

PATIENT INFORMATION

First name	MI	Last name	Date of birth (MM/DD/YYYY)	Biological sex <input type="radio"/> Male <input type="radio"/> Female	MRN (medical record number)
Email address		Mobile phone	Ancestry: <input type="radio"/> Asian <input type="radio"/> Black/African American <input type="radio"/> White/Caucasian <input type="radio"/> Ashkenazi Jewish <input type="radio"/> Hispanic <input type="radio"/> Native American <input type="radio"/> Pacific Islander <input type="radio"/> French Canadian <input type="radio"/> Sephardic Jewish <input type="radio"/> Mediterranean <input type="radio"/> Other: _____		
Address		City	State/Prov	Zip/Postal code	Country

ORGANIZATION INFORMATION

Organization name	Phone		Fax		
Address		City	State/Prov	Zip/Postal code	Country
Primary clinical contact name (contact for general inquiries)		Phone	Fax	Email address	
Ordering physician name	NPI	Phone	Fax	Email address	
Referring physician name		Phone	Fax	Email address	
Additional clinical or laboratory contact name		Fax	Email address		

INSURANCE BILLING (attach front and back of insurance card)

Attach applicable clinical notes and medical records. We do not accept insurance for patients outside the US. www.invitae.com/billing

Policyholder name	Patient relationship to policyholder <input type="radio"/> Self <input type="radio"/> Spouse <input type="radio"/> Child <input type="radio"/> Other: _____	Indicate ICD-10 code in Reason for testing	
Primary insurance company name	Primary member ID #	Primary insurance phone	Prior-authorization #
Secondary insurance company name	Secondary member ID #	Secondary insurance phone	Prior-authorization #

PATIENT PAY BILLING

Invitae will send an electronic invoice to the patient email address listed above.

INSTITUTIONAL BILLING

Invitae will send an invoice to the organization listed above.

Collection date: ____/____/____ # tubes: ____ Specimen ID#: _____

PRENATAL DIAGNOSIS
MICROARRAY ANALYSIS

- Invitae Prenatal Chromosomal Microarray Analysis (with follow-up FISH when required)
 Invitae Prenatal Targeted Chromosomal Microarray Analysis (with follow-up FISH when required)
 If microarray is normal, reflex to Invitae Prenatal Karyotype (Chromosome Analysis)

KARYOTYPE

- Invitae Prenatal Karyotype (Chromosome Analysis)
 If karyotype is normal, reflex to Invitae Prenatal Chromosomal Microarray Analysis (with follow-up FISH when required)
 If karyotype is normal, reflex to Invitae Prenatal Targeted Chromosomal Microarray Analysis (with follow-up FISH when required)

ADDITIONAL TESTING (must order microarray or karyotype first)

- Invitae Prenatal Interphase FISH (for 13, 18, 21, X, Y)
 Amniotic fluid AFP with reflex to AChE (if over 24 weeks gestational age only AChE will be performed)
 Culture cells for additional testing (provide details in special instructions below)

Specimen type:

- Chorionic villi Amniotic fluid DNA source: _____
 Cultured CVS Cultured amniocytes Other: _____

Reason for testing (select all that apply):

- O09.519 Advanced maternal age (1st pregnancy)
 O09.529 Advanced maternal age (subsequent pregnancy)
 O28.5 Abnormal chromosomal and genetic finding on antenatal screening of mother
 O28.1 Abnormal maternal serum screening suggestive of a neural tube defect
 O28.8 Other abnormal findings on antenatal screening of mother
 O28.3 Abnormal fetal ultrasound; Specify abnormality: _____
 Other ICD-10: _____ *Indicate abnormalities on Page 2 Phenotypic Checklist*

Pregnancy history:

Gestational age Wks: ____ Days: ____ Determined by LMP U/S
 Gravida: ____ Para: ____ SAB: ____ TAB: ____
 Is the pregnancy currently ongoing? Yes No, SAB/IUFD No, TAB
 How many fetuses? 1 2 3
 Fetal sex: Female Male Unknown
 Fetal karyotype: 46,XX 46,XY Not performed Pending Abnormal*
 NIPS results: Not performed Normal Abnormal* **enclose copy of report*
 If NIPS was performed at Invitae, provide Order ID: RQ _____

