

**BASIC3 GERMLINE SEQUENCING FORM**

**PATIENT INFORMATION** **REPORTING INFORMATION**

NAME: \_\_\_\_\_  
LAST NAME FIRST NAME MI

DATE OF BIRTH: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
MM DD YY

GENDER (Please select one):  FEMALE  
 MALE  
 UNKNOWN

MEDICAL RECORD #: \_\_\_\_\_

ETHNIC BACKGROUND (Select all that apply):

AFRICAN AMERICAN  
 ASIAN  
 ASHKENAZIC JEWISH  
 EUROPEAN CAUCASIAN  
 HISPANIC  
 NATIVE AMERICAN INDIAN  
 OTHER JEWISH  
 OTHER (Please specify): \_\_\_\_\_

ACCESSION #: \_\_\_\_\_

HOSPITAL #: \_\_\_\_\_

CONSENT DATE: \_\_\_\_\_

STUDY #: \_\_\_\_\_

\*BCM-MEDICAL GENETICS LABORATORIES HAS A FAX ONLY POLICY FOR REPORTING

PHYSICIAN: \_\_\_\_\_

INSTITUTION: \_\_\_\_\_

PHONE: \_\_\_\_\_ \*FAX: \_\_\_\_\_

**SAMPLE INFORMATION**

PERIPHERAL BLOOD (AS DNA SOURCE)

DATE OF COLLECTION: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
MM DD YY

TIME OF COLLECTION: \_\_\_\_\_

**TESTS REQUESTED**

Tumor Sample Available for Sequencing:  YES -- **9601 & 1590**  
 NO -- **1591**

**ADDITIONAL PROFESSIONAL REPORT RECIPIENTS**

NAME: \_\_\_\_\_

PHONE: \_\_\_\_\_ \*FAX: \_\_\_\_\_

NAME: \_\_\_\_\_

PHONE: \_\_\_\_\_ \*FAX: \_\_\_\_\_

INSTITUTION NAME: \_\_\_\_\_

INSTITUTION CODE: \_\_\_\_\_

**FAMILY HISTORY**

Family History of Cancer:  YES  NO

If yes, please provide detailed information on affected family members and/or a pedigree.

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LAST NAME FIRST NAME MI MM DD YY  MALE  
 UNKNOWN

Please provide the following clinical information regarding the patient to be tested. If answering "yes," please provide additional description as appropriate (e.g., percentiles for growth parameters, type of limb abnormality, etc.). Please also submit a clinic note and pedigree if available. This information is needed to facilitate interpretation of whole exome sequencing results. If the laboratory requires additional information, please indicate the health care provider to be contacted:

NAME: \_\_\_\_\_ PHONE/PAGER #: \_\_\_\_\_

		PATIENT DESCRIPTION	NO	UNKNOWN
Prematurity	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Intrauterine growth restriction	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Delayed motor milestones	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Delayed speech	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Developmental regression	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Autism/Autistic spectrum	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Intellectual disability	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Hearing loss	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Hypotonia	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Hypertonia/Spasticity	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Seizure disorder	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Ataxia	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Abnormal movements	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Dysmorphic features	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Short stature	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Tall habitus	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Microcephaly	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Macrocephaly	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Hyperextensibility	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Joint contractures	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Obesity/Overgrowth	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Failure to thrive	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Structural brain abnormalities	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Eye anomalies	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Vision loss	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>

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LAST NAME FIRST NAME MI MM DD YY  MALE  
 UNKNOWN

		PATIENT DESCRIPTION	NO	UNKNOWN
Congenital heart disease	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Kidney abnormalities	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Skeletal abnormalities	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Scoliosis	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Limb malformation	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Skin anomalies	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Genital anomalies	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Organomegaly	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Hemihypertrophy	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Cancer/tumor formation	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>
Family history of non-cancer disorder	<input type="checkbox"/>		<input type="checkbox"/>	<input type="checkbox"/>

Please read the below statements carefully and check the appropriate box and initial. Please note that if neither box is checked the lab will default to the YES/ reporting option.

Initial

Carrier Status for Autosomal Recessive Conditions

- \_\_\_\_\_  YES, please report carrier status. By checking this box, I choose to receive information regarding carrier status.  
 \_\_\_\_\_  NO, please do NOT report carrier status. By checking this box, I choose to NOT receive information regarding carrier status.