

# XOMEDX TEST REQUISITION FORM



PATIENT INFORMATION	
First Name	Last Name
Genetic 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: _____	
Email	
Address	
City	State Zip Code
Primary Phone	Is this patient deceased? <input type="radio"/> Yes <input type="radio"/> No Deceased Date: _____

SAMPLE INFORMATION	
Date Sample Collected (mm/dd/yy) (required):	Medical Record #
<input type="radio"/> Blood <input type="radio"/> Buccal Swab <input type="radio"/> Other (specify source): _____	
Patient has had a blood transfusion <input type="radio"/> Yes <input type="radio"/> No	Date of Last Transfusion: _____ (2-4 weeks of wait time is required for some testing)
Patient has had an allogenic bone marrow transplant <input type="radio"/> Yes <input type="radio"/> No	
Fibroblasts are recommended for patients who had an allogenic bone marrow transplant. See <a href="http://www.genedx.com/specimen-requirements">www.genedx.com/specimen-requirements</a> for details.	
<input type="radio"/> Treatment-Related <b>RUSH</b>	Date: _____

PATIENT CONSENTS	
<p>By signing this form, I acknowledge as the patient or relative being tested that I have read or have had read to me the GeneDx Informed Consent document at the end of this test requisition form, and understand the information regarding molecular genetics testing. I have had the opportunity to ask questions about the testing, the procedure, the risks, and the alternatives. By signing this form, I authorize GeneDx to perform genetic testing as ordered. I understand that, for tests that evaluate data from multiple family members concurrently, test results from these family members may be included in a single comprehensive report that will be made available to all tested individuals and their healthcare providers.</p> <p>More information, including the GeneDx Notice of Privacy Policies, is available on GeneDx's website: <a href="http://www.genedx.com">www.genedx.com</a></p> <p><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 test development studies.</p> <p><input type="radio"/> Check this box if you wish to opt out of being contacted for research studies.</p> <p><input type="radio"/> Check this box if you do not wish to receive ACMG secondary findings (Full Exome Sequencing and Genome Sequencing Tests ONLY; not for Xpanded® or Slice tests).</p>	
Signature of Patient/Legal Guardian (required)	Date
Signature of Relative A/Legal Guardian	Date
Signature of Relative B/Legal Guardian	Date
<p><b>OPTIONAL AND FOR COMMERCIAL INSURANCE ONLY:</b></p> <p>By entering my preferred contact information below, I give my permission to GeneDx to send me an email and/or text with a link to access my personalized Digital Patient Letter. Data rates may apply.</p>	
Mobile Number*	Email*
*Contact information provided must be for the individual authorizing the genetic testing.	

ACCOUNT INFORMATION	
GeneDx Account Number	Account Name
Phone	Fax
Address	City
State	Zip Code Country
Ordering Provider Name	Role/Title
NPI	Phone Number
Send Report Via <input type="radio"/> Fax <input type="radio"/> Email <input type="radio"/> Portal Fax #/Email: _____	
Additional Reporting Provider's Name	
Send Report Via <input type="radio"/> Fax <input type="radio"/> Email <input type="radio"/> Portal Fax #/Email: _____	
<b>SEND ADDITIONAL REPORT COPIES TO:</b>	
Provider Name	GeneDx Acct#
Fax #/Email: _____	

ICD-10 CODES (Required)	
ICD-10 Codes	
Clinical Diagnosis	Age of Onset

STATEMENT OF MEDICAL NECESSITY	
<p>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.</p>	
Signature of Provider (required)	Date