Special instructions/additional requests:
MATERNAL CELL CONTAMINATION

- Maternal Cell Contamination following Prenatal Testing **Specimen type:** Blood
 ICD-10: _____ 4-mL purple EDTA tube

PARENTAL FOLLOW-UP

- Invitae Parental Chromosomal Microarray Analysis **Specimen type:** Blood
 Invitae Parental FISH Analysis 4-mL purple EDTA and
 Invitae Parental Karyotype (Chromosome Analysis) 4-mL green NaHep tubes

Previous testing:

- Previous testing was performed at Invitae/Combimatrix - provide previously tested info:
 Invitae accession#: _____ Relationship to patient: _____
 Full name: _____ DOB: _____
 Previous testing was performed at another lab:
 Report is available - include copy with sample shipment
 Report is not available - call Invitae genetic counseling 800-436-3037 to confirm testing

Reason for testing (select all that apply):

- Evaluation of genetic disease carrier status for procreative management Z31.430 Female Z31.440 Male
 Z82.79 Family history of a chromosome abnormality Other ICD-10: _____

RECURRENT PREGNANCY LOSS OR INFERTILITY

- Invitae Reproductive Karyotype **Specimen type:** Blood
 (Chromosome Analysis) 4-mL purple EDTA and 4-mL green NaHep tubes

Reason for testing (select all that apply):

- N97.9 Female infertility, unspecified N96 Female recurrent pregnancy loss
 N46.9 Male infertility, unspecified Z31.441 Male partner of patient with recurrent pregnancy loss

By signing this form, the medical professional acknowledges that the individual/family member authorized to make decisions for the individual (collectively, the "Patient") has been supplied information regarding and consented to undergo genetic testing, substantially as set forth in Invitae's Informed Consent for Genetic Testing (www.invitae.com/cytogenomic-consent), has been informed that Invitae may notify them of clinical updates related to genetic test results (in consultation with the ordering medical professional as indicated), and for orders originating outside the US, has been informed that the Patient's personal information and specimen will be transferred to and processed in the US. The Patient has further been informed and authorizes Invitae Corporation ("Invitae") and its designees to release information concerning testing to their insurer, if applicable, in order to process and/or appeal claims on behalf of the Patient. If a letter of medical necessity (LMN) has not been provided, the medical professional agrees to allow Invitae to transfer the information from this requisition to a LMN and/or other documentation using the medical professional's name as the signature for insurance billing. For amounts received directly, the Patient has agreed to remit payment to Invitae for testing services rendered. I acknowledge that the Patient has agreed that if the Patient's insurer does not reimburse Invitae in full for any reason, including if the insurer considers the genetic test ordered to be a non-covered service or not medically necessary, then Invitae may bill the Patient directly for the services and the Patient will remit payment directly to Invitae. I acknowledge that I offered pre-test genetic counseling to the Patient, if required by their insurer. In addition to the above, I attest that I am the ordering physician, or I am authorized by the ordering physician to order this test, or I am authorized under applicable law to order this test.

Medical professional signature (required)

Date

The accurate interpretation and reporting of genetic results relies on the provision of pertinent clinical and family history information. To help ensure the most clinically appropriate interpretation of results, complete this clinical information form. Testing cannot be initiated without a reason for referral.

