

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

HOLOPROSENCEPHALY PANEL 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): _____

Does the patient/pregnancy have a diagnosis of holoprosencephaly? No Yes (check one of the following)

Alobar Semilobar Lobar Middle interhemispheric variant Microform Other: _____

Other findings:

Microcephaly

Single central incisor

Short stature/failure to thrive

Laterality defect (describe): _____

Craniofacial (describe): _____

Neurological (describe): _____

Cardiac defect (describe): _____

Urogenital anomalies (describe): _____

Endocrine issues (describe): _____

Gastrointestinal anomalies (describe): _____

Ecto-/Polydactyly (describe): _____

Other: _____

Has the patient undergone previous testing for holoprosencephaly? No Yes (check all that apply and describe results)

Chromosome analysis: _____

Genomic microarray: _____

Other genetic test(s): _____

Is there any relevant family history of holoprosencephaly? 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.

