

Highlighted fields are required.

Name \_\_\_\_\_  
Last First MI

Address \_\_\_\_\_

City State Zip \_\_\_\_\_

Male  Female Date of Birth / /

Home Phone Work Phone

Lab # Hospital #

I have obtained informed consent of the patient (or the patient's authorized representative) for the ordered genetic test(s) in accordance with applicable law.

Physician/Authorized Signature: \_\_\_\_\_

NPI#: \_\_\_\_\_ Taxonomy#: \_\_\_\_\_

Referring Physician (print): \_\_\_\_\_

Genetic Counselor (print): \_\_\_\_\_

Refer to [www.integratedgenetics.com](http://www.integratedgenetics.com) to access informed consent forms for genetic testing.

Date collected: / / Date sent: / / Collected by: \_\_\_\_\_

Pregnancy:  Yes  No Egg donor used:  Yes  No

Date of ultrasound: / / GA on date of US: weeks days

Sex of the fetus if known: Date of LMP: / / GA by LMP: weeks days

**Gestation History**

# of Fetuses:  1  2  >2 (submit separate requisitions)

Gravida \_\_\_\_\_ Para \_\_\_\_\_ SAB \_\_\_\_\_ TAB \_\_\_\_\_

Specimen Type: (check one)

Amniotic Fluid  CVS  PUBS  Other \_\_\_\_\_

Cord blood (specify prior to or after delivery): \_\_\_\_\_

POC/Fetal Tissue (GA: \_\_\_\_\_ tissue origin: \_\_\_\_\_)

POC formalin-fixed paraffin-embedded (FFPE) tissue

Parental blood for \_\_\_\_\_

Hold cells for \_\_\_\_\_

All diagnoses should be provided by the ordering physician or an authorized designee. Diagnosis/Signs/Symptoms in ICD-CM format in effect at Date of Service (Highest Specificity Required)

ICD-CM		ICD-CM		ICD-CM	
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**Indication(s) for Test (check all that apply)**

If ordering Reveal® SNP Microarray please submit Clinical Questionnaire

Advanced maternal age (≥35)  gravida 1  gravida 2+

Abnormal NIPS/NIPT (include report)  Abnormal maternal serum screen

Increased risk for:  NTD  Down Syndrome  Trisomy 18

Other (specify): \_\_\_\_\_

Abnormal fetal ultrasound:

CNS  Heart  Genitourinary

Growth/skeletal  Oligohydramnios  Polyhydramnios

Other/specify ultrasound finding(s): \_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

**Laboratory Test(s) Ordered (\*Reflex Policy on back)**

105  InSight® (FISH for 13,18,21, X and Y)

If InSight is normal, reflex to Reveal® SNP Microarray-Prenatal; if InSight is abnormal, reflex to chromosome analysis

See back  Chromosome analysis

477  Reflex to Reveal® SNP Microarray-Prenatal, if chromosomes are normal

478  Reflex to Reveal® SNP Microarray – POC, if POC/tissue fails to grow

See back  Reveal® SNP Microarray  Direct on amnio, CVS, or POC

add MCC analysis to Reveal® SNP Microarray (send 1 separate tube of blood)

See back  Reveal® SNP Microarray & Abbreviated Chromosome Analysis

451890  Noonan syndrome – prenatal (MCC required - send 1 separate tube of blood)

300  AF-AFP (alpha-fetoprotein)\*

330  AChE (acetylcholinesterase)\*

287  DiGeorge/VCF (FISH)

Parental follow-up to Reveal® SNP Microarray (additional charges may apply)

Test code on original report: \_\_\_\_\_

Other Testing – specify (call before sending) \_\_\_\_\_

Multiple SABs (Spontaneous abortion):  Pregnant  Not Pregnant

Fetal loss/Stillbirth (POC)  <20 wks  >20 wks

Parental chromosome analysis following abnormal prenatal results

Specify \_\_\_\_\_

Confirm prenatal analysis

Clarify abnormal fetal chromosomes - provide results and a copy of the karyotype

Other \_\_\_\_\_

**Family History**

Family History (include copy of report)

Specify relationship \_\_\_\_\_

Chromosome abnormality (specify) \_\_\_\_\_

Genetic disorders \_\_\_\_\_

Autism/Autism spectrum disorders  ID/DD

Birth defects (specify) \_\_\_\_\_

Parent(s) carrier(s) of \_\_\_\_\_

Parent has chromosome rearrangement/mosaicism (specify) \_\_\_\_\_

\_\_\_\_\_

Fetus at risk for \_\_\_\_\_

Other \_\_\_\_\_

Unless testing is clearly ordered as a reflex, all testing will be run concurrently when possible.

**BILLING INFORMATION**

Patient Hospital Status:  Inpatient  Outpatient  Non-hospital

Medicaid  Medicare  Insurance  Client Bill  CA XAFP  Self-Pay

Billing Information Attached (Please include a copy of insurance card or face sheet.)

Do not attach credit card information to this form for security purposes.

Insurance Company Name \_\_\_\_\_

Policy # \_\_\_\_\_ Group # \_\_\_\_\_

Relation to Insured:  Self  Spouse  Child  Other \_\_\_\_\_

Patient Signature \_\_\_\_\_ Date: \_\_\_\_\_

**INTEGRATED GENETICS INTERNAL USE ONLY**

By signing this form, I hereby authorize Laboratory Corporation of America® Holdings (LCAH), its subsidiaries and affiliated companies to furnish my designated insurance carrier the information on this form if necessary for reimbursement. I also authorize benefits to be payable to LCAH.

I understand that I am responsible for any amounts not paid by insurance for reasons including, but not limited to, non-covered and non-authorized services. I permit a copy of this authorization to be used in place of the original.

Bill Codes:	Chromosome Analysis	Abbreviated Chromosome Analysis	Reveal® SNP Microarray
	100 Amniotic Fluid	101 Amniotic Fluid	477 Prenatal
	110 CVS	111 CVS	478 POC
	123 PUBS	124 PUBS	
	180 POC/Fetal Tissue	181 POC/Fetal Tissue	

\*REFLEX POLICY: The following will be performed by reflex at an additional charge. AChE when AF-AFP is elevated &/or gestational age is out of range of normative values. Fetal Hemoglobin when AF-AFP is elevated and amniotic fluid is bloody.