



CUSTOM FAMILY SEQUENCING REQUISITION

PATIENT INFORMATION (COMPLETE ONE FORM FOR EACH PERSON TESTED)

Patient Last Name		Patient First Name		MI	Date of Birth (MM / DD / YYYY)	
Address		City	State	Zip	Phone	
Accession #	Hospital / Medical Record #		Patient discharged from the hospital/facility: ☐ Yes ☐ No		Genetic Sex: ☐ Female ☐ Male ☐ Unknown Gender identity (if different from above):	

REPORTING RECIPIENTS

Ordering Physician		Institution Name		
Email (Required for International Clients)	Phone	Fax	Client ID	NPI #

ADDITIONAL REPORTING RECIPIENTS

Name	Email	Fax
Name	Email	Fax

PAYMENT (ONLY REQUIRED IF ORDERING TEST CODE 1580)

The payment section is **ONLY** required to complete when ordering test code 1580, Custom Family Sequence Analysis (Billed). Payment information is not needed for test codes 1593, 1594, or 6989.

☐ SELF PAYMENT

☐ INSTITUTIONAL BILLING

Institution Name	Institution Code	Institution Contact Name	Institution Phone	Institution Contact Email
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☐ **INSURANCE**

☐ Do Not Perform Test Until Patient is Aware of Out-Of-Pocket Costs (excludes prenatal testing)

REQUIRED ITEMS	1. Copy of the Front/Back of Insurance Card(s)	2. ICD10 Diagnosis Code(s)	ICD10 Diagnosis Code(s) (Required)
	3. Name of Ordering Physician	4. Insured Signature of Authorization	

☐ Commercial ☐ Medicaid ☐ Medicare Advantage ☐ Medicare Fee-For-Service*

*A completed Advance Beneficiary Notice (ABN) is required for Medicare patients that do not meet Medicare criteria.

Primary Insurance Co. Name	Primary Insurance Co. Phone	Secondary Insurance Co. Name	Secondary Insurance Co. Phone		
Primary Member Policy #	Primary Member Group #	Secondary Member Policy #	Secondary Member Group #		
Name of Insured	Insured Date of Birth (MM / DD / YYYY)	Name of Insured	Insured Date of Birth (MM / DD / YYYY)		
Patient's Relationship to Insured	Phone of Insured	Patient's Relationship to Insured	Phone of Insured		
Address of Insured		Address of Insured			
City	State	Zip	City	State	Zip

STATEMENT OF MEDICAL NECESSITY (REQUIRED)

This requisition hereby incorporates the Terms and Conditions of the Laboratory Services found at <https://www.baylorgenetics.com/lab-terms-conditions/> or, in the case of international entities, <https://www.baylorgenetics.com/terms-conditions-of-the-laboratory-services-international/>. This test is medically necessary for the risk assessment, diagnosis, or detection of a disease, illness, impairment, symptom, syndrome, or disorder. The results will determine my patient's medical management and treatment decisions. The person listed as the Ordering Physician is authorized by law to order the test(s) requested herein. I confirm that I have provided genetic testing information to the patient, and they have consented to genetic testing.

Physician's Printed Name	Physician's Signature	Date (MM / DD / YYYY)
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CUSTOM FAMILY SEQUENCING REQUISITION

Patient Last Name _____ Patient First Name _____ MI _____ Date of Birth (MM / DD / YYYY) _____ Genetic Sex _____

SAMPLE

Date of Collection: _____ **SAMPLE TYPE** _____
_____/_____/_____
MM DD YYYY ☐ Blood ☐ Buccal Swab ☐ Skin Biopsy† ☐ Saliva ☐ DNA (Specify Source): _____ ☐ Cultured Skin Fibroblast

NOTE: Extracted DNA/RNA will only be accepted if the isolation of nucleic acids for clinical testing occurs in a CLIA-certified laboratory or a laboratory meeting equivalent requirements as determined by the CAP and/or the CMS.

CUSTOM FAMILY SEQUENCING INFORMATION

The test codes below are for targeted analysis of a known familial variant(s) for which Baylor Genetics does not have a separate test code. These test codes should only be used when the variant(s) have already been identified in the proband. These test codes should only be used for single nucleotide variants (SNVs) or insertion and deletions (indels) within the nuclear genome. If proband testing was performed at an outside laboratory, contact gc@baylorgenetics.com to discuss coverage prior to sending familial samples.

If the proband was tested at an outside laboratory, a positive control is highly recommended to be provided from the proband if possible. Test code 6989 below should be ordered for their positive control sample.

