**Short Communication**

BRCA2 gene mutations in families with aggregations of breast and stomach cancers

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Stomach cancer ranks second to lung cancer in the global cancer burden. It is estimated that 25% of families meeting the criteria for hereditary diffuse gastric carcinoma (HDGC) will have germline mutations in the E-cadherin gene (Hizawa et al, 1993). It has also been shown that the frequency of stomach cancers in BRCA2 carriers may be sex related, as it occurs primarily in males (BCLC, 1999; Johannsson et al, 1999).

In Polish families, with a strong aggregation of breast and ovarian cancers and no other cancer, the frequency of BRCA2 mutations is very low and does not exceed 5–10% (van Der Looij et al, 2000; Grzybowska et al, 2000; Górski unpublished data).

Given the high frequency of breast and stomach cancer families observed in our familial breast cancer data base, we analysed the BRCA2 gene by direct DNA sequencing to determine the frequency and nature of BRCA2 germline mutations in families where there was a clear aggregation of breast and male stomach cancers occurring at early ages.

**MATERIALS AND METHODS**

Twenty-nine families with an aggregation of at least one female breast cancer diagnosed before the age of 50 years and one male stomach cancer diagnosed under the age of 55 years were available for study. In 28 of these families at least one additional relative was diagnosed with a malignant tumour (Table 1). In 12 families stomach cancer was diagnosed in a first-degree relative of an early onset breast cancer proband (families 1–12). In 17 breast cancer families, stomach cancer was diagnosed among second degree relatives through an unaffected woman (10 cases, families 13–22) or through an unaffected male (seven cases, families 23–29). A control population of 100 healthy unaffected unrelated individuals was used for the assessment of potential pathogenic missense changes. For a comparison of the clinical features and age of diagnosis of malignancies of the 29 families see Table 1.
Peripheral blood for DNA isolation was taken from all index patients affected by breast cancer and from the 100 healthy controls (Lahiri and Schnabel, 1993).

None of the cancer patients showed signs of the five common Polish founder BRCA1 mutations (Gorski et al., 2000). The entire coding sequence of the BRCA2 gene was amplified in 27 separate PCR reactions, using primers and conditions described previously (BCLC, http://www.nhgri.nih.gov/Intramural_research/Lab_transfer/Bic/Member/BRCA2.html), with minor modification. Instead of subdividing exons 11, 10, 14 and 27 due to their size, each were amplified in one fragment, using primers located in the adjacent introns. After purification, all PCR products were analysed on an ABI 377 DNA Sequencer with the primers described previously (BCLC,http://www.nhgri.nih.gov/Intramural_research/Lab_transfer/Bic/Member/BRCA2.html).

Standard incidence rates (SIRs) and confidence intervals (CIs) for the various cancer sites were calculated by the conventional method, which assumes all events are independent, and by a method which adjusts for intrafamilial correlation, which is to fit the same model but to calculate robust standard errors that allow for this extra source of variation. The results for the adjusted approach are affected mainly via their standard errors, but if the number of observed cases is small, the SIR estimate will also be affected. All other statistical tests were performed using the STATA statistical package.

### RESULTS

From the 29 families, three frameshifts and three variants, potentially missense mutations, in six unrelated probands representing – 20.7% of families were identified. Two of the detected frameshift mutations (886delGT, 6696delTC) have been previously reported in BIC (http://www.nhgri.nih.gov/Intramural_research/Lab_transfer/Bic/index.html). One mutation (9174delA) localised in exon 22 and the three potential missense changes represented novel variations in the BRCA2 gene (Table 1). None of these changes were present in the controls (Lahiri and Schnabel, 1993). This suggests an approach for adjusting for intrafamilial correlation, which is to fit the same model but to calculate robust standard errors that allow for this extra source of variation. The results for the adjusted approach are affected mainly via their standard errors, but if the number of observed cases is small, the SIR estimate will also be affected. All other statistical tests were performed using the STATA statistical package.

