

INSURANCE AND PAYMENT INFORMATION

PATIENT INFORMATION

Patient Name (Last, First, Middle):	Maiden Name:	Date of Birth (MM-DD-YYYY):
_____	_____	_____
Social Security #:	Driver's License #:	Phone Number:
_____	_____	_____
Home Address:	City:	State: ZIP Code:
_____	_____	_____
First time here? <input type="radio"/> Yes <input type="radio"/> No	Communication Preference: <input type="radio"/> Mail <input type="radio"/> Phone <input type="radio"/> No Preference	
Smoking Status: <input type="checkbox"/> Current Everyday Smoker <input type="checkbox"/> Current Some Day Smoker <input type="checkbox"/> Former Smoker	<input type="checkbox"/> Light Tobacco Smoker <input type="checkbox"/> Never Smoker	
<input type="checkbox"/> Heavy Tobacco Smoker		
Employer:	Work Address:	Work Phone:
_____	_____	_____
Name of Nearest Relative (other than spouse/partner):	Relationship to Patient:	Phone Number:
_____	_____	_____

SPOUSE/PARTNER INFORMATION

Spouse/Partner Name (Last, First, Middle):	Date of Birth (MM-DD-YYYY):	
_____	_____	
Social Security #:	Driver's License #:	Work Phone:
_____	_____	_____
Employer:	Work Address:	
_____	_____	

INSURANCE INFORMATION

Primary Insurance:	_____		
Address:	_____		
Policy Number:	_____	Subscriber Name:	_____
Secondary Insurance:	_____		
Address:	_____		
Policy Number:	_____	Subscriber Name:	_____

Your signature below confirms that the information submitted above is true and correct to your knowledge and that you have read, understood, and accept our Insurance, Payment, and Other Terms on a separate page.

Patient's Signature: _____ **Date:** _____



GENETIC SCREENING QUESTIONNAIRE

Name: _____ DOB: _____ Ethnic Origin: _____

Partner's Name: _____ DOB: _____ Ethnic Origin: _____

Family and Patient History

- 1. Is your family or your partner's family...
a. Southeast Asian, Taiwanese, Chinese, or Filipino?
b. Italian, Greek, Middle Eastern, East Indian, or Pakistani
c. African or African-American (Black)?
d. Jewish?
e. Cajun or French Canadian?

- 2. Have you, your partner, or anyone in either of your families ever had any of the following disorders?
a. Down syndrome
b. Other chromosomal abnormalities
c. Neural tube defect
d. Cystic fibrosis
e. Bleeding disorder
f. Sickle cell/Thalassemia
g. Tay-Sachs/Canavan
h. Neurofibromatosis
i. Muscular dystrophy
j. Other nerve or muscle disorder
k. Bone or skeletal disorder
l. Polycystic kidney disease
m. Heart defect
n. Cleft lip/palate
o. Developmental delay
p. Any birth defect or genetic condition
q. Needed surgery before age 1 year
r. Cancer in childhood
s. Non-age-related blindness or deafness

- 3. Are you and your partner related by blood (such as cousins)?
4. Have you, your partner, or anyone in either of your families had a baby who died shortly after birth or in childhood?
5. Have you, your partner, or anyone in either of your families had a stillbirth or two or more pregnancy losses?
6. Have you or your partner had any genetic tests (such as chromosomes, cystic fibrosis, Tay-Sachs, or sickle cell screening)?
7. Have you ever been diagnosed with diabetes, cancer, another serious medical condition, or genetic condition?
8. Do you or your partner have a history of infertility?
9. If yes to any question above, please explain:

Pregnancy History

- 1. Was this pregnancy achieved through in-vitro fertilization (IVF) or other assisted reproductive methods?
2. Have you had the California Prenatal Screening Program blood test?
3. Have you had Non-invasive Prenatal Testing (NIPT)?
4. During this pregnancy, have you had any of the following?
a. Cramping, vaginal bleeding or vaginal leakage of fluid
b. Infections, rashes, or other illness, fever over 101
c. X-rays, hospitalizations, or surgery
d. Cigarettes, alcoholic beverages, or "street" drugs
e. Ultrasound ("sonogram")
f. Occupational, chemical, or other exposures
g. Prescription or non-prescription medication
h. Prenatal vitamins

My signature below indicates that the above family and pregnancy history information provided is complete and correct.

Completed by: _____ Date: _____

Reviewed by: _____ Date: _____

Recommended Prenatal Genetic Screening Tests

Patient Name: _____ **Date of Birth:** _____
(MM-DD-YYYY)

The following tests are recommended to be offered to every patient who is planning a pregnancy or who is already pregnant. The testing can determine if the pregnancy is at a higher risk than the general population of having these conditions. Currently, there is not a cure for these conditions prior to or after birth. If the risk of these conditions is higher, you would be offered further diagnostic testing. The purpose of having this information about your developing baby is so that you can prepare yourself to care for a child with special health care needs or choose to not continue an affected pregnancy. Please review the basic information regarding these conditions. If you would like more information or have further questions, your genetic counselor can answer them for you. We can also verify insurance coverage for this testing.

