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RESEARCH ARTICLE

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# Prevalence of *PALB2* mutations in Australian familial breast cancer cases and controls

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## Abstract

**Introduction:** *PALB2* is emerging as a high-penetrance breast cancer predisposition gene in the order of *BRCA1* and *BRCA2*. However, large studies that have evaluated the full gene rather than just the most common variants in both cases and controls are required before all truncating variants can be included in familial breast cancer variant testing.

**Methods:** In this study we analyse almost 2000 breast cancer cases sourced from individuals referred to familial cancer clinics, thus representing typical cases presenting in clinical practice. These cases were compared to a similar number of population-based cancer-free controls.

**Results:** We identified a significant excess of truncating variants in cases (1.3 %) versus controls (0.2 %), including six novel variants ( $p = 0.0001$ ; odds ratio (OR) 6.58, 95 % confidence interval (CI) 2.3–18.9). Three of the four control individuals carrying truncating variants had at least one relative with breast cancer. There was no excess of missense variants in cases overall, but the common c.1676A > G variant (rs152451) was significantly enriched in cases and may represent a low-penetrance polymorphism ( $p = 0.002$ ; OR 1.24 (95 % CI 1.09–1.47).

**Conclusions:** Our findings support truncating variants in *PALB2* as high-penetrance breast cancer susceptibility alleles, and suggest that a common missense variant may also lead to a low level of increased breast cancer risk.

## Introduction

Partner and localizer of *BRCA2* (*PALB2*) plays a central role in homologous recombination-mediated repair of double-strand DNA breaks [1] and biallelic mutations are responsible for Fanconi anemia complementation group N [2]. Monoallelic inactivating germline mutations in *PALB2* were subsequently shown to be associated with familial breast cancer [3] and numerous studies supported this association in various populations and established a mutation prevalence of approximately 1 % among familial breast cancer cases (varying from 0.1 % to 2.7 % as reviewed by Southey et al. [4]). Most recently, Antoniou et al. used a modified segregation analysis approach to determine that the age-specific risk of

breast cancer among female mutation carriers overlaps the risk conferred by *BRCA2* mutations [5] establishing that, despite the rarity of mutations, *PALB2* is the most significant breast cancer predisposition gene after *BRCA1* and *BRCA2*.

In Australia, early studies identified *PALB2* c.3113G > A (p.Trp1038\*) as a recurring truncating mutation among familial breast cancer index cases, and established the enrichment of c.3113G > A in cases compared to controls [6]. Further studies have identified a spectrum of truncating variants among breast cancer cases [7–10], the collective frequency of which has not been compared to Australian controls. Indeed few studies of *PALB2* mutations have analysed significant numbers of family cancer clinic-ascertained cases or matched controls. Because early studies focused on screening just for the presumed common pathogenic mutations, in Australia (eviQ Cancer Treatments Online; [11]) it is not recommended to test for *PALB2* truncating mutations aside from the recurring c.3113G > A variant, however, it is likely that all truncating

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mutations confer an equivalent loss of gene function and consequent breast cancer risk. Other guidelines, such as National Comprehensive Cancer Network [12], have made no specific distinction between different *PALB2* mutations but do raise a general caution around the interpretation of testing for mutations in *PALB2* and other “moderate penetrance” breast cancer predisposition genes, especially as part of panel tests. Identification of genetic risk factors is critical for individual risk assessment and reduction strategies, and in the near future may provide avenues for personalised therapy [4]. Therefore it is important to continue to amass the necessary data to support the implementation of whole gene testing of *PALB2* in breast cancer families. In this study, we performed germline mutation analysis of the entire coding region of *PALB2* in a cohort of 1996 breast cancer index cases referred to familial cancer clinics for genetic testing and tested negative for *BRCA1* and *BRCA2* mutations as well as 1998 Australian cancer-free female controls. This represents the largest single case/control screen of germline *PALB2* mutations to date.

## Methods

### Samples for mutation analysis

Cancer-affected women in the study were referred by their physician to a specialist Familial Cancer Centre (FCC) for genetic testing of *BRCA1* and *BRCA2* between 1997 and 2014, and were identified as being at “high risk” of carrying a predisposing allele. The criteria for high risk included a personal history of breast cancer, two or more first- or second-degree relatives with breast and/or ovarian cancer, and an additional risk factor (additional affected close relatives, diagnosis before 40 years, multiple primary breast or ovarian cancers in one individual, or Ashkenazi Jewish ancestry). From 2003, individuals with a  $\geq 10$  % risk of carrying a *BRCA1* or *BRCA2* mutation, as estimated by BRCAPro, including tumour pathology, were also eligible [13].

Our final case cohort (Additional file 1) included 997 breast (95 %) or ovarian (5 %) cancer-affected index cases from the Hunter Area Pathology Service (HAPS), Newcastle, Australia [9]. Family history information was available for a subset of this cohort only. A further 999 breast cancer-affected index cases each with detailed family history available were obtained from the combined Victorian Familial Cancer Centres (FCCs) through the Variants in Practice (ViP) study. For all cases, clinical genetic testing of *BRCA1* and *BRCA2*, including for large rearrangements by multiplex ligation probe-dependent amplification (MLPA), returned negative results.

A cohort of 1998 participants in the LifePool study [14] were utilised as cancer-free population control samples for this analysis. LifePool recruits female participants through the Australian population mammography

screening program (BreastScreen) for research studies utilising prospectively collected epidemiological, genetic and mammographic data with ongoing clinical follow-up obtained through the Victorian Cancer Registry. Participants provided breast cancer family history information for close relatives only. The average age of the participants recruited to this study was  $58.84 \pm 9.9$  years (range 19–91).

All cases and controls provided informed consent for genetic analysis of their germline DNA. This study was approved by the Human Research Ethics Committees at each participating ViP centre (see Acknowledgements), the Peter MacCallum Cancer Centre, Hunter New England Health and The University of Newcastle. This study was carried out in accordance with all relevant regulations and guidelines.