PAYMENT OPTIONS (Select One)				
<input type="radio"/> <b>INSURANCE BILL</b> (select all that applies)  <input type="radio"/> Commercial <input type="radio"/> Medicaid <input type="radio"/> Medicare <input type="radio"/> Tricare  FOR ALL INSURANCE CARDS PROVIDE FRONT AND BACK COPY OF CARD(S)	Patient Status	<input type="radio"/> Hospital outpatient <input type="radio"/> Hospital inpatient; Date of Discharge _____		
	Name of Insurance Carrier	Insurance ID#:		
	Relationship to Insured	Policy Holder's Name		
	<input type="radio"/> Self <input type="radio"/> Spouse <input type="radio"/> Child <input type="radio"/> Other: _____	Policy Holder's Date of Birth		
	Referral/Prior Authorization # (please attach)	GeneDx Benefit Investigation #		
	Secondary Insurance Type:			
	Insurance Carrier	Insurance ID #	Subscriber Name	Date of Birth
	Relationship to Insured: <input type="radio"/> Self <input type="radio"/> Spouse <input type="radio"/> Child <input type="radio"/> Other: _____			
	<input type="radio"/> <b>PATIENT BILL</b>	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.		
	Amount Due: _____	Authorized Patient/Guardian Signature		
<input type="radio"/> <b>INSTITUTIONAL BILL</b>	GeneDx Account #	Place Sticker/Stamp Here		
	Hospital/Lab Name			

GeneDx Account #	Account Name	
First Name	Last Name	Date of Birth

**CLINICAL INFORMATION (DETAILED MEDICAL RECORDS MUST BE ATTACHED)**  
*Relevant clinical records are required at the time of sample submission to ensure the information is included in data analysis.*

**REQUESTED GENES AND DIFFERENTIAL DIAGNOSIS**

Genes of interest: \_\_\_\_\_

Differential Diagnosis: \_\_\_\_\_

**Pre/Perinatal History**

- Cystic hygroma
- Diaphragmatic hernia
- Encephalocele
- Growth delay
- Increased nuchal translucency
- Intrauterine Growth Retardation
- Nonimmune hydrops fetalis
- Oligohydramnios
- Omphalocele
- Polyhydramnios
- Prematurity GA: \_\_\_\_\_
- Prolonged neonatal jaundice

**Structural Brain Abnormalities**

- Abnormal myelination
- Abnormality of basal ganglia
- Abnormality of brainstem
- Abnormality of periventricular white matter
- Abnormality of the corpus callosum
- Aplasia/hypoplasia of cerebellar vermis
- Aplasia/hypoplasia of cerebellum
- Arnold Chiari malformation
- Cerebellar atrophy
- Heterotopia (Periventricular nodular heterotopia)
- Holoprosencephaly
- Hydrocephalus
- Leukodystrophy
- Lissencephaly
- Pachygyria
- Polymicrogyria
- Ventriculomegaly

**Developmental/Behavioral Findings**

- Absent speech
- Aggressive behavior
- Anxiety
- Autistic Behavior
- Cognitive impairment
- Delayed speech & language development
- Developmental regression
- Dysarthria
- Gait disturbance
- Global developmental delay
- Hyperactivity
- Incoordination
- Intellectual disability
- Learning disability
- Memory impairment
- Sleep disturbance
- Stereotypy

**Neurological Findings**

- Abnormality of nervous system
- Ataxia
- Cerebral palsy
- Chorea
- Cortical Visual Impairment
- Dementia
- Dysarthria
- Dyskinesia
- Dysphasia
- Dystonia
- Encephalopathy
- Headaches
- Hemiplegia
- Infantile Spasms
- Migraines
- Myoclonus
- Parkinsonism
- Peripheral neuropathy
- Seizures
- Sensory neuropathy
- Spasticity
- Syncope
- Tremors
- Vertigo

**Craniofacial/Dysmorphism**

- Abnormal facial shape (Dysmorphic features)  
Specify: \_\_\_\_\_
- Brachycephaly
- Cleft lip and/or palate
- Coarse facial features
- Craniosynostosis
- Macrocephaly
- Microcephaly
- Short neck
- Synophrys

**Eye Defects/ Vision**

- Abnormality of Vision
- Anophthalmia
- Cataracts
- Coloboma
- Corneal opacity
- Ectopia lentis
- External ophthalmoplegia
- Microphthalmia
- Myopia
- Nystagmus
- Optic atrophy
- Optic neuropathy
- Ptosis
- Retinal detachment
- Retinitis pigmentosa
- Strabismus

**Hearing Impairment**

- Abnormal Newborn Screen: \_\_\_\_\_
- Conductive hearing impairment
- Sensorineural hearing impairment

**Endocrine Findings**

- Delayed puberty
- Diabetes Insipidus
- Diabetes Mellitus
- Hyperthyroidism
- Hypophosphatemia
- Hypothyroidism
- Maturity-onset diabetes of the young
- Rickets

