



Knight Diagnostic Laboratories

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 Questions? (855) 535-1522

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

Primary Insurance Name
 Primary Policy #
 Primary Group #
 Preauthorization #

Relation to Insured Medicaid Medicare
 Self Child Spouse Other _____

Bill Referring Provider or Institution

Invoice will be sent to Client Account and Address Listed Above

Secondary Insurance Name
 Secondary Policy #
 Secondary Group #
 Preauthorization #

Relation to Insured Medicaid Medicare
 Self Child Spouse Other _____

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.

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 ContinuedICD-10 (required)

Clinical Diagnosis 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	6100	Chorionic Villus Sampling: Full Chromosome Analysis
6054	Blood: High Resolution Chromosome Study reflexed to FISH	6754	Solid Tissue/Fibroblasts (includes POC): Full Chromosome Study reflexed to FISH
6078	Breakage Analysis: Blood Chromosome Study	6500	Chromosomal Microarray
6080	Breakage Analysis: Skin Chromosome Study	6510	Chromosomal Microarray - Prenatal Diagnosis
6550	SNP Microarray POC		

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

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-Hirschhorn 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	Freeze and Store Cells
6242	Fibroblast Retrieval	N/A	Primary Culture Sendout