

# Characterization of three alternative transcripts of the *BRCA1* gene in patients with breast cancer and a family history of breast and/or ovarian cancer who tested negative for pathogenic mutations

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**Abstract.** The study of *BRCA1* and *BRCA2* genes and their alterations has been essential to the understanding of the development of familial breast and ovarian cancers. Many of the variants identified have an unknown pathogenic significance. These include variants which determine alternative mRNA splicing, identified in the intronic regions and those are capable of destroying the splicing ability. The aim of this study was to detect *BRCA1/BRCA2* aberrant transcripts resulting from alternative splicing, in women with a known family history and/or early onset of breast and/or ovarian cancer, tested wild-type for *BRCA1* and *BRCA2*. The identification and characterization of aberrant transcripts through the analysis of mRNA levels in blood lymphocytes may help us to recognize families otherwise misclassified as wild-type *BRCA1* and *BRCA2*. Blood samples were collected from 13 women that had a family history of breast and/or ovarian cancer and tested negative for pathogenic mutations in the *BRCA1* and *BRCA2* genes. Total RNA was analyzed for the presence of *BRCA1* and *BRCA2* naturally occurring and pathological transcripts using RT-PCR. In 2 out of the 13 samples, 2 alternative transcripts of the *BRCA1* gene were identified. These were probably pathogenic as they lacked exon 17 and exon 15, respectively, giving rise to a truncated protein. In addition to these, we identified the  $\Delta 17-19$  transcript in 1 patient, which gives rise to a protein with an in-frame deletion of 69 amino acids. In conclusion, this study on alternative transcripts of the *BRCA1* and *BRCA2* genes

revealed the presence of isoforms (prevalence of 15%) in blood samples from women with breast and ovarian cancer that were probably pathogenic, that were not detected by conventional methods of mutation screening based on direct sequencing of all coding regions, intron-exons junctions and MLPA analysis.

## Introduction

Inactivating mutations in the *BRCA1* (MIM 113705) and *BRCA2* (MIM 600185) genes confer a high risk of developing breast and ovarian cancer (1,2). Both genes have a contribution of approximately 16% to the risk of familial breast cancer (3). Genetic testing for *BRCA1* and *BRCA2* provides valuable information for determining the clinical management of patients with breast/ovarian cancer. However, the data provided are difficult to interpret due to the identification of many DNA variants of unknown pathological significance or unclassified variants (UVs) that hamper genetic counseling in hereditary breast and ovarian cancer (HBOC) (4).

UVs have the potential to alter protein function by altering the coding sequence of a transcript, or the level of the gene transcript by disrupting regulatory regions in promoters, untranslated regions, exons or introns (5). Such regulatory variants include those affecting the normal splicing of *BRCA1* and *BRCA2*, many of which have been shown to be clinically significant using cDNA studies and multifactorial likelihood analysis methods that combine bioinformatics, as well as pathological and clinical information (6,7).

Assessing the impact of UVs on splicing is key to determining their pathogenicity. The accuracy of pre-mRNA splicing is determined by the recognition of well known 5' and 3' splice site consensus sequences. However, more discrete elements are also involved, such as exonic splicing enhancers (ESEs) that enhance pre-mRNA splicing when present in exons (8). As a result, each UV may potentially affect normal pre-mRNA splicing and be deleterious through the disruption of consensus sequences, or through the creation of *de novo* sequences or the alteration of splicing regulatory

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elements (9). Several *BRCA1* isoforms have been identified in different tissues; however, their functional significance is not yet fully understood (10-15).

A recent study on the comprehensive annotation of *BRCA1* splice junctions identified 63 independent alternative splicing events in RNA samples from healthy control individuals. Among these, 10 were predominant ( $\Delta 1Aq$ ,  $\Delta 5$ ,  $\Delta 5q$ ,  $\Delta 8p$ ,  $\Delta 9$ ,  $\Delta 9-10$ ,  $\Delta 9-10-11$ ,  $\Delta 11q$ ,  $\Delta 13p$  and  $\Delta 14p$ ) and represented 5-30% of the full-length signal; 48 were minor and 5 were non-classifiable events (16).