**Respiratory Findings**

- Asthma
- Bronchiectasis
- Hyperventilation
- Hypoventilation
- Pneumothorax
- Pulmonary fibrosis
- Respiratory insufficiency

**Hematologic or Immunologic Findings**

- Allergic rhinitis
- Anemia
- Immunodeficiency
- Neutropenia
- Pancytopenia
- Recurrent infections
- Thrombocytopenia

**Skin/Hair Findings**

- Abnormal blistering of the skin
- Abnormality of nail
- Alopecia
- Anhidrosis
- Café-Au-Lait Macules
- Coarse hair
- Cutis Laxa
- Eczema
- Hemangiomas
- Hyperextensible skin
- Hyperpigmentation of the skin
- Hypohidrosis
- Hypopigmentation of the skin
- Ichthyosis
- Skin rash
- Sparse hair
- Telangiectasia
- Vascular skin abnormality
- Velvety skin

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

### Cardiac Findings

- Abnormal heart morphology
- Amyloidosis
- Aortic root dilation
- Arrhythmia
- Atrial septal defect
- Bicuspid aortic valve
- Bradycardia
- Coarctation of aorta
- Dilated cardiomyopathy
- Heterotaxy
- Hypertension
- Hypertrophic cardiomyopathy
- Mitral valve prolapse
- Noncompaction cardiomyopathy
- Patent ductus arteriosus
- Patent foramen ovale
- Prolonged QTc interval
- Sudden death
- Tetralogy of Fallot
- Ventricular septal defect
- Ventricular tachycardia

### Gastrointestinal Findings

- Constipation
- Diarrhea
- Duodenal stenosis/atresia
- Exocrine pancreatic insufficiency
- Failure to thrive
- Feeding difficulties
- Gastroesophageal reflux
- Hepatomegaly
- Inflammatory bowel disease
- Intrahepatic biliary atresia
- Laryngomalacia
- Nausea
- Pancreatitis
- Pyloric stenosis
- Splenomegaly
- Tracheoesophageal fistula
- Vomiting

### Genitourinary Findings

- Ambiguous genitalia
- Cryptorchidism
- Cystic renal dysplasia
- Horseshoe kidney
- Hydronephrosis
- Hypospadias
- Inguinal hernia
- Micropenis

- Nephrolithiasis
- Polycystic kidney disease
- Renal agenesis
- Umbilical hernia

### Musculoskeletal Findings

- Abnormal connective tissue
- Abnormal form of the vertebral bodies
- Abnormality of the ribs
- Arachnodactyly
- Arthralgia
- Arthrogryposis
- Bruising susceptibility
- Clinodactyly
- Decreased muscle mass
- Ectrodactyly
- Exercise intolerance
- Fatigue
- Hemihypertrophy
- Hypertonia
- Hypotonia
- Joint hypermobility
- Muscle weakness
- Myalgia
- Myopathic facies
- Myopathy
- Osteoarthritis
- Osteopenia
- Pain
- Pectus carinatum
- Pectus excavatum
- Polydactyly
- Recurrent fractures
- Rhabdomyolysis
- Scoliosis
- Short stature
- Skeletal dysplasia
- Syndactyly
- Tall stature

### Cancer

- Type: \_\_\_\_\_
- Location: \_\_\_\_\_
- Age of onset: \_\_\_\_\_

### Metabolic Findings (Attached relevant lab reports/values)

- Abnormal activity of mitochondrial respiratory chain
- Abnormal Newborn Screen: \_\_\_\_\_
- Abnormality of mitochondrial metabolism
- Elevated CPK
- Elevated hepatic transaminase
- Hyperammonemia

- Hyperglycemia
- Hypoammonemia
- Hypoglycemia
- Increased serum pyruvate
- Lactic acidosis
- Plasma AA: \_\_\_\_\_
- Urine OA: \_\_\_\_\_

### Vascular System

- Aneurysm
- Arterial calcification
- Arterial dissection
- Arterial tortuosity
- Arteriovenous malformation
- Epistaxis
- Lymphedema
- Pulmonary hypertension
- Stroke