### Germline mutation analysis

Germline mutation analysis of the *PALB2* gene was performed as part of a custom sequencing panel. All coding *PALB2* exons were amplified from 225 ng of germline DNA extracted from blood or saliva using the HaloPlex Targeted Enrichment Assay (Agilent Technologies, Santa Clara, CA, USA) according to the manufacturer’s protocol using an Agilent Bravo Automated Liquid Handling System. Paired-end 100 or 150 bp sequence reads were generated from the indexed, pooled libraries on a HiSeq2500 Genome Analyzer (Illumina, San Diego, CA, USA). Sequence reads were trimmed of adapter using Cutadapt [15] and aligned using either BWA or BWA MEM [16]. Genome Analysis Toolkit (GATK) v3.1 was used to perform indel realignment and Unified Genotyper was used for variant calling [17, 18]. Protein consequence and additional annotations were added using Ensembl v73 Variant Effect Predictor [19]. Variant positions were determined by reference to GenBank reference sequence NM\_024675.3 according to Human Genome Variation Society (HGVS) guidelines [20]. All novel variants were validated by Sanger resequencing of germline DNA using primers from Tischkowitz et al. [21]. The following in silico prediction tools were used to assess the possible pathogenicity of missense mutations: Combined Annotation-Dependent Depletion (CADD) [22], Condel [23], SIFT [24] and PolyPhen2 [25]. CADD scores evaluate both missense and indel variants, integrating conservation measures, regulatory, transcriptional and protein effects to estimate the relative deleteriousness of the variants.

## Results

### Coverage

A total of 1996 breast cancer index cases and 1998 non-cancer controls were screened for germline mutations in the coding regions of *PALB2*. These coding regions were well covered by sequence reads in both cases and controls.

The mean read depth across the entire gene for all samples was 217 (192 for cases, 242 for controls), with an average of 98.66 % of the coding regions covered by at least 20 reads (98.12 % for cases and 99.20 % for controls).

### Truncating mutations

Nineteen different truncating variants were identified in 30 individuals in this study, 26 of these variants were detected among 1996 breast cancer index cases (1.3 %) and four among 1998 controls (0.2 %), demonstrating a significant enrichment in cases ( $p = 0.0001$ , chi-squared test; odds ratio (OR) 6.58, 95 % confidence interval (CI) 2.3–18.9) (Table 1). Five previously reported variants were detected recurrently (c.196C > T (p.Gln66\*) and c.758dupT (p.Ser254Ilefs\*3) each in two cases, c.3113G > A (p.Trp1038\*) in seven cases and one control, and c.3116delA (p.Asn1039Ilefs\*2) and c.3362delG (p.Gly1121Valfs\*3) each in one case and one control) with the remaining variants detected in single cases or controls only. Six truncating variants have not previously been reported (c.522\_523delAA (p.Arg175Thrfs\*9), c.577dupA (p.Thr193Asnfs\*2), c.693dupA (p.Gly232Argfs\*3), c.860dupT (p.Ser288Lysfs\*15), c.1947\_1966dup (p.Pro65

6Glnfs\*11) and c.2966\_2967insCAACAAGT (p.Glu990Asnfs\*3)). Novel variant p.Glu990Asnfs\*3 was detected in a control only.

The personal and family history information for carriers of the *PALB2* truncating variants are given in Table 2 and Additional file 2. As expected, the cases generally have a strong family history of cancer, especially breast cancer. In the controls, four individuals were identified with truncating variants. One individual had a maternal aunt diagnosed with breast cancer at under 40 years of age, and her mother, father and brother all had cancer although not of the breast. The mothers of both of the other individuals had a breast cancer diagnosis aged over 70 years of age, and one of these individuals also had two second-degree relatives with breast cancer. The final and youngest carrier (aged 48) did not report any breast cancer in her family. Thus, 3/4 carriers have some family history of breast cancer.

### Missense and synonymous variants

A large number of missense variants ( $n = 54$ ) were detected in the cohort (Table 3). There was a slight

**Table 1** Truncating variants

| Exon | Nucleotide change <sup>a</sup> | Predicted protein change <sup>a</sup> | dbSNP ID    | First reported                     | HAPS cases (n = 997) <sup>b</sup> | FCC-ViP cases (n = 999) <sup>b</sup> | Controls (n = 1998) <sup>b</sup> | CADD scaled C score |
|------|--------------------------------|---------------------------------------|-------------|------------------------------------|-----------------------------------|--------------------------------------|----------------------------------|---------------------|
| 3    | c.172_175delTTGT               | p.Gln60Argfs*7                        | .           | Jones (2009)                       | -                                 | 1                                    | -                                | 19.0                |
| 3    | c.196C > T                     | p.Gln66*                              | rs180177083 | Casadei (2011) [29]                | 2                                 | -                                    | -                                | 35.0                |
| 4    | c.522_523delAA                 | p.Arg175Thrfs*9                       | .           | -                                  | -                                 | 1                                    | -                                | 23.2                |
| 4    | c.577dupA                      | p.Thr193Asnfs*2                       | .           | -                                  | 1                                 | -                                    | -                                | 11.6                |
| 4    | c.693dupA                      | p.Gly232Argfs*3                       | .           | -                                  | 1                                 | -                                    | -                                | 11.2                |
| 4    | c.758dupT                      | p.Ser254Ilefs*3                       | .           | Zheng (2012) [40]                  | 1                                 | 1                                    | -                                | 16.2                |
| 4    | c.860dupT                      | p.Ser288Lysfs*15                      | .           | -                                  | 1                                 | -                                    | -                                | 17.6                |
| 5    | c.1947dupA                     | p.Glu650Argfs*13                      | .           | Teo (2013) [8]                     | -                                 | 1                                    | -                                | 24.1                |
| 5    | c.1947_1966dup                 | p.Pro656Glnfs*11                      | .           | -                                  | 1                                 | -                                    | -                                | 13.0                |
| 5    | c.2386G > T                    | p.Gly796*                             | rs180177112 | Rahman (2007) [3]                  | -                                 | 1                                    | -                                | 32.0                |
| 5    | c.2391delA                     | p.Gln797Hisfs*54                      | .           | Wong-Brown (2013)                  | 1                                 | -                                    | -                                | 23.5                |
| 9    | c.2966_2967insCAACAAGT         | p.Glu990Asnfs*3                       | .           | -                                  | -                                 | -                                    | 1                                | 20.7                |
| 9    | c.2982dupT                     | p.Ala995Cysfs*16                      | rs180177127 | Rahman (2007) [3]                  | 1                                 | -                                    | -                                | 31.0                |
| 10   | c.3113G > A                    | p.Trp1038* <sup>c</sup>               | rs180177132 | Rahman (2007) [3]                  | 2                                 | 5                                    | 1                                | 42.0                |
| 11   | c.3116delA                     | p.Asn1039Ilefs*2                      | rs180177133 | Reid (2007) [2]; Rahman (2007) [3] | -                                 | 1                                    | 1                                | 40.0                |
| 12   | c.3256C > T                    | p.Arg1086*                            | .           | Jones (2009)                       | -                                 | 1                                    | -                                | 40.0                |
| 13   | c.3362delG                     | p.Gly1121Valfs*3                      | .           | Blanco (2013) [27]                 | -                                 | 1                                    | 1                                | 22.1                |
| 13   | c.3507_3508del                 | p.His1170Phefs*19                     | .           | Antoniou (2014) [5]                | 1                                 | -                                    | -                                | 40.0                |
| 13   | c.3549C > G                    | p.Tyr1183*                            | rs118203998 | Reid (2007) [2]                    | 1                                 | -                                    | -                                | 37.0                |

