

PATIENT HISTORY FOR BIOCHEMICAL GENETIC TESTING

PATIENT INFORMATION

Clinic Number: _____

Patient's Last Name: _____ First Name: _____ MI: _____

Birth Date: _____ Gender: Male Female

Referring Physician (last, first): _____

Referring Physician Phone: (_____) _____ Referring Physician Fax*: (_____) _____

Genetics Counselor (last, first): _____

Genetics Counselor Phone: (_____) _____ Genetics Counselor Fax*: (_____) _____

**Fax number given must be from a fax machine that complies with applicable HIPAA regulations.*

COMMENTS OR SPECIAL INSTRUCTIONS

REFERRING DIAGNOSIS

PATIENT'S ETHNICITY (check all that apply)

- | | | | |
|---|------------------------------------|---|--|
| <input type="checkbox"/> African American | <input type="checkbox"/> Asian | <input type="checkbox"/> Hispanic | <input type="checkbox"/> Native American |
| <input type="checkbox"/> Ashkenazi Jewish | <input type="checkbox"/> Caucasian | <input type="checkbox"/> Middle Eastern | <input type="checkbox"/> Other _____ |

PATIENT'S SYMPTOMS (check all that apply)

- | | | |
|--|--|---|
| <input type="checkbox"/> Acidosis | <input type="checkbox"/> Failure to thrive | <input type="checkbox"/> Organomegaly |
| <input type="checkbox"/> Cardiomyopathy | <input type="checkbox"/> Hyperammonemia | <input type="checkbox"/> Seizures |
| <input type="checkbox"/> Coarse features | <input type="checkbox"/> Hypoglycemia | <input type="checkbox"/> Skeletal anomalies |
| <input type="checkbox"/> Corneal clouding | <input type="checkbox"/> Macrocephaly | <input type="checkbox"/> Other _____ |
| <input type="checkbox"/> Developmental delay | <input type="checkbox"/> Microcephaly | |

List the patient's medications, including antibiotics, anticonvulsants and enzyme replacement therapy:

List the patient's specific diet or formula:

Are the patient's parents related to one another? No Yes Unknown

If yes, please describe: _____