

**REFERRAL FORM FOR WHOLE EXOME SEQUENCING**

Child's Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Parents/Guardians' Names: \_\_\_\_\_

Contact Phone Number(s): \_\_\_\_\_

**This form is confidential and will become part of the patient's medical record.**

Referring MD: \_\_\_\_\_  
Phone: \_\_\_\_\_  
Fax: \_\_\_\_\_

Primary Care Physician: \_\_\_\_\_  
Phone: \_\_\_\_\_  
Fax: \_\_\_\_\_

**To refer a patient for whole exome sequencing, please complete the following:**

- **Complete this 2 page form in its entirety**
- **Provide printed copies of the patient's pertinent medical records:**
  - **Most recent clinic note from any specialists**
  - **Radiology reports**
  - **Previous laboratory studies (e.g. metabolic and biochemical testing, CSF studies)**
    - **Note: please do not send copies of routine labs such as CBCs and CMPs unless you believe they are relevant to the patient's underlying diagnosis.**
  - **Other diagnostic studies (e.g. EMG, EEG, muscle biopsy, pathology)**
  - **Copies of all previous genetic testing reports**
- **A copy (front and back) of the patient's current insurance card(s)**

**Please note that we are unable to schedule the patient for whole exome sequencing until we have received all of the above in addition to this form. These may be faxed to our office at (314) 454-2075, Attn: Genetic Counselors, or delivered to our office by hand or campus mail (Marisa Vineyard, Campus Box 8116).**

☐ Please check this box to indicate that you are requesting a consultation for whole exome sequencing for this patient and sign below.

\_\_\_\_\_  
Referring physician signature

\_\_\_\_\_  
Date

**For Genetics office use only:**

- ☐ *Form completed*
- ☐ *Printed records received*
- ☐ *Insurance card*
- ☐ *Reviewed by GC/MD*
- ☐ *Insurance BI submitted*

- ☐ *Insurance approval received*
- ☐ *Family notified of out of pocket*
- ☐ *Applied for financial assistance program*
- ☐ *Appointment scheduled*
- ☐ *Both parents to attend the appt*

*Out of pocket amount: \$\_\_\_\_\_*

### **Basic Information**

Have you already discussed whole exome sequencing with the family? ☐ Yes ☐ No

Will both biological parents be available to give blood samples? ☐ Yes ☐ No

If not, please explain: \_\_\_\_\_

If not, does the patient have any full siblings or other family members who might be available to provide a sample? ☐ Yes ☐ No If yes, please list: \_\_\_\_\_

### **Clinical Information**

Please list your patient's main features and most pertinent clinical history, including a summary of any relevant test results:

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Do you have any specific genetic diagnoses or genes in mind for this patient? ☐ Yes ☐ No

If yes, please list: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

### **Prior Genetic Testing – Please provide copies of all genetic test reports**

Has the patient had chromosome microarray (CMA)? This is *required* prior to exome sequencing, except in exceptional circumstances.

- ☐ Yes, and the results were normal. Please list the year CMA was performed: \_\_\_\_\_
- ☐ Yes, but the findings did not explain the patient's phenotype.
- ☐ No, I do not feel it is indicated. Please state reason: \_\_\_\_\_
- ☐ No, the patient's insurance would not cover CMA.

For each genetic test below, please check the box if the test was completed and list the results, if abnormal.

<u>Test</u>	<u>Abnormal results, including variants of uncertain significance</u>
<input type="checkbox"/> Karyotype/chromosomes	_____
<input type="checkbox"/> Fragile X	_____
<input type="checkbox"/> Epilepsy panel	_____
<input type="checkbox"/> Prader-Willi/Angelman syndrome	_____
<input type="checkbox"/> Rett/atypical Rett testing	_____
<input type="checkbox"/> X-linked intellectual disability panel	_____
<input type="checkbox"/> Other genetic tests:	_____
	_____
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