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REVIEW

Variants of uncertain clinical significance in hereditary breast and ovarian cancer genes: best practices in functional analysis for clinical annotation

Alvaro N Monteiro ¹, Peter Bouwman ², Arne N Kousholt ²,
Diana M Eccles ³, Gael A Millot ⁴, Jean-Yves Masson ⁵,
Marjanka K Schmidt ², Shyam K Sharan ⁶, Ralph Scully ⁷,
Lisa Wiesmüller ⁸, Fergus Couch ⁹, Maaikje P G Vreeswijk ¹⁰

For numbered affiliations see end of article.

Correspondence to

Dr Alvaro N Monteiro, Cancer Epidemiology, H. Lee Moffitt Cancer Center and Research Institute, Tampa, FL 33612, USA; Alvaro.Monteiro@moffitt.org and Dr Maaikje P G Vreeswijk, Human Genetics, Leiden University Medical Center, Leiden, 2300RC, The Netherlands; M.P.G.Vreeswijk@lumc.nl

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INTRODUCTION

Since the identification and cloning of *BRCA1* in 1994,¹ and shortly thereafter of *BRCA2*,² genetic tests of germline DNA to identify pathogenic variants in genes linked to hereditary breast and ovarian cancer (HBOC) have become mainstream.³ These tests are critical to identify women at increased risk relative to the general population. Women at moderate risk ($2 \leq \text{relative risk (RR)} < 4$) may benefit from enhanced screening and chemoprevention while those at high risk ($\text{RR} \geq 4$), including those with *BRCA1* and *BRCA2* pathogenic variants, may also benefit from preventive surgery. Germline mutation testing is also becoming increasingly relevant in the cancer treatment setting because carriers of pathogenic variants in *BRCA1/2* may benefit from poly-ADP ribose polymerase (PARP) inhibitors.^{4,5} Importantly, genetic tests can identify individuals in HBOC families who do not carry the relevant predisposing allele and are not at elevated risk of cancer.⁶

A significant fraction of documented variants in *BRCA1* and *BRCA2* are considered variants of uncertain clinical significance (VUS), for which cancer association has not been assessed or could not be determined due to insufficient information (table 1). In ClinVar (<https://www.ncbi.nlm.nih.gov/clinvar/>), a clinically oriented database, currently ~37% of *BRCA1* and ~45% of *BRCA2* unique variants recorded are VUS. Thus, there is a critical need to classify variants according to their pathogenicity.

Over the past decade, functional assays have emerged that can be included as a source of evidence to classify variants according to their pathogenicity, with the potential to greatly accelerate classification.⁷ Here, we discuss several technical and conceptual aspects relevant for the use of functional assays in the classification of variants. We present best practice recommendations to improve annotation quality and accuracy, and to provide a basis for the comparison and integration of functional data from different laboratories (box 1). For the coming years, we anticipate that recent technological developments such as VAMP-Seq (variant abundance by massively parallel sequencing) or high-throughput Clustered Regularly Interspaced

Short Palindromic Repeats (CRISPR)-based saturation mutagenesis will enable the functional assessment of every missense variant for all moderate-risk and high-risk HBOC genes.^{8–10} Once established and validated, these catalogues of functional data will provide valuable information for clinical annotation. The recommendations proposed here are the result of a discussion that started at a Netherlands Cancer Institute workshop on Functional Analysis of Sequence Variants in Hereditary Breast and Ovarian Cancer Genes (Amsterdam, The Netherlands) and was followed by additional discussion and extensive refinement. It represents a consensus view that was self-developed by an international group of investigators (the authors) who have been active in this field.

ASSESSMENT OF THE EVIDENCE FOR ASSOCIATION OF EACH GENE WITH HBOC RISK

The first step in developing or interpreting results from functional assays is to understand the level of evidence that links a particular gene to breast and ovarian cancer risk.³ To date, there are nine genes for which an association between protein-truncating variants and breast cancer risk has been established (*ATM*, *BRCA1*, *BRCA2*, *CDH1*, *CHEK2*, *PALB2*, *PTEN*, *STK11* and *TP53*) and several more (*BARD1*, *FANCM*, *NBN*, *NF1*, *MLH1*, *MSH2*, *MSH6*, *PMS2*, *RAD51C* and *RAD51D*)^{3 11–13} for which association has been suggested but not yet firmly established. At least 12 genes have been implicated in ovarian cancer risk (*ATM*, *BRCA1*, *BRCA2*, *BRIP1*, *MLH1*, *MSH2*, *MSH6*, *PALB2*, *PMS2*, *RAD51C* and *RAD51D*).^{14 15}

Development of functional tests for emerging genes provides opportunities to uncover new mechanistic aspects of their biology and identify functional domains. However, developers of functional assays should consider that clinical recommendations are unlikely to be made based on variants in genes for which the association has not been robustly established. Thus, a detailed understanding of the strength of evidence for association between each gene (and its variant alleles) and HBOC risk should be sought to evaluate the clinical utility of a proposed functional assay.



