

**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](http://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#: \_\_\_\_\_

**CHROMOSOMAL DEVELOPMENTAL DISORDERS**

Invitae Chromosomal Microarray Analysis (with follow-up FISH when required)

Fragile X Syndrome

Invitae Karyotype (Chromosome Analysis)

**Specimen type:**

Blood (4-mL purple EDTA and 4-mL green NaHep tubes)  Buccal swab (Microarray and Fragile X only)

**Reason for testing (select all that apply):**

F84 Autistic disorder

F80.9 Developmental disorder of speech & language, unspecified

F82 Specific developmental disorder of motor function

R62.0 Delayed milestone in childhood

F81.9 Developmental disorder of scholastic skills, unspecified

Intellectual disability:  F70 Mild  F71 Moderate  F72 Severe

Other ICD-10: \_\_\_\_\_

**CLINICAL PHENOTYPE**

Indicate abnormalities on the Chromosomal Developmental Disorders Checklist on page 2

**PARENTAL FOLLOW-UP**

Invitae Parental Chromosomal Microarray Analysis

Invitae Parental FISH Analysis

Invitae Parental Karyotype (Chromosome Analysis)

**Specimen type:** Blood  
4-mL purple EDTA and  
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: \_\_\_\_\_

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](http://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.

<b>Medical professional signature (required)</b>	<b>Date</b>
--	-------------

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		
<b>Perinatal history</b> <ul style="list-style-type: none"> <li><input type="radio"/> Prematurity</li> <li><input type="radio"/> IUGR</li> <li><input type="radio"/> Oligohydramnios</li> <li><input type="radio"/> Polyhydramnios</li> <li><input type="radio"/> Other fetal anomaly: _____</li> </ul>	<b>Cardiac</b> <ul style="list-style-type: none"> <li><input type="radio"/> Atrial septal defect</li> <li><input type="radio"/> AVSD/AV canal defect</li> <li><input type="radio"/> Coarctation of the aorta</li> <li><input type="radio"/> Hypoplastic left heart</li> <li><input type="radio"/> Tetralogy of Fallot</li> <li><input type="radio"/> Ventricular septal defect</li> <li><input type="radio"/> Dilated aortic root</li> <li><input type="radio"/> Cardiomyopathy: _____</li> <li><input type="radio"/> Arrhythmia</li> </ul>	<b>Musculoskeletal</b> <ul style="list-style-type: none"> <li><input type="radio"/> Contractures (arthrogryposis)</li> <li><input type="radio"/> Club foot</li> <li><input type="radio"/> Diaphragmatic hernia</li> <li><input type="radio"/> Limb anomaly</li> <li><input type="radio"/> Polydactyly (hands)</li> <li><input type="radio"/> Polydactyly (feet)</li> <li><input type="radio"/> Syndactyly (hands)</li> <li><input type="radio"/> Syndactyly (feet)</li> <li><input type="radio"/> Vertebral anomaly</li> </ul>
<b>Developmental/behavioral</b> <ul style="list-style-type: none"> <li><input type="radio"/> Fine motor delay</li> <li><input type="radio"/> Gross motor delay</li> <li><input type="radio"/> Speech delay</li> <li><input type="radio"/> Learning disability</li> <li><input type="radio"/> Intellectual disability</li> <li><input type="radio"/> Autistic features</li> <li><input type="radio"/> Autism spectrum disorder</li> <li><input type="radio"/> Oppositional-defiant disorder</li> <li><input type="radio"/> Psychiatric illness:                             <ul style="list-style-type: none"> <li><input type="radio"/> Bipolar disorder</li> <li><input type="radio"/> Anxiety</li> <li><input type="radio"/> Depression</li> <li><input type="radio"/> ADHD</li> <li><input type="radio"/> Other: _____</li> </ul> </li> </ul>	<b>Cutaneous</b> <ul style="list-style-type: none"> <li><input type="radio"/> Hyperpigmentation</li> <li><input type="radio"/> Hypopigmentation</li> <li><input type="radio"/> Café au lait spots</li> <li><input type="radio"/> Eczema</li> </ul>	<b>Neurological</b> <ul style="list-style-type: none"> <li><input type="radio"/> Ataxia</li> <li><input type="radio"/> Cerebral palsy</li> <li><input type="radio"/> Dystonia</li> <li><input type="radio"/> Hypotonia</li> <li><input type="radio"/> Hypertonia</li> <li><input type="radio"/> Developmental regression</li> <li><input type="radio"/> Muscle weakness</li> <li><input type="radio"/> Neural tube defect</li> <li><input type="radio"/> Seizures</li> <li><input type="radio"/> Abnormal brain MRI: _____</li> </ul>
<b>Craniofacial</b> <ul style="list-style-type: none"> <li><input type="radio"/> Cleft lip</li> <li><input type="radio"/> Cleft palate</li> <li><input type="radio"/> Coloboma</li> <li><input type="radio"/> Craniosynostosis</li> <li><input type="radio"/> Dysmorphic facial features</li> <li><input type="radio"/> Macrocephaly</li> <li><input type="radio"/> Microcephaly</li> <li><input type="radio"/> Hearing loss</li> <li><input type="radio"/> Ear malformation</li> <li><input type="radio"/> Cataracts</li> </ul>	<b>Gastrointestinal</b> <ul style="list-style-type: none"> <li><input type="radio"/> Gastroschisis</li> <li><input type="radio"/> Hirschprung disease</li> <li><input type="radio"/> Omphalocele</li> <li><input type="radio"/> Pyloric stenosis</li> <li><input type="radio"/> Tracheoesophageal fistula</li> <li><input type="radio"/> Meconium ileus</li> <li><input type="radio"/> Anal atresia</li> <li><input type="radio"/> Diaphragmatic hernia</li> </ul>	<b>Family history</b> <ul style="list-style-type: none"> <li><input type="radio"/> Patient with <math>\geq 2</math> miscarriages</li> <li><input type="radio"/> Relative with known chromosome abnormality: _____</li> <li><input type="radio"/> Other relatives with clinical history similar to patient (please describe) _____</li> </ul>
	<b>Genitourinary</b> <ul style="list-style-type: none"> <li><input type="radio"/> Ambiguous genitalia</li> <li><input type="radio"/> Hydronephrosis</li> <li><input type="radio"/> Kidney anomaly: _____</li> <li><input type="radio"/> Hypospadias</li> </ul>	<b>Other (describe below)</b> _____ _____ _____ _____
	<b>Genitourinary</b> <ul style="list-style-type: none"> <li><input type="radio"/> Short stature</li> <li><input type="radio"/> Tall stature</li> <li><input type="radio"/> Obesity</li> <li><input type="radio"/> Failure to thrive</li> <li><input type="radio"/> Delayed puberty</li> <li><input type="radio"/> Precocious puberty</li> </ul>	

### 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.