HAPS Hunter Area Pathology Service, FCC Familial Cancer Centre, ViP Variants in Practice

<sup>a</sup>Variant positions are reported in reference to NCBI RefSeq NM\_024675.3 (mRNA) and NP\_078951.2 (protein)

<sup>b</sup>Number of individuals carrying the variant

<sup>c</sup>c.3113G > A produces three different *PALB2* mRNA sequences: complete deletion of exon 10 (117 bp); use of an alternative splice site within exon 10, and deletion of 31 bp; and an immediate stop at codon 1038 [29]

**Table 2** Family history of carriers of truncating variants

| Case            | Variant <sup>a</sup> | Diagnosis                              | Family history first-degree relatives                                        | Family history other relatives                                                                                                                 |
|-----------------|----------------------|----------------------------------------|------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------|
| HAPS-102285     | p.Gln66*             | Breast 41                              | NA                                                                           | NA                                                                                                                                             |
| HAPS-60382      | p.Gln66*             | Breast 43                              | Mother breast 61; father prostate 65                                         | Paternal: 2 x 2 <sup>nd</sup> cousins breast 42, 37; 3 x great aunt breast 42, 47, 58; grandfather prostate 71. Maternal: grandmother other 60 |
| HAPS-90978      | p.Thr193Asnfs*2      | Bilateral breast 61                    | Sister breast 60s; sister breast 60s                                         |                                                                                                                                                |
| HAPS-90809      | p.Gly232Argfs*3      | Bilateral breast 41, 47                | Father lung 68                                                               | Cousin breast 45, maternal grandmother bowel 60                                                                                                |
| HAPS-120272     | p.Ser254Ilefs*3      | Breast 35, ovarian 58                  | 2 x Sisters breast 40s; brother renal 52; brother HNSCC 50s; father other    | Cousin breast 50s.                                                                                                                             |
| HAPS-102573     | p.Ser288Lysfs*15     | Breast 63                              | NA                                                                           | NA                                                                                                                                             |
| HAPS-114269     | p.Pro656Glnfs*11     | Breast 46, ovarian 49                  | Mother unknown primary 45                                                    | Maternal aunt breast 50s; grandmother breast 60s; maternal uncle prostate 70s                                                                  |
| HAPS-120953     | p.Gln797Hisfs*54     | Breast 46                              | Father melanoma 62                                                           | half-brother NHL 64; half-brother bladder 55; maternal cousin breast 41; maternal aunt breast 55                                               |
| HAPS-81242      | p.Ala995Cysfs*16     | Bilateral breast 46, 70                | Sister breast 38; daughter breast 47; daughter cervical 24; mother breast 39 | Paternal cousin breast 60s; maternal cousin other                                                                                              |
| HAPS-121549     | p.Trp1038*           | Breast 56                              | Sister breast 44                                                             | Maternal aunt breast 50s; paternal grandmother breast                                                                                          |
| HAPS-110283     | p.Trp1038*           | Breast 46                              | Mother breast 57                                                             | Maternal: aunt bowel 50; great-grandmother ovarian. Paternal: grandmother breast 85, grandfather bowel 60                                      |
| HAPS-100240     | p.His1170Phefs*19    | Breast 48                              | Mother breast 68                                                             | Maternal grandmother AML 72; paternal cousin ovarian 36.                                                                                       |
| HAPS-110583     | p.Tyr1183*           | Breast 39                              | Mother breast 80; father bowel 69                                            | Maternal aunt bilateral breast 50, 70; maternal grandfather prostate 80                                                                        |
| FCC-681-000     | p.Gln60Argfs*7       | Breast 47                              |                                                                              | Breast >40                                                                                                                                     |
| FCC-2121-000    | p.Gly796*            | Breast 62                              | Lung/prostate                                                                | Prostate, other x 2                                                                                                                            |
| FCC-1423-000    | p.Trp1038*           | Breast 44                              | Breast >40, bowel, other                                                     | Breast >40                                                                                                                                     |
| FCC-2104-000    | p.Trp1038*           | Breast 38                              | Breast >40, other x 2                                                        | Other                                                                                                                                          |
| FCC-2677-000    | p.Trp1038*           | Cervix 55, breast 65, 67, 68, bowel 67 | Breast >40, ovarian                                                          | Lung, other                                                                                                                                    |
| FCC-3527-000    | p.Trp1038*           | Breast 37                              |                                                                              | Other                                                                                                                                          |
| FCC-60-000      | p.Trp1038*           | Thyroid 42, breast 48, 51              | Breast >40, bowel                                                            | Breast x 2 ovarian, prostate x 2, lung, other x 2                                                                                              |
| FCC-905-000     | p.Asn1039Ilefs*2     | Melanoma 39, breast 47                 | Lung, prostate, other                                                        | Breast x 3, bowel x 2, ovarian                                                                                                                 |
| FCC-2965-000    | p.Arg1086*           | Bilateral breast 44                    | Prostate                                                                     |                                                                                                                                                |
| FCC-317-000     | p.Gly1121Valfs*3     | Breast 31                              | Other x 2                                                                    | Breast, other                                                                                                                                  |
| FCC-1322-000    | p.Glu650Argfs*13     | Breast 50                              | Breast, other                                                                | Breast x 3, other                                                                                                                              |
| FCC-3397-000    | p.Ser254Ilefs*3      | Melanoma 53, Breast 54, 61             | Breast                                                                       |                                                                                                                                                |
| FCC_2431-000    | p.Arg175Thrfs*9      | Breast 42                              | Breast, other x 3                                                            | Breast x 3, lung, other x 2                                                                                                                    |
| <b>Controls</b> | <b>Variant</b>       | <b>Age enrolled</b>                    | <b>Family history first-degree relatives</b>                                 | <b>Family history other relatives</b>                                                                                                          |
| LP-12031915     | p.Trp1038*           | 65                                     | Mother, other 58; father lung 64; brother other                              | Maternal aunt breast >40                                                                                                                       |