Name of First Patient Studied _____ Relationship to Patient Studied _____ Baylor Genetics Lab # _____ Family # _____

This Family Member is Currently:

- ☐ **ASYMPTOMATIC**
☐ **SYMPTOMATIC**

☐ Copy of Original Results
Attached (REQUIRED)

If SYMPTOMATIC, please provide details. Please attach additional pages, if needed.

Include a pedigree showing familial relationships.

CUSTOM FAMILY SEQUENCING TESTS

Please select one test code per gene for which targeted sequencing is being ordered:

TEST CODE	TEST NAME	GENE NAME (REQUIRED)	MUTATION/UNCLASSIFIED VARIANT (REQUIRED)
☐ 1580	Custom Family Sequence Analysis (Billed)	 	
☐ 1593	Known Familial Variant*	 	
☐ 1594	Proband Variant Resolution†	 	
☐ 6989	Positive Control Sample for Custom Sequencing	A separate Custom Family Sequencing Requisition should be used for the positive control sample.	

NOTE: Orders that do not meet these requirements may be delayed to clarify the order, or cancelled. Baylor Genetics will contact you if a test change is required.

† Available for test code 1580 only. This sample incurs an additional fee and typically adds 14 days to the turnaround time, depending on sample quality. Baylor Genetics will store this sample for up to 14 days after the report is issued, allowing for follow-up testing if needed.

‡ Please review <https://www.baylorgenetics.com/variant-resolution-testing-policy> for eligibility requirements. Place this order only if a VUS was identified by Baylor Genetics in the proband.

* Please review <https://www.baylorgenetics.com/familial-variant-testing-policy> for eligibility requirements. Place this order only if a pathogenic or likely pathogenic variant was identified by Baylor Genetics in the proband.



INFORMED CONSENT FOR CUSTOM FAMILY SEQUENCING

Patient Last Name

Patient First Name

MI

Date of Birth (MM / DD / YYYY)

Genetic Sex

GENERAL GENETIC TESTING CONSENT

This consent form cannot be used for whole exome sequencing (WES), whole genome sequencing (WGS), biochemical testing, or Huntington disease testing. Consent forms for other tests are located at Baylor Genetics' website (<https://www.baylorgenetics.com/consent/>).

For the purposes of this consent, "I", "my", "you", and "your" can refer to you, your child, your unborn child, or other individual you are the legal representative of.

TEST INFORMATION

Your healthcare provider (doctor, genetic counselor, or other person with medical training) wants to order one or more tests to find a cause for your health issues. This testing can see if there is a cause for your health issues or if there is an increased chance for a health issue to happen to you or your family. Some of these tests look for changes, called variants, in a person's DNA. DNA is our genetic material. You might have testing for variants in one or more genes, specific parts of DNA that are needed for our health. Variants can also be found in other places in the genome (all of the DNA that a person has). Some tests might look for changes in proteins or analytes that cause health issues. The testing ordered will depend on your health issues as well as what is already known about you and your family's genetics. These tests may also explain health issues that your family may have. Even if this test finds the cause of your health issues, this may not help treat or manage those issues.

Before you sign this consent form, you should speak with your healthcare provider. They can help you understand this testing and what it means for your health.

TEST RESULTS

There are several types of test results that may be reported including:

- **Positive:** A variant in the DNA was found that is related to your health issues or a health issue that you are at an increased risk of having in the future. These changes that cause disease are also known as pathogenic variants.
- **Negative:** No variants in the DNA were found that are related to your health issues or that would increase your risk of a health issue in the future.
- **Variant of Uncertain Clinical Significance (VUS):** A variant in the DNA was found that we do not know its effect, if any, on health. More testing may be needed for you or your family if a VUS is found that may be associated with your health issues.
- **Secondary and Incidental Findings:** Testing can sometimes find a variant in the DNA not related to the reason for testing. If this result is expected to affect your health, it is called a secondary or incidental finding.

CONSIDERATIONS AND LIMITATIONS

- You should speak with your provider before signing this consent form to understand the risks, benefits, and alternatives to testing.
- Testing may show you have, or are at increased chance of having, a health issue. It may show that you have an increased chance of having a child with a health issue.
- Even if the variant(s) causing your health issues are found, how these issues might progress or improve with treatment might not be known. Affected family members with the same variant might not be affected like you are.
- Depending on the results of testing, more testing may be needed to understand these results. This testing might be needed for you and/or other family members.
- A negative result does not rule out the chance for health issues. Our knowledge of variants and how they cause disease may change over time as we learn more about genetics. Testing has limitations to what it can find as well.
- Certain factors may lead to incorrect results. These include mislabeled samples, incorrect information in the test order, and rare technical errors.
- More sample may be needed from you if the first sample is not sufficient to complete testing.