Alternative splicing is a highly coordinated process. Mutations destroying the 5' and 3' splice site consensus sequences may alter the splicing patterns of one or more transcripts, interrupting the production or function of the encoded protein (17). A large number of pathological transcripts of the *BRCA1* and *BRCA2* genes, deriving from a mutation in the consensus regions, have been described in the literature (18).

The aim of this study was to identify aberrant transcript variants resulting from the alternative splicing of *BRCA1* and *BRCA2* genes in RNA extracted from blood lymphocytes from women with a family history of and/or early onset breast and/or ovarian cancer, in which genomic pathogenic alterations in *BRCA1* and *BRCA2* have not been detected by conventional analysis.

The analysis of all possible transcripts of the *BRCA1/BRCA2* genes may allow us to uncover mRNA splicing defaults overlooked by conventional protocols and to confirm that the alternative splicing of the *BRCA1* and *BRCA2* genes plays an important role in *BRCA1/2*-driven tumorigenesis. This approach has the potential to complete the process of the characterization of mutations of *BRCA1* and *BRCA2* in HBOC.

## Materials and methods

**Sample acquisition.** A total of 13 blood samples were collected from women with a family history of and/or early-onset breast and/or ovarian cancer and who tested negative for pathogenic mutations in the *BRCA1* and *BRCA2* genes. Sample data details and characteristics are presented in Table I.

The BRCAPRO (<http://www4.utsouthwestern.edu/breast-health/cagene/>) and BOADICEA (<https://pluto.srl.cam.ac.uk/cgi-bin/bd2/v2/bd.cgi>) (19,20) programs were used, calculating *a priori* mutation carrier risk of >10% for each patient.

Ten blood samples from healthy women, aged 25-45 years, with no family history of breast and/or ovarian cancer, were used as the controls. All patients and healthy donors were recruited at the Interdepartmental Centre for Cancer Genetics, University Hospital of Santa Chiara in Pisa, Italy between 2004 and 2011. Prior to enrollment, informed consent for genetic analysis was obtained from all patients. All the experiments carried out complied with the current laws of the country in which they were performed (Italy).

**RNA extraction and cDNA synthesis.** All samples were subjected to total RNA extraction in 2 steps: peripheral blood mononuclear cells (PBMCs) were isolated from ethylenediaminetetraacetic acid (EDTA)-treated peripheral blood by standard density gradient centrifugation (Ficoll 15%; Bio-Rad Medical Diagnostics GmbH, Dreieich, Germany) and RNA extraction was obtained using a TRI Reagent kit (Molecular

Research Center, Inc., Cincinnati, OH, USA) as recommended by the manufacturer. Isolated RNA was quantified using a NanoDrop spectrophotometer (NanoDrop Technologies, Inc., Wilmington, DE, USA) and examined for integrity on a 1.5% agarose/formaldehyde gel containing 0.5  $\mu\text{g/ml}$  ethidium bromide (RNA Analysis Notebook; Promega Corp., Madison, WI, USA). First-strand cDNA was synthesized from at least 1,000 ng of total RNA using an oligo(dT) primer or random primer and SuperScript III reverse transcriptase (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions.

**Reverse transcription-polymerase chain reaction (RT-PCR).** To perform the amplification of naturally occurring transcripts of the *BRCA1* and *BRCA2* genes, we used multiple combinations of forward and reverse primer pairs to amplify overlapping regions of the mRNA and to cover the entire open reading frame (Figs. 1 and 2). For each PCR reaction, 2  $\mu\text{l}$  of cDNA were used. The PCR conditions were as follows: 5 min at 95°C followed by 40-45 cycles at 95°C for 1 min, melting temperature according to primer pair for 30 sec, 72°C for 1 min followed by 72°C for 1 min. To encompass multiple exons for large regions, Long Range PCR (Expand Long Template PCR system; Roche, Indianapolis, IN, USA) was used. The PCR products, eventually isolated on agarose gels, were sequenced on both strands using the BigDye Terminator v3.1 Cycle Sequencing kit and the 3130xl Genetic Analyzer (both from Life Technologies, Foster City, CA, USA). The electropherograms were analyzed using SeqScape Software v2.6 (Life Technologies).