### Table 1 Occurrence of malignant tumours and detected germline alterations in tested families

| Family | No of breast cancers (age at diagnosis) | No of stomach cancers (age at diagnosis) | Other cancers | Detected alteration in DNA and consequence at protein level |
|--------|---------------------------------------|----------------------------------------|--------------|----------------------------------------------------------|
| 1      | 4 (41–46)                            | 1 (44)                                 | –            |                                          |
| 2      | 3 (45, 48)                           | 1 (36)                                 | lar, lung    |                                          |
| 3      | 3 (35–45)                            | 1 (49)                                 | –            | 886delGT, V220fs223X |
| 4      | 3 (40–50)                            | 1 (49)                                 | 2 × kid, leu |                                          |
| 5      | 3 (30–48)                            | 1 (54)                                 | Leu          |                                          |
| 6      | 2 (42)                               | 1 (50)                                 | –            |                                          |
| 7      | 2 (48, 71)                           | 2 (38, 35)                             | –            |                                          |
| 8      | 2 (47, 56)                           | 2 (40, 64)                             | pan, lung, 2 × leu |                                          |
| 9      | 1 (39)                               | 2 (44, 30)                             | –            |                                          |
| 10     | 1 (39)                               | 1 (50)                                 | liv          |                                          |
| 11     | 1 (49)                               | 2 (27, 30)                             | Panc         | 6509A/G, Y2094C*                                |
| 12     | 1 (42)                               | 4 (48–60)                              | –            |                                          |
| 13     | 3 (44–64)                            | 1 (27)                                 | –            |                                          |
| 14     | 2 (35, 36)                           | 1 (54)                                 | –            | 9174delA, K2982fs2987X*                          |
| 15     | 2 (42, 52)                           | 1 (45)                                 | FGT          |                                          |
| 16     | 2 (30, 76)                           | 1 (50)                                 | –            |                                          |
| 17     | 1 (49)                               | 2 (42, 44)                             | col, lar     |                                          |
| 18     | 1 (47)                               | 2 (48, 61)                             | pancre, lar  |                                          |
| 19     | 1 (43)                               | 2 (52, 68)                             | –            |                                          |
| 20     | 1 (39)                               | 2 (22, 52)                             | –            |                                          |
| 21     | 1 (25)                               | 1 (52)                                 | –            | 6696delTC, S2156fs2174X                         |
| 22     | 1 (41)                               | 1 (50)                                 | Leu          |                                          |
| 23     | 3 (46–67)                            | 1 (52)                                 | Lung         | 6612AC, K2128N*                                |
| 24     | 2 (47, 50)                           | 1 (42)                                 | Brain        |                                          |
| 25     | 2 (46, 64)                           | 1 (45)                                 | lar          | 6443C/G, S2072C*                                |
| 26     | 1 (36)                               | 1 (52)                                 | lung         |                                          |
| 27     | 1 (41)                               | 1 (55)                                 | bl           |                                          |
| 28     | 1 (45)                               | 1 (51)                                 | col, CSU     |                                          |
| 29     | 1 (45)                               | 1 (54)                                 | kid, pan     |                                          |

Col, colon; panc, pancreas; lar, larynx; bl, urinary bladder; kid, kidney; leu, leukaemia; liv, liver; CSU, cancer site unknown; FGT, female genital tact. *Alteration reported for the first time.
**Figure 1** Pedigrees of three families carrying the BRCA2 alterations: 3 (A), 23 (B) and 11 (C). st, stomach cancer; br, breast cancer; lu, lung cancer; pan, pancreatic cancer; numbers in brackets indicate age at diagnosis of cancer; (+), (−) indicate presence or absence of BRCA2 alteration.

In families where the occurrence of breast and stomach cancers was among 1° relatives, BRCA2 abnormalities were detected in two out of 12 cases (one frameshift, one assumed missense variation; 16.7%). In families with stomach cancers among 2° relatives, BRCA2 changes were identified in four out of 17 cases (two frameshifts, two assumed missense variations; 23.5%).

To further verify the uniqueness of the population under investigation, a statistical comparison was performed to determine if the observed cancers in the families were due to chance or most probably a result of a genetic predisposition. The relative frequency of the reported malignancies, compared to those of the general population, revealed several significant differences as shown in Table 2.

As expected in the study population breast and stomach cancers are over-represented with a high degree of confidence. Interestingly, leukaemia is also significantly over-represented followed by pancreatic cancer. All other cancers shown in Table 2 (except brain and bladder) are over-represented, but the confidence intervals are altered indicating that the statistical significance of the results are affected by familial relationships. Overall, all but two cancer types (brain and bladder) are significantly over-represented.

