

CYTOGENETICS TEST REQUISITION FORM

PATIENT INFORMATION		ACCOUNT INFORMATION	
First name _____		Last name _____	
Sex <input type="radio"/> Male <input type="radio"/> Female Gender identification (optional): _____		Date of birth (mm/dd/yy) _____	
Ancestry <input type="radio"/> White/Caucasian <input type="radio"/> Hispanic <input type="radio"/> Black/African American <input type="radio"/> Native American <input type="radio"/> East Asian <input type="radio"/> South Asian <input type="radio"/> Middle Eastern <input type="radio"/> Ashkenazi Jewish <input type="radio"/> Other: _____		Address _____ City _____	
Email _____		State _____ Zip code _____ Country _____	
Address _____		Ordering provider	
City _____ State _____ Zip code _____		Name _____ Role/Title _____	
Primary phone _____		Phone _____ NPI _____	
Is this patient deceased? <input type="radio"/> Yes <input type="radio"/> No Deceased date: _____		Email address (for report access) _____	
Reporting Preference: <input type="radio"/> Portal <input type="radio"/> Fax <input type="radio"/> Email <input type="radio"/> Care Evolve <small>If unmarked, we will use the account's default preferences or fax to new clients.</small>			
Additional Reporting Providers <input type="radio"/> Same as ordering provider			
Name _____		Role/Title _____	
Phone _____		NPI _____	
Email address (for report access) _____			
Additional clinical or laboratory contact (optional)			
Name _____		Email address (for report access) _____	
SEND ADDITIONAL REPORT COPIES TO			
Healthcare provider/Acct # _____		Fax #/Email _____	
STATEMENT OF MEDICAL NECESSITY			
By submission of this test requisition and accompanying sample(s), I: (i) authorize and direct GeneDx to perform the testing indicated; (ii) certify that the person listed as the ordering provider is authorized by law to order the test(s) requested; (iii) certify that any custom panel and/or ordered test(s) requested on this test requisition form are reasonable and medically necessary for the diagnosis and/or treatment of a disease, illness, impairment, symptom, syndrome or disorder; (iv) the test results will determine my patient's medical management and treatment decisions of this patient's condition on this date of service; (v) have obtained this patient's and relatives', when applicable, written informed consent to undergo any genetic testing requested; and (vi) that the full and appropriate diagnosis code(s) are indicated to the highest level of specificity.			
Signature of provider (required) _____		Date _____	
ICD-10 codes (required):			
Clinical diagnosis: _____		Age at initial presentation: _____	
PATIENT CONSENT			
By signing this form I acknowledge as the patient that I have read the attached informed consent document and that I authorize GeneDx to perform genetic testing as described. I have been informed that GeneDx may contact me or my healthcare provider about research opportunities in the future. For the insurance bill, I understand and authorize GeneDx to share information with the designated insurance carrier for reimbursement. I understand that GeneDx will attempt to contact me if my estimated out-of-pocket responsibility will be greater than \$100 per test. If GeneDx is unsuccessful in its attempts to contact me, it will be my responsibility to contact GeneDx to determine and pay the out-of-pocket cost. More information, including the GeneDx Notice of Privacy Policies, is available on GeneDx's website: www.genedx.com			
<input type="radio"/> By checking this box, I confirm that I am a New York state resident, and I give permission for GeneDx to retain any remaining sample longer than 60 days after the completion of testing, and to be used as a de-identified sample for test development and improvement, internal validation, quality assurance, and training purposes. Otherwise, New York law requires GeneDx to destroy my sample after 60 days, and it cannot be used for the studies listed above.			
<input type="radio"/> Check this box if you wish to opt out of being contacted for research studies.			
Signature of patient (required) _____		Date _____	
PATIENT STATUS – ONE MUST BE CHECKED: <input type="radio"/> Hospital outpatient <input type="radio"/> Hospital inpatient Date of discharge: _____ <input type="radio"/> Not a hospital patient			
PAYMENT OPTIONS			
<input type="radio"/> Insurance Bill		<input type="radio"/> Copy of front and back of primary and secondary (if applicable) insurance card(s) included	
Insurance carrier _____		Referral/Prior authorization number (please attach) _____	
Insurance ID # _____		GeneDx benefit investigation # _____	
Relationship to insured: <input type="radio"/> Self <input type="radio"/> Spouse <input type="radio"/> Child <input type="radio"/> Other _____		Policyholder's name _____	
Policyholder's date of birth _____		Policyholder's date of birth _____	
<input type="radio"/> Patient Bill		<input type="radio"/> Institutional Bill	
Amount (\$): _____		GeneDx account # _____	
Hospital/Lab name _____		Hospital/Lab name _____	
If Patient Bill is selected, I am electing to be treated as a self-pay patient for this testing. I agree that neither GeneDx nor I will submit a claim to my insurance for this testing, if I have insurance. GeneDx will send an invoice to the patient listed above.			
<input type="radio"/> GeneDx Affiliate Code: _____			
Place sticker/stamp here			