### Other Testing/Imaging (Please provide copy or report if possible)

- Echo: \_\_\_\_\_
- EEG: \_\_\_\_\_
- EMG: \_\_\_\_\_
- Gene Panel: \_\_\_\_\_  
Results: \_\_\_\_\_  
Performed at: \_\_\_\_\_
- Gene Sequencing\*: \_\_\_\_\_  
Results: \_\_\_\_\_  
Performed at: \_\_\_\_\_  
\*If you would like us to comment on the presence/absence of previously identified variants, including parental status (if included), provide complete variant information or a copy of the original report.
- Microarray: \_\_\_\_\_
- MRI: \_\_\_\_\_
- Muscle Biopsy: \_\_\_\_\_
- Ultrasound: \_\_\_\_\_
- X-rays: \_\_\_\_\_

### Additional Clinical Findings:

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

Signature of Provider (required)	Date
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# XOMEDX TEST REQUISITION FORM



GeneDx Account #	Account Name	
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## REASON FOR EXPEDITED TESTING (REQUIRED)

Pregnancy (gestational age \_\_\_\_\_ weeks)
  Transplantation
  Other: \_\_\_\_\_

## TARGETED VARIANT TESTING AND SPECIAL SERVICES

Individual to be Tested:
  Affected/Symptomatic
  Unaffected/Asymptomatic  
 Known Familial Variant(s) in a Nuclear Gene
  Confirmation of Variant Identified in Research Lab
  Targeted Mosaic Variant Testing  
 Known Familial Copy Number Variant(s)
  Known mtDNA Variant(s) Testing
 **(Insurance Billing NOT Accepted; Patient Bill or Institutional Bill MUST be selected on page 1)**

Proband Name	Relationship to Proband	Proband GeneDx Accession #
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Non-GeneDx Test:
  Family member test report included (recommended if previous test was performed at another lab)  
 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)

## VARIANT INFORMATION (please fill out the below information if family member report is not included)

Gene	Coding DNA (c./m.)	Amino Acid (p.)	Transcript (NM#)	Number of Variants: _____
Gene	Coding DNA (c./m.)	Amino Acid (p.)	Transcript (NM#)	

## COPY NUMBER VARIANTS (CNV(s) require coordinates and genome build or transcript # and exon #)

Gene(s)	Exon #	Coordinates	Genome Build	Number of Variants: _____
Gene(s)	Exon #	Coordinates	Genome Build	

## TESTING OPTIONS

### CUSTOM DEL/DUP TESTING

906 Deletion/Duplication Analysis of ONE Nuclear Gene
  703 Deletion/Duplication Analysis of 2-20 Nuclear Genes

Write-in Desired Gene(s) to be Tested: \_\_\_\_\_

**WRITE-IN TEST SELECTION:**
 Test Code: \_\_\_\_\_
  Test Name: \_\_\_\_\_

## HISTORY

**FAMILY HISTORY:**
 No Known Family History
  Pedigree Attached
  Adopted

Relationship	Maternal	Paternal	Relevant History	Age at Dx
1	<input type="radio"/>	<input type="radio"/>		
2	<input type="radio"/>	<input type="radio"/>		
3	<input type="radio"/>	<input type="radio"/>		

**TESTING HISTORY:**
 Test Report Included (recommended)

Other relevant results (clinical, laboratory/biochemical or research): \_\_\_\_\_

# XOMEDX TEST REQUISITION FORM



GeneDx Account #	Account Name	
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## XOMEDX TESTING OPTIONS

TEST CODE	TEST NAME	TEST CODE	TEST NAME
<input type="radio"/> 561a	XomeDx Trio*	<input type="radio"/> 690a	XomeDxPlus Trio* (XomeDx + 690c mtDNA)
<input type="radio"/> 561e	XomeDx Duo*	<input type="radio"/> 690e	XomeDxPlus Duo* (XomeDx + 690c mtDNA)
<input type="radio"/> 561b	XomeDx Proband	<input type="radio"/> 690b	XomeDxPlus Proband (XomeDx + 690c mtDNA)

\*If a Trio or Duo test is ordered, please fill out the Biological Parent Sample Information section below.