Cystic Fibrosis	<ul style="list-style-type: none"> CF is a life long illness that causes problems with digestion and breathing. CF does not affect intelligence or appearance. In California, all infants are tested for this condition at birth through the State Newborn Screening Program. CF is most common in individuals of European, Caucasian, or Jewish descent. However, CF has been reported in other populations as well. To have a child with CF, both parents must carry a genetic change or mutation. CF carrier screening does not detect all CF carriers. The detection rate varies by specific ethnic group.
<i>I would like more information regarding this testing:</i> <input type="checkbox"/> Yes <input type="checkbox"/> No	
Spinal Muscular Atrophy (SMA)	<ul style="list-style-type: none"> SMA is a serious condition that causes progressive muscle weakness and paralysis. Children with SMA type 1 usually die from respiratory failure by 2 years of age. Other types of SMA are less severe, but are also disabling. SMA affects all ethnic groups equally. The chance of being a carrier is approximately 1/50 in the general population. To have a child with SMA, both parents must carry a genetic change or mutation. SMA carrier screening does not detect all SMA carriers. 6-9% of carriers of SMA are not detected by current technology. The severity of SMA cannot be predicted by genetic testing.
<i>I would like more information regarding this testing:</i> <input type="checkbox"/> Yes <input type="checkbox"/> No	

Continued on next page

Recommended Prenatal Genetic Screening Tests

Continued from previous page

Fragile X syndrome	<ul style="list-style-type: none">• This condition is the most common cause of inherited mental retardation and developmental delay in males. Behavioral abnormalities, including autism spectrum disorder, are also common.• Only females can carry the genetic mutation that causes Fragile X syndrome. Males with Fragile X syndrome are more severely affected than females with this condition.• The incidence in the general population is approximately 1/2000 to 1/4000 live births.
<p><i>I would like more information regarding this testing:</i> <input type="checkbox"/> Yes <input type="checkbox"/> No</p>	

Patient's Signature: _____ **Date:** _____

**I acknowledge that the values and limitations of DNA testing have been explained to me, and I give my consent for testing.*

FOR COUNSELOR USE ONLY

I have received and understand the information on Spinal Muscular Atrophy & Fragile X syndrome.

- I do not want Fragile X carrier testing.
- I want Fragile X carrier testing. *

- I do not want Spinal Muscular Atrophy carrier testing.
- I want Spinal Muscular Atrophy carrier testing. *

Insurance, Payment, and Other Terms

Patient Name: _____ Date of Birth: _____
(MM-DD-YYYY)

AUTHORIZATION TO RELEASE INFORMATION FOR BILLING
I authorize the Genetics Center and its medical affiliates to release any information acquired in the course of my examination and treatment to my insurance company for billing purposes.
AUTHORIZATION TO RELEASE PAYMENT(S) TO GENETICS CENTER
I irrevocably assign and transfer insurance payment(s) directly to the Genetics Center.
INSURANCE ELIGIBILITY
I certify that I am eligible with my insurance company. I understand that if this is not true or if I am not eligible for some or all of the Genetics Center services under the terms of my insurance contract, I am liable for any and all charges for services rendered. Also, if I am not eligible, I agree to pay in full for all services rendered within thirty days of receiving a bill from the Genetics Center.
INSURANCE AND PAYMENT TERMS
I acknowledge that all medical bills are due and payable at the time services are rendered. However, as a courtesy to me the patient, Genetics Center will submit my claim to my insurance company for me. I understand that my insurance coverage is a contract between me and my insurance carrier. If it is my desire to have Genetics Center bill my insurance carrier for these services, I will present my insurance card. I also acknowledge that all co-pays and unmet deductibles are due and must be paid at the time of service . In certain cases, Genetics Center may also require some deposit in advance. If my insurance company pays more than was collected, Genetics Center will promptly reimburse me that amount of the deposit. In some cases, my insurance will only cover a portion of the fees. If I have made an initial payment, it will then be applied to my balance. If Genetics Center does not receive payment from my insurance carrier within 60 days from the date of my service, Genetics Center may look to me for payment in full. A monthly 1.5% service charge will be added to balances over 30 days old, and a \$10 statement fee will be added to balances over 60 days old. The charges for Genetics Center services are ultimately my responsibility.
BENEFITS AND COVERAGE CHECK IS SUBJECT TO CHANGE
Genetics Center cannot accept responsibility for any differences between what was quoted to them by my insurance during their courtesy benefits and coverage check (copay, deductible, etc), and the final benefit determination performed by my insurance when my claim is processed. Therefore, I may owe a different amount than what was quoted to me prior to services.
ACKNOWLEDGEMENT OF INDEPENDENT CONTRACTORS
I acknowledge that some providers involved here are not employees, but are independent contractors, specifically including the NT practitioners, sonographers, and perinatologists.
ACKNOWLEDGEMENT OF POTENTIAL BILLING BY OTHER PROVIDERS
I acknowledge that there could be other providers involved, such as ultrasound, hospital, perinatologist, etc., which will have their own billing.
AUTHORIZATION TO RECEIVE VOICE MESSAGES
I authorize the doctor and/or facility and/or staff to identify themselves as being from Genetics Center when calling to leave a message regarding my appointment, results, or other medical information on any answering device or with another person answering the phone
ACKNOWLEDGEMENT OF NOTICE OF PRIVACY PRACTICES
I acknowledge that I was offered a copy of the Genetics Center's Notice of Privacy Practices.

My signature confirms that I have read, understand, and accept these terms.

Signature of Patient: _____ Date: _____
(or parent if minor)