Identification of a founder BRCA2 mutation in Sardinia

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Summary Sardinian population can be instrumental in defining the molecular basis of cancer, using the identity-by-descent method. We selected seven Sardinian breast cancer families originating from the northern-central part of the island with multiple affected members in different generations. We genotyped 106 members of the seven families and 20 control nuclear families with markers flanking BRCA2 locus at 13q12–q13. The detection of a common haplotype shared by four out of seven families (60%) suggests the presence of a founder BRCA2 mutation. Direct sequencing of BRCA2 coding exons of patients carrying the shared haplotype, allowed the identification of a ‘frame-shift’ mutation at codon 2867 (8765delAG), causing a premature termination-codon. This mutation was found in breast cancer patients as well as one prostate and one bladder cancer patient with shared haplotype. We then investigated the frequency of 8765delAG in the Sardinian breast cancer population by analysing 270 paraffin-embedded normal tissue samples from breast cancer patients. Five patients (1.7%) were found to be positive for the 8765delAG mutation. Discovery of a founder mutation in Sardinia through the identity-by-descent method demonstrates that this approach can be applied successfully to find mutations either for breast cancer or for other types of tumours. © 2000 Cancer Research Campaign

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Breast cancer is the most common malignancy in women, with an incidence that varies between 40 and 90 per 100 000 (standardized rate) worldwide. Breast cancer is the most frequent female tumour in Italy, representing about 25% of all female tumours as reported in Italian registries (Zanetti et al, 1997).

A positive family history is known to be a high risk factor for developing the disease: 5–10% of all breast cancers arise in individuals carrying a germline mutation and are usually considered hereditary forms (Claus et al, 1991). Two major breast cancer-susceptibility genes, BRCA1 and BRCA2, have been cloned (Miki et al, 1994; Wooster et al, 1995) and both are thought to account for 30–60% of hereditary breast cancer (Serova et al, 1997; Szabo et al, 1997; Vehmanen et al, 1997a, 1997b). However, large-scale mutation analyses conducted in several populations suggest the existence of additional breast cancer-susceptibility gene(s). BRCA1 mutations are responsible for the majority of familial breast cancer associated with ovarian carcinoma, for about 50% of cases with breast cancer alone and for very few male breast cancer cases (Easton et al, 1993; Stratton et al, 1994; Narod et al, 1995). It has been estimated that women carrying a germline mutation in BRCA1 have a risk ranging from 80 to 90% for developing breast cancer and from 44 to 63% for developing ovarian cancer (Easton et al, 1993, 1995; Ford et al, 1994; Miki et al, 1994; Wooster et al, 1994). BRCA2 mutations account for a similar proportion of inherited breast cancer and are frequently associated with male breast cancer (Wooster et al, 1995). Breast cancer risk in females carrying BRCA2 mutations is calculated to be similar to that conferred by BRCA1 mutations (Easton et al, 1993, 1997; Ford et al, 1994; Miki et al, 1994; Wooster et al, 1994). BRCA1 and particularly BRCA2 families are often affected by other tumours such as prostate, liver, pancreas, lung, stomach and colorectum (Wooster et al, 1995; Gudmundsson et al, 1996; Phelan et al, 1996; Thorlacius et al, 1996; Vehmanen et al, 1997b; Tonin et al, 1998). Except for higher incidences of ovarian cancer in families with mutations in a 3.3-kb region of exon 11 of BRCA2 (the so-called ovarian cancer cluster region [OCCR]; Gayther et al, 1997), no other significant association between genotype and phenotype was described. BRCA1 and BRCA2 mutations are for the most part frame-shifts due to small deletions leading to premature translation termination (Wooster et al, 1995; Phelan et al, 1996; Tavtigian et al, 1996; Gayther et al, 1997).

