

PRENATAL GENETICS 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		
Specimen ID	Medical record #	Date Sample Collected (mm/dd/yy)
<input type="radio"/> CVS <input type="radio"/> Amniotic Fluid <input type="radio"/> Fetal Blood (PUBS) <input type="radio"/> Cultured CV <input type="radio"/> Cultured Amniocytes <input type="radio"/> Products of Conception (POC), specify tissue: _____ <input type="radio"/> DNA: tissue source _____ <input type="radio"/> Maternal blood for MCC (please send ALL studies) <input type="radio"/> Paternal blood (please send for ALL molecular studies)		
Father's Last Name	Father's First Name	Father's Date of Birth (mm/dd/yy)

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

PATIENT CONSENT FOR GENETIC TESTING, FINANCIAL AGREEMENT AND GUARANTEE:	
<p>By signing this form, I acknowledge as the patient or relative being tested that I have read the GeneDx Informed Consent document available from my healthcare provider or at genedx.com/forms, and I authorize GeneDx to perform genetic testing as ordered. I understand that, for tests that evaluate data from multiple family members concurrently, 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. By my signature below, I accept full and complete financial responsibility for all genetic testing ordered by my healthcare provider. 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 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 and agree to be financially responsible for any and all amounts as indicated on the explanation of benefits issued by my health insurance plan. I am aware that my insurance provider may send 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. 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 and I agree to pay any associated collection costs, including attorney fees.</p> <p>More information, including the GeneDx Notice of Privacy Policies, is available on GeneDx's website: www.genedx.com</p> <p>Medicare: A completed Advance Beneficiary Notice (ABN) is required for Medicare patients. Please visit our website, www.genedx.com/billing for more information.</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 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.</p> <p><input type="radio"/> Check this box if you wish to opt out of being contacted for research studies.</p>	
Signature of Patient (required)	Date

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

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"/> INSURANCE BILL (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 _____			
	<input type="radio"/> Not a hospital patient			
	Name of Insurance Carrier		Insurance ID#:	
	Relationship to Insured <input type="radio"/> Self <input type="radio"/> Spouse <input type="radio"/> Child <input type="radio"/> Other: _____		Policy Holder's Name	
	Referral/Prior Authorization # (please attach)		Policy Holder's Date of Birth	
	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"/> PATIENT BILL	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"/> INSTITUTIONAL BILL	GeneDx Account #		Place Sticker/Stamp Here	
	Hospital/Lab Name			

CLINICAL INFORMATION

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

CLINICAL INFORMATION

Relevant clinical records are required at the time of sample submission to ensure the information is included in data analysis.

Fetal Gender: M F Discrepant: _____ Chromosome Analysis (if known): _____ Diagnosis: _____
 ICD-10 codes: _____ IVF Pregnancy Maternal Diabetes mellitus Egg Donor Sperm Donor

ULTRASOUND INFORMATION/FINDINGS

Date of Ultrasound: ___/___/___ GA at time of Ultrasound: ___Weeks ___Days
 Date of collection: ___/___/___ Last menstrual period: ___/___/___

Pre/Perinatal History

- Cystic hygroma
- Diaphragmatic hernia
- Encephalocele
- Increased nuchal translucency
- Intrauterine growth retardation
- Neural tube defect
- Nonimmune hydrops fetalis
- Oligohydramnios
- Omphalocele
- Polyhydramnios
- Increased nuchal fold (____ mm)

Structural Brain Abnormalities

- Abnormal myelination
- Abnormality of periventricular white matter
- Abnormality of the corpus callosum
- Arnold Chiari malformation
- Cerebellar atrophy
- CNS hypomyelination
- Cortical dysplasia
- Cortical tubers
- Holoprosencephaly
- Hydrocephalus
- Lissencephaly
- Pachygyria
- Polymicrogyria
- Ventriculomegaly

Craniofacial/Dysmorphism

- Cleft lip
- Cleft palate
- Macrocephaly
- Microcephaly

Eye Defects/ Vision

- Anophthalmia
- Microphthalmia

Cardiac Findings

- Atrial septal defect
- Cardiac rhabdomyoma
- Heterotaxy
- Tetralogy of Fallot
- Ventricular septal defect

Gastrointestinal Findings

- Congenital diaphragmatic hernia
- Duodenal stenosis/atresia
- Gastroschisis
- Omphalocele

