

# The clonal and mutational evolution spectrum of primary triple-negative breast cancers

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**Primary triple-negative breast cancers (TNBCs), a tumour type defined by lack of oestrogen receptor, progesterone receptor and *ERBB2* gene amplification, represent approximately 16% of all breast cancers<sup>1</sup>. Here we show in 104 TNBC cases that at the time of diagnosis these cancers exhibit a wide and continuous spectrum of genomic evolution, with some having only a handful of coding somatic aberrations in a few pathways, whereas others contain hundreds of coding somatic mutations. High-throughput RNA sequencing (RNA-seq) revealed that only approximately 36% of mutations are expressed. Using deep re-sequencing measurements of allelic abundance for 2,414 somatic mutations, we determine for the first time—to our knowledge—in an epithelial tumour subtype, the relative abundance of clonal frequencies among cases representative of the population. We show that TNBCs vary widely in their clonal frequencies at the time of diagnosis, with the basal subtype of TNBC<sup>2,3</sup> showing more variation than non-basal TNBC. Although *p53* (also known as *TP53*), *PIK3CA* and *PTEN* somatic mutations seem to be clonally dominant compared to other genes, in some tumours their clonal frequencies are incompatible with founder status. Mutations in cytoskeletal, cell shape and motility proteins occurred at lower clonal frequencies, suggesting that they occurred later during tumour progression. Taken together, our results show that understanding the biology and therapeutic responses of patients with TNBC will require the determination of individual tumour clonal genotypes.**

To understand the patterns of somatic mutation in TNBC, we enumerated genome aberrations at all scales from 104 cases of primary TNBC (Affymetrix SNP6.0, 104 cases; RNA-seq, 80 cases; genome/exome sequencing, 65 cases: 54 exomes, 15 genomes with 4 overlapping) (Supplementary Table 1 and Supplementary Fig. 1), annotated with clinical information (Supplementary Table 2). We revalidated 2,414 somatic single nucleotide variants<sup>4,5</sup> (SNVs) (Supplementary Table 3) with targeted deep sequencing to a median of 20,000×

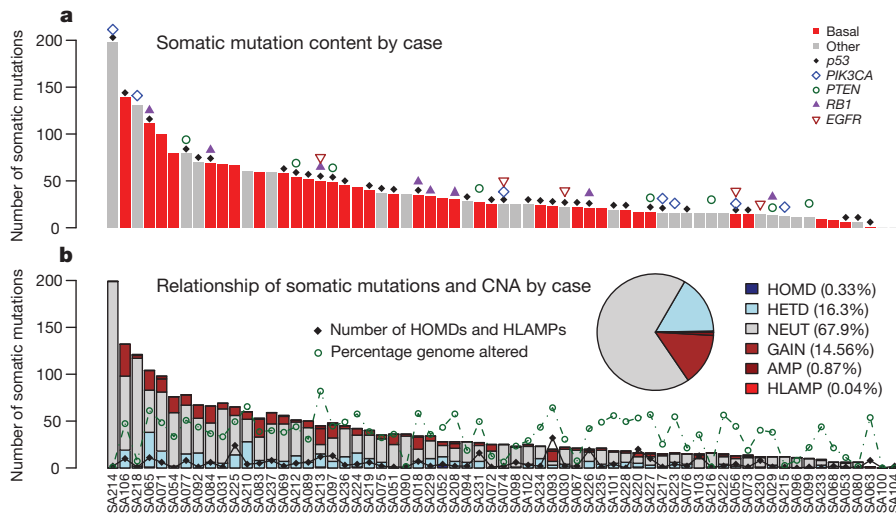
coverage, including 43 non-coding splice site dinucleotide mutations (Supplementary Table 4) and 104 genes with 107 indels (Supplementary Table 5 and Supplementary Methods). Notably, the distribution of somatic mutation abundance varies in a continuous distribution among tumours (Fig. 1a) and seems to be unrelated to the proportion of the genome altered by copy number alterations (CNAs) (Fig. 1b) or tumour cellularity (Supplementary Fig. 2b). Although this distribution could be partially explained by a false-negative rate in mutation discovery, others have noted similar distributions in epithelial cancers<sup>6</sup>, suggesting that the total mutation content of individual tumours may be shaped by biological processes or differential exposure to mutagenic influences in the population.

