14	(0.73-1.80)	0.553	25.53 (12)	0.49	(0.20-1.17)	0.106
Risk genotype analysis.														
	Controls			Hereditary cases			Familial cases			Sporadic cases				
	RG (No.)	% (No.)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)	% (No.)	Corr OR	(95% CI)	p-value (α=0.05)
<i>Rad51</i> <sup>a</sup>	0	75.32 (116)	68.60 (59)	Ref.	-	-	50.28 (89)	Ref.	-	-	55.56 (25)	Ref.	-	-
	>1	24.68 (38)	31.40 (27)	1.37	(0.76-2.48)	0.293	24.86 (44)	1.46	(0.87-2.47)	0.152	22.22 (10)	1.13	(0.44-2.92)	0.804
<i>XRCC3</i> <sup>b</sup>	0	91.67 (154)	81.03 (94)	Ref.	-	-	87.57 (155)	Ref.	-	-	93.62 (44)	Ref.	-	-
	>1	8.33 (14)	18.97 (22)	<b>2.48</b>	<b>(1.20-5.15)</b>	<b>0.015</b>	12.43 (22)	1.61	(0.79-3.30)	0.193	6.38 (3)	0.70	(0.23-5.32)	0.902
<i>Rad51</i> and <i>XRCC3</i> <sup>c</sup>	0	70.67 (106)	57.14 (48)	Ref.	-	-	42.25 (79)	Ref.	-	-	55.56 (25)	Ref.	-	-
	>1	29.33 (44)	42.86 (36)	<b>1.80</b>	<b>(1.03-3.15)</b>	<b>0.041</b>	<b>28.88 (54)</b>	<b>1.62</b>	<b>(0.99-2.68)</b>	<b>0.057</b>	22.22 (10)	1.02	(0.40-2.63)	0.965
*c.-98 G>T and c.-61 G>T; <sup>b</sup> c.562-14 and c.-316 A>G; <sup>c</sup> c.-98 G>T and c.-316 A>G ( <i>Xrcc3</i> ). Age-corrected ORs (Corr OR) for each individual single nucleotide polymorphism (SNP) and for grouped 'risk genotypes' (RGs) are shown, together with 95% confidence intervals (95% CI). Significant and borderline-significant findings are highlighted in bold.														

<sup>a</sup>c.-98 G>T and c.-61 G>T; <sup>b</sup>c.562-14 and c.-316 A>G; <sup>c</sup>c.-98 G>T and c.-61 G>T (*Rad51*) and c.562-14 and c.-316 A>G (*XRCC3*). Age-corrected ORs (Corr OR) for each individual single nucleotide polymorphism (SNP) and for grouped 'risk genotypes' (RGs) are shown, together with 95% confidence intervals (95% CI). Significant and borderline-significant findings are highlighted in bold.

Table V. Genotype frequencies in control subjects and in breast cancer patients divided on the basis of presence vs. absence of a *BRCA1* and/or *BRCA2* mutation.

