

Uncovering Mutations in Tumor DNA

& providing clinically actionable data for
choice of the most optimal treatment

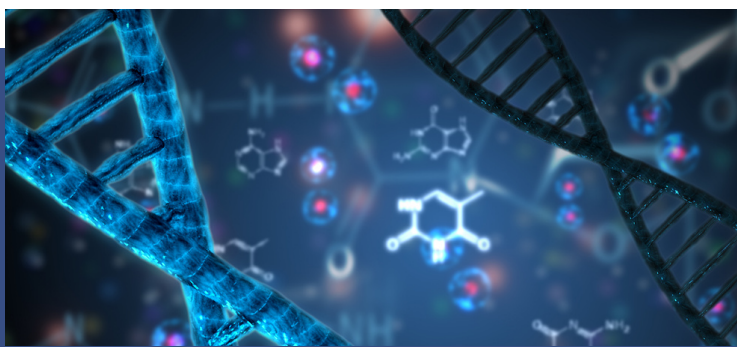


BRCA Extended Panel

BRCA extended panel contains 15 genes associated with hereditary Breast and Ovarian cancer, pancreatic and prostate cancer

PANEL PROVIDES

- Profiling of 15 genes associated with hereditary Breast and Ovarian cancer, pancreatic and prostate cancer
- Guidance for PARPi therapy
- Detect heritable, disease-related gene mutations which indicate predisposition to risk of certain cancers.
- Identify at-risk family members for early detection and improved outcomes



WHO CAN BENEFIT

- Strong family history of breast, uterine, and/or other related cancers
- Family history pointing both hereditary breast and ovarian cancer and Lynch syndrome
- Diagnosis of breast cancer with negative BRCA 1/2 test and Positive family history
- Diagnosis of Breast, Ovarian, Pancreatic, male breast, or metastatic prostate cancer
- Or as evaluated by your oncologist

GENES COVERED IN THIS PANEL

ATM BARD1 BRCA1 BRCA2 BRIP1 CDK12 CHEK2
FANCD2 MRE11 NBN PALB2 PPP2R2A RAD51B
RAD54L TP53

WHAT ARE THE BENEFITS OF GENETIC TESTS?

- Genetic testing can help in making better medical and lifestyle choices for some women while relieving the anxiety of not understanding their genetic history. With both drugs and prophylactic surgery, you may also decide about prevention.
- Knowledge of genetic mutation in the patient can help in early or timely detection of mutations in close family members.

OUR OTHER PANELS



Homologous Recombination Repair Gene Panel (>15 genes)



Comprehensive Cancer Panel



Inherited Cancer Panel (>140 genes)



Somatic panel with >400 genes



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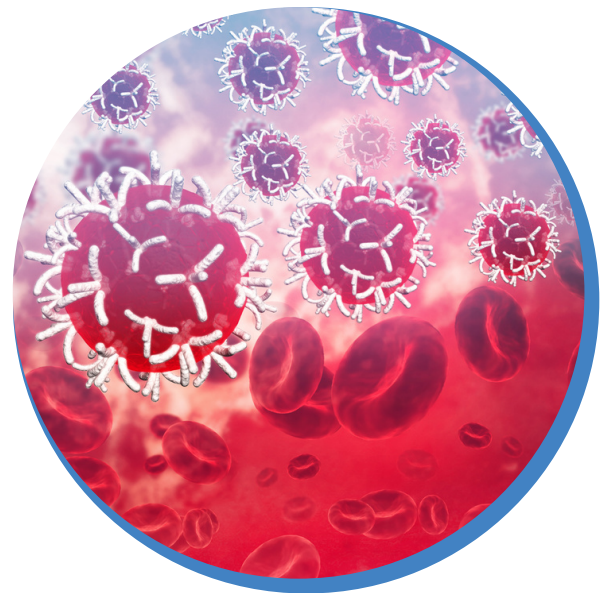
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SPECIMEN REQUIREMENTS

Blood in EDTA tubes

WHY BIOSERVE - REPROCELL INDIA?

- High precision and accuracy of the BRCA1 and BRCA2 genes are guarded with the sensitivity and specificity to detect SNVs and InDels.
- We provide a free pre-test and post-test genetic therapy session with trained experts who provide impartial insights into the risk, incidence, and recurrence of genetic disorders in individuals/families.
- The test has been validated in house and offers coverage range of 500-1000X.



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