Musculoskeletal Findings

- Abnormal form of the vertebral bodies
- Abnormality of the ribs
- Abnormality of the upper limb
- Arthrogryposis
- Bowing of the long bones
- Ectrodactyly
- Fractures of the long bones
- Limb joint contracture
- Multiple prenatal fractures
- Polydactyly
- Scoliosis
- Small chest circumference
- Syndactyly
- Talipes equinovarus
- Thoracic hypoplasia

Genitourinary Findings

- Ambiguous genitalia
- Cystic renal dysplasia
- Horseshoe kidney
- Polycystic kidney dysplasia
- Renal agenesis
- Umbilical hernia

Signature of Provider (required) _____ Date _____

Other: _____

Previous pregnancy or family history of: ONTD Down Syndrome Genetic disorders (please explain below)
 Other: _____ (Attach Pedigree if Available)

Please explain pregnancy or family history: AMA Abnormal maternal serum screen for _____ Abnormal NIPT for _____

Please choose test(s) and provide clinical information in the appropriate section above. GeneDx performs maternal contamination studies for prenatal tests, so a maternal blood sample is requested for prenatal tests. All tests will be performed concurrently unless order of testing is specified.

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REASON FOR EXPEDITED TESTING (REQUIRED)

Pregnancy (gestational age _____ weeks)
 Transplantation
 Other: _____

TARGETED VARIANT TESTING AND SPECIAL SERVICES

902 Prenatal Testing for Known Familial Pathogenic Variant
 Fetus to be tested: Known Abnormalities Unknown

Affected Relative Name	Relationship to Fetus	GeneDx Accession #
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Known Familial Sequence Variant
 Known Familial Copy Number Variant

Non-GeneDx Test:
 Relative tested at GeneDx (provide GeneDx ID number above)
 Positive control included/will be sent - Positive control is REQUIRED if previous test was performed at another lab.
 Family member test report included (REQUIRED if previous test was performed at another lab)

Variant Information (please fill out the below information if family member report is not included)
 Number of Variants: _____

Gene	Coding DNA (c./m.)	Amino Acid (p.)	Transcript (NM#)
Gene	Coding DNA (c./m.)	Amino Acid (p.)	Transcript (NM#)

Copy Number Variants (CNV(s) require coordinates and genome build or transcript # and exon #)
 Number of Variants: _____

Gene(s)	Exon #	Coordinates	Genome Build
Gene(s)	Exon #	Coordinates	Genome Build

WRITE-IN TEST SELECTION

Test Code: _____ Test Name: _____

HISTORY

FAMILY HISTORY:
 No Known Family History
 Pedigree Attached
 Adopted

Relationship to Fetus to be Tested	Maternal	Paternal	Relavant 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): _____