	Controls	Patient group without <i>BRCA1/BRCA2</i> mutation				Patient group with <i>BRCA1</i> mutation				Patient group with <i>BRCA2</i> mutation				
		% (No.)	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )
<i>Rad51</i> c.-98 G>C	GG	91.18 (155)	87.65 (220)	Ref.	-	-	89.47 (51)	Ref.	-	-	88.24 (30)	Ref.	-	-
	GC	8.82 (15)	12.35 (31)	1.42	(0.74-2.75)	0.293	10.53 (6)	1.09	(0.38-3.14)	0.867	11.76 (4)	1.13	(0.34-3.74)	0.842
	CC	0.00 (0)	0.00 (0)	-	-	-	0.00 (0)	-	-	-	0.00 (0)	-	-	-
	GC+CC	8.82 (15)	12.35 (31)	1.42	(0.74-2.75)	0.293	10.53 (6)	1.09	(0.38-3.14)	0.867	11.76 (4)	1.13	(0.34-3.74)	0.842
<i>Rad51</i> c.-61 G>T	GG	32.47 (50)	32.43 (60)	Ref.	-	-	43.59 (17)	Ref.	-	-	29.63 (16)	Ref.	-	-
	GT	52.60 (81)	47.57 (88)	0.98	(0.60-1.61)	0.950	41.03 (16)	0.68	(0.31-1.53)	0.353	40.74 (11)	1.00	(0.37-2.71)	0.992
	TT	14.94 (23)	20.0 (37)	1.39	(0.72-2.67)	0.323	15.38 (6)	0.89	(0.29-2.70)	0.887	29.63 (8)	2.37	(0.77-7.28)	0.133
	GT+TT	67.53 (104)	67.57 (125)	1.08	(0.68-1.72)	0.756	56.41 (22)	0.73	(0.34-1.55)	0.408	70.37 (19)	1.32	(0.53-3.31)	0.550
<i>XRCC3</i> c.562-14 A>G	AA	48.21 (81)	47.83 (121)	Ref.	-	-	56.90 (33)	Ref.	-	-	36.11 (13)	Ref.	-	-
	AG	44.64 (75)	43.08 (109)	0.92	(0.61-1.40)	0.697	36.21 (12)	0.54	(0.27-1.09)	0.085	55.56 (30)	1.59	(0.73-3.46)	0.241
	GG	7.14 (12)	9.09 (23)	1.30	(0.60-2.80)	0.504	6.9 (4)	0.81	(0.23-2.92)	0.750	8.33 (3)	1.60	(0.39-6.53)	0.513
	AG+GG	51.79 (87)	52.17 (132)	0.97	(0.65-1.45)	0.889	43.10 (25)	0.58	(0.30-1.12)	0.103	63.89 (23)	1.59	(0.75-2.28)	0.227
<i>XRCC3</i> Thr241 Met	CC	32.14 (54)	37.29 (88)	Ref.	-	-	32.69 (17)	Ref.	-	-	40.63 (13)	Ref.	-	-
	CT	50.00 (84)	46.61 (110)	0.81	(0.52-1.27)	0.359	57.69 (30)	1.04	(0.50-2.19)	0.908	53.13 (17)	0.80	(0.35-1.80)	0.588
	TT	17.86 (30)	16.10 (38)	0.89	(0.49-1.62)	0.692	9.66 (5)	0.61	(0.18-2.09)	0.428	6.25 (2)	0.34	(0.07-1.62)	0.175
	CT+TT	67.86 (114)	62.71 (148)	0.83	(0.54-1.27)	0.388	67.31 (35)	1.05	(0.51-2.16)	0.889	59.38 (19)	0.69	(0.32-1.51)	0.353
<i>XRCC3</i> c.-316 A>G	AA	68.02 (117)	65.20 (163)	Ref.	-	-	51.79 (29)	Ref.	-	-	73.53 (25)	Ref.	-	-
	AG	30.23 (52)	30.40 (76)	1.00	(0.65-1.54)	0.999	42.86 (24)	1.86	(0.99-3.50)	0.054	23.53 (8)	0.61	(0.25-1.47)	0.269
	GG	1.74 (3)	4.40 (11)	2.43	(0.65-9.06)	0.187	5.36 (3)	4.03	(0.77-21.03)	0.098	2.94 (1)	1.59	(0.15-16.63)	0.700
	AG+GG	31.98 (55)	34.80 (87)	1.08	(0.71-1.65)	0.727	48.21 (27)	1.94	(0.99-3.80)	0.053	26.47 (9)	0.65	(0.28-1.53)	0.329
Risk genotype analysis.														
	Controls	Patient group without <i>BRCA1/2</i> mutation				Patient group with <i>BRCA1</i> mutation				Patient group with <i>BRCA2</i> mutation				
		% (No.)	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )	% (No.)	Corr OR	(95% CI)	p-value ( $\alpha=0.05$ )
<i>Rad51</i> <sup>a</sup>	0	75.32 (116)	69.73 (129)	Ref.	-	-	74.36 (29)	Ref.	-	-	55.56 (15)	Ref.	-	-
	>1	24.68 (38)	30.27 (56)	1.38	(0.85-2.24)	0.2	25.64 (10)	1.03	(0.44-2.39)	0.9	<b>44.44 (12)</b>	<b>2.23</b>	<b>(0.94-5.26)</b>	<b>0.06</b>
<i>XRCC3</i> <sup>b</sup>	0	91.67 (154)	86.35 (215)	Ref.	-	-	83.64 (46)	Ref.	-	-	87.88 (29)	Ref.	-	-
	>1	8.33 (14)	13.65 (34)	1.75	(0.90-3.42)	0.1	<b>16.36 (9)</b>	<b>2.77</b>	<b>(1.04-7.36)</b>	<b>0.04</b>	12.12 (4)	1.63	(0.49-5.41)	0.4
<i>Rad51</i> and <i>XRCC3</i> <sup>c</sup>	0	70.67 (106)	60.96 (114)	Ref.	-	-	61.54 (24)	Ref.	-	-	53.85 (14)	Ref.	-	-
	>1	29.33 (44)	<b>39.04 (73)</b>	<b>1.57</b>	<b>(0.98-2.50)</b>	<b>0.059</b>	38.46 (15)	1.58	(0.73-3.43)	0.2	46.15 (12)	1.94	(0.82-4.60)	0.1
a,c,c.-98 G>T; b,c.562-14 and c.-316 A>G; c.-98 G>T and c.-316 A>G ( <i>Xrcc3</i> ). Age-corrected ORs (Corr OR) for each individual single nucleotide polymorphism (SNP) and for grouped 'risk genotypes' (RGs) are shown, together with 95% confidence intervals (95% CI). Significant and borderline-significant findings are highlighted in bold.														