The overall pattern (Supplementary Fig. 3a, b) of CNA abundance appears similar (Supplementary Fig. 4) to that seen in a larger, independent series of ~2,000 SNP6.0 profiled breast tumours<sup>7</sup>. Among the most frequently observed CNA events (Supplementary Table 6) are the tumour suppressor and oncogenes *PARK2* (6%), *RB1* (5%), *PTEN* (3%) and *EGFR* (5%). Here we report intragenic deletions (Supplementary Fig. 5) in the *PARK2* tumour suppressor<sup>8,9</sup>, specifically linking *PARK2* with TNBC for the first time. Consistent with previous reports in breast cancer<sup>10</sup>, we did not observe frequent recurrent structural rearrangements (Supplementary Fig. 3d and Supplementary Table 7), although we revalidated many individual fusion events involving known oncogenes or tumour suppressors (for example, *KRAS*, *RB1*, *IDH1*, *ETV6*) (Supplementary Tables 8–10).

A comparison of RNA-seq data with genomes/exomes data revealed that only 36% of validated somatic SNVs were observed in the transcriptome sequence (Supplementary Table 3 and Supplementary Fig. 2b). In a recent lymphoma study, similar proportions were observed (137 of 329 somatic mutations expressed in RNA-seq)<sup>11</sup>. As expected, the proportion of low-abundance somatic SNVs observed in RNA is reflected in the distribution of wild-type, heterozygous and homozygous expressed mutations (Supplementary Fig. 2b), consistent with the notion that

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**Figure 1 | Distribution of number of validated somatic mutations by case over 65 cases.**

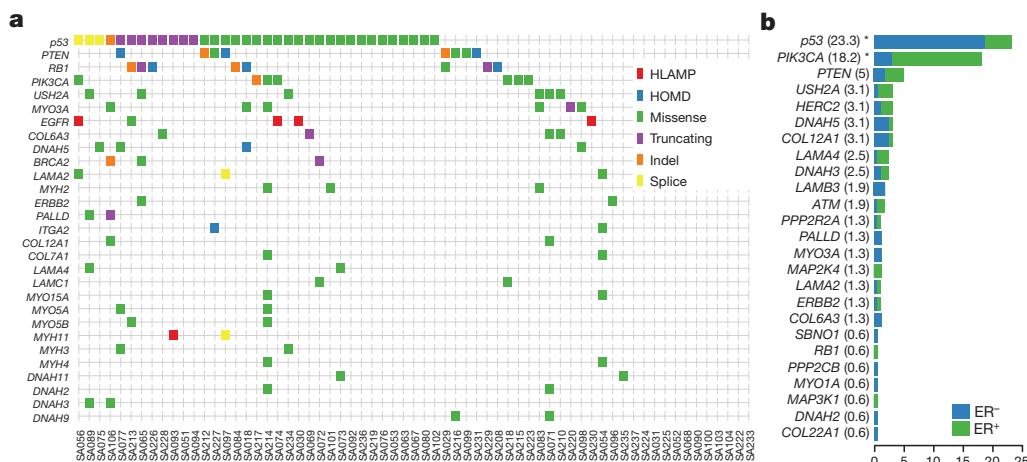
**a**, Mutation frequency (basal, red; other, grey). Patients harbouring known driver gene mutations are indicated. **b**, Case-specific and overall (inset) distributions of mutations in CNA classes. AMP, amplification; GAIN, single copy gain; HETD, hemizygous deletion; HLAMP, high-level amplification; HOMD, homozygous deletion; NEUT, no copy number change. The number of (HOMD, HLAMP) CNAs (black diamonds) and percentage genome altered (green circles) are indicated.

low-abundance alleles may represent rarer clones in the primary tumour. We found 43 splice junction mutations with evidence for an impact on splicing patterns (Supplementary Table 4), encompassing several known tumour suppressors (*p53*, *PIK3R1*; Supplementary Fig. 6) as well as many genes not yet implicated in carcinogenesis. Analysis of 72 somatic mutations in the non-coding space of experimentally determined human regulatory regions<sup>12</sup> showed (Supplementary Table 11) a significant overrepresentation (31.9% versus expected 2.5%, Fisher exact test  $P = 2 \times 10^{19}$ ) of mutations within retinoblastoma-associated protein (RB)-binding sites. Six mutations were predicted to be damaging to RB binding (Supplementary Methods and Supplementary Fig. 7). This is consistent with observations of frequent functional disruption of the RB-regulated cell cycle network<sup>13</sup> in TNBC.