Some of these mutations are prevalent in genetically homogeneous populations as a consequence of a founder effect. A single BRCA2 mutation accounts for the majority of hereditary breast cancer in Iceland (Gudmundsson et al, 1996; Thorlacius et al, 1996) and for 40% of male breast cancer cases (Johannesdottir et al, 1996), whereas three different founder mutations (185delAG and 5382insC in BRCA1, and 6174delT in BRCA2) have a high frequency in Ashkenazi Jews (Roa et al, 1996). Although at different rates, BRCA1 and BRCA2 founder mutations have been detected in other genetically homogeneous populations, such as the Finns (Vehmanen et al, 1997a) and the French-Canadians.
(Tonin et al, 1998). In Sardinia, epidemiological data from the Regional Tumor Registry (accounting for the northern part of the island) indicate that breast carcinoma is the principal death-causing malignancy, with an incidence of 93 per 100 000 inhabitants (standardized rate) (Budroni et al, 1998). Sardinian population is genetically separated from that of the rest of Italy as well as from other European populations, due to strong genetic drift. All the monogenic disorders analysed, such as thalassemia (Pirastu et al, 1987), seem to be associated with a single founder mutation throughout the island. Therefore, it seems possible that such a founder effect could also be identified for a complex disease like cancer. This can be done by tracing back the mutation by linkage disequilibrium with genetic markers which give rise to a shared haplotype among the patients. On this basis, we decided to analyse the \textit{BRCA2} gene using the identity-by-descent method, which allowed the identification of a mutation with founder effect in Sardinian breast cancer families.

**MATERIALS AND METHODS**

**Breast cancer patients**

Collaborating physicians at both the Department of Medical Oncology and the Institute of Histo-Pathology at Sassari University collected seven Sardinian families. Family ascertainment was carried out using the following criteria: (a) families with at least two affected members in different generations (either a first-degree relative or relative affected before age 50), or (b) families with at least three affected members. Clinical information was obtained from medical records.

Families were all apparently unrelated and originated from different small villages located in the northern-central part of the island; none of them presented cases of ovarian cancer. Three families had other forms of cancer (Figure 1). No breast cancer was detected in male members of the pedigrees. Blood samples were collected from 17 affected (15 breast cancer, one prostate cancer and one bladder cancer) and 89 unaffected members.

Twenty unrelated nuclear families, originating from the same geographical area with no history of breast cancer, were used as controls for the haplotype study.

Paraffin-embedded normal tissues were obtained from 270 breast cancer patients consecutively collected during 1997. No additional selection criteria were used to enrol patients in the screening; all cases were included regardless of age of onset. Sardinian origin was ascertained in all cases through genealogical studies. Informed consent was obtained from each family member before drawing blood.

**DNA analysis**

DNA was isolated from blood samples using standard methods (Sambrook et al, 1989). DNA extraction from paraffin-embedded tissue was performed by a modification of the Jackson et al (1989) procedure. Briefly, single 7- to 8-mm tissue sections, cut from paraffin blocks, were stirred for 30 min with 1 ml of xylene in 1.5-ml tubes and centrifuged. The pellet was washed with ethanol, air-dried and resuspended in lysis solution (0.5% sodium dodecyl sulphate (SDS), 0.5 mg ml\(^{-1}\) proteinase K in 1× TE buffer). After incubation at 37°C overnight and inactivation of proteinase K for 15 min, DNA was extracted with 1 vol of phenol, phenol–chloroform and chloroform. The supernatant was precipitated at −20°C overnight. The DNA was washed with 70% ethanol, air dried and resuspended in 10 mM Tris–HCl pH 7.5, 0.1 mM EDTA.

Polymorphic microsatellite markers used for haplotype analysis are linked to \textit{BRCA2} gene at 13q12–q13 as reported in published genetic and physical maps: cen-D13S1246-D13S289-D13S260-D13S1698-\textit{BRCA2}-D13S1701-D13S171-D13S267-D13S263-tel (Couch et al, 1996; Vehmghan et al, 1997a; Neuhausen et al, 1998; Marshmed map at http://www.marshmed.org). Polymerase chain reactions (PCR) were carried out as suggested in the Human Genome Database. PCR products were end-labelled with \(\gamma\)-\[^{32}\text{P}\]dATP and electrophoresed on 6% acrylamide/7M urea sequencing gels. Alleles, visualized by X-ray autoradiography, were numbered according to size for each microsatellite repeat marker.

**\textit{BRCA2} sequence analysis**

Nucleotide sequencing of the entire \textit{BRCA2} coding regions was initially performed in two patients with an identical haplotype belonging to two different families. DNA was amplified with primers specific for \textit{BRCA2} exons (sequences and conditions are reported in the Human Genome Database). PCR products were gel purified using Qiaquick spin columns (Qiagen) and sequenced by Thermo Sequenase \(^{32}\text{P}\)-labelled terminator cycle sequencing kit (Amersham Pharmacia Biotech).