<sup>a</sup>c.-98 G>T and c.-61 G>T; <sup>b</sup>c.562-14 and c.-316 A>G; <sup>c</sup>c.-98 G>T and c.-61 G>T (*Rad51*) and c.562-14 and c.-316 A>G (*XRCC3*). Age-corrected ORs (Corr OR) for each individual single nucleotide polymorphism (SNP) and for grouped 'risk genotypes' (RGs) are shown, together with 95% confidence intervals (95% CI). Significant and borderline-significant findings are highlighted in bold.

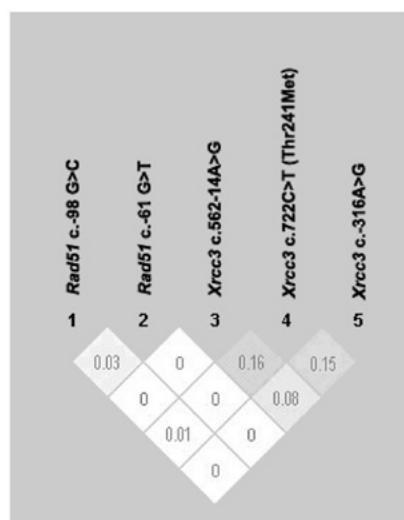


Figure 1. Linkage disequilibrium (LD) plot of the c.-98 G>C (rs1801320) and c.-61 G>T (rs1801321) variations in the 5' UTR of *Rad51*, and the c.562-14 A>G (rs1799796), c.722 C>T (rs861539) and c.-316 A>G (rs1799794) variations in *Xrcc3*. Analysis was performed in 172 healthy control individuals. The LD coefficient  $r^2$  is shown in the plot squares.

Since the five SNPs investigated in this study are not in LD with each other (Fig. 1), we analyzed whether the combination of different DSB repair risk genotypes (RGs) may result in a significant association with BC risk. For this specific analysis, the He genotype 'GC' of the c.-98 G>C SNP in *Rad51* and the HV genotypes 'TT' of the c.-61 G>T variant (*Rad51*), as well as 'GG' of c.562-14 A>G (*Xrcc3*) and 'GG' of c.-316 A>G (*Xrcc3*), were considered as putative breast cancer susceptibility genotypes (Table III).

