

THIS IS NOT A TEST REQUEST FORM.
The information below is required to perform molecular genetic testing.
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** ____/____/____ **Gender** F M

Physician _____ **Physician Phone** (____) _____ **Practice Specialty** _____

Genetic Counselor _____ **Counselor's Phone #** (____) _____

Patient's Ethnicity (check all that apply)

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

Does the patient have a clinical diagnosis of CHARGE syndrome? Yes Suspected Unknown

Does the patient have SYMPTOMS of CHARGE syndrome? No Yes (please check all that apply)

- | | |
|---|---|
| <input type="checkbox"/> Ocular coloboma | <input type="checkbox"/> Cranial nerve dysfunction/anomaly |
| <input type="checkbox"/> Choanal atresia/stenosis | <input type="checkbox"/> Ear abnormalities, describe: _____ |
| <input type="checkbox"/> Growth deficiency | <input type="checkbox"/> Congenital heart defect, describe: _____ |
| <input type="checkbox"/> Developmental delay | <input type="checkbox"/> Characteristic facial features |
| <input type="checkbox"/> Genital hypoplasia | <input type="checkbox"/> Tracheoesophageal (TE) fistula |
| <input type="checkbox"/> Cleft lip/palate | <input type="checkbox"/> Other: _____ |

Does the patient have a FAMILY HISTORY of CHARGE syndrome or related findings? No Yes Unknown
If yes, attach a PEDIGREE or specify the relatives' RELATIONSHIP to the patient. List their symptoms & age of onset:

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

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

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

Circle the test below 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