**Multiplex ligation-dependent probe amplification (MLPA).** To exclude large genomic deletions in the *BRCA1* and *BRCA2* genes, MLPA (MRC-Holland, Amsterdam, The Netherlands) was performed as recommended by the manufacturer's instructions. The electropherograms were analyzed using GeneScan (Life Technologies) and Coffalyser software (MRC-Holland).

## Results

The aim of this study was to identify alternative transcripts of the *BRCA1* and *BRCA2* genes resulting from aberrant splicing events. The analysis was conducted on 13 total RNA samples extracted from PBMCs from women with a family history of and/or early-onset breast and/or ovarian cancer, more specifically, 5 cases of hereditary breast cancer (HBC), 4 cases of HBOC, 1 case of bilateral carcinoma, 1 case of breast/ovarian cancer and 2 cases of early-onset breast cancer (Table I).

All women were affected by breast cancer, 1 women by breast and ovarian cancer, 3 by bilateral breast cancer and 2 by early-onset breast cancer. A total of 10 control RNA samples from healthy women, aged between 25 and 45 years with no family history of any form of cancer, were also included in this study.

The genomic DNA of each patient was analyzed by direct sequencing of the entire open reading frame, 5' and 3'UTRs and exon/intron junctions of both genes. Approximately 100 bps from the 5' and 3' end of each intron were sequenced. All cases tested negative for the presence of germline mutations in the *BRCA1* and *BRCA2* genes. The presence of variants

Table I. Summary of patient data and family history of cancer.

Patient	Personal history (age at onset, years)	Family history		
		No. of breast cancer cases	No. of ovarian cancer cases	No. of cases of other types of cancer
P1	Br (57)	4	-	1
P2	Br (34)	1	-	1
P3	Br (43)	8	1	5
P4	Br (37)	1	1	4
P5	Br (55), Ov (61)	-	-	2
P6	Br (35)	3	-	1
P7	Br (55)	3	1	4
P8	Br bil (43)	4	1	3
P9	Br (56)	3	-	2
P10	Br bil (45 and 50)	4	-	9
P11	Br (25)	-	-	1
P12	Br (32)	-	-	1
P13	Br bil (38 and 48)	-	-	6

Br, breast cancer; Ov, ovarian cancer; bil, bilateral.

EXON	PRIMERS 5'→3'	Amplicons bp	Annealing (C°)
BR1 2F BR1 7R	GCTCTTCGCGTTGAAGAAGT3' 5'TCTTTTGGCACGCGTTTCTGT	396	58
BR1 6F BR1 9R	5'TCAGCTTGACACAGGTTTGG3' 5'CTTGATCTCCCACACTGCAA	329	57
BR1 8F BR1 11aR	GAGGACAAAGCAGCGGATAC3' 5'GCTGTAATGGGCTGGCATGA	359	57
BR1 8F BR1 12R	GGAAACCAAGTCTCAGTGCCA3' 5'CTGAGAGGATAGCCCTGAGC	3800	57
BR1 11bF BR1 15R	5'GAACGGGCTTGGAAAGAAA3' 5'GAGCAACTGTGCATGTACCAC	499	57
BR1 13F BR1 16R	GACTCTTCTGCCCTTGAGGA3' 5'GGTCATCAGAGAAGAGGCTGA	421	54
BR1 15F BR1 17R	5'GCACAGTTGCTCTGGGAGTC 5'GGCAAAGTTGACACGAGCA	478	57
BR1 16F BR1 21R	AGAGTCCAGTGTCTGTCTCAT3' GGCCATAGCAACAGATTTTC	600	59
BR1 20F BR1 24R	GAAGAAACCAAGGTTCA3' AAGCTCATCTTGGGGTCTCT	413	54

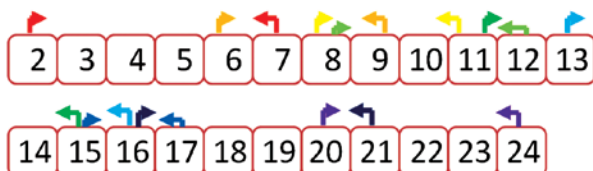


Figure 1. Primers used for the scanning of exons in the *BRCA1* gene. Primer pairs were used to amplify the cDNA covering the entire coding region of the *BRCA1* gene overlapping the exonic region.