**Table 2** Family history of carriers of truncating variants (*Continued*)

|             |                  |    |                  |                                                   |
|-------------|------------------|----|------------------|---------------------------------------------------|
| LP-13099711 | p.Asn1039Ilefs*2 | 73 | Mother breast 72 | Maternal: grandmother breast >40 yrs; aunt breast |
| LP-13243620 | p.Glu990Asnfs*3  | 73 | Mother breast 82 | NA                                                |
| LP-12025195 | p.Gly1121Valfs*3 | 48 | None             | Yes, no details                                   |

NA not applicable, HNSCC head and neck squamous cell carcinoma, NHL non-Hodgkin lymphoma

<sup>a</sup>Novel variants in italics

enrichment for missense variants overall in cases (39.6 % versus controls (36.1 %,  $p = 0.025$ , OR 1.15, 95 % CI 1.02–1.32). The four most common variants (>3 % carrier frequency) were tested for association with breast cancer. Surprisingly, the most common variant, c.1676A > G (Gln559Arg; rs152451), which is predicted by CADD, Condel, PolyPhen2 and SIFT to be benign, was significantly more common in cases (19.9 % carried at least one non-wild-type allele) than controls (16.8 % non-wild-type) with a per-allele OR of 1.24 (95 % CI 1.09–1.47,  $p = 0.002$ , logistic regression). There were 20 cases homozygous for this variant versus ten controls ( $p = 0.058$  logistic regression, OR 2.09, 95 % CI 0.98–4.48). The overall trend for an effect had a  $p$  value of 0.0018 (Cochrane-Armitage trend test). The minor allele frequency was 0.105 in cases and 0.086 in controls, compared to other databases where the minor allele frequency of this variant in European populations was 0.09 (1000 Genomes), 0.096 (ExAC) and 0.09 (EVS), but showed increased frequency in African and Asian populations.

Considering only those rare variants present in fewer than five carriers among 3994 cases and controls (approximately 0.1 %), a similar number of missense variants were detected in both groups (40 in cases (2 %), 28 in controls (1.4 %)), which does not suggest any association of rare missense variants with risk. There was also no significant enrichment in cases when limited to rare variants that were predicted to be deleterious by any of Condel, SIFT or Polyphen2 (28/1996 cases, 18/1998 controls) or with a CADD score of >10 (29/1996 cases, 20/1998 controls).

We detected 23 synonymous variants (Table 4). Neither the most common alone (c.3300 T > G) nor all together were significantly enriched in cases or controls.

## Discussion

This study screened Australian individuals with breast cancer who had been referred to a Familial Cancer Centre for genetic testing and in whom no pathogenic *BRCA1* and *BRCA2* variant could be identified. The frequency of *PALB2* truncating variants in this cohort (1.1 %) is similar to other studies analysing high-risk breast cancer individuals (0.64–3.4 %, 1.35 % overall [3, 6–9, 26–41]) or triple-negative breast cancer (0.9–2.5 % [10, 42, 43]) but is the largest to include an analysis of the full gene in both cases and controls. However, we would not be able to detect any large deletions or rearrangements. The low

frequency of truncating variants in controls supports *PALB2* as a high-penetrance breast cancer predisposing gene. The diversity of truncating mutations identified, comprising 16 different variants in eight of the 13 exons including five novel variants, highlights the need for full gene screening, not just the most common variant c.3113G > A (rs180177132). These data will enable evidence-based clinical guidelines to include full *PALB2* screening if previously they had advised testing limited to the specific common variant only.

The prevalence of truncating variants in cancer-free controls was 0.15 % in the LifePool cohort. These individuals were ascertained from women attending population-based mammographic screening, which in Australia is targeted towards women over 50, although some younger women are included. Thus, this volunteer cohort may not be entirely representative of the general population, although all were cancer-free at the time of analysis. Nonetheless, the frequencies of missense and synonymous variants are consistent with those reported in large databases such as 1000 Genomes [44], Exome Aggregation Consortium [45] and Exome Variant Server [46].