We next searched for mutation enrichment patterns in three ways: by single gene mutation frequency over multiple cases; by the mutation frequency over multiple members of a gene family; and by correlating mutation status with expression networks. First, similar to other studies<sup>14,15</sup>, *p53* is the most frequently mutated gene (Supplementary Table 12) with 62% of basal TNBC (determined by gene expression classification with PAM50 (ref. 16) analysis on RNA-seq expression profiles) and 43% of non-basal TNBC cases harbouring a validated somatic mutation. We also observed frequent mutations in *PIK3CA* at 10.2% (7/65), *USH2A* (Usher syndrome gene, implicated in actin cytoskeletal functions) at 9.2% (6/65), *MYO3A* at 9.2%, *PTEN* and *RB1* at 7.7% (5/65) and a further eight genes (including *ATR*, *UBR5* (also known as *EDD1*), *COL6A3*) at 6.2% (4/65) of cases in the cohort

(Fig. 2a). Considering background mutation rates<sup>17</sup>, *p53*, *PIK3CA*, *RB1*, *PTEN*, *MYO3A* and *GHI* showed evidence of single gene selection ( $q < 0.1$ ) (Supplementary Table 13). Additional recurrent mutations of note occurred in the synuclein genes (*SYNE1* and *SYNE2*, 9.2% 6/65, recently implicated in squamous head and neck cancers<sup>18,19</sup>), *BRCA2* (three cases), and several other well known oncogenes (*BRAF*, *NRAS*, *ERBB2* and *ERBB3*) with mutations in two cases each. Approximately 20% of cases contained examples of potentially ‘clinically actionable’ somatic aberrations, including *BRAF* V600E, high-level *EGFR* amplifications and *ERBB2* and *ERBB3* mutations.

In the second approach we searched for statistically overrepresented gene families and protein functions using the Reactome functional protein interaction database<sup>20</sup> (Supplementary Methods). This analysis quantifies gene family involvement through sparse mutation patterns in functionally connected genes, which would be statistically underrepresented by single gene recurrent mutation analysis. The overrepresented pathways (false discovery rate (FDR) < 0.001) included *p53*-related pathways along with chromatin remodelling, PIK3 signalling, ERBB signalling, integrin signalling and focal adhesion, WNT/cadherin signalling, growth hormone and nuclear receptor co-activators, and ATM/RB-related pathways (Fig. 3a and Supplementary Table 14). We note that the candidate ‘driver’ *MYO3A*, a cytoskeleton motor protein involved in cell shape and motility, relates to several pathways upstream and downstream of integrin signalling. The mutated genes include extracellular matrix (ECM) interactions (laminins, collagens), ECM receptors (integrins), several proteins



**Figure 2 | Population patterns of co-occurrence and mutual exclusion of genomic aberrations in TNBC.**

**a**, Case-specific mutations in known driver genes, plus genes from integrin signalling and ECM-related proteins (laminins, collagens, integrins, myosins and dynein) derived from all aberration types: high-level amplifications (HLAMP), homozygous deletions (HOMD), missense, truncating, splice site and indel somatic mutations are depicted in genes with at least two aberrations in the population. **b**, Distribution of somatic mutations in 25 genes across all exons of 159 additional breast cancers (relative proportion of ER<sup>+</sup> cases in green, and ER<sup>-</sup> in blue), shown as a percentage of cases (in parentheses) with one or more mutations. \* $P < 0.05$ .