CLINICAL INFORMATION

Account #	Account Name	
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CLINICAL INFORMATION (DETAILED MEDICAL RECORDS MUST BE ATTACHED)

Diagnosis: _____

Dysmorphic features: _____

Other medical problems: _____

Suspected syndrome(s): _____

Previous cytogenetics (attach copy): _____

Clinical and family history: _____

Pre/Perinatal History

- Cystic hygroma
- Diaphragmatic hernia
- Encephalocele
- Growth delay
- Increased nuchal translucency
- Intrauterine growth retardation
- Neural tube defect
- Omphalocele

Structural Brain Abnormalities

- Abnormality of the corpus callosum
- Holoprosencephaly
- Hydrocephalus
- Lissencephaly
- Pachygyria

Developmental/Behavioral Findings

- Attention deficit hyperactivity disorder
- Autistic behavior
- Behavioral abnormality
- Delayed fine motor development
- Delayed gross motor development
- Delayed speech & language development
- Developmental regression
- Gait disturbance
- Global developmental delay
- Hyperactivity
- Intellectual disability
- OCD
- Sleep disturbance
- Specific learning disability

Neurological Findings

- Abnormality of nervous system
- Anosmia, congenital
- Ataxia
- Cerebral palsy
- Encephalopathy
- Epileptic encephalopathy
- Peripheral neuropathy
- Seizures
- Spasticity

Craniofacial/Dysmorphism

- Abnormal facial shape (Dysmorphic features)
- Brachycephaly
- Cleft lip
- Cleft palate
- Craniosynostosis
- External ear malformation
- Macrocephaly
- Microcephaly

Eye Defects/ Vision

- Abnormality of vision
- Aniridia
- Coloboma
- Esotropia
- Microphthalmia
- Retinitis pigmentosa

Hearing Impairment

- Conductive hearing impairment/bilateral
- Sensorineural hearing impairment/bilateral

Cardiac Findings

- Abnormal heart morphology
- Atrial septal defect
- Cardiomyopathy
- Coarctation of aorta
- Tetralogy of fallot
- Ventricular septal defect

Gastrointestinal Findings

- Aganglionic megacolon
- Failure to thrive
- Tracheoesophageal fistula

Musculoskeletal Findings

- Clinodactyly
- Craniosynostosis
- Hemihypertrophy
- Hypertonia
- Hypotonia
- Pectus excavatum
- Polydactyly
- Scoliosis
- Short stature

- Syndactyly
- Talipes equinovarus
- Tall stature

Skin/Hair Findings

- Café-au-lait macules
- Cutis laxa
- Hyperpigmentation of the skin
- Hypopigmentation of the skin
- Ichthyosis
- Sparse hair

Genitourinary Findings

- Ambiguous genitalia
- Cryptorchidism
- Horseshoe kidney
- Hydronephrosis
- Hypospadias
- Polycystic kidney disease

Metabolic Issues/Mito (Attach relevant lab reports/values)

- Elevated CPK_____

Endocrine Findings

- Delayed puberty



Signature of provider (required)

Date

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EXPEDITED TESTING

I would like expedited testing for the following reason: Pregnancy (gestational age: _____ weeks) Transplantation Other: _____