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Table 1 Fraction of VUS in *BRCA1* and *BRCA2*

Databases	BIC*		ClinVar†		BRCA Exchange‡		gnomAD§	
	N	% (%VUS)	N	% (%VUS)	N	% (%VUS)	N	%
<i>BRCA1</i> unique variants	1781	100	5821	100	7898	100	2936	100
<i>BRCA1</i> VUS	891	50.0 (100)	2146	36.9 (100)	5186	65.7 (100)	n/a	n/a
<i>BRCA1</i> missense	607	34.1 (68.1)	1715	29.5 (79.9)	1892	24.0 (36.5)	938	31.9
<i>BRCA1</i> missense VUS	569	31.9 (63.9)	1633	28.1 (76.1)	1714	21.7 (33.1)	n/a	n/a
<i>BRCA2</i> unique variants	2000	100	8119	100	10 422	100	4262	100
<i>BRCA2</i> VUS	1065	53.3 (100)	3615	44.5 (100)	6980	67.0 (100)	n/a	n/a
<i>BRCA2</i> missense	891	44.6 (83.7)	3111	38.3 (86.1)	3484	33.4 (49.9)	1909	44.8
<i>BRCA2</i> missense VUS	838	41.9 (78.7)	3011	37.1 (83.3)	3190	30.6 (45.7)	n/a	n/a

*BIC (<https://research.nhgri.nih.gov/bic/>) is a locus-specific database established in 1995 for *BRCA1* and *BRCA2* variants, including loci primarily found in clinical or research testing.

†ClinVar (<https://www.ncbi.nlm.nih.gov/clinvar/>) is a public archive of reports of the relationships among human variations and phenotypes and includes submissions reporting variants found in patient samples from clinical or research testing, and from the literature (note: ClinVar also includes BIC data).⁶⁹ VUS counts also include conflicting assessments.

‡BRCA Exchange (<http://brcaexchange.org/>) pools data on *BRCA1/2* genetic variants and corresponding clinical data from around the world (including BIC, ClinVar, 1000 Genomes Project). BRCA Exchange is part of the Global Alliance for Genomics and Health. VUS counts also include 'not yet reviewed'.

§gnomAD (<http://gnomad.broadinstitute.org/>), initially released as ExAC aggregates and harmonises both exome and genome sequencing data from a wide variety of large-scale sequencing projects. It does not contain pathogenicity assessments. All searches were conducted in December 2017.

BIC, Breast Cancer Information Core; gnomAD, The Genome Aggregation Database; VUS, variants of uncertain clinical significance.

An additional aspect to consider when developing a functional assay is the proportion of missense VUS that are probably pathogenic. Missense variation is unlikely to significantly affect the overall protein function when located in disordered regions or

in repeat motifs. Therefore, functional assays for these regions (or for a protein with a large portion of its coding sequence composed by these regions) may not be a priority.

Box 1 Summary recommendations for the development, reporting and interpretation of functional assays

- ▶ Assess the strength of evidence for association between each gene and hereditary breast and ovarian cancer risk to evaluate the potential clinical utility of a proposed functional assay.
- ▶ Consider the assumptions, biological characteristics, controls and limitations of each assay.
- ▶ Choose genomic DNA/cDNA/protein sequences that correspond to the coding sequence of the most commonly found haplotype in non-affected individuals to be used as a reference (wild-type).
- ▶ A minimal set of non-pathogenic and pathogenic variants should be used as internal reference for each run of an assay. Larger sets of reference variants should be used to assess the overall sensitivity and specificity of an assay.
- ▶ Verify that elements of the assay (reagents and data) have been through quality control, including periodical verification of cell line and strain identity (eg, identity by short tandem repeat analysis for mammalian cells and phenotyping for yeast strains) and quality (mycoplasma testing).
- ▶ Due to protein stability issues, exercise caution when developing and interpreting results from model organisms that are cultured at temperatures lower than 37°C.
- ▶ Inspect data to identify and correct batch effects.
- ▶ Do not assume that intermediate levels of activity necessarily reflects intermediate risks.
- ▶ In addition to loss-of-function effects also consider dominant negative and gain-of-function effects.
- ▶ When reporting results use explicitly defined terminology and aim for the development of a controlled vocabulary.
- ▶ Be explicit about assay's limitations, performance metrics and thresholds used to classify variants.

ASSESSMENT OF VARIANT PATHOGENICITY

Genes implicated in HBOC are tumour suppressor genes and therefore variants leading to disruption of function(s) are usually considered pathogenic for clinical purposes. Notable exceptions of variants with dominant negative or gain-of-function have also been reported.¹⁶ Loss-of-function genetic alterations include frameshift and nonsense variants leading to truncation of a functionally important segment of the protein, alterations of donor and acceptor splice sites and large genomic rearrangements altering segments of the coding region. Conversely, synonymous changes without effect on mRNA splicing are considered non-pathogenic. These variants can be reliably classified by a rule-based system that incorporates general DNA/RNA/protein rules and takes into account exceptions specific to each gene (ENIGMA rules for classification of *BRCA1* and *BRCA2* variants: <https://enigmaconsortium.org/library/general-documents/enigma-classification-criteria/>).

For a significant fraction of rare variants pathogenicity cannot be predicted based on DNA changes alone. Primarily, these variants include small deletions or insertions that do not disturb the reading frame, missense changes, intronic and exonic variants that may lead to altered mRNA splicing and in-frame exon deletions or duplications. Missense variants represent the largest contributor to this class, making up to 79.9% and 86.1% of all *BRCA1* and *BRCA2* VUS in ClinVar, respectively (table 1).

