

OUTLINED AREAS MUST BE COMPLETED

| | | | |
|--|---------------------------|-------|---------------------|
| P A T I E N T | PATIENT IDENTIFIER | | |
| | NAME, LAST (Please Print) | | FIRST M.I. |
| | BIRTHDATE | M/F | DATE/TIME COLLECTED |
| | STREET | | PHONE # |
| | CITY | STATE | ZIP |

| | | | |
|--|--|--|--|
| B I L L I N G | INSURANCE CARRIER NAME | | ADDRESS |
| | INSURED NAME | INSURED ID# | PT. RELATIONSHIP TO INSURED: <input type="checkbox"/> Self <input type="checkbox"/> Spouse <input type="checkbox"/> Dependent |
| | GROUP # or NAME | | <input type="checkbox"/> INSURANCE CHANGE |
| | <input type="checkbox"/> MEDICARE # | <input type="checkbox"/> MEDICAID # | <input type="checkbox"/> SELF-PAY |

RACE/ETHNICITY NATIVE AMERICAN ASIAN AFRICAN-AMERICAN CAUCASIAN
 ASHKENAZI JEWISH PACIFIC ISLANDER HISPANIC OTHER _____

AFFIX TO SPECIMEN CONTAINER

I attest that this patient has been informed about and has given consent for the test(s) I have ordered below under applicable law.
PHYSICIAN SIGNATURE: (required)

X

REPORT COPY TO:

| | |
|-----------|---------|
| DIAGNOSIS | DX CODE |
| DX CODE | DX CODE |
| DX CODE | DX CODE |

ORDER COMMENTS

| | | |
|-------------------------|---|--|
| INFORMED CONSENT | Patient has read the appropriate consent form on the following pages and has been informed of the required information regarding testing for inherited genetic disorders or predisposition. | |
| | Patient Signature: _____ | Healthcare Provider Signature: _____ Date: _____ |

| | |
|----------------|---|
| ACCOUNT | S4762 Northwell Health Genomics Alliance T: (516) 719-1100 F: (516) 719-1220 |
|----------------|---|

OncoGeneDx Tests

Breast/Gynecologic Cancers

| | |
|--------------|---|
| BRCA 1/2 | BRCA1/BRCA2 Ashkenazi Founder Panel |
| BRCA 1/2 Seq | BRCA1 BRCA2 Sequencing and DEL/Dup Analysis |
| BC High/Mod | Breast Cancer Management Panel |
| B/O Cancer | Breast/GYN Cancer Panel |

Colorectal Cancer

| | |
|------------|-------------------------------|
| Colorectal | Colorectal Cancer Panel |
| LYNCH | GI/Colorectal High Risk Panel |

Multiple Cancers

| | |
|-------------|-----------------------------------|
| CompCancer | Comprehensive Common Cancer Panel |
| Hi/Mod Risk | Common Cancer Management Panel |

Tumor Specific Panels

| | |
|------------|-------------------------|
| MaligMelan | Melanoma Panel |
| PancCancer | Pancreatic Cancer Panel |
| PedTumor | Pediatric Tumor Panel |
| PGL/PCC | PGL/PCC Panel |
| RenCancer | Renal Cancer Panel |

Patient Clinical Information

Personal History of Cancer: _____

Breast Cancer(s) Age at Dx: _____ ER _____ PR _____ HER2 _____ triple negative
 Bilateral Two primaries Invasive Ductal Invasive Lobular DCIS LCIS Other: _____
 Colorectal Cancer(s) Age at Dx: _____ Pathology: _____
 Location: Right Left Transverse Rectum

Patient has had a blood transfusion Yes No Date of last transfusion ____/____/____ (2-4 weeks of wait time is required for some testing)
 Specimens are not accepted for patients who have had allogeneic bone marrow transplants.