EXON	PRIMERS 5'→3'	Amplicons bp	Annealing (C°)
BR2 2F BR2 6R	5'CTTATTTTACCAAGCATTGGAGAA3' 5'AGCTAAAGAACTTGACCAAGAC	632	55
BR2 2F BR2 9R	5'CCGCTGTACCAATCTCCTGT3' 5'ATCATTITTTTCTCAGACTTTCATCA3'	468	55
BR2 8F BR2 10R	5'TCAGAAATGAAGAAGCATCTGA3' 5'GCGTTTGGCTTCATGGAAAAT3'	382	50
BR2 9F BR2 11R	5'TCAAAGAGAAGCTGCAAGTCA3' 5'AAAGAGCTAGTTAAGGACAAAGTTGG3'	1218	54
BR2 10F BR2 12R	5'CAGCCAGTTTGAAGCAAAT3' GATTTTCTATTATCCTGTCAAATTCAT3'	5317	51
BR2 11F BR2 14R	5'CGAAAATGAGGAAATGGTTTTG3' 5'CTTGTAGCAGAACTTGATAAAATGG3'	396	51
BR2 13F BR2 15R	5'TTCTTTAGAGCCGATTACCT3' 5'GCATTCTGAAGACTTGTAAT3'	486	50
BR2 14F BR2 20R	CATTGATGGACATGGCTCTG3' 5'TTGCTGCTTCTTTCTCC3'	1250	52
BR2 19F BR2 24R	TTCTCTGCCCTTATCATCG3' 5'CCTCAGAACAAAGATGGCTGA3'	813	53
BR2 23F BR2 27R	5'TGAGTATTTGGCGTCCATCA3' 5'CCTTTGGCCATACAAAGTGA3'	752	53



Figure 2. Primers used for the scanning of exons in the *BRCA2* gene. Primer pairs were used to amplify the cDNA covering the entire coding region coding of the *BRCA2* gene overlapping the exonic region.

of unknown pathological significance was also excluded from our analysis. The presence of large genomic deletions or rearrangements was excluded by MLPA of DNA extracted from the peripheral blood lymphocytes of all patients and the controls.

The *BRCA1* and *BRCA2* mRNA in each patient was analyzed by dividing the cDNA into 9 amplicons for *BRCA1* and 10 amplicons for *BRCA2*. The size of the full-length transcripts of the *BRCA1* (5,592 bp) and *BRCA2* (10,987 bp) genes prevents the analysis of the cDNA as a single amplicon.

*Naturally occurring transcripts of BRCA1.* The cDNA of the *BRCA1* gene was amplified with primers localized in exonic sequences so that partially overlapping amplified products were obtained. The primers were selected in order to highlight all predominant naturally occurring alternative splicing isoforms and 28 of the minor transcripts, starting from exon 2, as previously reported by Colombo *et al* (16).

Under our experimental conditions, the following naturally occurring alternative transcripts were observed: the  $\Delta 9$ -10 transcript was detected in 3 samples, the  $\Delta 9$  transcript was detected in 1 patient and  $\Delta 14p$  was detected in all the samples. Using the long range PCR approach, the following transcripts were detected:  $\Delta 11q$  (transcript lacking exon 11 except the initial 121 bp) in 10 samples and  $\Delta 9$ -10-11q (transcript lacking exons 9, 10 and 11 except the initial 121 bp) in 2 cases only.