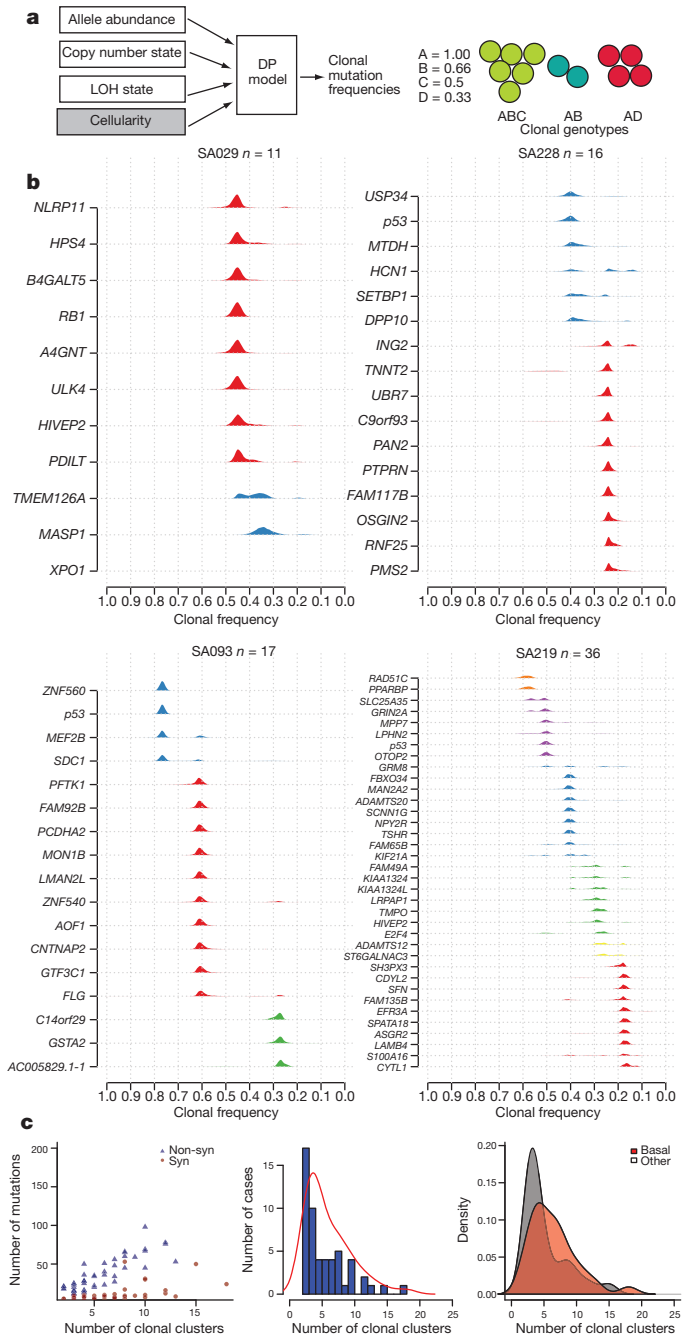


seen to be only partially related to the total number of mutations in a case, groups 1 and 2 having on average fewer mutations per case. The frequent involvement of pathways with *p53*, *PTEN* and *PIK3CA* as members, is noted (Supplementary Fig. 9); however, the case groupings also vary by the progressive inclusion of additional pathways (for example, WNT signalling, integrin signalling, ERBB signalling, hypoxia and PI3K). More than two thirds of cases contained one or more mutations in the actin/cytoskeletal functions group of genes (Supplementary Fig. 9). Some 12% of cases did not contain somatic aberrations in any of the frequent drivers or cytoskeletal genes (Supplementary Table 12). This suggests that primary TNBCs are mutationally heterogeneous from the outset, with some patients' tumours having a small number of implicated pathways and few mutations, whereas other patients present with tumours containing extensive mutation burdens and multiple pathway involvement.

Motivated by the observation that early primary TNBCs show a wide variation of mutation content, we asked whether the clonal composition of these primary cancers is similarly varied. We and others have shown<sup>22,23</sup> how deep-frequency measurements of allelic abundance can be used to study tumour clonal evolution. Clonal mutation frequency, a compound measure of clonal complexity, (Fig. 4a) can be estimated from allele abundance, once the influence of copy number states, regional loss of heterozygosity (LOH state) and tumour cellularity have been considered (although we note that approximately 68% of SNVs in this study are in diploid, neutral regions). To extend allelic abundance measurements to estimation of clonal frequencies, we implemented a Dirichlet process clustering model (pyclone; Supplementary Methods and Supplementary Fig. 10) that simultaneously estimates the genotype and clonal frequency given a list of deeply sequenced mutations and their local copy number and heterozygosity contexts.

