



# Knight Diagnostic Laboratories

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## Constitutional/Prenatal Cytogenetics Requisition

### Patient Information

Full Name   
 Street Address   
 City, State, Zip   
 Phone  DOB  /  /   
 Fax  Male  Female   
 ID/MRN #   
 Hospital In-Patient Yes  No

### Ordering Physician Information

Full Name   
 NPI   
 Office/Facility Name   
 Address   
 City, State, Zip   
 Phone  Fax   
 Account #   
 Notes

Physician Signature \_\_\_\_\_ Date \_\_\_\_\_

### Send additional copies of test results to:

Physician Name   
 Physician Name

Physician Phone  Fax   
 Physician Phone  Fax

### Billing Information - Select One Billing Method

Self Pay

Bill Insurance

Attach Copy of Insurance Card or Billing Face Sheet

Bill Referring Provider or Institution

Invoice will be sent to Client Account and Address Listed Above

Billing is done in accordance with the information provided below and OHSU policy. Appropriate areas must be completed or referring laboratory/physician will be billed.

Primary Insurance Name   
 Primary Policy #   
 Primary Group #   
 Preauthorization #   
 Relation to Insured Medicaid Medicare  
 Self Child Spouse Other \_\_\_\_\_

Secondary Insurance Name   
 Secondary Policy #   
 Secondary Group #   
 Preauthorization #   
 Relation to Insured Medicaid Medicare  
 Self Child Spouse Other \_\_\_\_\_

### Clinical Information

**Specimen Type**

Amniotic Fluid  
 Blood, EDTA  
 Blood, Sodium Heparin  
 CVS  
 Fibroblasts  
 Skin Biopsy  
 Other \_\_\_\_\_

Date of Specimen Collection  /  /   
 Time of Specimen Collection  :

### Pregnancy History

G: \_\_\_\_\_ TAB: \_\_\_\_\_  
 P: \_\_\_\_\_ SAB: \_\_\_\_\_  
 Gestational Age: \_\_\_\_\_  
 Determined By: \_\_\_\_\_  
 WBC: \_\_\_\_\_

**Clinical Information Continued**ICD-10 (required) **REQUIRED:** Reason for Referral Description (fetal abnormalities, provisional diagnosis, family history of chromosome abnormalities, etc):**\*Chromosome Assays**

Code	Test Name	Code	Test Name
6020	Amniotic Fluid: Full Chromosome Analysis	6754	Solid Tissue/Fibroblasts (includes POC): Full Chromosome Study reflexed to FISH
6054	Blood: High Resolution Chromosome Study reflexed to FISH	6500	Chromosomal Microarray
6078	Breakage Analysis: Blood Chromosome Study	6510	**Chromosomal Microarray - Prenatal Diagnosis
6080	Breakage Analysis: Skin Chromosome Study	N/A	**Prenatal FISH With Reflex Cytogenetics, Amnio
6550	SNP Microarray POC	N/A	**Prenatal FISH With Reflex Cytogenetics, CVS
6100	Chorionic Villus Sampling: Full Chromosome Analysis	N/A	**Prenatal FISH With Reflex Cytogenetics, POC

\* Chromosome studies will reflex to FISH if clinically relevant abnormalities are detected; appropriate charges will apply.

\*\* Prenatal studies with microarray testing will need maternal blood collected in EDTA for MCRO testing.

**FISH Assays**

Code	Test Name	Code	Test Name
7018	AneuVysion (chromosomes 13, 18, 21, X and Y)	8756	SHOX-related Haploinsufficiency Disorders, SHOX (Xp22.33)
7020	Angelman Syndrome / Prader Willi (SNRPN/D15S10) (15q11-13)	7750	Smith-Magenis Syndrome
8080	CEP X and CEP Y FISH	8762	SNRPN Dup(15) in autism
8105	Cri-du-Chat (5p-) Syndrome (5p15.2)	8772	SRY-related disorders of sex development, SRY (Yp11.3)
7140	DiGeorge Syndrome (TUPLE1) (22q11.2)	8775	Steroid Sulfatase (STS) (Xp22.3)
8395	Kallman Syndrome (KAL) (Xp22.3)	7870	Velocardiofacial Syndrome (TUPLE1)
7510	Miller Dieker Syndrome, (LIS1)(17p13.3)	7900	Williams Syndrome (ELN) (7q11.23)
7020	Prader-Willi Syndrome (SNRPN/D15S10) (15q11-13)	7920	Wolf-Hirshhorn Syndrome, (WHS) (4p-) (WHSCR) (4p16.3)
8692	RB1/D13S319 (13q14)		

**Non-Testing Services**

Code	Test Name	Code	Test Name
6240	Fibroblast Primary Culture	N/A	Primary Culture Sendout

**Additional Comments**

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**Result Release**

**Results will be immediately available to the patient unless you mark the box below**

Do not release (I reasonably believe that an Information Blocking exception applies)

Comments / Requests: