

## RUTGERS CLINICAL GENOMICS REQUISITION: **PEDIATRIC Precision Medicine**

<b>Patient Information</b>			
Name: _____ <small style="display: flex; justify-content: space-between; width: 100%;"> <span>LAST NAME</span> <span>FIRST NAME</span> <span>MI</span> </small>			
Medical Records#: _____ Date of Collection: ____/____/____ <small style="display: flex; justify-content: space-between; width: 100%;"> <span>MM</span> <span>DD</span> <span>YY</span> </small>			
Sex: <input type="checkbox"/> Female <input type="checkbox"/> Male Sample Type: <input type="checkbox"/> Blood <input type="checkbox"/> Amniotic Fluid <input type="checkbox"/> POC <input type="checkbox"/> Skin Biopsy <input type="checkbox"/> Other(Specify) _____			
<b>Reporting Information</b>			
Physician: _____ Institution: _____ Phone: _____ Email/Fax: _____	Additional Professional Report Recipients Name: _____ Institution: _____ Phone: _____ Email/Fax: _____		
<b>Indication for Study</b>			
<b>Pediatric Precision Medicine Testing (Orderable in EPIC)</b> <small>*Purple/EDTA vacutainer except for Chromosomal analysis</small>			
<input type="checkbox"/> NEURO	Comprehensive Neurology Sequencing Panel	<input type="checkbox"/> MEDEX	Medical exome
<input type="checkbox"/> CGH	SNP/CNV Microarray Analysis, Constitutional	<input type="checkbox"/> EPI	Epilepsy and Seizure Disorder Panel
<input type="checkbox"/> CBR	Chromosome Analysis, Constitutional <small>*Dark Green/Heparin tube</small>	<input type="checkbox"/> AUT	Autism and Intellectual Disability Panel
<input type="checkbox"/> IRDP	Inherited Recessive Disease Panel	<input type="checkbox"/> MITO	Mitochondrial DNA Sequencing
<b>Single Gene Tests (Orderable in EPIC)</b> <small>*Purple/EDTA vacutainer</small>		<b>FISH Tests (Orderable in EPIC)</b> <small>*Dark Green/Sodium Heparin Vacutainer</small>	
<input type="checkbox"/> DMD	Deletion/duplication for DMD	<input type="checkbox"/> FINT03	Trisomy 21 (LSI 21), 21q22
<input type="checkbox"/> FXMO	Fragile X FMR1 molecular analysis	<input type="checkbox"/> FINT201	Sex Determination, CEPX/CEPY
<input type="checkbox"/> CF2000	CFTR Gene Sequencing	<input type="checkbox"/> FINT01	Trisomy 13 (RB1), 13q34
<input type="checkbox"/> SC	Sickle Cell Disease	<input type="checkbox"/> FINT02	Trisomy 18 (D18Z1), 18cen
<input type="checkbox"/> MYO	Myotonic Dystrophy (DM1)	<input type="checkbox"/> FINT03	Trisomy 21 (LSI 21), 21q22
<input type="checkbox"/> SMA	Spinal Muscular Atrophy (SMN1 & SMN2)		
<input type="checkbox"/> HCR	Hereditary Hemochromatosis		
Other Tests (Please Specify)			
<b><u>PARENTAL STUDIES</u></b>			
Name of Proband: _____		DOB: _____ MR# _____	
Testing Requested: _____			
Name of Parent (MOTHER): _____		DOB: _____ MR# _____	
Sample submitted: <input type="checkbox"/> SALIVA <input type="checkbox"/> BLOOD <input type="checkbox"/> BUCCAL			
Name of Parent (FATHER): _____		DOB: _____ MR# _____	
Sample submitted: <input type="checkbox"/> SALIVA <input type="checkbox"/> BLOOD <input type="checkbox"/> BUCCAL			

# PEDIATRIC Precision Medicine

## Guidelines for Sample Collection and Submission

### Prior to Collection

#### Order Testing:

- **EPIC**
  - Order testing in EPIC. Simultaneously, submit the filled in paper requisition form.
  - If testing does not appear – please order as an MSO and clearly type the test requested in the paper requisition form with test code- with a notation in EPIC “please send to Clinical Genomics lab at MSB F656”
- **Patients:**
  - All patients must be identified positively, by a minimum of two patient identifiers prior to specimen collection for clinical testing.
  - Patients with a recent history of blood transfusion (<30 days) are not able to submit whole blood specimens, but can submit saliva samples.

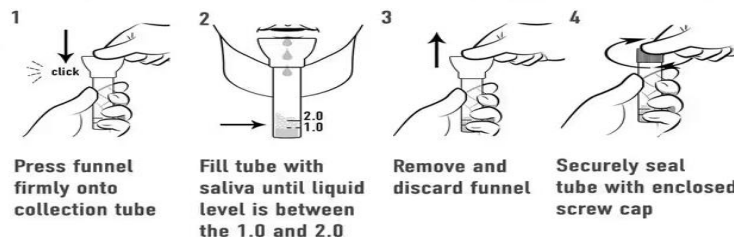
### Collection of Specimens

#### **Peripheral blood**

- 3-5cc of whole blood in Lavender/EDTA vacutainer
  - minimum blood volumes of 0.5cc. for newborn/pediatric patients.
  - Label tube with Patient Name, DOB, Date of collection and Medical record Number. Place the labeled specimen tube into the biohazard bag with the appropriate intake form

#### **Saliva**

- Please contact the laboratory for saliva collection devices, to be provided upon request. All kits are to be stored according to the manufacturer’s requirements and only to be used before the expiration date.



- **Steps**
  - Hold the tube and remove the stopper.
  - Place funnel into tube.
  - Check tube indicator to fill with saliva until this mark
  - Expel saliva into tube until mark
  - Remove and discard the funnel, replace the stopper
  - Label tube with Patient Name, DOB, Date of collection and Medical record Number. Place the labeled specimen tube into the biohazard bag with the appropriate intake Form.

### Specimen Submission

- Send the blood or saliva samples to UH C107 Accessions lab or contact the Rutgers Clinical Genomics (973-972-3170) to arrange pick up of the specimen.
  - If the specimen cannot be picked up same day, please keep saliva specimen at room temperature until send for accessioning. Please store blood at 2-6C or at room temperature.