We did not observe any significant enrichment in missense mutations overall, although the frequency was slightly higher in the cases when only rare, deleterious mutations were considered. The contribution of rarer variants to breast cancer risk will need to be evaluated in larger case-control cohorts. Surprisingly, the common variant c.1676A > G (Gln559Arg; rs152451) was significantly enriched in cases versus controls, although with only a modest odds ratio (1.24). There was a trend towards homozygous carriers of this variant being enriched in cases versus controls with an OR of 2.08. This variant was shown to be associated with an increased breast cancer risk in multiple-case breast cancer families in Chile compared to population controls [47] with an OR of 2.0 when at least three family members were breast or ovarian cancer-affected. No association was found for individuals diagnosed at a young age (<50) and with no affected relatives. In a small Malaysian case-control study, there was a trend towards enrichment for carriers of the variant in non-familial breast cancer cases (286/871, 33 %) versus controls (70/257, 27 %, OR 1.3 [38]), however, cases and controls were not well matched for ethnicity, with an excess of Indian and Malay women over Chinese in the controls compared to cases. Larger numbers of cases and

**Table 3** Missense variants

| Exon | Nucleotide change <sup>a</sup> | Protein change <sup>a</sup> | dbSNP ID    | First reported          | 1000 G MAF <sup>b</sup> | NHLBI GO ESP MAF <sup>c</sup> | ExAC <sup>d</sup> | HAPS cases (n = 997) <sup>e</sup> | FCC-ViP cases (n = 999) <sup>e</sup> | Controls (n = 1998) <sup>e</sup> | CADD scaledC score | Condel      | PolyPhen2         | SIFT        |
|------|--------------------------------|-----------------------------|-------------|-------------------------|-------------------------|-------------------------------|-------------------|-----------------------------------|--------------------------------------|----------------------------------|--------------------|-------------|-------------------|-------------|
| 1    | c.11C > T                      | p.Pro4Leu                   | rs45619737  | Rahman (2007) [3]       | .                       | 0.00038                       | 0.0000999         | 1                                 | 1                                    | 1                                | 12.9               | Deleterious | Probably damaging | Tolerated   |
| 2    | c.53A > G                      | p.Lys18Arg                  | rs138789658 | Tischkowitz (2008)      | 0.0032                  | 0.00523                       | 0.0000552         | -                                 | -                                    | 1                                | 18.0               | Deleterious | Possibly damaging | Deleterious |
| 2    | c.94C > G                      | p.Leu32Val                  | rs151316635 | Teo (2013) [8]          | .                       | 0.00023                       | 0.0000184         | -                                 | 1                                    | 1                                | 16.3               | Deleterious | Possibly damaging | Deleterious |
| 3    | c.194C > T                     | p.Pro65Leu                  | rs62625272  | Adank (2011) [26]       | .                       | 0.00015                       | 0.00006           | -                                 | -                                    | 1                                | 4.6                | Neutral     | Benign            | Tolerated   |
| 4    | c.232G > A                     | p.Val78Ile                  | .           | Tischkowitz (2012) [39] | .                       | .                             | 0.000325          | .                                 | 2                                    | .                                | 0.4                | Neutral     | Benign            | Tolerated   |
| 4    | c.298C > T                     | p.Leu100Phe                 | rs61756147  | Wong (2011) [9]         | 0.0005                  | 0.00023                       | 0.000037          | -                                 | -                                    | 2                                | 10.3               | Neutral     | Probably damaging | Tolerated   |
| 4    | c.344G > T                     | p.Gly115Val                 | rs145598272 | Foulkes (2007) [34]     | .                       | 0.00015                       | 0.0000921         | 1                                 | -                                    | 1                                | 6.5                | Neutral     | Benign            | Tolerated   |
| 4    | c.353 T > C                    | p.Ile118Thr                 | .           | -                       | .                       | 0.00008                       | 0.000037          | -                                 | -                                    | 1                                | 5.4                | Neutral     | Benign            | Tolerated   |
| 4    | c.400G > A                     | p.Asp134Asn                 | rs139555085 | Zheng (2011)            | 0.0005                  | 0.00184                       | 0                 | .                                 | .                                    | 1                                | 4.8                | Neutral     | Benign            | Tolerated   |
| 4    | c.508A > G                     | p.Arg170Gly                 | .           | -                       | .                       | .                             | .                 | -                                 | -                                    | 1                                | 4.4                | Neutral     | Benign            | Tolerated   |
| 4    | c.557A > T                     | p.Asn186Ile                 | .           | -                       | .                       | .                             | .                 | -                                 | -                                    | 1                                | 13.1               | Deleterious | Probably damaging | Deleterious |
| 4    | c.571C > G                     | p.Pro191Ala                 | .           | -                       | .                       | .                             | .                 | -                                 | -                                    | 1                                | 9.4                | Neutral     | Probably damaging | Tolerated   |
| 4    | c.629C > T                     | p.Pro210Leu                 | rs57605939  | Rahman (2007) [3]       | 0.0174                  | 0.02216                       | 0.0000736         | 1                                 | 1                                    | 2                                | 10.6               | Deleterious | Probably damaging | Tolerated   |
| 4    | c.899C > T                     | p.Thr300Ile                 | .           | Ding (2011)             | .                       | .                             | 0                 | 1                                 | -                                    | -                                | 15.1               | Deleterious | Probably damaging | Tolerated   |
| 4    | c.656A > G                     | p.Asp219Gly                 | rs45594034  | Rahman (2007) [3]       | .                       | 0.00015                       | 0.000221          | .                                 | 1                                    | .                                | 1.8                | Neutral     | Benign            | Tolerated   |
| 4    | c.740C > G                     | p.Thr247Arg                 | .           | .                       | .                       | .                             | .                 | .                                 | 1                                    | .                                | 15.5               | Deleterious | Probably damaging | Tolerated   |
| 4    | c.925A > G                     | p.Ile309Val                 | rs3809683   | Rahman (2007) [3]       | 0.0087                  | 0.00970                       | 0.0000184         | 4 (1)                             | -                                    | -                                | 0.5                | Neutral     | Benign            | Tolerated   |
| 4    | c.1010 T > C                   | p.Leu337Ser                 | rs45494092  | Rahman (2007) [3]       | 0.0133                  | 0.01424                       | 0.0197            | 44 (1)                            | 44                                   | 93 (1)                           | 8.9                | Deleterious | Probably damaging | Tolerated   |
| 4    | c.1085 T > C                   | p.Leu362Pro                 | .           | -                       | .                       | .                             | .                 | 1                                 | -                                    | -                                | 14.4               | Deleterious | Probably damaging | Tolerated   |
| 4    | c.1145G > T                    | p.Ser382Ile                 | .           | Tischkowitz (2012) [39] | .                       | .                             | 0.0000184         | -                                 | 1                                    | -                                | 15.2               | Deleterious | Possibly damaging | Deleterious |