Classification of these *BRCA1/2* variants for clinical use can be based on the American College of Medical Genetics and Genomics/Association for Molecular Pathology (ACMG/AMP) guidelines in which pathogenicity is determined by the entire body of evidence.⁷ In this proposed five-tier classification, variants with >90% certainty of being pathogenic (which includes likely pathogenic and pathogenic variants), are considered actionable, and carriers are managed as high risk (RR ≥4). Evidence is qualitatively weighed as strong, moderate, supporting or not used. Functional data provide strong (PS3: well-established functional studies show a deleterious effect) and moderate (PM1:

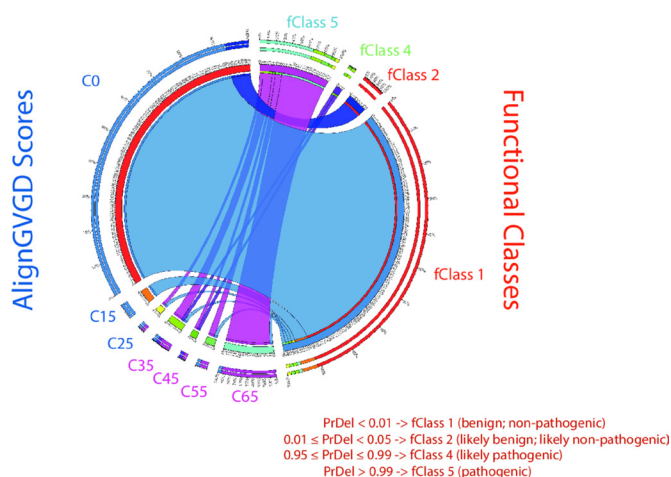


Figure 1 Circos plot illustrating the concordance between alignGVGD predictions and experimental data derived from analysis of BRCA1 C-terminal variants (aa 1396–1863). Variants were analysed using the transcription activation assay²⁹ and assigned to functional classes by VarCall.⁵⁷ Blue ribbons show that all variants scoring C0 and C15 in alignGVGD and predicted to have no or little functional impact score as non-pathogenic or likely non-pathogenic (fClass 1 or 2) in a validated functional assay. Conversely, most variants scoring as C65 and predicted to have a functional score as fClass 5 (thick purple ribbon). Despite the strong concordance between alignGVGD and the transcriptional assay, a small but significant fraction of variants scoring as fClass 1 were incorrectly predicted (C45–C65).

mutational hot spot or well-studied functional domain without benign variation) evidence for pathogenicity; and strong (BS3: well-established functional studies show no deleterious effect) evidence for benign impact.⁷ Reproducible and robust functional assays that have been validated are considered the most well-established source.

Alternatively, classification of *BRCA1/2* variants can be based on a multifactorial statistical model that incorporates data on family history, co-segregation and co-occurrence with another allele with a known pathogenic variant in the same gene (because biallelic inactivation is embryonic lethal while biallelic partial loss of function leads to Fanconi anaemia).¹⁷ In the International Agency for Research on Cancer (IARC) proposed five-tier classification, variants with >95% certainty of being pathogenic are considered actionable.¹⁸ Variants that reach >1.0% allele frequency in the population are considered unlikely to be pathogenic on their own but there is simply insufficient clinic and family-based genetic information to determine the likelihood of pathogenicity of many uncommon (<1.0%) variants. Currently, functional data are not integrated in these multifactorial statistical models.

The effects of these rare VUS can be predicted by a wide variety of publicly available in silico tools with variable performance.^{19–20} For tools that use multiple sequence alignments, performance has been tied to the choice of alignments and calibration.^{21–23} Reliance on multiple sequence alignment and evolutionary approaches may also generate false negatives. For example, Kondrashov *et al* have estimated that approximately 10% of variants that are classified as ‘tolerated’, because a corresponding amino acid residue is found in the cognate position in another species, only score as ‘tolerated’ because of compensatory variation elsewhere in the protein sequence.²⁴ Despite these limitations, algorithms are constantly improving and the concordance between some predictors and empirical data is

sufficiently high (figure 1) to guide prioritisation of variants for functional assessment. However, empirical functional data will be necessary for the robust clinical annotation of uncommon variants.

THE SPECTRUM OF LOW-RISK, MODERATE-RISK AND HIGH-RISK ALLELES IN HBOC GENES

When developing or interpreting a functional assay for VUS classification, the level of risk conferred by pathogenic variants should be considered. Findings of pathogenic variants in a low penetrance gene ($RR < 2$) currently do not trigger clinical recommendations making the development of a functional assay a low priority. It is also plausible that variants within the same gene may span the spectrum of low ($RR < 2$), moderate ($2 \leq RR < 4$) and high ($RR \geq 4$) risk. In other words, distinct ‘pathogenic’ variants in the same gene may carry significantly different levels of risk.

Currently, the multifactorial statistical model for classification of *BRCA1* or *BRCA2* variants determines whether a variant is likely to be pathogenic. The clinical inference of the IARC classification is based on variants that typically are associated with a high cancer risk comparable to a truncating variant in *BRCA1* or *BRCA2* ($RR \geq 4$).¹⁸ However, it is now clear that some pathogenic missense variants in *BRCA1* (p.R1699Q and p.V1736A) and *BRCA2* (p.Y3035S and p.G2508S) confer only moderate breast cancer risk ($2 \leq RR < 4$).^{25–27} On the other hand, the *BRCA2* p.K3326X, classified by the model as non-pathogenic, was shown to confer a mildly increased risk ($RR = 1.4$) of breast and ovarian cancer.²⁸ Although finding this variant would not trigger a change in clinical recommendation currently, this variant can contribute to polygenic risk scores based on common genetic variants that are now being used for risk stratification, and may prove effective for selection of screening and prevention options.

