

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 CHARGE SYNDROME TESTING

Patient Name _____ Date of Birth _____ Sex F M
 Physician _____ Physician Phone _____
 Practice Specialty _____ Physician Fax _____
 Genetic Counselor _____ Counselor Phone _____

Patient's Ethnicity (check all that apply)

African-American Asian Hispanic Native American
 Ashkenazi Jewish Caucasian Middle Eastern Other: _____

Clinical Diagnosis of CHARGE syndrome? Yes Suspected Unknown

Does the patient have symptoms? No Yes (check all that apply and describe)

Characteristic facial features Cranial nerve dysfunction/anomaly Growth deficiency
 Choanal atresia/stenosis Developmental delay Ocular coloboma
 Cleft lip/palate Genital hypoplasia Tracheoesophageal (TE) fistula
 Congenital heart defect; describe: _____
 Ear abnormalities; describe: _____
 Other symptom(s): _____

Has the patient undergone previous DNA testing for CHARGE syndrome? No Yes Unknown

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

Is there any relevant family history? 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 mutation testing.)

Check the test you intend to order.

- 2012609 CHARGE Syndrome (CHD7) Sequencing:** Detects 90% of CHARGE syndrome.
- 2012717 CHARGE Syndrome (CHD7) Sequencing, Fetal:** Prenatal testing for CHARGE syndrome.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a previously identified familial mutation only. A copy of relative's DNA laboratory result is REQUIRED.

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

Master Label