**Table 3** Missense variants (Continued)

|   |             |             |             |                            |        |         |           |         |          |          |      |             |                   |             |
|---|-------------|-------------|-------------|----------------------------|--------|---------|-----------|---------|----------|----------|------|-------------|-------------------|-------------|
| 4 | c.1189A > T | p.Thr397Ser | .           | Rahman (2007) [3]          | .      | 0.00008 | 0.0000184 | -       | 1        | 1        | 22.9 | Deleterious | Possibly damaging | Deleterious |
| 4 | c.1250C > A | p.Ser417Tyr | rs45510998  | Rahman (2007)              | .      | .       | 0.000203  | -       | -        | 1        | 20.6 | Deleterious | Probably damaging | Deleterious |
| 4 | c.1478C > T | p.Pro493Leu | .           | -                          | .      | .       | 0.0000184 | -       | -        | 1        | 12.4 | Neutral     | Benign            | Tolerated   |
| 4 | c.1492G > T | p.Asp498Tyr | rs75023630  | Phuah (2013) [38]          | 0.0014 | .       | 0         | -       | -        | 1        | 16.2 | Deleterious | Benign            | Deleterious |
| 4 | c.1544A > G | p.Lys515Arg | .           | Tischkowitz (2012) [39]    | .      | .       | 0.000037  | -       | 1        | -        | 16.2 | Deleterious | Possibly damaging | Tolerated   |
| 4 | c.1610C > T | p.Ser537Leu | rs142103232 | -                          | .      | 0.00015 | 0.000166  | -       | 1        | -        | 11.2 | Neutral     | Possibly damaging | Tolerated   |
| 4 | c.1676A > G | p.Gln559Arg | rs152451    | Rahman (2007) [3]          | 0.1465 | 0.13483 | 0.0961    | 216 (9) | 182 (11) | 335 (10) | 0.0  | Neutral     | Benign            | Tolerated   |
| 5 | c.1699C > T | p.His567Tyr | .           | Tischkowitz (2012) [39]    | .      | 0.00008 | 0.000094  | -       | -        | 1        | 0.9  | Neutral     | Benign            | Tolerated   |
| 5 | c.1931G > A | p.Gly644Glu | .           | -                          | .      | .       | .         | -       | -        | 1        | 16.0 | Deleterious | Probably damaging | Deleterious |
| 5 | c.2014G > C | p.Glu672Gln | rs45532440  | Rahman (2007) [3]          | 0.0142 | 0.02324 | 0.0278    | 64 (1)  | 68 (4)   | 123 (1)  | 11.4 | Neutral     | Possibly damaging | Tolerated   |
| 5 | c.2106A > G | p.Ile702Met | .           | -                          | .      | .       | 0.0000184 | 1       | -        | -        | 11.1 | Neutral     | Probably damaging | Tolerated   |
| 5 | c.2135C > T | p.Ala712Val | rs141458731 | Dansonka-Meiskowska (2010) | 0.0014 | 0.00062 | 0.00039   | -       | -        | 1        | 12.2 | Neutral     | Benign            | Tolerated   |
| 5 | c.2200A > T | p.Thr734Ser | rs45543843  | Rahman (2007) [3]          | .      | .       | 0.0000368 | -       | -        | 3        | 25.3 | Deleterious | Possibly damaging | Deleterious |
| 5 | c.2228A > G | p.Tyr743Cys | rs141749524 | -                          | 0.0009 | .       | 0.0000184 | 1       | -        | -        | 8.5  | Neutral     | Benign            | Tolerated   |
| 5 | c.2289G > C | p.Leu763Phe | .           | Phuah (2013) [38]          | .      | .       | 0         | 2       | -        | -        | 15.5 | Neutral     | Probably damaging | Tolerated   |
| 5 | c.2360C > T | p.Thr787Ile | rs201042302 | -                          | 0.0005 | .       | 0         | -       | 1        | -        | 9.7  | Neutral     | Probably damaging | Tolerated   |
| 5 | c.2417C > T | p.Pro806Leu | rs45464991  | Rahman (2007) [3]          | .      | 0.00008 | 0.000037  | -       | 1        | 1        | 0.5  | Neutral     | Benign            | Tolerated   |
| 7 | c.2590C > T | p.Pro864Ser | rs45568339  | Rahman (2007) [3]          | 0.0018 | 0.00239 | 0.00396   | 6       | 10       | 19       | 12.0 | Neutral     | Benign            | Tolerated   |
| 7 | c.2606C > G | p.Ser869Cys | .           | .                          | .      | .       | 0.0000184 | .       | 1        | .        | 19.9 | Deleterious | Probably damaging | Deleterious |
| 7 | c.2674G > A | p.Glu892Lys | rs45476495  | Rahman (2007) [3]          | .      | 0.00008 | 0.000092  | 1       | 2        | -        | 20.8 | Deleterious | Possibly damaging | Deleterious |
| 8 | c.2755G > A | p.Val919Ile | .           | -                          | .      | .       | .         | -       | -        | 1        | 15.7 | Neutral     | Benign            | Tolerated   |