Treatment-Related RUSH: _____ (If known, please provide date)

Family History of Cancer(s) Tumor(s) or Relevant History

No Known Family History Pedigree Attached Adopted

Please include clinical details, such as bilateral, pathology (including triple negative breast cancer), premenopausal breast cancer, and Gleason score for prostate cancer. For pancreatitis history, please indicate acute or chronic, if available.

| Relationship | Maternal | Paternal | Cancer/Tumor Site or Relevant History | Age at Dx |
|--------------|--------------------------|--------------------------|---------------------------------------|-----------|
| _____ | <input type="checkbox"/> | <input type="checkbox"/> | _____ | _____ |
| _____ | <input type="checkbox"/> | <input type="checkbox"/> | _____ | _____ |
| _____ | <input type="checkbox"/> | <input type="checkbox"/> | _____ | _____ |
| _____ | <input type="checkbox"/> | <input type="checkbox"/> | _____ | _____ |

| OncoGeneDx Panel Components | |
|--|--|
| Breast Cancer Management Panel (9 genes) | ATM, BRCA1, BRCA2, CDH1, CHEK2, NBN, PALB2, PTEN, TP53 |
| Breast/Gyn Cancer Panel (23 genes) | ATM, BARD1, BRCA1, BRCA2, BRIP1, CDH1, CHEK2, EPCAM*, FANCC, MLH1, MSH2, MSH6, MUTYH, NBN, NF1, PALB2, PMS2, POLD1, PTEN, RAD51C, RAD51D, RECQL, TP53 |
| Lynch/Colorectal High Risk Panel (7 genes) | APC, EPCAM*, MLH1, MSH2, MSH6, MUTYH, PMS2 |
| Colorectal Cancer Panel (20 genes) | APC, ATM, AXIN2, BMPR1A, CDH1, CHEK2, EPCAM*, MLH1, MSH2, MSH6, MUTYH, NTHL1, PMS2, POLD1, POLE, PTEN, SCG5/GREM1*, SMAD4, STK11, TP53 |
| Common Cancer Management Panel (37 genes) | APC, ATM, AXIN2, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDKN2A, CHEK2, EPCAM*, FH, FLCN, MLH1, MSH2, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, PTEN, RAD51C, RAD51D, SCG5/GREM1*, SDHB, SDHC, SDHD, SMAD4, STK11, TP53, TSC1, TSC2, VHL |
| Comprehensive Common Cancer Panel (46 genes) | APC, ATM, AXIN2, BAP1, BARD1, BMPR1A, BRCA1, BRCA2, BRIP1, CDH1, CDK4, CDKN2A, CHEK2, EPCAM*, FANCC, FH, FLCN, HOXB13, MET, MITF*, MLH1, MSH2, MSH6, MUTYH, NBN, NF1, NTHL1, PALB2, PMS2, POLD1, POLE, POT1, PTEN, RAD51C, RAD51D, RECQL, SCG5/GREM1*, SDHB, SDHC, SDHD, SMAD4, STK11, TP53, TSC1, TSC2, VHL |
| Melanoma Panel (9 genes) | BAP1, BRCA2, CDK4, CDKN2A, MITF*, POT1, PTEN, RB1, TP53 |
| Pancreatic Cancer Panel (15 genes) | APC, ATM, BRCA1, BRCA2, CDK4, CDKN2A, EPCAM*, MLH1, MSH2, MSH6, PALB2, PMS2, STK11, TP53, VHL |
| Pediatric Tumor Panel (27 genes) | ALK, APC, CDC73, DICER1, EPCAM*, MEN1, MLH1, MSH2, MSH6, NF1, NF2, PHOX2B*, PMS2, PRKAR1A, PTCH1, PTEN, RB1, RET*, SMARCA4, SMARCB1, STK11, SUFU, TP53, TSC1, TSC2, VHL, WT1 |
| PGL/PCC Panel (12 genes) | FH, MAX, MEN1, NF1, RET*, SDHA*, SDHAF2, SDHB, SDHC, SDHD, TMEM127, VHL |
| Renal Cancer Panel (18 genes) | BAP1, EPCAM*, FH, FLCN, MET, MITF*, MLH1, MSH2, MSH6, PMS2, PTEN, SDHB, SDHC, SDHD, TP53, TSC1, TSC2, VHL |