

2015 PRENATAL DIAGNOSIS REFERRAL FORM FOR CHROMOSOME ANALYSIS

Medical Genetics – Wake Forest University School of Medicine – Winston Salem NC

www.wfubmc.edu/medicalgenetics

Phone: 336-716-4321 Fax: 336-716-2554

Collection Date: _____ **Time:** _____ am/pm **WFU LAB #:** _____

Name: _____ / _____ / _____ / _____
(Please print) Last First Middle Maiden

Address: _____ / _____ / _____ / _____ **Daytime Phone:** (____) _____
Mailing Address City State Zip

Birth Date: _____ **SS# :** _____ **Patient's Mother's first name:** _____

Hospital : _____ **Hospital/Unit #:** _____ **Race:** _____

Type of Specimen: Amniotic Fluid CVS PUBS Tissue Blood Other: _____

COLLECTION TECHNIQUE: AF- Discard the first 2 cc of fluid. Draw 20 -30 ml of fluid. CVS: >20mg tissue.
 PUBS: 1-2 mls in a green stoppered sodium heparin vacutainer. **KEEP ALL SAMPLES AT ROOM TEMPERATURE**

Physician/Provider Order	Statement of Financial Responsibility
Physician: Last, First / Phone/beeper	I authorize any holder of medical or other information about me to release to my healthcare provider, third party processor, the Centers for Medicare and Medicaid Services or its intermediaries or carriers any information needed for this health care encounter or related claim. I permit a copy of this authorization to be used in place of the original, and request payment of authorized insurance benefits be made on my behalf to the WFU Physicians. I understand I am responsible for payment of these charges. I am also responsible for payment if my insurance carrier decides this is a non-covered service or requires prior authorization, which I did not obtain.
1. _____	
X Physician Signature Required	
2. _____	
Tap by: _____	Patient Signature: _____ Date: _____

Billing Information

Bill: Forsyth Novant Moses Cone Hospital CFMFC
 Solstas Women's Hospt of Greensboro Wesley Long Hospital Other : _____
 Medicare # _____ Medicaid #: _____ Carolina Access# _____
 Insurance: _____ Employer: _____ Policy #: _____
(Enclose copy of both sides of insurance card)

OBSTETRIC / PATIENT INFORMATION

G _____ **P** _____ **A** _____ [**SAB** _____ **TAB** _____] **Gestation (wks):** LMP _____ or U/S _____

Genetic Counselor: _____ **Does patient want to know sex of the fetus? :** Yes / No

SIGNS/SYMPTOMS/INDICATION (ICD-9 CODES) FOR CHROMOSOME STUDY

Indicate all that apply. Codes here do not represent entire listing of ICD-9 codes available. Please consult current ICD-9 codebook for complete listing.

Advanced Maternal Age (659.63) _____ Abnormal findings in amniotic fluid NOS (792.3)

Suspected Fetal chromosome abnormality (655.13) Habitual aborter w/o current pregnancy (629.9)

Fetal abnormality antepartum (655.93) Pregnancy with history of abortion (V23.2)

Inc. Down Syndrome Risk {AFP/hCG/uE3} (796.5) → → → { } by WFUSM { } by outside lab: _____

Inc. Trisomy 18 Risk {AFP/hCG/uE3} (796.5) → → → { } by WFUSM { } by outside lab: _____

Neural Tube Defect Risk {Elev. MSAFP}(796.5) → → → { } by WFUSM { } by outside lab: _____

Abnormal abdomen Ultrasound (793.6)- specify: _____

Family history of genetic/chromosome disorder (V19.8) -specify: _____

Other Clinical/ICD-9 code specify: _____

Test Requested Note: When ordering tests for which Medicare reimbursement will be sought, it is recommended that the Provider consult any Local Medical Review Policies (LMRP) or National Coverage Decisions (NCD) that may be applicable to the test(s) being ordered. Based on guidance issued in either of these policies it may be necessary to obtain an Advanced Beneficiary Notice (ABN) from the Medicare Patient. For Medicaid and other carriers a signed **Statement of Financial Responsibility** from the patient may be necessary. (See Statement of Financial Responsibility at top of form.)

TEST(s) REQUESTED	FISH Specific Probes
<input type="checkbox"/> Routine chromosome / karyotype + AF-AFP + ACHe <small>(88235, 88269, 88280, 88285;82105;82013)</small> <input type="checkbox"/> add Microarray	<input type="checkbox"/> Prader-Willi 15q12 <input type="checkbox"/> Sex - X&Y
<input type="checkbox"/> Routine chromosome / karyotype + AneuVysion FISH (13/18/21/X/Y) <small>(88235, 88269, 88280, 88285, 88271x5, 88274 ;82105;82013)</small> + AF-AFP + ACHe	<input type="checkbox"/> DiGeorge/VCF 22q11 <input type="checkbox"/> SRY Yp
<input type="checkbox"/> Routine chromosome / karyotype + specific FISH → → →	<input type="checkbox"/> STS Xp22.3 <input type="checkbox"/> KAL Xp
<input type="checkbox"/> Culture and freeze cells + AF-AFP + ACHe	<input type="checkbox"/> Angelman 15q11
<input type="checkbox"/> AF-AFP <input type="checkbox"/> ACHe <input type="checkbox"/> DNA testing: _____	<input type="checkbox"/> Miller Dieker 17p13 <input type="checkbox"/> other _____
<input type="checkbox"/> Culture & Hold cells per physician <input type="checkbox"/> PCR Testing for:infec /micro/viral/etc _____	

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Collection Date: _____ Time: _____ am/pm WFU LAB #: _____

CYTOGENETIC LAB USE ONLY

Name: _____ Lab #: _____
last first middle maiden

Date Received: _____/_____/_____ Time Received: _____

Sample Type: Amniotic Fluid CVS PUBS

Fluid appearance: clear cloudy bloody brown green clotted

Amount of fluid: _____ mls Size of pellet: tiny small medium large

Number of Tubes: 1 2 3 4

Additional Specimen Evaluation: _____

Primary Cultures: A B C D

Date culture initiated: _____/_____/_____ Tech: _____

Media: Amnio Max other: _____

SENT OUT: to referring institution / # flasks sent: _____ flasks frozen down / # flasks sent: _____

REPORT OF RESULTS / SPECIMEN SUMMARY

Final Preliminary Read Back Date _____ Tech _____

To: _____

KARYOTYPE: 46,XY 46,XX Sex: Yes or No

INTERPRETATION: normal male normal female

abnormal: _____

AF on GEL

Additional Studies / Results: NOR C-band R-band

FISH: normal male normal female
 abnormal +13 +18 +21 +/-X/Y Other: _____

To: _____ By: _____ Date _____