TEST MENU

TEST CODE	TEST NAME
<input type="radio"/> 910	GenomeDx: Whole-Genome Chromosomal Microarray (CMA)
<input type="radio"/> 905	Known Familial Deletion/Duplication Testing
<input type="radio"/> 725	Parental follow up testing for copy number variant(s) (for eligible VUS identified by GenomeDx)
<input type="radio"/> J542	FISH Follow-up after GenomeDx (Test 910) Please specify chromosomal region: 1: _____ 2: _____
<input type="radio"/> 336	FISH Please specify chromosomal region: 1: _____ 2: _____
<input type="radio"/> 0559	Chromosome Analysis, Peripheral blood (routine study)
<input type="radio"/> T982	Rule out Mosaicism Chromosome Analysis, Peripheral blood (Must indicate suspected mosaic chromosomal abnormality)

VARIANT INFORMATION (CNV(s) require coordinates and genome build or transcript # and exon #)

Variant(s): _____
 Genome Build: _____ Coordinates: _____
 Gene: _____ Exon #: _____ Transcript #: _____
 Proband Name: _____
 Relationship to Proband: _____
 Proband GeneDx Accession #: _____

PROBAND TESTED AT ANOTHER LABORATORY

- Positive control included/will be sent - **Positive control is recommended if previous test was performed at another lab.**
- Positive control not available (Caveat language will be included on a negative report)
- Family member test report included (recommended if previous test was performed at another lab)

INFORMED CONSENT

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General Information About Genetic Testing

What is genetic testing?

DNA provides instructions for our body's growth and development. Genes are distinct sequences of DNA, and are arranged on chromosomes. The DNA in a gene contains instructions for making proteins, which determine things like growth and metabolism as well as traits like eye color and blood type. Genetic disorders are caused by certain changes in DNA affecting the structure or number of chromosomes. Genetic testing is a laboratory test that tries to identify these changes in chromosomes or the DNA. Genetic testing can be a diagnostic test, which is used to identify or rule out a specific genetic condition. Genetic screening tests are used to assess the chance for a person to develop or have a child with a genetic condition. Genetic screening tests are not typically diagnostic and results may require additional testing.

The purpose of this test is to see if I, or my child, may have a genetic variant or chromosome rearrangement causing a genetic disorder or to determine the chance that I, or my child, will develop or pass on a genetic disorder in the future. 'My child' can also mean my unborn child, for the purposes of this consent.

If I/my child already know the specific gene variant(s) or chromosome rearrangement that causes the genetic disorder in my family, I will inform the laboratory of this information.

What could I learn from this genetic test?

The following describes the possible results from the test:

1) Positive: A positive result indicates that a genetic variant has been identified that explains the cause of my/my child's genetic disorder or indicates that I/my child am at increased risk to develop the disorder in the future. It is possible to test positive for more than one genetic variant.

2) Negative: A negative result indicates that no disease-causing genetic variant was identified by the test performed. It does not guarantee that I/my child will be healthy or free from genetic disorders or medical conditions. If I/my child test negative for a variant known to cause the genetic disorder in other members of my/my child's family, this result rules out a diagnosis of the same genetic disorder in me/my child due to this specific change.

3) Inconclusive/Variant of Uncertain Significance (VUS): A finding of a variant of uncertain significance indicates that a genetic change was detected, but it is currently unknown whether that change is associated with a genetic disorder either now or in the future. A variant of uncertain significance is not the same as a positive result and does not clarify whether I/my child is at increased risk to develop a genetic disorder. The change could be a normal genetic variant or it could be disease-causing. Further analysis may be recommended, including testing parents and other family members. Detailed medical records or information from other family members also may be needed to help clarify results.

4) Unexpected results: In rare instances, this test may reveal an important genetic change that is not directly related to the reason for ordering this test. For example, this test may tell me about the risk for another genetic condition I/my child is not aware of or it may indicate differences in the number or rearrangement of sex chromosomes. This information may be disclosed to the ordering health care provider if it likely impacts medical care.

Result interpretation is based on currently available information in the medical literature, research and scientific databases. Because the literature, medical and scientific knowledge are constantly changing, new information that becomes available in the future may replace or add to the information GeneDx used to interpret my/my child's results. Providers can contact GeneDx at any time to discuss the classification of an identified variant. In addition, I or my/my child's health care providers may monitor publicly available resources used by the medical community, such as ClinVar (www.clinvar.com), to find current information about the clinical interpretation of my/my child's variant(s).

For tests that evaluate data from multiple family members, my spouse, or partner concurrently, results may be included in a single comprehensive report.

What are the risks and limitations of this genetic test?