Using the set of deeply sequenced (median 20,000 $\times$ ), validated SNVs, our analysis revealed (Fig. 4b) that groups of mutations within individual cases have different clonal frequencies, indicative of distinct clonal genotypes. Remarkably, the tumours exhibit a wide spectrum of modes over clonal frequencies (Fig. 4b and Supplementary Fig. 11), with some cases showing only one or two frequency modes (Fig. 4b), indicating a smaller number of clonal genotypes, whereas other tumours exhibit multiple clonal frequency modes, indicating more extensive clonal evolution. Consistent with early 'driver gene' status, mutations in known tumour suppressors such as *p53* tend to occur in the highest clonal frequency group in most tumours. However, in some cases (for example, SA219, SA236; Fig. 4b, Supplementary Fig. 11) *p53* resides in lower-abundance clonal frequency groups (Supplementary Fig. 12 and Fig. 3a), suggesting that it was not the founding event. Although the number of clonal frequency modes tends to increase with the number of mutations, the relationship is not strictly linear (Fig. 4c). To determine whether basal and non-basal cancers differ in their clonality, we compared the distribution of clonal modes (clusters) by case and as an overall distribution, and note that basal TNBCs have more clonal frequency modes than non-basal TNBCs (Fig. 4c). Both of these distributions emphasize a key observation; namely, that at the time of diagnosis TNBCs already display a widely varying clonal evolution that mirrors the variation in mutational evolution.

Finally, we asked where key pathways appear in the distribution of clonal frequency groups. We examined the clonal frequency of genes in each pathway and ascertained if there was a deviation away from the distribution of clonal frequency for all mutations. As expected, pathways involving *p53* and *PIK3CA* showed significantly skewed distributions (Wilcoxon,  $q < 0.01$ ; Fig. 3b and Supplementary Fig. 12) towards higher clonal frequencies, consistent with their roles in early tumorigenesis (Fig. 3a and Supplementary Table 17). Intriguingly, pathways with cytoskeletal genes such as myosins, laminins, collagens and integrins tend to have lower median clonal frequencies, suggesting that somatic mutations in these genes are acquired much later (Fig. 3b). Notably, the median clonal frequency for Reactome pathway 'p53



**Figure 4 | Clonal evolution in TNBC.** **a**, Schematic representation of integration of CNA, LOH, allelic abundance measurements and normal cell contamination for clonal frequency estimation using a Dirichlet process (DP) model (left). Example of a mixture of three clonal genotypes composed of four mutations (A, B, C, D) and their resulting clonal frequencies. **b**, Estimated clonal frequencies for four cases are shown as the distribution of posterior probabilities from the pycclone model (Supplementary Methods). Clonal frequency distributions are coloured by their frequency group membership. **c**, Left, relationship of mutation abundance (synonymous (Syn) and non-synonymous (Non-syn)) and the inferred number of clonal clusters. Middle, distribution and kernel density (red line) of the number of inferred clonal clusters over 54 TNBCs. Right, kernel density distribution of clonal clusters for basal (red) and non-basal (grey) tumours.

pathway feedback loops', including 46 mutations in *ATM*, *ATR*, *NRAS*, *PIK3CA*, *PTEN*, *SIAH1* and *p53*, was 73% (Wilcoxon,  $q = 0.0007$ ), whereas 'integrin cell surface interactions', including 23 mutations in integrin, laminin and collagen genes, had a median clonal frequency of 42% (Wilcoxon,  $q = 0.9569$ ).

Primary TNBCs are still treated as if they were a single disease entity, yet it is clear they do not behave as a single entity in response to current therapies. Here we show for the first time, using next-generation sequencing mutational profiling methods, that treatment-naïve TNBCs display a complete spectrum of mutational and clonal evolution, with some patients' tumours showing only a few somatic coding sequence point mutations with a limited number of molecular pathways implicated, whereas other patients' tumours exhibit considerable additional mutational involvement. Moreover, the clonal heterogeneity of these cancers is also a continuum, with some patients presenting with low-clonality cancers and other cases exhibiting more extensive clonal evolution at diagnosis. In this respect, the basal expression subtype of TNBCs also tends to show higher clonality at diagnosis, although the relationship is not exact.