**Table 2** Standard incidence rates of the cancers identified in the 29 families

| Site     | Unadjusted | Adjusted |
|----------|------------|----------|
|          | SIR        | 95% CI   | SIR        | 95% CI   |
| Breast   | 576.8      | 439.5,756.9 | 576.8      | 473.7,702.2 |
| Breast   | 584.1      | 445.1,766.3 | 584.1      | 479.9,710.8 |
| Stomach  | 597.7      | 429.1,832.5 | 597.7      | 449.7,794.4 |
| Laryngeal| 134.7      | 59.3,588  | 160.1      | 449.7,571.5 |
| Kidney   | 84.8       | 27.4,263.0 | 94.8       | 242,371.8  |
| Leukaemia | 260.3     | 108.3,625.4 | 256.2      | 98.4,666.9  |
| Pancreas | 137.1      | 51.4,365.2 | 137.1      | 52.6,375.7  |
| Colon    | 40.4       | 10.1,161.6 | 40.4       | 10.3,158.7  |
| Brain    | 35.8       | 5.0,254.2  | 20.2       | 2.9,140.8   |
| Liver    | 60.1       | 8.5,426.6  | 183.4      | 25.7,131.1  |
| Lung     | 29.2       | 13.1,65.1  | 29.2       | 13.7,62.2   |
| Bladder  | 27.2       | 3.8,193.2  | 28.2       | 3.9,202.3   |
| FGT      | 212.2      | 29.9,1506.6 | 212.2      | 29.9,1504.2 |

*Expected count includes both males and females. aExpected number of cases based on lymphatic and myeloid leukaemia rates. Unadjusted SIRs do not take into account the relation between groups of patients whereas the adjusted SIRs do. The overall difference in rates is not changed, but the confidence intervals are altered indicating that the statistical significance of the results is affected by familial relationships. Overall, all but two cancer types (brain and bladder) are significantly over-represented.

**DISCUSSION**

Currently there is little information about the nature and frequency of BRCA2 constitutional mutations in families selected for the coexistence of breast and stomach cancers. In an earlier study it was reported that male stomach cancer was over-represented in BRCA2 mutation positive families. Verification of this relationship between breast and stomach cancer in our series of cases suggests that this disease constellation can be used as a phenotypic indicator for the pre-selection of families for BRCA2 testing. Unfortunately, the stomach cancer patients had succumbed to their disease and it was impossible to obtain mutation data from any of them. Notwithstanding, the stomach cancer cases were on the same transmittting lineage of the mutation, were diagnosed with disease under the age of 55 and it remains highly likely that they were gene carriers.

The SIRs for breast and stomach cancer in the families presented in this report is highly significant, indicating that it is a real entity and not a chance association, and it provides additional evidence that male stomach cancer is part of the spectrum of disease in BRCA2 families. Similarly, other cancers are over-represented providing additional evidence that BRCA2 mutations result in a less restricted disease phenotype than BRCA1 mutations. A significant difference in cancer incidence was observed between the two groups, which could be accounted for by the increased frequency of breast cancer within the mutation negative group accounted for the difference in cancer incidence between the two groups. The frequency of stomach cancer was identical in both groups (7%). There were too few other cancers to determine if there were differences between the two groups.

Finally, a comparison was made between the frequency of BRCA2 mutations in this study population compared to the frequency of BRCA2 mutations in breast ovarian cancer families. Collectively 248 breast/ovarian cancer families have thus far been screened for BRCA2 mutations, and 17 have been identified (van der Looij et al, 2000, Grzybowska et al, 2000 and Górski unpublished data). Using Fishers exact test we observed a significant difference between the incidence of BRCA2 mutations in the breast/male stomach cancer families compared to the breast/ovarian cancer families (P<0.025).
of breast cancer in the mutation negative group. This observation is most likely due to fragmentary pedigree information (due to the decimation of this population in recent history) and is unlikely to represent an actual difference between the two groups.

Three mutations in this study can be clearly identified as causative, whilst the remaining three represent missense changes which can not unequivocally be assigned as causative. The missense changes occurred in a small region of the BRCA2 gene (from position 6443 to 6612) within 169 base pairs of one another and were all identified in patients with breast cancer and not in 100 control subjects. The amino acid changes conferred by the missense mutations were not conservative, one change was a tyrosine to cysteine, the second a serine to cysteine and the third a lysine to asparagine. The introduction of a cysteine is likely to interact with other cysteines, which occur within this region of the gene and thereby disrupt the structural and functional activity of the protein. Since we do not know of any functional domains in this region of the gene it is impossible to determine the functional significance of these changes. Nevertheless, the evidence suggests that these missense mutations are likely to affect the function of the protein.

It has been shown that BRCA2 mutations are characterised by a lower level of penetrance than in BRCA1 mutation carriers (Boyd, 1996; Levy-Lahad et al, 1997). The results presented here support these findings, since we detected BRCA2 changes almost twice as frequently in patients without cancers among first degree relatives compared to patients with cancers diagnosed in parents or siblings (Boyd, 1996; Roa et al, 1996; Levy-Lahad et al, 1997; Risch et al, 2001).

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