Some assays may have the ability to reflect different levels of risk depending on the dynamic range of the read-out and on the specific biological assay being performed. It is important to stress that it should not be assumed that intermediate levels of activity in a biochemical or biological assay necessarily reflects intermediate risks. Several reference variants with known intermediate risks should be used to determine the ability of an assay to reflect the continuum of risk. While the transcription activation assay for *BRCA1* does not seem to discriminate between variants with intermediate risks from variants associated with high risk,²⁹ the *BRCA2* homologous recombination (HR) assay may be able to distinguish high from moderate and low/neutral as suggested by functional assessment of variant p.Y3035S.³⁰ For genes in which pathogenic variants are clearly associated with disease risk, a two-stage reporting system has been proposed, that is, the first stage would establish pathogenicity of the variant based on multiple criteria and the second stage would denote the likely severity or clinical consequence for that variant (high, moderate or low risk).³¹ Capturing the full spectrum of risk associated with distinct pathogenic variants is a critical challenge for assay development and for reporting laboratories.

Classification from both ACMG/AMP and multifactorial models are designed to distinguish high-risk variants ($RR \geq 4$; actionable) from not high-risk ($RR < 4$) variants and are currently not suitable to identify moderate-risk variants. From a clinical standpoint, while these models have a binary outcome (actionable vs non-actionable), carriers of moderate-risk variants ($2 \leq RR < 4$) may also benefit from enhanced screening.³²

FUNCTIONAL ASSAYS FOR HBOC GENE VARIANT CLASSIFICATION

For the purposes of our discussion, 'function' is considered as any aspect defined by the Gene Ontology Consortium³³ molecular function, cellular component and biological processes. A 'functional assay' is generally defined as any *in vitro* and *in vivo* system able to determine the impact of a variant by assessing its effect on protein stability, conformation and function. Thus, assays for splicing alterations are not considered functional assays for the purposes of this manuscript (for assessment of splice variants please refer to Thomassen *et al* and Walker *et al*^{34,35}).

Several characterised functions of the BRCA1 and BRCA2 proteins have been exploited in the development of functional assays.^{36–38} Reflecting their central role in DNA damage repair, many assays revolve around measuring the ability of the variant to promote survival following DNA insults, such as treatment with ionising radiation or DNA damaging compounds. In addition to these viability assays, specific biochemical assays such as those measuring HR or ubiquitylation are also rooted in the known biology of BRCA1 and BRCA2. Finally, more limited biochemical assays, measuring binding to specific interacting proteins have also been applied to the functional analysis of variants.^{36–38}

In general, there is enough evidence to tie each of these functions to the aetiology of tumours arising in carriers. However, their individual contribution to cancer risk is unclear, making it difficult to determine which assay is more or less biologically appropriate, or to assign different weights to results obtained from different assays. Rather, the determination of which assays should be used for clinical annotation relies on their accuracy, and not on their biological properties. Preliminary analysis has shown that these functional assays display high (>80%) sensitivity and specificity (table 2).³⁹

After the development of a large number of functional assays for high-risk genes (BRCA1, BRCA2 and TP53),^{36–37,40} significant attention has been focused on developing assays for other high-risk/moderate-risk genes such as PALB2, ATM and CHEK2.^{41–46} However, here we will focus on functional assays for missense variants of BRCA1 and BRCA2 as exemplars from which we have derived general guidelines.

Functional assays can be defined by three broad categories according to their experimental set-up (cell-free or cell-based), host (human or model organism) and read-out. Result interpretation requires careful consideration of the assumptions, the biological characteristics and limitations of each assay.

Cell-free systems either test a specific biochemical activity *in vitro* (eg, phosphopeptide binding, ubiquitin ligase activity, DNA combing, DNA binding, DNA recombination), protein-protein interactions (eg, yeast two-hybrid screening, co-immunoprecipitations) or the effect of different factors on protein structure and stability (eg, protease sensitivity, calorimetry). Interpretation of results from cell-free assays should consider that they are restricted to specific functions, sometimes limited to specific regions of the protein, and may be particularly sensitive to temperature, buffer conditions and concentrations of exogenous substrates.

Cell-based systems use a human or model organism (eg, yeast, bacteria or mouse) host cell as the basis for the assay. Cell-based systems can be further distinguished as *in cellulo* (when the assay context is a single cell) or *in vivo* (in the context of a whole metazoan organism), although there are currently no established *in vivo* functional assays for VUS. We recommend periodical authentication of cell line and strain identity by short tandem

repeat analysis and phenotyping, respectively. Cell lines should be checked regularly for mycoplasma infection. Interpretation of results from assays performed in model systems should consider the degree of divergence of proteins from the host involved in the assay, differences in biology and in growth conditions.