**Mutation screening**

A new set of primers (F, 5′-GTGTAACACATTATTACAGT-3′ and Rv, 5′-AATTCTCCCTGAATTTTAGT-3′) was generated in order to bracket the region containing the mutation. Amplification conditions were: 94°C for 5 min, 30 cycles of 94°C for 30 s, 53°C for 30 s, 72°C for 1 min and 72°C for 10 min. Samples were electrophoresed on a 6% denaturing polyacrylamide gel for 2 h and the 2-bp deletion mutation was visualized as the faster migrating fragment after silver staining.

**RESULTS**

Among the seven unrelated Sardinian families, 106 members, including 15 breast cancer cases, were genotyped with markers flanking the \textit{BRCA2} locus at 13q12–q13. Pedigrees of all selected families are shown in Figure 1. The haplotypes generated with markers from D13S1246 to D13S1701 showed a pattern that was constant within each family for all affected members, as indicated in Figure 1. Four families shared the same haplotype in a 6.5 cm region from D13S1246 to D13S267. The haplotype shared by families 2 and 4 extended over a 15-cm interval reaching D13S263, the most telomeric marker (Figure 1). Interestingly, this haplotype was not detected in 80 control chromosomes except for alleles 11 and 2 of D13S171 and D13S267, respectively, found to be in linkage disequilibrium in the general Sardinian population (data not shown and Figure 1).

Patients III:18 of family 2 and patient III:6 of family 4 were affected by bladder and prostate cancer respectively. They shared the common extended haplotype of breast cancer patients. DNA samples from other family members with different tumours, including prostate, bladder, colorectal, gastric and brain (Figure 1) were unfortunately not available for analysis.

Two unrelated affected members carrying the common extended haplotype (patient III:7 of family 1 and II:2 of family 2; Figure 1) were chosen for direct sequencing of the \textit{BRCA2} coding
Table 1  Mutational screening of unselected breast cancer patients.

| Age | No. of patients | Patients positive to 8765delAG (age at diagnosis) | Additional cancer cases in positive families |
|-----|----------------|-------------------------------------------------|---------------------------------------------|
| ≤ 40 | 42             | 1 (39)                                          | 5 breast (1 male), 1 lung                    |
| 41–60 | 120            | 1 (45)                                          | 5 breast, 1 lung, 1 liver                    |
| > 60  | 108             | 1 (57)                                          | 1 breast, 1 lung, 1 colon                    |
|      |                 | 1 (69)                                          | 2 breast, 1 colon, 1 lung, 1 larynx          |
|      |                 | 1 (80)                                          | 2 breast, 1 rectum                          |

The 270 consecutively collected patients are grouped according to age at diagnosis. For the five positive cases we indicate the age of onset and the number and types of cancer present in their families.

region. An AG deletion was found in exon 20 at codon 2846 (8765delAG), in both patients (Figure 2). This mutation is predicted to produce a truncated protein at codon 2867. The 8765delAG mutation was found in the remaining patients of family 2 (patients III:2 and III:19) and family 1 (patient III:11) (Figure 1). Interestingly, in patient III:19 of family 1 (Figure 1), who carries a different haplotype (Figure 1) this mutation was not detected. After screening the other five families, patient III:5 of family 6 and patients III:7, III:12 and III:6 (prostate cancer) of family 4 as well as patient III:18 (bladder cancer) of family 2 were found positive for the presence of 8765delAG mutation as expected from the shared haplotype (Figure 1). Patients from families 3, 5 and 7 were negative for the 8765delAG mutation, confirming the haplotype results. Altogether this mutation was detected in four out of seven families (60%).

To investigate the frequency of the 8765delAG mutation, we analysed 270 paraffin-embedded normal tissues from breast cancer patients coming from the northern-central part of Sardinia. All cases were collected regardless of family history and age of onset. We found that five out of 270 patients (1.7%) carried this mutation (Table 1).

DISCUSSION

Sardinia has a relatively small, isolated, and genetically homogeneous population with a high rate of inbreeding making it ideal for genetic studies on either monogenic or multifactorial disorders. Several founder effects have already been demonstrated for monogenic diseases in this population. Therefore it seemed possible that founder mutations could also be detected in cancer patients. Analysis of family pedigrees over several generations and use of polymorphic markers may identify a common identical-by-descent haplotype, in affected individuals. This strategy could help restricting the number of cases for mutation screening, avoiding extensive analysis principally when the candidate gene is as large as BRCA2.