### XOMEDX REFLEX TESTING

Test code(s): \_\_\_\_\_

Reflex to (choose one from above): \_\_\_\_\_

### XOMEDX XPERT

*This is applicable for patients with previous exome analysis performed by an outside clinical laboratory where a second opinion is desired. Outside lab report required along with new specimen(s). Institutional/self-pay only.*

TG56 XomeDx Xpert\* (Proband or Trio)

### CUSTOM SLICE TESTING OPTIONS

<input type="radio"/> TG70	Slice - Single Gene (1 gene)	Approved Slice ID: _____
<input type="radio"/> 706	Slice - Multi-Gene (2-150 genes)	Approved Slice ID: _____
<input type="radio"/> 706H	Slice - Homozygous (no gene list)*	Approved Slice ID: _____
<input type="radio"/> J757	Slice - Xpanded (>150 genes, Proband or Trio*)	Approved Slice ID: _____

\*If a Trio or Duo test is ordered, please fill out the Biological Parent Sample Information section below.

### SKIN DISORDER SLICES

- 707 Slice - Epidermolysis Bullosa (EB)
- 708 Slice - Congenital Ichthyosis

### REANALYSIS OF XOMEDX TESTING OPTIONS

*These test options are only appropriate if the patient previously had a XomeDx test (full exome analysis) at GeneDx. We recommend waiting at least one year from original/prior analysis before ordering a Reanalysis.*

<input type="radio"/> 660	XomeDx First Time Reanalysis (no charge)	Reason for Reanalysis Is there new clinical information available? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Other _____
<input type="radio"/> 947	XomeDx Subsequent Reanalysis (charged)	

## BIOLOGICAL PARENT SAMPLE INFORMATION

**ADDITIONAL SAMPLES MUST BE RECEIVED WITHIN 3 WEEKS.** Ordered test codes may require adjusting to appropriately correspond with relative samples received. A change in the ordered test will impact billing, including prior benefits investigations.

<b>Mother</b>	First Name	Last Name	DOB	<input type="radio"/> Asymptomatic	<input type="radio"/> Symptomatic
				<input type="radio"/> At GeneDx	<input type="radio"/> Not Available <input type="radio"/> To be Sent Within 3 Weeks*
<b>Father</b>	First Name	Last Name	DOB	<input type="radio"/> Asymptomatic	<input type="radio"/> Symptomatic
				<input type="radio"/> At GeneDx	<input type="radio"/> Not Available <input type="radio"/> To be Sent Within 3 Weeks*
<b>Other</b>	Relationship to Proband				
	First Name	Last Name	DOB	<input type="radio"/> Asymptomatic	<input type="radio"/> Symptomatic
				<input type="radio"/> At GeneDx	<input type="radio"/> Not Available <input type="radio"/> To be Sent Within 3 Weeks*

ACMG secondary findings, as discussed in the Informed Consent and Authorization Form, are only returned for the patient if an XomeDx test (full exome analysis) is completed.

GeneDx tests are frequently updated and improved based upon the most recent scientific evidence. The test codes, genes, and gene quantities listed on this test requisition are subject to change by GeneDx at any time. The most current test menu and list of genes included for a specific test panel may be found on our website, [genedx.com](http://genedx.com). Please note that GeneDx reserves the right to modify and upgrade any ordered panel to the version currently listed on our website.

GeneDx Account #	Account Name	
First Name	Last Name	Date of Birth

For the purposes of this consent, “I”, “my”, and “your” will refer to me or to my child, including my unborn child, if my child is the person for whom the healthcare provider has ordered testing.

### PURPOSE OF THIS TEST

The purpose of this test is (a) to see if I may have a genetic variant or chromosome rearrangement causing a genetic disorder; or (b) to evaluate the chance that I will develop or pass on a genetic disorder in the future. If I already know the specific gene variant(s) or chromosome rearrangement that causes the genetic disorder in my family, I agree to inform the laboratory of this information.

