

# BRCAcare®

DNA-based Next Generation Sequencing (NGS) technologies provide high sensitivity and accuracy for the detection of BRCA and other gene mutations.

## Tests Available:

- **BRCA1/2: Comprehensive BRCA Analysis**
- **BRCA1/2: Ashkenazi Jewish 3-site Mutation Analysis**
- **BRCA1 & BRCA 2: Specific Site Analysis**
- **Testing for Hereditary Breast & Ovarian Cancer Syndrome (HBOC)**
  - **Breast Cancer High Risk Extended Panel Plus: 14 genes** (BRCA1, BRCA2, CDH1, PTEN, TP53, STK11, ATM, CHEK2, PALB2, BARD1, BRIP1, MUTYH, RAD51C, RAD51D)
  - **Breast Cancer High Risk Extended Panel (No BRCA1, BRCA2): 12 genes** (CDH1, PTEN, TP53, STK11, ATM, CHEK2, PALB2, BARD1, BRIP1, MUTYH, RAD51C, RAD51D)
  - **BRCA1/2: Ashkenazi Jewish 3-site Mutation Analysis** (Reflex to Breast Cancer High Risk Extended Panel Plus if negative)
- **Testing for Lynch Syndrome genes:**
  - **Comprehensive Hereditary Breast and Gynecologic Cancer Panel: 19 genes** analyzed by Gene Sequencing and/or Deletion/Duplication Analysis (BRCA1, BRCA2, ATM, BARD1, BRIP1, CDH1, CHEK2, MUTYH, PALB2, PTEN, RAD51C, RAD51D, STK11, TP53, EPCAM, MLH1, MSH2, MSH6, PMS2).

**Now Available**

*Targets genes of **elevated risk** for the development of breast, ovarian, and endometrial uterine cancer.*

## Advantages:

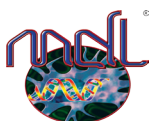
- Non-invasive
- Ease of collection
- Rapid turnaround time
- High sensitivity and specificity
- Genetic counseling can be arranged
- Patient friendly pricing

UNLOCK YOUR

 **BRCA**  
CODE®



Upd: 4/2017



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