Cell-based assays can be further defined by read-out. Reporter systems include those in which the read-out for functional impact is an ectopic reporter (eg, transcription activation or HR assays) or in which ectopic overexpression in a heterologous system leads to a defined phenotype (eg, small colony phenotype in yeast). Limitations of reporter systems based on ectopic expression may include artefacts of over expression. Alternatively, assays in which the full length variant allele/protein replaces the endogenous gene/protein and defined biological processes are assessed are considered complementation/perturbation assays. Some assays may combine reporter systems and complementation.

Ultimately, the value of an assay will depend on its performance, defined using a set of known non-pathogenic and pathogenic control variants as reference (see below). Given the complexity of the interaction of multiple biochemical functions and breast and ovarian cancer phenotype, it is uncertain that there will be a single comprehensive and highly accurate functional assay. Rather, the combination of approaches using diverse sources of data obtained with transparent methodology and careful interpretation is likely to solve the challenges of VUS in HBOC genes.

Mouse models, although not suitable for high throughput analysis, can be helpful in determining the effect of such variants on tumour predisposition and treatment response. *Brca1* mutant mice expressing the p.L26A variant showed that the E3 ubiquitin ligase activity of BRCA1 is dispensable for tumour suppression and mice expressing 185delAG (c.68_69delAG; p.E23VfsTer17) revealed the hypomorphic nature of this pathogenic variant in response to therapy.^{47–49} A knock-in mouse model of the BRCA2 p.G25R variant, which had no effect on ES cell viability but had subtle defect in HR, showed a significant increase in tumour formation in mutant animals.⁵⁰ Similarly, the effect on tumour predisposition of an alternatively spliced *Brca2* transcript lacking exons 4–7 was revealed in mutant mice lacking these exons.⁵¹

REQUIREMENTS FOR A CLINICALLY RELEVANT FUNCTIONAL ASSAY

The analytical validity is the degree of accuracy with which a functional assay correctly classifies variants as pathogenic or non-pathogenic. For each assay performance metrics (true positive rate or sensitivity; true negative rate or specificity; false positive and false negative rates; positive and negative likelihood ratios; false discovery rate; false omission rate; positive predicted value or precision; negative predictive value; accuracy and diagnostic OR) should be derived from testing a panel of known pathogenic and non-pathogenic variants. The recommendation is to choose a set of pathogenic and non-pathogenic missense controls whose likelihood of pathogenicity has been established by the multifactorial statistical model and can be found in a recent ENIGMA publication.^{36–37,52}

There are no specific recommendations about which threshold of sensitivity or specificity should be used to consider the inclusion of data from a functional assay for variant classification. Plon *et al*¹⁸ have pointed out that clinical decisions based on predictive values of 80%–85% are normally used in oncology. A more strict approach would require that the lower bound of the 95% CI be above the suggested 80%–85% threshold but that

Table 2 Categories and performance of functional assays for *BRCA1* and *BRCA2*

Gene	Assay	Set-up	Read-out	# of variants assessed (# of known non-pathogenic; pathogenic)*	Sensitivity (95% CI)†	Specificity (95% CI)†	Reference
<i>BRCA1</i>	Colony size	Cell-based (yeast)	Complementation/perturbation	40 (15; 25)	0.96 (0.80 to 1.00)	0.93 (0.68 to 1.00)	58
<i>BRCA1</i>	Yeast localisation	Cell-based (yeast)	Complementation/perturbation	40 (15; 25)	0.84 (0.64 to 0.95)	0.93 (0.68 to 1.00)	58
<i>BRCA1</i>	Transcription activation	Cell-based (HEK293T)	Reporter system	204 (25; 40)	1.00 (0.75 to 1.00)	1.00 (0.83 to 1.00)	29
<i>BRCA1</i>	BARD1 binding	Cell-based (yeast)	Reporter system	35	n/a	n/a	70
<i>BRCA1</i>	UbcH5a binding	Cell-based (yeast)	Reporter system	35	n/a	n/a	70
<i>BRCA1</i>	Ubiquitin ligase activity	Cell-free (in vitro)	In vitro enzymatic activity	35	n/a	n/a	70
<i>BRCA1</i>	Protease sensitivity	Cell-free (in vitro)	In vitro binding activity	117 (10; 14)	0.79 (0.49 to 1.00)	0.80 (0.44 to 0.98)	56
<i>BRCA1</i>	Phosphopeptide binding activity	Cell-free (in vitro)	In vitro binding activity	117 (10; 14)	0.86 (0.57 to 0.98)	1.00 (0.69 to 1.00)	56
<i>BRCA1</i>	Phosphopeptide binding specificity	Cell-free (in vitro)	In vitro binding activity	117 (10; 14)	1.00 (0.77 to 1.00)	0.99 (0.56 to 1.00)	56
<i>BRCA1</i>	ES cell survival	Cell-based (mouse ES cells)	Complementation/perturbation	86 (25; 9)‡	n/a	n/a	62
<i>BRCA1</i>	Cisplatin sensitivity	Cell-based (mouse ES cells)	Complementation/perturbation	86 (25; 9)	1.00 (0.63 to 1.00)	1.00 (0.83 to 1.00)	62
<i>BRCA1</i>	BARD1 binding	Cell-based (yeast)	Reporter system	1287 (3; 19)	n/a	n/a	71
<i>BRCA1</i>	Ubiquitin ligase activity	Cell-free (in vitro)	In vitro enzymatic activity	1287 (3; 19)	n/a	n/a	71
<i>BRCA1</i>	Haploid cell survival	Cell-based (HAP1 cells)	Complementation/perturbation	3893 (22; 162)§	0.967	0.982	10
<i>BRCA1</i>	Homologous recombination	Cell-based (RG37-shBRCA1 cells)	Complementation/perturbation	78 (6; 7)	1.00	1.00	72
<i>BRCA1</i>	Localisation	Cell-based (RG37-shBRCA1 cells)	Complementation/perturbation	78 (6; 7)	0.714	1.00	72
<i>BRCA1</i>	Protein expression and stability	Cell-free (in vitro)	In vitro solubility and thermostability	78 (6; 7)	0.714	0.83	72
<i>BRCA1</i>	Phosphopeptide binding activity	Cell-free (in vitro)	In vitro binding activity	42 (5; 2)	n/a	n/a	72
<i>BRCA1</i>	Homologous recombination	Cell-based (HeLa-DR-FRT)	Complementation/perturbation	1056 (5; 8)	0.875	1.00	9
<i>BRCA1</i>	Homologous recombination	Cell-based (HEK293T)	Complementation/perturbation	35 (23; 5)¶	1.00	1.00	73
<i>BRCA2</i>	Homologous recombination	Cell-based (V-C8 cells)	Complementation/perturbation	64 (18; 13)	1.00 (0.75 to 1.00)	1.00 (0.82 to 1.00)	30
<i>BRCA2</i>	Homologous recombination	Cell-based (V-C8 cells)	Complementation/perturbation	139 (12; 13)	1.00 (0.75 to 1.00)	1.00 (0.69 to 1.00)	74
<i>BRCA2</i>	Homologous recombination	Cell-based (mouse ES cells)	Complementation/perturbation	43 (20; 15)	1.00 (0.78 to 1.00)	1.00 (0.83 to 1.00)	75