**Table 3** Missense variants (*Continued*)

|    |              |              |             |                         |        |         |           |    |        |        |      |             |                   |             |
|----|--------------|--------------|-------------|-------------------------|--------|---------|-----------|----|--------|--------|------|-------------|-------------------|-------------|
| 8  | c.2794G > A  | p.Val932Met  | rs45624036  | Rahman (2007) [3]       | 0.0009 | 0.00431 | 0.00869   | 15 | 8      | 23     | 18.3 | Deleterious | Probably damaging | Tolerated   |
| 8  | c.2816 T > G | p.Leu939Trp  | rs45478192  | Rahman (2007) [3]       | 0.0009 | 0.00154 | 0.0015    | 2  | 3 (1)  | 8      | 20.9 | Deleterious | Probably damaging | Deleterious |
| 9  | c.2993G > A  | p.Gly998Glu  | rs45551636  | Rahman (2007) [3]       | 0.0105 | 0.01785 | 0.0213    | 42 | 46 (2) | 92 (1) | 22.7 | Deleterious | Probably damaging | Deleterious |
| 10 | c.3054G > C  | p.Glu1018Asp | rs183489969 | Tischkowitz (2012) [39] | 0.0009 | .       | 0         | 1  | -      | 1      | 16.4 | Deleterious | Possibly damaging | Deleterious |
| 10 | c.3106G > C  | p.Val1036Leu | .           | -                       | .      | .       | 0.000037  | -  | 1      | -      | 13.3 | Neutral     | Benign            | Tolerated   |
| 11 | c.3128G > C  | p.Gly1043Ala | .           | Hellebrand (2011) [36]  | 0.0009 | .       | 0.000037  | 1  | -      | -      | 20.9 | Deleterious | Probably damaging | Deleterious |
| 11 | c.3146 T > C | p.Met1049Thr | rs138273800 | -                       | .      | 0.00008 | .         | -  | 1      | -      | 18.0 | Deleterious | Probably damaging | Deleterious |
| 12 | c.3307G > A  | p.Val1103Met | rs201657283 | Casadei (2011) [29]     | .      | 0.00015 | 0.0000184 | -  | 1      | -      | 13.0 | Neutral     | Benign            | Tolerated   |
| 13 | c.3366C > A  | p.Asp1122Glu | .           | .                       | .      | .       | .         | -  | .      | 1      | 16.0 | Neutral     | Possibly damaging | Tolerated   |
| 13 | c.3367G > A  | p.Val1123Met | .           | .                       | .      | .       | .         | -  | 1      | -      | 18.7 | Deleterious | Probably damaging | Deleterious |
| 13 | c.3428 T > A | p.Leu1143His | rs62625284  | Balia (2010)            | .      | 0.00008 | 0.00031   | 2  | -      | -      | 20.0 | Deleterious | Possibly damaging | Tolerated   |
| 13 | c.3448C > T  | p.Leu1150Phe | .           | .                       | .      | .       | .         | -  | 1      | -      | 15.5 | Deleterious | Possibly damaging | Deleterious |
| 13 | c.3449 T > G | p.Leu1150Arg | rs45566737  | .                       | .      | .       | 0.0000368 | -  | 1      | -      | 18.2 | Deleterious | Possibly damaging | Deleterious |

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<sup>a</sup>Variant positions are reported in reference to NCBI RefSeq NM\_024675.3 (mRNA) and NP\_078951.2 (protein)<sup>b</sup>Minor allele frequency (MAF) reported in the 1000 Genomes (1000 G) cohort Phase 1<sup>c</sup>MAF reported in the Exome Variant Server, NHLBI GO Exome Sequencing Project (ESP) [46] (data release ESP6500SI-V2)<sup>d</sup>MAF reported in ExAC [45] from non-Finnish Europeans, excluding individuals in the database who were part of The Cancer Genome Atlas and therefore known to have had cancer<sup>e</sup>Number of individuals carrying the variant. Where applicable, the number of homozygous carriers is indicated in parentheses