*Naturally occurring transcripts of BRCA2.* The cDNA of the *BRCA2* gene was amplified with primers localized in exonic sequences so that partially overlapping amplified products were obtained. A total of 5 predominant naturally occurring transcripts have been previously identified:  $\Delta 4$ ,  $\Delta 4$ -7,  $\Delta 17$ -18,  $\Delta 18$  and  $\Delta 20$  (7,18,21). Under our experimental conditions, all 5 alternative transcripts were detected: More specifically, the  $\Delta 4$  transcript was detected in 2 patients, the  $\Delta 4$ -7 transcript was detected in 2 patients, the  $\Delta 17$ -18 transcript was detected in 2 patients and the  $\Delta 18$  transcript was detected in 7 patients. The  $\Delta 20$  transcript was also detected in 1 case. All these *BRCA1* and *BRCA2* naturally occurring transcripts were detected in the controls.

*Abberant transcripts.* In addition to the above-mentioned predominant naturally occurring transcripts, we detected 3 aberrant transcripts in the *BRCA1* gene in 2 patients (2 in patient P6 and 1 in patient P7). No aberrant transcripts were detected in the *BRCA2* gene.

In patient P6, following the amplification of the cDNA of *BRCA1* with primers localized in exons 16 and 21 in addition to the full-length transcript (600 bp), 2 additional transcripts of approximately 500 and 400 bp were obtained. These transcripts were not detected in the 10 healthy control samples. The aberrant transcripts were gel-purified and then sequenced as described in the Materials and methods. Sequence analysis allowed us to detect the presence of a transcript lacking exon 17 and another transcript lacking exons 17, 18 and 19 (Fig. 3). The transcript containing the deletion of exon 17 produced an abnormal stop signal at codon 1673 (HGVS codification: p.Val1665Serfs\*8) and then a truncated protein lacking the last 192 amino acids. The transcript containing the deletion of exons 17, 18 and 19 did not produce an abnormal stop signal, lost 207 nucleotides and retained the open reading frame, producing a protein lacking 69 amino acids.

In patient P7, following the amplification of the cDNA of the *BRCA1* gene with primers localized in exon 13 and 16 in addition to the full-length transcript of approximately 420 bp, an additional transcript of approximately 220 bp was obtained (Fig. 4). Direct sequencing of this transcript allowed us to detect an exon 15 deletion. The transcript containing the deletion of exon 15 produced an abnormal stop signal at codon 1510 (HGVS codification: p.Ser1496Glyfs\*14) and then a truncated protein lacking the last 405 amino acids.

*Phenotype-genotype correlation.* Patient P6, at the age of 35 years, developed an infiltrating ductal carcinoma of the right breast, which was estrogen receptor-positive, progesterone receptor-negative and Her2/neu 3+, with loco-regional lymph node metastasis. The patient's sister had been diagnosed with breast cancer of a similar phenotype at the age of 34. These were not the only cases of breast cancer in the family; the grandmother and maternal aunt (their mother's twin sister) had also been afflicted by the disease. Segregation analysis of the aberrant transcript in this family was not possible as DNA samples were not available.

Patient P7 was diagnosed with breast cancer at the age of 55 years. The tumor was an infiltrating breast carcinoma of the right breast, and was estrogen receptor-positive, progesterone receptor-negative and Her2/neu-negative, without any lymph node metastasis. The patient's mother had been affected by ovarian cancer at 77 years of age and 3 cases of breast cancer were reported in the family: 2 sisters of the mother and the mother's cousin (36 years of age). On the mother's side, an uncle had been affected by malignant melanoma, and an aunt by a brain tumor. These relatives were not available for co-segregation analysis, but this aberrant transcript was not detected in the 42-year-old healthy daughter.

## Discussion

This study focused on the possibility that a proportion of patients with HBOC have mutations in the *BRCA1* or *BRCA2* genes that affect splicing, but are not detectable through sequencing of the gene exons or intronic sequences near the intron-exon boundaries. Several studies have demonstrated that, due to variations in splice sites, *BRCA1/BRCA2* may generate truncated, non-functional proteins that may be associated with a predisposition to breast and ovarian cancer (22). However, to the best of our knowledge, no studies have evaluated the presence of *BRCA1/BRCA2* pathological transcripts in patients without mutations identified in the canonical splice sites or regulatory sequences.