The ORs calculated for various combinations of DSB repair RGs and BC are shown in Tables III-V.

Combination of *Xrcc3* RGs resulted in high and significant ORs in the hereditary patient group ( $OR_{>IRG}=2.48$ ; 95% CI, 1.20-5.15;  $p=0.015$ ; Table IV) and *BRCA1* mutation carriers ( $OR_{>IRG}=2.77$ ; 95% CI, 1.04-7.36;  $p=0.041$ ; Table V).

The combined *Rad51* risk genotype analysis yielded ORs of  $>1$  in most cases, indicating a possible enhancement of BC risk in the presence of at least 1 risk genotype. In particular, *BRCA2* mutation carriers showed a borderline-significant  $OR_{>IRG}$  of 2.23 (95% CI, 0.94-5.26;  $p=0.06$ ; Table V).

When all of the DSB repair genotypes were combined, a significant positive association between the putative risk alleles listed in the previous paragraph and BC was found in the total patient population ( $OR_{>IRG}=1.59$ ; 95% CI, 1.02-2.47;  $p=0.041$ ; Table III) and in hereditary cases ( $OR=1.80$ ; 95% CI, 1.03-3.15;  $p=0.04$ , Table IV). Familial non-*BRCA1* or non-*BRCA2* cases showed borderline-significant ORs (Table V).

**Variations in *Rad51* and *Xrcc3* and in vitro CRS.** To examine the potential effect of the 5 variations under study on CRS, we previously determined the *in vitro* CRS of 62 patients and 100 healthy female individuals using the  $G_2$  chromatid-break or  $G_0$  MN assays (4). In this study, the 75 percentiles of the  $G_0$  MN or  $G_2$  chromatid-break values calculated in the healthy female population were used as cut-off points for radiosensitivity, as previously described (8). Age-corrected ORs

were calculated by logistic regression. The genotype frequencies in radiosensitive patients were compared to frequencies assessed in the non-radiosensitive patient group (Table VI). The two SNPs in *Rad51* and the c.-316 A>G variation in *Xrcc3* showed ORs of  $<1$ , suggesting a potential protective effect of the variant alleles towards CRS. High but non-significant ORs were assessed for the c.562-14 A>G SNP in *Xrcc3* ( $OR_{He}=3.45$ ,  $OR_{HV}=1.65$ ) and for the He genotype of the c.722 C>T variation in *Xrcc3* ( $OR_{He}=1.73$ ).

A risk allele (RA) analysis was performed to assess the effect of combining different DSB repair variant alleles on CRS (Table VI). ORs of  $<1$  were obtained by combining variant alleles of the SNPs in *Rad51*. The protective effect for radiosensitive patients carrying two *Rad51* variant alleles was significant ( $OR_{2VA}=0.10$ , 95% CI, 0.01-0.96;  $p=0.046$ ; Table VI). Non-significant results were obtained by analyzing the combined effect of the c.562-14 A>G and c.722 C>T SNPs in *Xrcc3*.

## Discussion

Misrepair or inefficient repair of DSBs may lead to genetic instability and ultimately to carcinogenesis. This process is confirmed by the fact that different cancer-prone genetic disorders such as Ataxia-telangiectasia and Nijmegen Breakage syndrome, are linked to the HR-DSB repair pathway, and by the finding that genes involved in HR, such as *BRCA1* and *BRCA2*, are tumor-suppressor genes (11). However, BC-predisposing genes such as *BRCA1* and *BRCA2* account for only 15-20% of inherited susceptibility (48,49). Epidemiological analysis suggests that the remaining 80-85% can mainly be explained by a polygenic model whereby the combined effects of numerous individual weak genetic variants are responsible for an enhanced breast cancer risk (50). This model is also supported by the results obtained in this study, where a significant association between the genetic variants and BC risk was observed only after combined risk genotype analysis.

