2016 PERIPHERAL BLOOD REFERRAL FORM FOR CHROMOSOME ANALYSIS 2016

Medical Genetics – Wake Forest University School of Medicine – Winston Salem NC
www. wfubmc.edu/medical genetics **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# :** _____ **Sex:** ☐ male / ☐ female

Hospital : _____ **Hospital/Unit #:** _____ **Patient's Mother's first name:**_____

Type of Specimen: ☐ **Blood** ☐ **PUBS** ☐ **Blood (POC)** ☐ **Cancer**

Collection Technique: 5 – 10 mls blood collected in a green stoppered sodium heparin tube. **Keep at room temperature**

Physician/Provider Order Physician: Last, First / Phone / Beeper	
1.	
X	
X Physician Signature Required	
2.	
3.	

Statement of Financial Responsibility

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.

Patient Signature: _____ **Date:** _____

Billing Information

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

SIGNS/SYMPTOMS/INDICATION (ICD-10 CODES) FOR CHROMOSOME STUDY

Indicate **all** that apply. **Codes do not represent entire listing of ICD-10 codes available**. Please consult current ICD-10 code book for complete listing.

☐ Other chromosome conditions (Q99.9)
☐ Other sex chromosome anomaly (Q97.8)
☐ Ambiguous genitalia (Q56.4)
☐ Multiple congenital anomalies (Q89.7)
☐ Unspec anom congen anomaly (Q89.9)
☐ Other Specified anomalies (Q87.89) congenital
☐ Developmental delay (F81.9) NOS
☐ Delayed learning (F81.81) NOS
☐ MR Moderate (F71) ☐ Severe (F72) ☐ Profound (F73)
☐ Infertility -> ☐ female (N97.8) NOS / ☐ male (N46.9) NOS)
☐ Multiple Miscarriages (habitual abortion) {G ____ P ____ A ____} [SAB ___ TAB ____]

☐ Anomalies face/neck (Q18.8)
☐ Eye anomaly (Q15.9) NOS
☐ Cleft lip/palate (Q37.9) NOS
☐ Hypotonia, cong. (P94.2)
☐ Obesity (E66.9)
☐ Delay in sexual development (E30.0)
☐ Abnormal ears (Q17.8)
☐ Short Stature (R62.52)

☐ Lack of nml physiol dev (R62.50)
☐ Brain anomalies (Q04.8) NOS
☐ Failure to thrive (R62.51)
☐ Mixed develop disorder (F82) NOS
☐ Other spec cond. perinatal (P96.89)
☐ Fam hx other cong anomal (Z82.79)
☐ Family hx, mental retard (Z81.0)
☐ Fam hx other conditions (Z84.89)
☐ Congen heart anomaly (Q24.9)
☐ Language Delay (F80.1)

☐ **Additional Clinical Information/ICD-10 codes**:_____

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.)

SUSPECTED DIAGNOSIS OR CHROMOSOME ABNORMALITY *(not for billing purposes)*

☐ Trisomy 21 (Down Syndrome)
☐ Trisomy 13
☐ Trisomy 18
☐ Turner Syndrome (45,X)
☐ Habitual Aborter
☐ Prenatal confirmation
☐ Prader Willi Synd. (15q-)
☐ Angelman Synd. (15q-)
☐ Smith Magenis Synd. (17p-)
☐ Di-George/ VCF Synd. (22q-)
☐ Miller-Dieker Synd. (17p-)
☐ Klinefelter Syn. (47,XXY)
☐ Mental Retardation ***(for Fragile X Syndrome – purple top also required)***
☐ Other genetic/chromosome abnormality: _____

TEST REQUESTED

☐ **Routine Chromosome / karyotype** *(7 - 10 days)*
(88230, 88262, 88280, 88285)
☐ **STAT Chromosomes** *(48 hrs. - additional charges)*
(88230 88262,88280, 88285, 88261)
☐ **Chromosome / karyotype +Stat FISH (*13/18/21/X/Y*)**
(88230, 88262, 88280, 88285,88271x5, 88274x5)
☐ **Routine Chromosome / karyotype +FISH** **select → →**
☐ **FISH Only - chromosome previously done** **select → → →**
☐ **Culture Only – Microarray testing**

FISH Specific Probes

☐ Angelman **15q11** ☐ +13 ☐ +18 ☐+21
☐ Prader-Willi **15q12** ☐ Sex - X&Y
☐ DiGeorge/VCF **22q11.2** ☐ SRY **Yp11.3**
☐ Miller-Dieker **17p13** ☐ 4p-
☐ STS **Xp22.3** ☐ 5p-
☐ Kallmann **Xp22.3** ☐ _____
☐ Smith-Magenis **17p11** ☐ Subtelomere
☐ Williams **7q11** ☐ SHOX **Xp22.3**

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

CYTOGENETIC LAB USE ONLY

Name:_____ **Lab #:** _____

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

Additional Specimen Evaluation:_____

_____

Additional Samples Received: [ ] DNA: _____ [ ] FRAX [ ] FISH [ ] other_____

LABORATORY REPORT SUMMARY

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

Sample type: [ ] PB [ ] FISH [ ] FRAX [ ] PUBS [ ] BM stat

Culture: [ ] 48h. [ ] 72h. col. [ ] 72h.EtBr [ ] DBM [ ] 24h [ ] 96h.

Media: [ ] RPMI 1640 less folic acid [ ] RPMI 1640

REPORT OF RESULTS / SPECIMEN SUMMARY

[ ] Final [ ] Preliminary [ ] Read Back Date_____Tech_____

To:_____

KARYOTYPE: [ ] 46,XY [ ] 46,XX

INTERPRETATION: [ ] normal male [ ] normal female

[ ] Abnormal:_____

_____

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

_____

FISH: [ ] normal male [ ] normal female

[ ] +13 [ ] +18 [ ]+21 [ ] -X [ ] +X [ ] +Y

[ ] Williams 7q22
[ ] DiGeorge/VCF 22q11
[ ] Prader-Willi 15q12
[ ] Angelman 15q12
[ ] Smith-Magines 17p11
[ ] Miller-Dieker 17p13
[ ] STS Xp22.3
[ ] chromosome paint #_____

To:_____By:_____ Date_____