Only assays in which >30 variants were tested are listed.

*Known pathogenic and non-pathogenic variants used for estimating sensitivity and specificity are those classified using the multifactorial model as IARC classes 1, 2, 4 or 5,¹⁸ unless otherwise indicated.

†As originally published, unless otherwise stated.

‡Used missense variants classified by multifactorial model as IARC classes 1, 2, 4 or 5¹⁸ plus the recently classified G1770V variants as pathogenic.

§Used ClinVar as a source of known pathogenic and non-pathogenic variants.

¶Used as non-pathogenic variants alignGVGD grade of C0 and IARC class 1, and as pathogenic variants alignGVGD grade of C35–C65 and IARC class 5. IARC, International Agency for Research on Cancer.

may be difficult to achieve for genes for which there are very few known pathogenic and non-pathogenic variants to use in a validation set, which will be reflected as wider 95% CIs.

Controls are critical for validation of assays, assessment of dynamic range, and to determine metrics of performance such as sensitivity and specificity. Some thought should be given to decide on genomic DNA/cDNA/protein sequence that corresponds to the coding sequence in the most commonly found haplotype in non-affected individuals to be used as a reference (wild-type). Note that differences in frequency of common alleles may exist across different populations. This reference cDNA or

genomic sequence must be included in every experiment. Variants are scored as having functional impact or not depending on how much they differ from the reference.

It is recommended that within each run of the assay, in addition to the reference sequence, at least one known missense pathogenic and one missense non-pathogenic variant is included. If possible, known missense variants for each protein domain are recommended. To account for the range of variation of non-pathogenic variants, additional known non-pathogenic variants should also be included. Addition of hypomorphic (attenuated) variants with established intermediate risk may help calibration

of assay results. Concerning VUS in genes for which there are no known missense non-pathogenic and pathogenic variants, alternative approaches, such as the use of missense variants with greater than 1.0% allele frequency and truncating variants as benign and pathogenic controls, respectively, might provide a yardstick. Results from assays using only truncating variants as pathogenic control should be interpreted with caution as truncating variants may not produce detectable protein, with implications to measuring baseline activity.

LESSONS FROM *BRCA1* AND *BRCA2* FUNCTIONAL ASSESSMENT

BRCA1 and *BRCA2* have 1863 and 3418 codons, respectively. If we consider all possible single nucleotide changes in these codons, 11 015 and 20 169 unique missense variants are generated (some changes will result in the same amino acid changes), respectively. Because many have never or only sparsely been observed, we expect that most are rare (<0.01%) such that data from functional assays will be required to assess their likelihood of pathogenicity. In order to maximise the chances of identifying pathogenic variants, investigators have focused on functional domains and motifs in which it seems more likely that variants affect protein function. Thus far, most assays have focused on variants at the RING and BRCT domains of *BRCA1* and at the DSS1 and DNA interaction domain of *BRCA2*.^{9 10 36 37} Several functional assays have been described for *BRCA1* and *BRCA2*, but few have tested large (>30) sets of variants (table 2).

For specificity and sensitivity calculations, variants are classified according to a binary classification based on the functional data: functional impact versus no functional impact. Variants with intermediate scores are ignored. This classification is then compared with a binary classification of a reference panel which combines the non-actionable IARC classes 1 and 2 (benign and likely benign) or actionable classes 4 and 5 (likely pathogenic and pathogenic). This simplification allows for the estimation of the assay's ability to correctly identify actionable and non-actionable variants. Most published functional assays have reported high sensitivity and specificity, often close to 100% (table 2). However, these numbers partially reflect the relatively low numbers of known variants used to assess specificity and sensitivity. To obtain a better sense of an assay's performance, it is critical to record and report the lower bounds of the 95% CI.