Recently, a systematic description of 'naturally occurring' alternative splicing at the *BRCA1* locus was conducted by the Evidence-based Network for the Interpretation of Germline Mutant Alleles (ENIGMA) consortium (16). This led to the annotation of 63 splicing events, of which 35 were novel findings, even though most of them are rather minor, and it is likely that some do not qualify as 'naturally occurring' events, suggesting that the characterization of the full complexity of *BRCA1* splicing requires further investigation (16).

The present study was performed in accordance with a standard assay design and detection methods formulated by ENIGMA Consortium members (23) in order to detect the predominant naturally occurring alternative splicing isoforms of *BRCA1*, as reported by Orban *et al* (13,14), which were the only data available at the moment of the study design. There is no standard assay design for the detection of the predominant naturally occurring alternative splicing isoforms of *BRCA2*; thus we referred to the transcripts described in Ensembl (<http://www.ensembl.org/index.html>).

Using this strategy, we detected the  $\Delta 9$ ,  $\Delta 9$ -10,  $\Delta 11q$ ,  $\Delta 9$ -10-11q and  $\Delta 14p$  *BRCA1* isoforms and the  $\Delta 4$ ,  $\Delta 4$ -7,  $\Delta 17$ -18,  $\Delta 18$  and  $\Delta 20$  *BRCA2* isoforms. In addition these predomi-