PATIENT CONFIDENTIALITY AND SAMPLE RETENTION

- If several family members are tested, knowing the correct biological relationships among them is important. In rare cases, testing can show that family members are not related as expected. If this is found, we may contact the provider who ordered your testing.
- If this testing is requested to be cancelled after the order and sample are sent to the laboratory, please see our Test Cancellation Policy at www.baylorgenetics.com/cancel-test/.
- Only Baylor Genetics and its contracted partners will have access to your sample for the ordered testing. Results from testing will only be released to: (i) a licensed healthcare provider, (ii) those authorized in writing, (iii) the patient or their personal representative, and (iv) those allowed access to test results by law. You have the right to access your test results from Baylor Genetics by providing a written request. You also have the right to request raw data obtained from your sample by providing a written request or HIPAA Authorization Form.
- In rare cases, people with genetic diseases may have problems with health insurance and employment. The U.S. Federal Government has several laws that prohibit discrimination based on test results by health insurance companies and employers. These laws also prohibit unauthorized disclosure of this information. For more information, please visit www.genome.gov/10002077.
- Samples will be kept in the laboratory based on our retention policy. Once testing completes, de-identified sample may be used for test development, quality assurance, and training purposes. Samples are not returned to patients or providers unless requested prior to testing. You and your heirs will not receive payments, benefits, or rights to any resulting products or discoveries.
- The information from your testing may be used in scientific research, publications or presentations, but your specific identity will not be revealed. We may contact your provider to obtain more clinical information about you. Baylor Genetics also performs other types of scientific research and may contact you to see if you would like to be involved.
- Variants found may be submitted to databases. The medical community uses these databases to collect information about how variants might cause disease to improve testing and treatment for patients. An example is ClinVar, a free, public archive of reports on human genetics. Limited clinical information may need to be shared with these databases. In rare cases, this information may be enough to allow you or your family members to be identified.
- For more information on privacy practices at Baylor Genetics, please visit www.baylorgenetics.com/privacy-practices/.



INFORMED CONSENT FOR CUSTOM FAMILY SEQUENCING

Patient Last Name Patient First Name MI ____ / ____ / ____
Date of Birth (MM / DD / YYYY) Genetic Sex

FOR SAMPLES FROM NEW YORK STATE RESIDENTS

Samples from New York State residents shall not be included in research without written consent. Samples will not be retained for more than sixty (60) days after receipt by Baylor Genetics, unless authorized by marking below. No tests other than those authorized shall be performed on the samples.

☐ I authorize Baylor Genetics to retain sample(s) longer based on our retention policy for test development, quality assurance, and training purposes.

FINANCIAL AGREEMENT

By signing below, I hereby authorize Baylor Genetics to provide my insurance carrier any information necessary, including test results, for processing my insurance claim. I understand that I am responsible for any co-pay, co-insurance, and unmet deductible that the insurance policy dictates. I designate Baylor Genetics as my designated representative for purposes of appealing any denial of benefits by my insurance carrier. I irrevocably assign associated payment to Baylor Genetics, and direct that payment be made directly to Baylor Genetics. Please note, some payers may not cover certain screening tests.

If my health insurer does not cover the test or I do not have health insurance, I have received a good faith estimate of the cost for the genetic testing ordered by my provider and agree to pay for the cost of the genetic testing billed to me by Baylor Genetics based on that good faith estimate. More information is available in Baylor Genetics' No Surprises Act and Good Faith Estimate Notice located at <https://www.baylorgenetics.com/no-surprises-act/>.

A Medicare Advance Beneficiary Notice (ABN) is required for services Medicare identifies as not medically necessary.

PATIENT AUTHORIZATION

By signing this statement of consent, I acknowledge that I have read, understand, and hereby grant my informed consent for genetic testing. I have received appropriate explanations from my healthcare provider about the planned genetic test(s) and possible results. I have been informed by my healthcare provider about the availability and importance of genetic counseling and have been provided with written information identifying a genetic counselor or medical geneticist who can provide such counseling services. All my questions have been answered, and I have had the necessary time to make an informed decision about the genetic test(s).

I hereby give permission to Baylor Genetics to conduct genetic testing as recommended by my physician*.

Patient Name Patient's Signature ____ / ____ / ____
Date Signed (MM / DD / YYYY)

Patient's Parent / Personal Representative* Name Patient's Parent / Personal Representative Signature ____ / ____ / ____
Date Signed (MM / DD / YYYY)

*If you are signing on behalf of the patient as the parent(s) and/or person with legal authority to act on behalf of the patient or parent, you may be required to provide evidence of your authority.