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TEST MENU

SEQUENCING TESTS FOR FETUSES WITH ABNORMAL ULTRASOUND FINDINGS

TEST CODE	TEST NAME	# OF GENES	TEST CODE	TEST NAME	# OF GENES
Noonan Spectrum and Rasopathies			Brain Malformations (Continued)		
<input type="radio"/> 357	Prenatal Noonan Spectrum Disorders Panel	11	<input type="radio"/> J802	Prenatal Pontocerebellar Hypoplasia Panel*	19
Skeletal and Limb Abnormalities			<input type="radio"/> 2373	Holoprosencephaly Panel*	4
<input type="radio"/> 937	Prenatal Limb Abnormalities Panel*	5	<input type="radio"/> 2553E	X-linked Hydrocephalus/MASA/CRASH Syndrome (<i>L1CAM</i>): FEMALES*	1
<input type="radio"/> 949	Prenatal Skeletal Dysplasia Panel*	48	<input type="radio"/> 2553	X-linked Hydrocephalus/MASA/CRASH Syndrome (<i>L1CAM</i>): MALES	1
<input type="radio"/> 3383	Campomelic Dysplasia* (<i>SOX9</i>)	1	Disorders of Sex Differentiation		
<input type="radio"/> 738	Cornelia de Lange Syndrome* (<i>NIPBL</i> -related)	1	<input type="radio"/> J719-6	Prenatal 46, XY Disorders of Sex Development Panel*	19
<input type="radio"/> 408	Duane-Radial-Ray Syndrome* (<i>DRRS</i> ; <i>SALL4</i>)	1	<input type="radio"/> 702	Prenatal SRD5A2 Gene Sequencing (5-Alpha Reductase Deficiency)	1
<input type="radio"/> 407	Ectrodactyly-ED-Clefting/Hay-Wells Syndrome (<i>TP63</i> , <i>p63</i>)	1	<input type="radio"/> 2201	Androgen Insensitivity Syndrome (<i>AR</i>)	1
<input type="radio"/> 2363	Holt-Oram Syndrome (<i>TBX5</i>)	1	<input type="radio"/> 409	XY Female Gonadal Dysgenesis (<i>SRY</i>)	1
<input type="radio"/> 2523	Townes-Brocks Syndrome (<i>SALL1</i>)	1	<input type="radio"/> 746	46, XY Gonadal Dysgenesis (<i>NR5A1</i> -related)	1
Brain Malformations			Other Panels and Full Gene Testing for Multiple Congenital Anomalies		
<input type="radio"/> TF32	Prenatal <i>FMR1</i> CGG Repeat Analysis	1	<input type="radio"/> TG85	Prenatal Akinseia/Arthrogyrosis Panel*	28
<input type="radio"/> TF33	Prenatal Dosage Analysis of <i>SMN1</i> and <i>SMN2</i>	2	<input type="radio"/> 663	Adrenal Hypoplasia Congenita (AHC), X-linked (<i>NROB1</i>)	1
<input type="radio"/> TF71	Prenatal <i>AR</i> Repeat Analysis	1	<input type="radio"/> 428	Anophthalmia/Microphthalmia: Sequencing (<i>SOX2*</i> , <i>OTX2*</i> and <i>VSX2</i>)	3
<input type="radio"/> TG16	Prenatal <i>DMPK</i> Repeat Analysis	1	<input type="radio"/> 2262	Prenatal CHD7 Gene Sequencing (CHARGE Syndrome)	1
<input type="radio"/> J803	Prenatal Joubert Syndrome and Related Disorders Panel*	29	<input type="radio"/> 2503	Smith-Lemli-Opitz Syndrome* (<i>DHCR7</i>)	1
<input type="radio"/> J793	Prenatal Lissencephaly Panel*	26	<input type="radio"/> 934	Tuberous Sclerosis* (<i>TSC1</i> and <i>TSC2</i>)	2

* Entire panel/test or specific gene includes deletion/duplication testing.

CYTOGENETICS AND BIOCHEMICAL TESTS

Cytogenic Tests		Chromosome Analysis (Select Specimen Type)	
<input type="radio"/> 3582	Rapid aneuploidy FISH (13, 18, 21, X, Y)	<input type="radio"/> 2136	Karyotype: Amniotic Fluid
<input type="radio"/> 410	Prenatal Targeted Chromosomal Microarray (CMA) Run as REFLEX test if _____ test(s) are negative	<input type="radio"/> A587	Karyotype: CVS
<input type="radio"/> 460	Prenatal/POC Whole Genome Chromosomal Microarray (CMA) Run as REFLEX test if _____ test(s) are negative	<input type="radio"/> 1053	Karyotype: Products of Conception (POC)
<input type="radio"/> J542	FISH Follow-up after CMA (Test 410 or 460) Please specify chromosomal region: 1. _____ 2. _____	<input type="radio"/> 0559	Karyotype: Peripheral Blood
Biochemical and Other Tests		<input type="radio"/> T982	Karyotype to Rule out Mosaicism: Peripheral Blood (Must indicate suspected mosaic chromosomal abnormality)
<input type="radio"/> 2122	Amniotic fluid AFP (1952-1 automatic reflex to AChE testing if AF-AFP is elevated)		
<input type="radio"/> 437	Zygosity Testing		

Please write any special instructions, e.g., in which order to perform tests:

Notes:

- (1) If sufficient fetal material is submitted, most testing can be performed concurrently. If no other instructions are given, all tests will be performed concurrently.
- (2) If you choose to have the testing done in a particular order ('reflex testing'), indicate the order of tests by numbering the tests (Example: (1) Chromosome analysis, CVS; (2) Prenatal targeted array (if chromosomes normal); (3) Noonan syndrome testing (if array normal)).

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First Name	Last Name	Date of Birth

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 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 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/my child's test results. The patient report may include clinical and genetic information about a relative when it is relevant to the interpretation of the results. Relatives do not receive an independent analysis of data nor a separate 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.