- Genetic testing is an important part of the diagnostic process. However, genetic tests may not always give a definitive answer. In some cases, testing may not identify a genetic variant even though one exists. This may be due to limitations in current medical knowledge or testing technology.
- Accurate interpretation of test results may require knowing the true biological relationships in a family. Failing to accurately state the biological relationships in my/my child's family may result in incorrect interpretation of results, incorrect diagnoses, and/or inconclusive test results. In some cases, genetic testing can reveal that the true biological relationships in a family are not as they were reported. This includes non-paternity (the stated father of an individual is not the biological father) and consanguinity (the parents of an individual are related by blood). It may be necessary to report these findings to the health care provider who ordered the test.
- Genetic testing is highly accurate. Rarely, inaccurate results may occur for various reasons. These reasons include, but are not limited to: mislabeled samples, inaccurate reporting of clinical/medical information, rare technical errors, or unusual circumstances such as bone marrow transplantation, or the presence of change(s) in such a small percentage of cells that the change(s) may not be detectable by the test (mosaicism).

- This test does not have the ability to detect all of the long-term medical risks that I/my child might experience. The result of this test does not guarantee my health or the health of my child/fetus. Other diagnostic tests may still need to be done, especially when only a genetic screening test has been performed previously.
- Occasionally, an additional sample may be needed if the initial specimen is not adequate.

Patient Confidentiality and Genetic Counseling

It is recommended that I receive genetic counseling before and after having this genetic test. I can find a genetic counselor in my area here: www.nsgc.org. Further testing or additional consultations with a health care provider may be necessary.

To maintain confidentiality, the test results will only be released to the referring health care provider, to the ordering laboratory, to me, to other health care providers involved in my/my child's diagnosis and treatment, or to others as entitled by law. The United States Federal Government has enacted several laws that prohibit discrimination based on genetic test results by health insurance companies and employers. In addition, these laws prohibit unauthorized disclosure of this information. For more information, I understand that I can visit www.genome.gov/10002077.

International Specimens

If I/my child reside outside the United States, I attest that by providing a sample for testing, I am not knowingly violating any export ban or other legal restriction in the country of my/my child's residence.

Additional information about the specific test being ordered is available from my health care provider or I can go to the GeneDx website, www.genedx.com. This information includes the complete gene lists, the specific types of genetic disorders that can be identified by the genetic test, the likelihood of a positive result, the limitations of genetic testing, as well as information about how specimens and information are stored and used.

Specimen Retention

After testing is complete, the de-identified submitted specimen may be used for test development and improvement, internal validation, quality assurance, and training purposes. DNA specimens are not returned to individuals or to referring health care providers unless specific prior arrangements have been made.

I understand that samples from residents of New York State will not be included in the de-identified research studies described in this authorization and will not be retained for more than 60 days after test completion, unless specifically authorized by my selection. The authorization is optional, and testing will be unaffected if I do not check the box for the New York authorization language. No tests other than those authorized shall be performed on the biological sample.

Database Participation

De-identified health history and genetic information can help health care providers and scientists understand how genes affect human health. Though I/my child may not personally benefit, sharing this information helps health care providers to provide better care for their patients and researchers to make discoveries. GeneDx shares this type of information with health care providers, scientists and health care databases. No personal identifying information will be shared, as it will be replaced with a unique code.

Even though only a code is used for the reporting to the database, there is a risk that I/my child could be identified based on the genetic and health information that is shared. GeneDx believes that this is unlikely, though the risk is greater if I have already shared (my/my child's) genetic or health information with public resources, such as genealogy websites.

Recontact for Research Participation

Separate from the above, GeneDx may collaborate with scientists, researchers and drug developers to advance knowledge of genetic diseases and to develop new treatments. If there are opportunities to participate in research relevant to the disorder in (my/my child's) family, and if I have consented for recontact, GeneDx may allow my healthcare provider to be recontacted for research purposes, such as the development of new testing, drug development, or other treatment modalities. In some situations, such as if my health care provider is not available, I may be contacted directly.

Any research that results in medical advances, including new products, tests or discoveries, may have potential commercial value and may be developed and owned by GeneDx or the collaborating researchers. If any individuals or corporations benefit financially from these studies, no compensation will be provided to (my/my child) or (my/my child's) heirs.