Although the c.-98 G>C SNP in *Rad51* was initially shown to have no impact on BC risk in the general BC population (24,31,33,51), contradicting results were obtained concerning the effect of the variant 'C' allele of c.-98 G>C on BC risk in a population of BC patients with a familial history of the disease (25,26,28). Various studies reported that the c.-98 G>C variation appeared to modify BC risk in *BRCA1* or *BRCA2* carriers. In their study, Jakubowska *et al* (27) demonstrated a protective effect of this variation on BC susceptibility in cases carrying the *BRCA1* gene, one of the three most common Polish founder mutations. Notably, a protective effect was also shown against colon cancer risk in Polish cohorts (52).

In different studies involving *BRCA2* mutation carriers, the c.-98 G>C variation was shown to enhance BC susceptibility (29,30,32). In 2007, a meta-analysis of 19 studies confirmed that the c.-98 G>C SNP is a BC risk modifier in *BRCA2*, but not in *BRCA1* mutation carriers (23). A significant odds ratio was also found between BC risk and c.-98 G>C in *BRCA2* mutation carriers in a recent meta-analysis by Zhou and coworkers (53). Most recently, four meta-analysis studies were published concerning the role of the c.-98 G>C SNP in breast cancer predisposition (53-56). With the exception of the Yu *et al* study (55), the findings of the meta-analyses showed a



Table VI. Genotype frequencies, age-corrected OR and 95% confidence intervals (95% CI) assessed in both radiosensitive (RS) and non-radiosensitive (NRS) cases.

		NRS cases		RS cases		
		% (No.)	% (No.)	Corr OR	(95% CI)	p ( $\alpha=0.05$ )
<i>Rad 51</i>	GG	78.57 (22)	88.24 (15)	-	-	-
c.-98	GC	21.43 (6)	11.76 (2)	0.49	(0.09-2.78)	0.422
G>C	CC	0.00 (0)	0.00 (0)	-	-	-
	GC+CC	21.43 (6)	11.76 (2)	0.49	(0.09-2.78)	0.422
<i>Rad 51</i>	GG	16.67 (4)	33.33 (5)	-	-	-
c.-61	GT	45.83 (11)	46.67 (7)	0.50	(0.10-2.54)	0.402
G>T	TT	37.50 (9)	20.00 (3)	0.27	(0.04-1.70)	0.161
	GT+TT	83.33 (20)	66.67 (10)	0.39	(0.09-1.81)	0.230
<i>XRCC3</i>	AA	67.86 (19)	47.06 (8)	-	-	-
c.562-14	AG	21.43 (6)	41.18 (7)	3.45	(0.78-15.31)	0.104
A>G	GG	10.71 (3)	11.76 (2)	1.65	(0.23-12.06)	0.622
	AG+GG	32.14 (9)	52.94 (9)	2.73	(0.73-10.17)	0.135
<i>XRCC3</i>	CC	37.04 (10)	35.29 (6)	-	-	-
Thr241	CT	40.74 (11)	58.82 (10)	1.73	(0.44-6.83)	0.437
Met	TT	22.22 (6)	5.88 (1)	0.26	(0.02-2.76)	0.262
	CT+TT	62.96 (17)	64.71 (11)	1.12	(0.31-4.02)	0.863
<i>XRCC3</i>	AA	65.38 (17)	76.47 (13)	-	-	-
c.-316	AG	34.62 (9)	23.53 (4)	0.58	(0.15-2.31)	0.438
A>G	GG	0.00 (0)	0.00 (0)	-	-	-
	AG+GG	34.62 (9)	23.53 (4)	0.58	(0.15-2.31)	0.438