**Table 4** Synonymous variants

| Exon | Nucleotide change <sup>a</sup> | Protein change <sup>a</sup> | dbSNP ID    | First reported    | 1000 G MAF <sup>b</sup> | NHLBI GO ESP MAF <sup>c</sup> | ExAC <sup>d</sup> | HAPS cases (n = 997) <sup>e</sup> | FCC-ViP cases (n = 999) <sup>d</sup> | Controls (n = 1998) <sup>e</sup> | CADD scaled score |
|------|--------------------------------|-----------------------------|-------------|-------------------|-------------------------|-------------------------------|-------------------|-----------------------------------|--------------------------------------|----------------------------------|-------------------|
| 1    | c.12 T > C                     | p.(=)                       | rs145291423 | -                 | .                       | .                             | 0.00012           | -                                 | 1                                    | 1                                | 11.2              |
| 4    | c.768C > T                     | p.(=)                       | rs45487491  | -                 | .                       | .                             | 0.000037          | -                                 | 1                                    | -                                | 0.3               |
| 4    | c.1188C > T                    | p.(=)                       | .           | -                 | .                       | .                             | 0.00003           | -                                 | -                                    | 1                                | 7.6               |
| 4    | c.1194G > A                    | p.(=)                       | rs61755173  | Rahman (2007) [3] | 0.0009                  | 0.00154                       | 0.00114           | 3                                 | 2                                    | 8                                | 6.4               |
| 4    | c.1194G > T                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | 1                                 | -                                    | -                                | 6.1               |
| 4    | c.1242A > C                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | -                                 | -                                    | 1                                | 7.7               |
| 4    | c.1431C > T                    | p.(=)                       | .           | Teo (2013)        | .                       | .                             | 0                 | 2                                 | -                                    | -                                | 0.0               |
| 4    | c.1470C > T                    | p.(=)                       | rs45612837  | Rahman (2007) [3] | 0.0005                  | 0.00015                       | 0.000405          | 1                                 | -                                    | 2                                | 5.8               |
| 4    | c.1572A > G                    | p.(=)                       | rs45472400  | Rahman (2007) [3] | 0.0032                  | 0.00339                       | 0.0041            | 12                                | 12                                   | 27                               | 5.6               |
| 4    | c.1623G > A                    | p.(=)                       | .           | -                 | .                       | .                             | 0                 | -                                 | 1                                    | 2                                | 4.2               |
| 5    | c.2067G > A                    | p.(=)                       | .           | Phuah (2013) [38] | .                       | 0.00015                       | 0.000018          | 1                                 | -                                    | -                                | 4.1               |
| 5    | c.2082A > G                    | p.(=)                       | rs150569240 | -                 | .                       | .                             | 0.000055          | -                                 | -                                    | 1                                | 5.8               |
| 5    | c.2091C > A                    | p.(=)                       | .           | -                 | .                       | .                             | 0                 | -                                 | -                                    | 1                                | 4.7               |
| 5    | c.2244A > G                    | p.(=)                       | .           | -                 | .                       | .                             | 0.000037          | -                                 | -                                    | 1                                | 5.6               |
| 5    | c.2328C > T                    | p.(=)                       | rs45508997  | -                 | .                       | 0.00008                       | 0                 | 1                                 | -                                    | -                                | 4.0               |
| 5    | c.2337A > C                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | -                                 | 1                                    | -                                | 5.2               |
| 5    | c.2379C > T                    | p.(=)                       | .           | -                 | .                       | 0.00008                       | 0.000111          | 1                                 | 1                                    | -                                | 0.1               |
| 5    | c.2478C > T                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | -                                 | -                                    | 1                                | 4.2               |
| 5    | c.2484C > T                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | -                                 | -                                    | 1                                | 6.0               |
| 7    | c.2742C > T                    | p.(=)                       | rs115759702 | -                 | 0.0018                  | 0.00146                       | 0.00003           | 2                                 | -                                    | -                                | 0.7               |
| 12   | c.3294G > A                    | p.(=)                       | .           | -                 | .                       | .                             | .                 | -                                 | -                                    | 1                                | 9.9               |
| 12   | c.3300 T > G                   | p.(=)                       | rs45516100  | Rahman (2007) [3] | 0.0183                  | 0.02801                       | 0.0278            | 64 (1)                            | 68 (4)                               | 123 (1)                          | 7.9               |
| 13   | c.3495G > A                    | p.(=)                       | rs45439097  | Bogdanova (2010)  | .                       | 0.00108                       | 0.000994          | 4                                 | 7                                    | 9                                | 8.6               |

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<sup>a</sup>Variant positions are reported in reference to NCBI RefSeq NM\_024675.3 (mRNA) and NP\_078951.2 (protein)

<sup>b</sup>Minor allele frequency (MAF) reported in the 1000 Genomes (1000 G) cohort Phase 1

<sup>c</sup>MAF reported in the Exome Variant Server, NHLBI GO Exome Sequencing Project (ESP) [46] (data release ESP6500SI-V2)

<sup>d</sup>MAF reported in ExAC [45] from non-Finnish Europeans, excluding individuals in the database who were part of The Cancer Genome Atlas and therefore known to have had cancer

<sup>e</sup>Number of individuals carrying the variant. Where applicable, the number of homozygous carriers is indicated in parentheses

controls will be required to confirm whether the association of rs152451 with breast cancer is a robust finding. In addition, the wide variation in the frequency of the minor allele in different populations means that cases and controls will have to be carefully matched for ethnicity. This variant is not located in a known protein domain and was consistently found to have predicted benign effects on protein function by all algorithms tested. However, this base change is only 9 bp away from the exon 4 splice donor site and Human Splicing Finder (v3) found that rs152451 could alter an exonic splicing enhancer motif [48], offering a potential mechanism for how this variant could affect *PALB2* function. It should be noted that such a prediction was relatively common for the variants we detected in *PALB2* (35/77 missense or nonsynonymous variants had a similar prediction from at least three

algorithms) and any effect would need to be confirmed by an RNA-based assay.

There has been only one study to date that has examined the likely functional effect of missense variants in *PALB2*, which examined p.Leu939Trp, p.Leu1143Pro and p.Thr1030Ile [49]. The first two variants had subtle but significant effects on homologous recombination repair: p.Leu1143Pro in particular showed decreased repair capacity and binding to BRCA2 and RAD51C. *PALB2* p.Thr1030Ile was unstable, leading to decreased protein levels and this was assumed to impair homologous recombination repair. However, it should be noted that these functional assays were performed by overexpression of a retroviral transgene in a null cell line and may not reflect the heterozygote situation. In our study, p.Leu939Trp was not enriched in cases (four in cases, eight in controls),

p.Leu1143Pro was only seen in two cases and no controls, while p.Thr1030Ile was not observed in either cases or controls.

## Conclusions

Our data strongly support *PALB2* as a breast cancer predisposition gene when considering truncating mutations. We did not see any excess in missense mutations in cancer cases overall, although there may be individual variants that are associated with risk at low penetrance. We advise extreme caution in attributing risk to missense *PALB2* mutations when determining clinical management.

## Additional files

**Additional file 1: Cohort information (Table).** (DOCX 48 kb)

**Additional file 2: *PALB2* truncating variant carrier family pedigrees (Figure).** (PDF 1036 kb)

## Abbreviations

CI: confidence interval; FCC: Familial Cancer Centre; HAPS: Hunter Area Pathology Service; OR: odds ratio; VIP: Variants in Practice.

## Competing interests

The authors declare that they have no competing interests.

## Authors' contributions

IGC, PAJ, Lifepool Investigators, RJS, AHT and GM conceived of and designed the study. ERT, NL, LD, JL, RL, MAD, MWB, MBD, SM, PAJ and SMR carried out experiments, acquired and analysed data. MWB, LD, SM, RJS, MBD, Lifepool Investigators, AHT and PAJ provided data and samples. KLG interpreted data. IGC, PAJ, GM, KLG, SMR, RJS and ERT were involved in drafting the manuscript and all authors read and provided critical feedback on the manuscript. All authors had final approval of the submitted and published versions and are accountable for the work.

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