Several assays have been developed using yeast (*Saccharomyces cerevisiae*), which provides a cost-effective and practical platform to evaluate missense variants. However, caution is warranted when interpreting results from model organisms that are cultured at temperatures lower than 37°C. Some pathogenic variants are relatively stable at lower temperatures (30°C vs 37°C) and may score as false negatives,^{53–55} reflected in the assay's slightly lower sensitivity (table 2).

It is important to note that contradictory results, for example, a variant scoring pathogenic in a functional assay while being classified by clinical and family data as non-pathogenic, provide opportunities for discovery. *BRCA1* variant p.V1736A scored as pathogenic in several functional assays and by in silico prediction tools⁵⁶ despite being classified as non-pathogenic due to a co-occurrence with the known pathogenic p.D821Ifs*25 variant in the same patient. However, on further examination it was found that the carrier presented several features (eg, developmental delay, microcephaly, short stature, very early onset ovarian cancer) pointing at hypomorphic *BRCA1* activity.²⁵ Detailed genetic and functional investigation of the p.V1736A variant led to the discovery of the first documented carrier of biallelic

pathogenic variants in *BRCA1*. This analysis established the existence of variants with intermediate effects and highlighted the power of functional assessment.²⁵

Although there are several missense variants that have displayed intermediate effects in vitro or in mouse models, only three variants, in addition to p.V1736A, have been established as hypomorphic in humans.^{25 27} *BRCA1* variant p.R1699Q (c.5096G>A; OR=4.29) and *BRCA2* variants p.Y3035S (c.9104A>C; OR=2.52) and p.G2508S (c.7522G>A; OR=2.68), have been shown to confer moderate increased risks. There are currently no consensus guidelines about their clinical management but the ENIGMA consortium has recommended breast surveillance for female carriers of p.R1699Q based on mammogram annually from age 40 and bilateral salpingo-oophorectomy should be considered based on family history.³² Care should be exercised in the choice of statistical treatment of the data generated in functional assays. Results from assays are usually normalised using the mean of the activity of the wild-type or reference sequence. Normalisation allows for comparisons across multiple experiments and, in some cases across multiple assays since variant activity is being represented as a percentage of the wild-type activity. Batch effects may be problematic (variance of the wild-type activity across multiple batches should also be taken into account) and some statistical models have taken that in consideration.⁵⁷

A more difficult task is the decision of a specific threshold of activity to separate pathogenic from non-pathogenic variants. Several approaches use arbitrary thresholds (eg, 20% or 50% of wild-type activity; number of SD from the wild-type reference; highest activity of a pathogenic variant and lowest activity of a non-pathogenic variant) or linear regression. Recent approaches have moved towards providing a probabilistic interpretation (ie, likelihood of pathogenicity of the variant given the functional data).^{57 58}

Probabilistic approaches also provide a path for integration of functional data with other data sources used to classify variants. By generating likelihood ratios (LRs) from the raw or processed functional read-outs (eg, viability counts, luciferase activity, Green Fluorescent Protein intensity), these approaches allow for the incorporation of functional assays as a data source into traditional multifactorial models that have so far not integrated functional data.⁵⁷

Integration of functional data can also be achieved using the ACMG/AMP classification model. According to the ACMG/AMP, 'well-established assays' can be used to obtain a PS3 or BS3 (strong evidence) criterion but there are no specific guidelines, which are likely to be established by expert panels for each gene. For example, concordant results from at least three independent validated assays would be needed for PS3 or BS3 (strong) criteria, while concordant results in two independent validated assays would generate a PM1 (moderate) criterion.

FUNCTIONAL ASSAYS BASED ON SENSITIVITY TO THERAPEUTIC COMPOUNDS

Insight in the importance of *BRCA1* and *BRCA2* for HR led to the development of carrier-specific treatment modalities for breast and ovarian cancers,^{59–61} in particular with the use of PARP inhibitors (PARPi), which are synthetic lethal with *BRCA1* or *BRCA2* deficiency. The therapeutic window for these types of treatment is greatly increased by the fact that *BRCA1* and *BRCA2* mutation carriers are generally heterozygous for the pathogenic allele, while tumour cells frequently undergo loss of the wild-type allele. Thus, while the tumour cells are not viable

in the presence of PARPi, non-tumour cells survive, making the therapy highly effective yet well tolerated.

Because inactivation of the gene product is required both for disruption of HR and sensitivity to PARPi or platinum compounds, functional assays based on sensitivity to these compounds can be used as an indirect read-out for HR to classify germline variants according to their pathogenicity.⁶²

In addition to the classification of germline variants according to associated cancer risk, there is an emerging need to classify germline and somatic variants according to their response to PARPi or DNA damaging compounds to predict drug response. PARPi have been approved in the USA and Europe for treatment of advanced and metastatic breast and ovarian cancers. It is unclear the extent to which sensitivity to PARPi or platinum compounds measured in a functional assay predicts in vivo tumour sensitivity. Pathogenic variants conferring high cancer risk with hypomorphic activity towards PARPi response are known to exist. For example, mouse tumour cells carrying the pathogenic *Brca1* p.C61G variant showed a poor response to platinum compounds and to PARPi, and resistance rapidly emerged.⁶³ Importantly, determining whether a *BRCA1* or *BRCA2* variant found in tumour tissue confers sensitivity to a given drug may need further clinical information to calibrate the functional assays for this purpose.

