

A nonprofit enterprise of the University of Utah and its Department of Pathology

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THIS IS NOT A TEST REQUEST FORM.

Please fill out this form and submit it with the test request form or electronic packing list.

PATIENT HISTORY FOR BIOCHEMICAL GENETIC TESTING

| Patient Name: | | | Dat | e of Birth: | s | ex: □ Female □ Male |
|---|------------------------|------------------|----------------------|---------------------|------------------|---------------------------|
| Physician: Practice Specialty: | | | | Physician Phone: | | _ |
| | | | | Physician Fax: | | |
| Genetic Counselor: | | unselor Phone: | | | | |
| Patient's Ethnicity (chec | k all that apply) | | | | | |
| ☐ African-American | □ Asian | ☐ Hispan | ic | □ Native Amer | ican | |
| ☐ Ashkenazi Jewish | ☐ Caucasian | ☐ Middle | Eastern | □ Other: | | |
| Clinical Diagnosis / Reas | son for Referral: | | | | | |
| Newborn Screening Res | ults: | | | | Abnorma | ıl □ Normal □ N/A |
| Describe any abnormal r | esults: | | | | | |
| Does the patient have sy | mptoms? | | | | 🗆 No 🗆 Ye | es (check all that apply) |
| ☐ Acidosis | . ☐ Failure to thrive | | | nomegaly | | |
| ☐ Cardiomyopathy | ☐ Hyperamr | ☐ Hyperammonemia | | ures | | |
| ☐ Coarse features | ☐ Hypoglycemia | | ☐ Skeletal anomalies | | | |
| ☐ Corneal clouding | ☐ Macrocephaly | | ☐ Other symptom(s): | | | |
| ☐ Developmental Delay | ☐ Microcephaly | | | | | |
| List the patient's medica | itions, including an | ntibiotics, anti | convulsants | s, and enzyme repla | acement therapy: | |
| List the patient's specifi | | | | | | |
| Are the patient's parents | | | | | □ No | ☐ Yes ☐ Unknown |
| Comments or special ins | structions: | | | | | |
| | | | | | | |
| Please submit this form ARUP Biochemical Gene | | | | | | |
| For questions, contact the ARUP Biochemical Genetics Laboratory at 800-242-2787, ext. 3922. | | | | | Ма | ster Label |