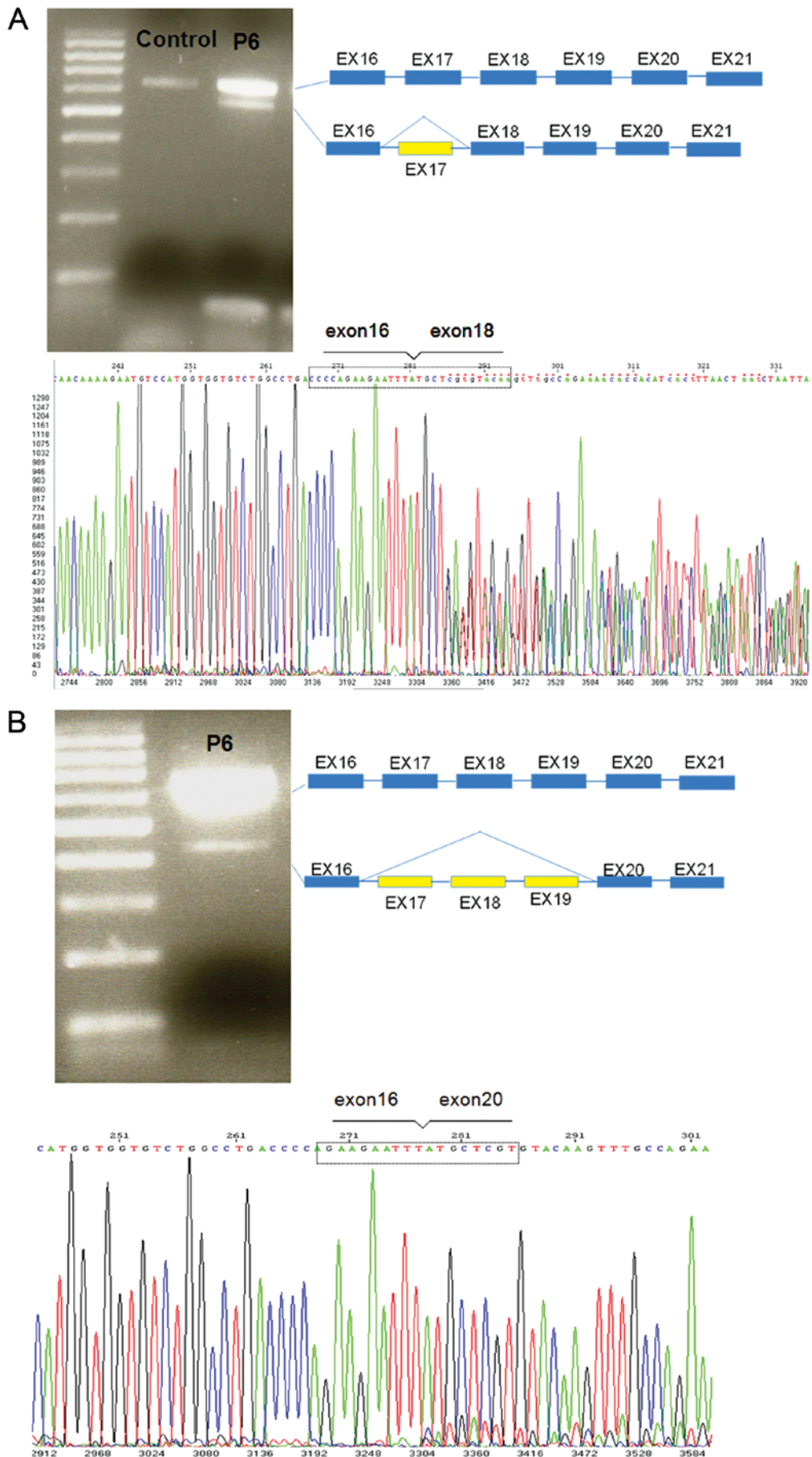


Figure 3. Alternative transcript isoforms in patient P6. Following the amplification of the cDNA of *BRCA1* with primers localized in exons 16-21, the wild-type (wt) transcript (~600 bp) and 2 additional bands of (A) ~500 bp and (B) ~400 bp were obtained. Sequencing analysis allowed us to detect the presence of a transcript lacking (A) exon 17 and another transcript lacking (B) exons 17, 18 and 19.