Please check all that apply		
Primary indication for testing <input type="radio"/> Abnormal screening results: <input type="radio"/> Abnormal NIPT result (please include report): _____ <input type="radio"/> Abnormal serum screen: _____ <input type="radio"/> Increased NT/cystic hygroma <input type="radio"/> Advanced maternal age <input type="radio"/> Fetal abnormality	Pulmonary <input type="radio"/> CCAM <input type="radio"/> Diaphragmatic hernia <input type="radio"/> Eventration of diaphragm <input type="radio"/> Pleural effusion <input type="radio"/> Pulmonary sequestration <input type="radio"/> Small thoracic cavity <input type="radio"/> Other: _____	Musculoskeletal <input type="radio"/> Contractures (arthrogryposis) <input type="radio"/> Club Foot <input type="radio"/> Limb anomaly (lower) <input type="radio"/> Limb anomaly (upper) <input type="radio"/> Polydactyly (feet) <input type="radio"/> Polydactyly (hands) <input type="radio"/> Rocker-bottom feet <input type="radio"/> Scoliosis <input type="radio"/> Shortened long bones <input type="radio"/> Skeletal dysplasia <input type="radio"/> Syndactyly (feet) <input type="radio"/> Syndactyly (hands) <input type="radio"/> Vertebral anomaly
Growth <input type="radio"/> Hydrops <input type="radio"/> IUGR <input type="radio"/> Macrosomia <input type="radio"/> Molar pregnancy <input type="radio"/> Oligohydramnios <input type="radio"/> Placental abnormality <input type="radio"/> Polyhydramnios <input type="radio"/> Single umbilical artery/2 vessel cord <input type="radio"/> Other: _____	Cardiac <input type="radio"/> Aortic atresia <input type="radio"/> Atrial septal defect (ASD) <input type="radio"/> AV canal defect <input type="radio"/> Coarctation of aorta <input type="radio"/> Ebstein anomaly <input type="radio"/> Echogenic intracardiac focus <input type="radio"/> Hypoplastic left heart <input type="radio"/> Hypoplastic right heart <input type="radio"/> Pericardial effusion <input type="radio"/> Pulmonary atresia <input type="radio"/> Tetralogy of Fallot <input type="radio"/> Transposition of the great vessels <input type="radio"/> Truncus arteriosus <input type="radio"/> Ventricular septal defect (VSD) <input type="radio"/> Other: _____	Genitourinary <input type="radio"/> Ambiguous genitalia <input type="radio"/> Hydronephrosis <input type="radio"/> Megacystis <input type="radio"/> Pyelectasis <input type="radio"/> Polycystic kidneys <input type="radio"/> Renal agenesis <input type="radio"/> Urethral/ureteral obstruction
Gastrointestinal <input type="radio"/> Anal atresia <input type="radio"/> Absent stomach <input type="radio"/> Duodenal atresia (double bubble sign) <input type="radio"/> Echogenic bowel <input type="radio"/> Gastroschisis <input type="radio"/> Omphalocele <input type="radio"/> Tracheoesophageal fistula <input type="radio"/> Other: _____	Neurological <input type="radio"/> Abnormal gyri/Lissencephaly <input type="radio"/> Agenesis of the corpus callosum <input type="radio"/> Cerebellar hypoplasia <input type="radio"/> Choroid plexus cyst(s) <input type="radio"/> Dandy Walker (posterior fossa abn.) <input type="radio"/> Decreased fetal movement <input type="radio"/> Holoprosencephaly <input type="radio"/> Open neural tube defect (ONTD) <input type="radio"/> Anencephaly <input type="radio"/> Spina bifida <input type="radio"/> Structural brain anomaly <input type="radio"/> Ventriculomegaly/hydrocephaly <input type="radio"/> Other: _____	Family history <input type="radio"/> Patient has had ≥ 2 miscarriages <input type="radio"/> Previous pregnancy with chromosome abnormality: _____ <input type="radio"/> Family history of known chromosome abnormality/genetic condition: _____ <input type="radio"/> Other: _____ _____
Craniofacial <input type="radio"/> Cleft lip and/or cleft palate <input type="radio"/> Hypertelorism <input type="radio"/> Hypotelorism <input type="radio"/> Macrocephaly <input type="radio"/> Microcephaly <input type="radio"/> Micrognathia <input type="radio"/> Pierre Robin sequence <input type="radio"/> Other: _____		Other (describe below) _____ _____ _____

Testing notes

Invitae will attempt to perform all tests ordered. If the sample size is insufficient to do so, Invitae will promptly notify the client and discuss what testing can and cannot be performed. If Invitae is unable to make contact with a client the same day (the day the sample was received) to determine what testing is desired, Invitae will run only the tests that are possible based on the sample received.