In clonally evolving tumours, identification of genes by single gene mutation frequency measurements will probably favour early driver genes, because the subsequent involvement of multiple additional pathways during tumour progression is unlikely to be observed as a frequent single gene mutation. The clonality analysis emphasizes this point: known drivers such as *p53*, *PIK3CA* and *PTEN* have among the highest clonal frequencies, whereas mutations in cell shape/motility and ECM-signalling genes appear in the lower clonal frequency groups, distributed over many genes. Although *p53* somatic mutations are clearly early events, the clonal frequencies observed in some TNBC suggest that they are not always the first event, raising a question about what drives early clonal expansion in some of these cancers. Our findings suggest that each TNBC at the time of primary diagnosis may be at a very different phase of molecular progression, with possible implications for approaches to the biology of low clonality versus high clonality primary tumours.

## METHODS SUMMARY

The genomes and transcriptomes of 104 TNBCs were profiled with Affymetrix SNP6.0 arrays (all cases), RNA-seq (80 cases; Illumina GAI), and whole exome/genome sequencing (65 cases; tumour and normal DNA). Exomes were obtained using Agilent's Human All Exon SureSelect Target Enrichment System v.1 followed by Illumina GAI sequencing, and whole genomes were sequenced using Life Technologies SOLiD system. Data were analysed using computational approaches to detect somatic SNVs<sup>4,5</sup>, indels, copy number alterations, gene fusions and gene expression patterns. Predictions were then validated using orthogonal experimental assays, including targeted ultra-deep amplicon sequencing of SNVs to ~20,000× redundancy. We determined single genes under selection using a statistical approach that considers patient-specific background mutation and transition/transversion rates. Mutations predicted to alter transcriptional profiles were determined using an integrated bipartite graph-based method (driverNet) that associates genomic aberrations with outlying expression patterns informed by pre-defined pathway gene sets. Disrupted pathways were determined using the Reactome FI Cytoscape plugin. Clonal analysis was performed (cases with >10 mutations) using a Dirichlet process statistical model that simultaneously estimates clonal frequencies and mutation genotype given deeply sequenced somatic SNVs and copy number estimates. Experimental assays and analytical methodology are detailed in the Supplementary Information.

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Supplementary Information is linked to the online version of the paper at [www.nature.com/nature](http://www.nature.com/nature).

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**Author Contributions** S.A., S.P.S., C. Caldas and M.A.M. designed and implemented the research plan and wrote the manuscript. S.P.S., A.R., R. Goya, G. Ha, J.D., G. Haffari, A. Bashashati, A. McPherson, K.S., A.C., R. Giuliani, A.H.-M., J.R., D.L., I.B., R.V., S.W.C., M.G., I.M.M., S.J., C. Curtis, O.M.R., P.D.P., V.B. and W.W.W. conducted bioinformatic analyses of the data and/or gave advice on analytic methodology. G.T. conducted histopathological review and immunohistochemistry. A.O., Y.Z., G.T., K.T., L.M.P., J.K., A.B., D.Y., A.T., N.D., T.Z., S.-F.C., K.M. and M.H. conducted sequencing or experimental validation of somatic aberrations. D.Y., A. Moradian, S.-W.G.C. and G.B.M. conducted proteome validation of splicing. P.W., K.G., S.C., S.-F.C., G.T., J.M., C. Caldas, P.D.P. and D.H. collected and interpreted clinical data. S.D., J.F.C., T.T., M.S., P.G. and C.J.E. contributed materials or reagents. K.H., V.T., T.H., M.H. and M.A.M. generated sequence data.

**Author Information** Aligned exome/genome sequence data, RNA-seq data and Affymetrix SNP6.0 data sets are available at the European Genome-phenome Archive (<http://www.ebi.ac.uk/ega/>) under study accession number EGAS00001000132. Normal reference RNA-seq datasets are available at the NCBI Short Read Archive (<http://www.ncbi.nlm.nih.gov/Traces>) under study accession number SRP000930. Reprints and permissions information is available at [www.nature.com/reprints](http://www.nature.com/reprints). The authors declare no competing financial interests. Readers are welcome to comment on the online version of this article at [www.nature.com/nature](http://www.nature.com/nature). Correspondence and requests for materials should be addressed to S.A. ([saparicio@bccrc.ca](mailto:saparicio@bccrc.ca)), C. Caldas ([carlos.caldas@cancer.org.uk](mailto:carlos.caldas@cancer.org.uk)), S.P.S. ([sshah@bccrc.ca](mailto:sshah@bccrc.ca)) or M.M. ([mmarra@bcgsc.ca](mailto:mmarra@bcgsc.ca)).