## Risk allele analysis.

Risk alleles		NRS cases		RS cases		
(No.)		% (No)	% (No)	Corr OR	(95% CI)	p ( $\alpha=0.05$ )
<i>Rad51</i> <sup>a</sup>	0	8.70 (2)	28.57 (4)	-	-	-
	1	47.83 (11)	57.14 (8)	0.37	(0.05-2.55)	0.313
	2	43.48 (10)	14.29 (2)	<b>0.10</b>	<b>(0.01-0.96)</b>	<b>0.046</b>
	3	4.35 (1)	7.14 (1)	0.56	(0.02-15.01)	0.732
<i>XRCC3</i> <sup>b</sup>	0	19.23 (5)	17.65 (3)	-	-	-
	1	26.92 (7)	41.18 (7)	1.72	(0.29-10.23)	0.551
	2	53.85 (14)	41.18 (7)	0.89	(0.16-4.94)	0.889

<sup>a</sup>c.-98 G>T and c.-61 G>T; <sup>b</sup>c.562-14 and Thr241Met. The odds ratios (ORs) for each individual single nucleotide polymorphism (SNP) and for the risk allele analysis are shown. Significant findings are highlighted in bold.

significant association with BC risk in the carriers of the c.-98 G>C variation. In particular, the study by Gao *et al* demonstrated a correlation between the CC genotype and elevated BC risk in sporadic patients and in the broader European population (54). Although no significant association of this polymorphism with BC was found by Wang *et al* in overall and European populations, the authors observed a significant increase in breast cancer risk (recessive model CC vs. GG/CG: OR=1.35, 95% CI, 1.05-1.74) following the exclusion of studies that did not show a Hardy-Weinberg equilibrium for the SNP (56). Finally, the Zhou study found significant ORs in both additive (1.34; 95% CI, 1.01-1.78) and recessive (1.37; 95% CI, 1.03-1.82) models (53).

The c.-61 G>T SNP in the 5' UTR of *Rad51* has been studied less extensively and no association with BC risk was found (32,57,58). Our results indicate that the He genotype of c.-98 G>C and the HV genotype of c.-61 G>T may be considered as putative RGs for BC since the combined effect of the two genotypes exhibited a borderline significant OR of 2.23 in the BC patient group with a *BRCA2* mutation (95% CI, 0.94-5.26). This combined effect on BC risk in *BRCA2* carriers may be due to the fact that the c.-98 G>C and c.-61 G>T variations are functional and result in an increased promoter activity (59).

The functionality of the two variations also substantiates the finding that the variant alleles of the two SNPs in *Rad51* have a protective effect towards *in vitro* CRS. The protective

effect became significant after analysis of the combined risk alleles ( $OR_{2RA}=0.10$ , 95% CI, 0.01-0.96;  $p=0.046$ ; Table VI). According to Hasselbach *et al* (59), the two variations are gain-of-function mutations. This observation explains the protection observed towards *in vitro* CRS as cells with an increased level of *Rad51* are expected to be more resistant to DNA damage. The enhanced BC risk observed in *BRCA2* mutation carriers in this study appear to be contradictory to this rationale. However, increased levels of *Rad51* have been found in cell lines derived from breast carcinoma. It is likely that a balanced relationship between *Rad51* expression and other repair factors is required. Additionally, the overexpression of *Rad51* may lead to increased genomic instability and therefore contribute to carcinogenesis (60). In contrast to Hasselbach *et al* (59), Antoniou *et al* (23) found that the c.-98 'C' allele may cause an overall lower abundance of the *Rad51* protein. The effects of these SNPs on *Rad51* protein levels, particularly in breast tissue, require further investigation.

The association between the c.562-14 A>G SNP in *Xrcc3* and BC is controversial. Some studies showed a protective effect of the HV genotype against BC (58,36), whereas other reports showed no association with BC risk (37).

