

THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

PEROXISOMAL DISORDERS PATIENT HISTORY FORM

Patient Name: _____ Date of Birth: _____

Sex Assigned at Birth: ☐ Female ☐ Male ☐ Intersex Gender Identity (optional): ☐ Female ☐ Male ☐ _____

Ordering Provider: _____ Provider's Phone: _____

Practice Specialty: _____ Provider's Fax: _____

Genetic Counselor: _____ Counselor's Phone: _____

Patient's Ethnicity/Ancestry (check all that apply)

☐ African American/Black ☐ Asian ☐ Hispanic ☐ White ☐ Other: _____

List country of origin (if known): _____

Did the patient have an abnormal newborn screen? ☐ No ☐ Yes ☐ Unknown

If yes, please provide details: _____

Suspected Diagnosis: ☐ Zellweger spectrum disorder ☐ Rhizomelic chondrodysplasia punctata (RCDP)

☐ Other: _____

Does the patient have symptoms? ☐ No ☐ Yes (check all that apply) ☐ Unknown

☐ Growth failure/feeding difficulties ☐ Seizures

☐ Hearing loss ☐ Hypotonia

☐ Visual impairment ☐ Brain MRI, abnormal findings: _____

☐ Liver dysfunction _____

☐ Renal insufficiency _____

☐ Adrenal insufficiency ☐ Other symptom(s): _____

☐ Skeletal abnormalities _____

Laboratory findings:

VLCFA levels (plasma) ☐ Normal ☐ Abnormal ☐ Not Performed

C26:0-lysophosphatidylcholine (LPC) (NBS/plasma) ☐ Normal ☐ Abnormal ☐ Not Performed

Plasmalogens (erythrocytes) ☐ Normal ☐ Abnormal ☐ Not Performed

Phytanic/pristanic acid levels (plasma) ☐ Normal ☐ Abnormal ☐ Not Performed

Pipecolic acid levels (plasma/urine) ☐ Normal ☐ Abnormal ☐ Not Performed

Bile acid intermediates (plasma/urine) ☐ Normal ☐ Abnormal ☐ Not Performed

Has the patient had an allogeneic bone marrow or umbilical cord blood transplant? ☐ No ☐ Yes ☐ Unknown

Has the patient undergone previous DNA testing? ☐ No ☐ Yes ☐ Unknown

If yes, describe the test(s) and results: _____

Is there any relevant family history of peroxisomal disorders? ☐ No ☐ Yes ☐ Unknown

If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset: _____

Has DNA testing been performed for the family member(s)? ☐ No ☐ Yes ☐ Unknown

If yes, attach a copy of the relative's DNA laboratory result (REQUIRED for familial variant testing).

For questions, contact an ARUP genetic counselor at 800-242-2787 ext. 2141.

Master Label