In the seven breast cancer families with multiple affected members in different generations, selected for our study, clinical phenotype, absence of ovarian cancer and late age-of-onset suggested BRCA2 as a candidate gene. Genotyping with markers flanking the BRCA2 gene at 13q12–q13 locus identified a large haplotype in four out of seven families, not found in control chromosomes from the same geographical area. A few patients from each family were additionally genotyped with markers closely linked to the BRCA1 gene at 17q21. We found no differences in haplotype frequency between patients and normal controls (data not shown). Presence of a founder mutation in the BRCA2 gene was confirmed by identification of a 2-bp (AG) deletion in exon 20. This mutation is located outside of the OCCR region of the BRCA2 gene, in agreement with ovarian cancer absence in our families. This AG deletion at nucleotide 8765 was already described as a founder mutation in Yemenite-Jews families (Lerer et al, 1998), as well as French-Canadian families (Phelan et al, 1996; Tonin et al, 1998). In order to understand if this mutation has a common ancestral origin we carried out a haplotype analysis of Sardinian and French-Canadian families (DNA samples of two French-Canadian 8765delAG carriers were kindly provided by P Tonin). This study showed that in the two populations the 8765delAG is associated to different haplotypes (data not shown). These results support the hypothesis that the 8765delAG mutation occurred at least twice in different populations because of its position in an AG-rich sequence which may be a mutational hot-spot.

The 8765delAG mutation was present in all affected individuals who shared the identical-by-descent haplotype. Unfortunately, some family members were not evaluated for such mutation due to individual refusal to undergo this analysis. Patient III:19 of family 1 and breast cancer patients from other families showing a different haplotype (Figure 1) were found to be negative for the 8765delAG mutation. Family 1 strikingly shows that two sisters (patients III:11 and III:19) do not have the same genotype: patient III:11, who carries the 8765delAG mutation, seems to have received it from the father’s side (because the same mutation is present in family member III:7). For patient III:19, who does not carry this mutation, we can hypothesize that she either received another mutation from the mother’s side (in which breast cancer is also present) or she is a phenocopy due to the high heterogeneity of the genetic and non-genetic factors causing the disease. Actually, the young age of the third-generation relatives and missing genotype data on some family members (i.e. the affected maternal aunt) are hindering clarification of this point.

As reported in Figure 1, two patients with other tumours (bladder carcinoma for patient III:18 of family 2 and prostate carcinoma for patient III:6 of family 4) shared the same haplotype with the breast cancer patients. In these two cases, we also found the 8765delAG mutation. The association of other tumours with breast cancer is reported by several authors (Phelan et al, 1996; Thorlacius et al, 1996; Serova et al, 1997; Tonin et al, 1998) and is confirmed in our families. It would be interesting to carry out a screening for this mutation in families with familial bladder and prostate cancers.

The frequency of 8765delAG in Sardinia was subsequently verified by screening 270 breast cancer patients consecutively...
Figure 1: Pedigrees and haplotype results. All family members for each pedigree are included. Symbol definitions are as follows: ○ unaffected, ■ breast cancer, ● prostate cancer, □ bladder cancer, △ colorectal cancer, ▼ liver cancer, ▲ brain cancer. Each selected individual is indicated with the generation identifier, age at diagnosis and specific haplotype. Arrows indicate the cases carrying the 8765delAG mutation.
Figure 1 continued
collected over a 1-year period, regardless of family history or age of onset, from the central-northern part of the island. We found five patients positive to the 8765delAG mutation with a frequency of 1.7% in this group of unselected patients (see Table 1). Preliminary results of a similar screening conducted in breast cancer patients from the southern part of the island indicate that 8765delAG mutation is present at a lower frequency (1/208; 0.5%).

The discrepancy in the frequency of the 8765delAG mutation between familial (60%) and unselected cases (1.7%) probably is due to the selection criteria used. Families were selected on the basis of the presence of several affected cases in different generations segregating a dominant trait with high penetrance. On the contrary, the mutation screening was carried out in unselected cases of which only 5–10% are expected to be familial (Claus et al., 1991). In addition, as already shown in several similar studies (Lerer et al., 1998; Tonin et al., 1998), the 8765delAG may be a founder in Italian breast cancer families. The evidence of a founder effect (De Benedetti et al., 1996; Montagna G, 1998) would be important for the detection of asymptomatic carriers. Searching for the 8765delAG mutation in these kind of families would be important for a prevention programme based on DNA analysis.

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