

PRENATAL/TISSUE TEST REQUISITION FORM



Cytogenetic Laboratories

Indiana University School of Medicine
975 W. Walnut, IB 350, Indianapolis, IN 46202
317/274-2243 (Office) 317/278-1616 (Fax)

317/274-2243 or 317/274-2246 (Lab)

CAP#: 16789-30 CLIA#: 15D0647198

Cytogenetic Lab Use Only

1) PHYSICIAN(S): **FOR LABORATORY USE ONLY:**

Referring Physician: _____
 Institution: _____
 Phone/Pager: _____ Fax: _____

Genetic Counselor: _____
 Phone/Pager: _____ Fax: _____

Additional Recipients: _____
 Phone/Pager: _____ Fax: _____

Date Received: ____/____/____ Family #: _____
 Time Received: ____:____ am/pm Proband: Yes No
 Received By _____

AM: CV: TI: CMA: CMA+5:
 AneuVysion FISH:
 Handling Charge: x _____ Handling **ONLY**:

Lab Comment(s): _____

2) PATIENT/CLINICAL INFORMATION

(Fetus of) _____ Sex: Male Female
 Patient Last Name Patient First Name Date of Birth (MM/DD/YY)

Address: _____ Hospital: _____ MRN: _____

Date of Collection (MM/DD/YY) ____/____/____ Time: _____ Estimated Gestational Age (EGA) _____ By LMP or Ultrasound

Was pregnancy achieved through ART? If so, how: Egg Donor Sperm Donor IVF ICSI

Sample Type:

Amniotic Fluid _____ cc Products of Conception (POC): Villi Placenta Fetal, Source: _____ Other: _____

CVS _____ mg (TA TC) Skin Biopsy (Non-fetal), Source: _____ Other, Describe: _____

Maternal Blood for MCC (3-5cc in EDTA tube, **Required** for prenatal CMA) _____ cc

Paternal Blood (3-5cc in EDTA tube, Requested for prenatal CMA) _____ cc Paternal Name/Date of Birth _____

3) REFERRING DIAGNOSES (please check all that apply):

Advanced maternal age

Abnormal NIPT (attach report): T13 T18 T21 MONX Other _____

Abnormal maternal serum screen: T13 T18 T21 MONX Other _____

Abnormal ultrasound, describe: _____

Family history of chromosome abnormality, describe: _____

Fetal demise Recurrent spontaneous abortion/miscarriage

Parental concern ICD10 Code/Other: _____

PATIENT LABEL

4) REQUESTED TESTING (please check all that apply):

Amniotic Fluid and CVS Test Options:

Chromosome analysis

Chromosomal Microarray Analysis (CMA) with abbreviated chromosome analysis

Maternal cell contamination studies included: Maternal blood sample (3-5cc in EDTA tube) required.

Aneuploidy FISH (13, 18, 21, X, Y). Chromosomes or CMA required.

AFP (amniotic fluid only)

ACHE reflex (amniotic fluid only)

Save cultured cells for send-out testing

Products of Conception (POC)/Skin Biopsy Test Options:

Chromosome analysis

Chromosomal Microarray Analysis (CMA) (POC only)

Chromosome analysis with Reflex to CMA (POC only)

Chromosome analysis to rule out mosaicism (skin biopsy only)

Save cultured cells for send-out testing

Maternal Cell Contamination Studies (RECOMMENDED for POC CMA):
 Maternal blood sample (3-5cc in EDTA tube) required

6) SPECIMEN COLLECTION REQUIREMENTS

Specimen	Collection	Container(s)	Instructions
Amniotic Fluid	20-25 mL of fluid at ≥ 16 weeks of gestation (30 mL if additional studies are ordered). Discard first 2-3 mL to avoid maternal cell contamination. Place remaining fluid in 3-4 aliquots, labeled 1 st , 2 nd , etc.	Sterile Corning centrifuge tubes can be provided upon request. For bloody specimens, use Dark Green-top sodium heparin tubes. These tubes are also available upon request by calling the lab.	Refrigerate. Do not centrifuge.
Chorionic Villus (CVS)	20-30 mg (50 mg if additional studies are ordered).	Transport media will be provided upon request by calling the lab.	Refrigerate.
Products of Conception (POC)	5-10 mm ³ Villi from the placenta is the preferred sample type. Fetal cartilage, membranes, and tendon will also be accepted.	Transport media will be provided upon request by calling the lab. If not available, use a sterile screw-top container with sterile media.	Refrigerate. Do not send entire fetus.
Skin Biopsy (non-fetal)	Skin punch or surgery skin specimen	Transport media will be provided upon request by calling the lab. If not available, use a sterile screw-top container with sterile media.	Refrigerate.

7) SPECIMEN HANDLING REQUIREMENTS

- Collect all specimens aseptically.
- **Do not freeze any specimen type.**
- **Do not place specimens in formalin or any other fixative.**
- Keep all specimens refrigerated until transport. **Specimens should be received within 24 hours of collection.**
- Label all containers and requisition forms with patient name, MRN, date of collection, and physician name.
- MATERNAL samples are REQUIRED to accompany the fetal sample for prenatal CMA. Paternal specimens are requested.
- MATERNAL samples are RECOMMENDED to accompany the fetal sample for POC CMA.

Shipping Instructions:

- Please enclose the cytogenetic test requisition and consent form with the sample. All required information must be completed before sample can be processed.
- Send specimens in a plastic zipper biohazard bag and place paperwork in the side pocket of the bag or in a separate bag or envelope. **Do not place paperwork in the same bag as the specimen.**
- Place specimen bag in Styrofoam mailer and ship at room temperature by overnight Federal Express to arrive Monday through Friday. Saturday delivery is available. Please call for specific information and instructions for Saturday delivery.
- Outside of Indianapolis, please ship by FedEx or other overnight courier to:
IU Cytogenetic Laboratories
Department of Medical and Molecular Genetics
975 W. Walnut St.
IB 350
Indianapolis, IN 46202
- Please call the laboratory and alert us that a specimen is en route: (317) 274-2243

8) PATIENT BILLING INFORMATION:

Bill Patient's Insurance: Policy #: _____ Group #: _____

Insurance/Managed Careplan: _____

Street Address: _____ City: _____ State: _____ Zip: _____

Relationship to Insured: Self Spouse Other: _____ Insured's Social Security #: _____

OR Copy of patient's insurance card attached

Bill Medicare: _____

Bill Medicaid: _____

Bill Patient/Self-Pay (*Please Attach Patient Demographic Sheet*)

Bill Hospital: _____