### WHAT TYPE OF TEST RESULTS CAN I EXPECT FROM GENETIC TESTING?

- Positive:** A change in your DNA was found, which is very likely the cause of your features/symptoms. This is the most straightforward test result, which can be used as the basis to test other family members to determine their chances of having either the disease or a child with the disease.
- Negative:** No variants were found to explain your symptoms. This does not mean that you do not have a genetic condition. It is still possible that there is a genetic variant not found by the test that was ordered. Your healthcare provider or genetic counselor may discuss more testing either now or in the future.
- Variant of Uncertain Significance (VUS):** A change in a gene was found. However, we are not sure whether this variant is the cause of your symptoms/features. More information is needed. We may suggest testing other family members to help figure out the meaning of the test result.
- 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 find you are at risk for another genetic condition I am not aware of or it may indicate differences in the number or rearrangement of sex chromosomes. We may disclose this information to the ordering healthcare provider if it likely affects medical care.

Because medical and scientific knowledge is constantly changing, new information that becomes available may supplement the information GeneDx used to interpret my results. Healthcare providers can contact GeneDx at any time to discuss the classification of an identified variant.

### WHAT IS TRIO/DUO-BASED GENETIC TESTING?

For some genetic tests, including samples from the biological parents and/or other biological relatives along with the patient’s sample can help with the interpretation of the test results. These tests are often referred to as “trio tests” since they typically include samples from the patient and both parents.

Samples from relatives should be submitted with the patient’s sample. Clinical information must be provided for the patient and any relative who submits a sample.

I understand that GeneDx will use the relative sample(s) when needed for the interpretation of my test results and that my test report may include clinical and genetic information about a relative when it is relevant to the interpretation of the test results. I further understand that relatives will not receive an independent analysis of data nor a separate report.

### RISKS AND LIMITATIONS OF GENETIC TESTING

- 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. I understand that if I fail to accurately state the biological relationships in my family, it could lead to incorrect interpretation of the test results, incorrect diagnoses, and/or inconclusive test results. If genetic testing reveals that the true biological relationships in a family are not as I reported them, including non-paternity (the reported father is not the biological father) and consanguinity (the parents are related by blood), I agree to have these findings reported to the healthcare provider who ordered the test.
- Although genetic testing is highly accurate, inaccurate results may occur. These reasons include, but are not limited to mislabeled samples, inaccurate reporting of clinical/medical information, rare technical errors, or other reasons.
- I understand that this test may not detect all of the long-term medical risks that I might experience. The result of this test does not guarantee my health and that additional diagnostic tests may still need to be done.
- I agree to provide an additional sample if the initial sample 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 at [www.nsgc.org](http://www.nsgc.org). Further testing or additional consultations with a healthcare provider may be necessary.

To maintain confidentiality, test results will only be released to the referring healthcare provider, the ordering laboratory, to me, to other healthcare providers involved in my care, diagnosis and treatment, or to others with my consent or as permitted or required by law. Federal laws prohibit unauthorized disclosure of this information. More information can be found at: [www.genome.gov/10002077](http://www.genome.gov/10002077)

### INTERNATIONAL SAMPLES

If I 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 residence.

### SAMPLE RETENTION

After testing is complete, my sample may be de-identified and be used for test development and improvement, internal validation, quality assurance, and training purposes. GeneDx will not return DNA samples to you or to referring healthcare 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 GeneDx will not retain them 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. GeneDx will not perform any tests on the biological sample other than those specifically authorized.

### DATABASE PARTICIPATION

De-identified health history and genetic information can help healthcare providers and scientists understand how genes affect human health. Sharing this de-identified information helps healthcare providers to provide better care for their patients and researchers to make new discoveries. GeneDx shares this type of information with healthcare providers, scientists, and healthcare databases. GeneDx will not share any personally identifying information and will replace the identifying information with a unique code not derived from any personally identifying information. Even with a unique code, there is a risk that I 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 genetic or health information with public resources, such as genealogy websites.

GeneDx Account #	Account Name	
First Name	Last Name	Date of Birth

## EXOME/GENOME SEQUENCING SECONDARY FINDINGS

- Applicable Only for Full Exome Sequencing and Genome Sequencing Tests.
- Does not pertain to Xpanded® or Slice tests

As many different genes and conditions are analyzed in an exome or genome sequencing test, these tests may reveal some findings not directly related to the reason for ordering the test. Such findings are called “incidental” or “secondary” and can provide information that was not anticipated.

Secondary findings are variants, identified by an exome or genome sequencing test, in genes that are unrelated to the individual’s reported clinical features.