Many other genes implicated in HBOC (such as *ATM*, *CHEK2*, *PALB2* and *TP53*) are involved in the DNA damage response, suggesting that associated tumours may also have a targetable DNA repair defect. Importantly, recent results from the NOVA, Study 19 and ARIEL3 studies have raised the possibility of a significant benefit of PARPi in ovarian cancer irrespective of *BRCA* mutation status.^{64–66}

FUTURE CHALLENGES AND OPPORTUNITIES

As we move forward, functional assays should be able to face a significant increase in the number of genes and variants to be analysed for clinical use. The number of alleles for genes predisposing to breast and ovarian cancer is expected to be very large. *BRCA1* and *BRCA2* have 7898 and 10422 unique alleles documented (table 1; BRCA Exchange: <http://brcaexchange.org/>) and other HBOC genes are expected to significantly add to the number of predisposing alleles. In addition, recent systematic germline sequencing efforts of breast and ovarian cancer cases are expected to reveal additional genes and variants associated with risk.

To face the rapid increase in variants, high-throughput functional assays that can generate and analyse large numbers of variants have been developed for *BRCA1*.^{9 10} Initially, these consisted of the large-scale generation and analysis of ectopically expressed cDNA constructs. Recent development of CRISPR-assisted gene targeting has now also allowed saturated mutagenesis at endogenous loci, exemplified by high-throughput mutagenesis of the *BRCA1* RING and BRCT domains in haploid cells.^{9 10} Likewise, assays based on human primary cells from carriers may also be used to capture more subtle functional defects that depend on the carrier's genetic background, but may be less amenable to high-throughput approaches.⁴² Such high-throughput approaches can provide catalogues of potentially pathogenic variants that can aid in the interpretation of newly observed VUS. CRISPR-assisted gene targeting is also expected to accelerated the pace at which mouse models can be generated.^{67 68}

The path to clinical implementation will necessarily involve the inclusion of clinically calibrated functional data into comprehensive multifactorial statistical models. Functional assays are

specialised tests that most diagnostic testing laboratories are currently not able to provide. In the USA, results from such tests cannot be directly used for clinical decisions unless they are conducted under Clinical Laboratory Improvement Amendments guidelines. However, if data have been validated (ie, evaluated for a comprehensive series of performance metrics such as sensitivity, specificity, accuracy, precision, repeatability and robustness), functional assays reported from research labs can be used as evidence for clinical annotation.⁷ It is important to consider that mistakes may occur in the processing of a sample in the absence of standard operating procedures. Those mistakes include clerical mistakes in reports (eg, reporting pathogenic when it should be non-pathogenic), sample swapping, errors in measurements due to lack of equipment calibration or improper staff training. Thus, evidence from functional studies performed in a research setting must be carefully verified to determine data quality, reliability and the degree of confidence in the results. In the context of risk assessment, we caution against the use of functional data as the sole source of information for clinical recommendations. Despite these challenges, the value of functional assessment of variants to improve cancer care, based on international and multidisciplinary collaborations, is expected to be high and therefore we envisage the clinical implementation of functional assays for classification of VUS to proceed at an accelerated pace.

Author affiliations

¹Cancer Epidemiology, H. Lee Moffitt Cancer Center and Research Institute, Tampa, Florida, USA

²Division of Molecular Pathology, Oncode Institute, The Netherlands Cancer Institute, Amsterdam, The Netherlands

³Cancer Sciences, University of Southampton Faculty of Medicine, Southampton, UK

⁴Hub-DBC, Institut Pasteur, USR 3756 CNRS, Paris, France

⁵CHU de Québec-Université Laval, Oncology Division, Laval University Cancer Research Center, Quebec City, Quebec, Canada

⁶National Cancer Institute at Frederick, Frederick, Maryland, USA

⁷Beth Israel Deaconess Medical Center, Boston, Massachusetts, USA

⁸Ulm University, Ulm, Baden-Württemberg, Germany

⁹Mayo Clinic, Rochester, Minnesota, USA

¹⁰Human Genetics, Leiden University Medical Center, Leiden, The Netherlands

Twitter Alvaro N Monteiro @AccidentalGenet

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ORCID iDs

Alvaro N Monteiro <http://orcid.org/0000-0002-8448-4801>

Peter Bouwman <http://orcid.org/0000-0003-0920-8433>

Arne N Kousholt <http://orcid.org/0000-0002-3972-1740>

Diana M Eccles <http://orcid.org/0000-0002-9935-3169>

Gael A Millot <http://orcid.org/0000-0002-0591-3509>

Jean-Yves Masson <http://orcid.org/0000-0002-4403-7169>

Marjanka K Schmidt <http://orcid.org/0000-0002-2228-429X>

Shyam K Sharan <http://orcid.org/0000-0002-9333-870X>
 Ralph Scully <http://orcid.org/0000-0002-5064-0175>
 Lisa Wiesmüller <http://orcid.org/0000-0002-2397-5041>
 Fergus Couch <http://orcid.org/0000-0001-9417-9985>
 Maaike P G Vreeswijk <http://orcid.org/0000-0003-4068-9271>

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