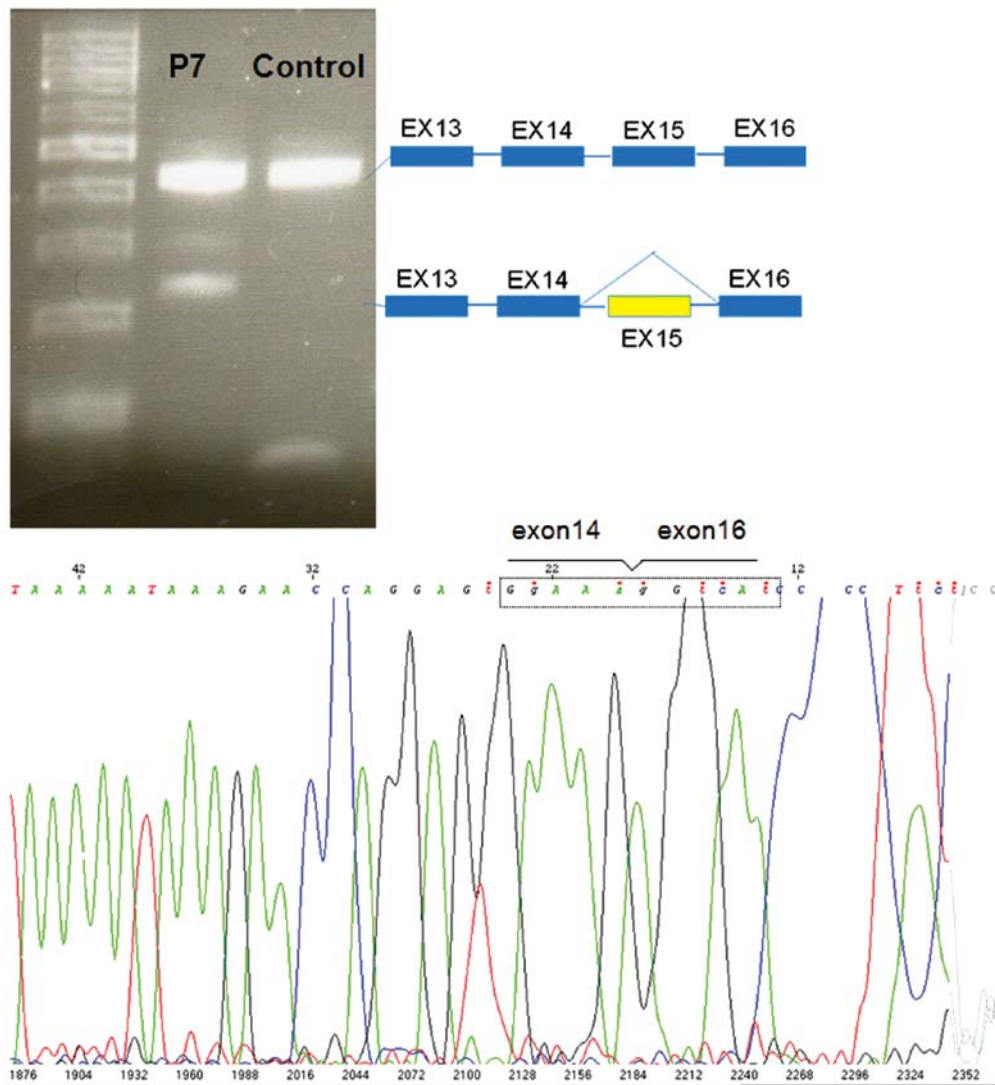


Figure 4. Alternative transcript isoform in patient P7. Following the amplification of the cDNA of *BRCA1* with primers localized in exons 13 and 16 with primers 13F and 16R, an additional band of ~420 bp was obtained. The fragment had a deletion of exon 15 as detected by direct sequencing.

nant *BRCA1* transcripts, we detected the  $\Delta 17$  and the  $\Delta 17-19$  isoforms in patient P6 and the  $\Delta 15$  isoform in patient P7. No minor transcripts were detected for *BRCA2*.

The  $\Delta 17$  transcript lost the open reading frame, leading to the formation of a truncated protein. The  $\Delta 17$  transcript was not detected in any of the 10 healthy controls in our study; however, it has previously been found in 28 out of 103 (27%) independent control samples or technical replicas in the ENIGMA Consortium study (16). The  $\Delta 17$  transcript, due to the genomic deletion of exon 17, has frequently been detected in Italian families (24) possibly as a result of a non-homologous rearrangement among 'Alu repeats'. The  $\Delta 17$  transcript is not derived from a genomic rearrangement as demonstrated by MLPA of the patient's DNA. The  $\Delta 17$  transcript, even if not derived from a genomic rearrangement or from aberrant mRNA maturation due to canonical splice site mutations, it has to be considered as pathogenetic as it encodes for a truncated protein.

The  $\Delta 17-19$  transcript gave rise to a protein with an in-frame deletion of 69 amino acids. The  $\Delta 17-19$  isoform was not detected in either our 10 healthy control samples or the

223 independent control samples and technical replicas in the ENIGMA Consortium study (16). Although the  $\Delta 17-19$  transcript had an in-frame deletion, a recent study demonstrated that the expression of the human *BRCA1* alternative splicing variant,  $\Delta 17-19$ , in the MCF-7 cells resulted in an impaired assembly of DNA repair complexes and aberrant DNA damage response (25), thus contributing to an increased risk of developing cancer in the carrier subjects.

In patient P7, the *BRCA1* alternative transcript  $\Delta 15$  was detected. The alternative  $\Delta 15$  transcript lost the open reading frame and produced an abnormal stop signal to position p.Ser1510Glyfs\*14, leading to a truncated protein lacking the last 405 amino acids. To evaluate the presence of  $\Delta 15$  at the genomic level, we performed MLPA, which did not reveal any deletion. This isoform was not detected in our 10 healthy PBMC control samples or in the 8 PBMC samples analyzed by Colombo *et al* (16). They found this transcript in only 4 of 10 replicas derived from samples of leukocytes, PHA-stimulated peripheral blood leukocytes and lymphoblastoid cell lines (they did not specify if these were independent samples or replicas of the same sample).

In conclusion, the present study on 13 patients with a family history of breast and/or ovarian cancer detected 3 alternative transcripts, probably pathogenic and not predictable by genomic screening, in 2 patients. A number of studies have described these 3 transcripts as consequences of large genomic rearrangements or mutations in canonical splice sites (24-28). These aberrant alternative transcripts may undergo nonsense-mediated decay (NMD) as they give rise to truncated proteins (29); therefore, the accurate quantification of these alternative transcripts is crucial to discriminating what is considered pathological and what is neutral for clinical relevance. The relative abundance of the minor transcripts detected in this study may prove to be useful in evaluating their possible role in cancer predisposition and risk modification.

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