Kuschel *et al* (58) revealed that the c.-316 A>G variation in *Xrcc3* did not correlate with BC susceptibility. The Breast Cancer Consortium pooled data from 5 and 6 studies focusing on c.562-14 A>G and c.-316 A>G, respectively, and found no correlation between the frequency of the variant alleles and BC risk (34). In our study, positive but non-significant ORs were obtained for the rare HV genotype of the two SNPs. However, the analysis performed by combining the two HV RGs, resulted in significant ORs in hereditary ( $OR_{>IRG}=2.48$ ; 95% CI, 1.20-5.15;  $p=0.015$ ; Table IV) and in *BRCA1* patient groups ( $OR_{>IRG}=2.77$ ; 95% CI, 1.04-7.36;  $p=0.041$ ; Table V), and borderline-significant-positive ORs for the total patient population ( $OR_{>IRG}=1.79$ ; 95% CI, 0.94-3.41;  $p=0.076$ ; Table III). Houlston and Peto (61) estimated that, depending on the risk exhibited by a variant allele, genotype studies in familial populations may be 2-5 times more powerful than studies involving unselected patients. The genotyping explains the occurrence of the strongest associations between SNPs and BC in the hereditary patient group. The strong association observed in *BRCA1* patients indicates that low-penetrant variations are capable of modifying BC risk in patients already carrying a high penetrant mutation. Although the *BRCA1* and *Xrcc3* proteins do not physically interact, both are indispensable components of the same HR pathway. *Xrcc3* is essential for the *Rad51* filament formation (20), whereas *BRCA1* plays a role as a 'molecular scaffold' for the assembly of the HR machinery, and recruits checkpoint factors to initiate a signaling cascade that halts cell cycle progression (10).

The c.722 C>T variation in *Xrcc3*, encoding the missense mutation Thr241Met, has been extensively studied. Published data have produced varying results, ranging from i) a protective effect towards BC (37), particularly in post-menopausal women carrying at least one 'T' allele (57); to ii) no association with breast (or colon) cancer risk (26,33,38,42,62); to iii) an enhanced susceptibility to BC, (58,25,35-37,39-41,43); or iv) to an increased risk of onset of local metastases (51). A pooled analysis of nine studies revealed no association of the variant 'T' allele with breast cancer (34). Our results confirm the

absence of an association between the presence of the variant 'T' allele and BC risk (Tables III-V). The lack of an association between this variation and BC is further supported by the fact that no significant difference in DNA repair activity was found between cells expressing the wild-type Thr protein of *Xrcc3* and the Met variant (63).

*Xrcc3* has also been extensively tested for its association with *in vivo* radiosensitivity after radiotherapy in a variety of cancers (reviewed in ref. 64). The majority of the studies failed to note a correlation between the c.722 C>T variant and radiotherapy-induced acute complications (64). However, in non-cancer subjects the protein encoded by the gene harboring the c.722 C>T variant has been found to repair radiation-induced damage with significantly less efficiency than its wild-type counterpart (65). In patients treated for gynaecological tumors, the c.-316 A>G SNP exhibited an association with incidence of cancer, but not with the development of acute reactions following radiotherapy. In the same study, the c.562-14 A>G variation exhibited no correlation with cancer incidence, but the variant 'G' allele was predisposed to more severe acute complications following radiotherapy (66). When genotype distributions were compared in radiosensitive vs. non-radiosensitive BC patients (Table VI), the positive ORs exhibited by c.562-14 A>G and c.722 C>T may indicate that these SNPs are correlated with *in vitro* CRS. However, risk allele analysis results were inconclusive.

In conclusion, the effects of SNPs in *Rad51* and *Xrcc3* on BC risk remain to be elucidated. Nevertheless, findings of this study revealed that low-penetrant variations in *Rad51* and *Xrcc3*, two proteins belonging to the HR repair pathway, may modify BC risk in patients already carrying a pathological mutation in the highly penetrant BC genes *BRCA1* and *BRCA2*. Combined risk genotype analysis also revealed that *Rad51* SNPs enhanced BC risk in *BRCA2* patients, whereas *Xrcc3* SNPs enhanced BC risk in *BRCA1* patients and in patients with hereditary BC. When the four putative RGs of *Rad51* and *Xrcc3* were combined, a positive significant OR was observed for the total patient population. This finding is supportive of a polygenic model whereby combinations of weak variations are responsible for enhanced breast cancer risk. Since the different subpopulations of BC patients used in this report were relatively small, larger sample sizes are required to confirm the conclusions (study in progress).

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