The American College of Medical Genetics and Genomics (ACMG) has recommended that secondary findings identified in a specific subset of medically actionable genes associated with various inherited disorders be reported for all probands undergoing exome or genome sequencing. Please refer to the latest version of the ACMG recommendations for reporting of secondary findings in clinical exome and genome sequencing for complete details of the genes and associated genetic disorders. Reportable secondary findings will be confirmed by an alternate test method when needed.

### WHAT WILL BE REPORTED FOR THE PATIENT?

All pathogenic and likely pathogenic variants associated with specific genotypes identified in the genes (for which a minimum of 10X coverage was achieved by exome sequencing or a minimum of 15X coverage was achieved by genome sequencing), as recommended by the ACMG.

### WHAT WILL BE REPORTED FOR RELATIVES?

The presence or absence of any secondary finding(s) reported for the proband will be provided for all relatives analyzed by an exome or genome sequencing test.

### LIMITATIONS

Pathogenic and/or likely pathogenic variants may be present in a portion of the gene not covered by this test and therefore are not reported. The absence of reportable secondary findings for any particular gene does not mean there are no pathogenic and/or likely pathogenic variants in that gene. Pathogenic variants and/or likely pathogenic variants that may be present in a relative, but are not present in the proband, will not be identified, or reported. Only changes at the sequence level will be reported in the secondary findings report. Larger deletions/duplications, abnormal methylation, triplet repeat or other expansion variants, or other variants not routinely identified by clinical exome and genome sequencing will not be reported.

## FINANCIAL AGREEMENT AND GUARANTEE

For insurance billing, I understand and authorize GeneDx to bill my health insurance plan on my behalf, to release any information required for billing, and to be my designated representative for purposes of appealing any denial of benefits. I irrevocably assign to and direct that payment be made directly to GeneDx.

I understand that my out-of-pocket costs may be different than the estimated amount indicated to me by GeneDx as part of a benefit investigation. I agree to be financially responsible for any and all amounts as indicated on the explanation of benefits issued by my health insurance plan. If my insurance provider sends a payment directly to me for services performed by GeneDx on my behalf, I agree to endorse the insurance check and forward it to GeneDx within 30 days of receipt as payment towards GeneDx’s claim for services rendered.

If I do not have health insurance, I agree to pay for the full cost of the genetic testing that was ordered by my healthcare provider and billed to me by GeneDx. I further understand and agree that, if I fail to make payment for genetic testing, in accordance with the payment policies of GeneDx, my account may be turned over to an external collection agency for non-payment. I agree to pay any associated collection costs, including attorney fees. By my signature on the GeneDx Test Requisition Form or at the bottom of this form, I accept full and complete financial responsibility for all genetic testing ordered by my healthcare provider.

## MEDICARE

A completed Advance Beneficiary Notice (ABN) is required for Medicare patients. Please visit our website, [www.genedx.com/billing](http://www.genedx.com/billing) for more information.

## DIGITAL PATIENT LETTER CONSENT

- Applicable Only for Commercial Insurance
- Estimate is provided by your health insurance company and therefore NO estimate will be sent for any orders placed with federal or state-funded insurance plans (e.g. Medicare, Medicaid, Tricare, etc.), institutional bill, or patient bill (self-pay).

To provide you with the estimated out-of-pocket expenses related to your test, GeneDx will send you an email and/or text with the link to access your personalized Digital Patient Letter. In order to send this information, we need your consent and agreement to the following items:

1. GeneDx can use your email address or mobile phone number solely for the purpose of GeneDx sending your estimated financial obligation. Text message data rates may apply. GeneDx is not responsible for undelivered messages due to incorrect or illegible contact information.
2. GeneDx will send you an email and/or text message containing a link to view your personalized Patient Letter that includes the test out-of-pocket estimate. The link is time-sensitive and will only be available for 72 hours from the time the message is sent. In order to view the estimate, you must click the link in the message.
3. If you take no action, GeneDx will assume that you agree to move ahead with testing and will bill your health insurance. You can approve testing with insurance, switch to self-pay, or cancel the test via the link within the given 72-hour window. In turn, if GeneDx receives your sample(s) and the billing method hasn’t been changed, or the test hasn’t been cancelled, we will move ahead with testing as ordered, and you will be responsible for any out-of-pocket costs for the completion of the test(s).