Mental Retardation, Stereotypic Movements, Epilepsy, and/or Cerebral Malformations: **MEF2C** Gene Deletion/Duplication

**Test Code:** DMEF2  
**Turnaround time:** 2 weeks  
**CPT Codes:** 81228 x1

### Condition Description
Le Meur et al. (2010) describe six unrelated children with a deletion or mutation of the **MEF2C** gene (5q14). All six patients had a similar phenotype including severe intellectual disability, developmental delay, hypotonia, absent speech, and the inability to walk unaided. Variable features include stereotypic movements, epilepsy and/or cerebral malformations, and dysmorphic features. Mutation of the **MEF2C** gene causes autosomal dominant mental retardation, stereotypic movements, epilepsy, and/or cerebral malformations.

For patients with suspected mental retardation, stereotypic movements, epilepsy, and/or cerebral malformations, sequence analysis is recommended as the first step in mutation identification. For patients in whom mutations are not identified by full gene sequencing, deletion/duplication analysis is appropriate.

### References:
- OMIM #600662: **MEF2C** gene
- OMIM #613443: Mental retardation, stereotypic movements, epilepsy, and/or cerebral malformations

### Genes
**MEF2C**

### Indications
This test is indicated for:
- Confirmation of a clinical diagnosis of mental retardation, stereotypic movements, epilepsy, and/or cerebral malformations in an individual in whom sequence analysis was negative.
- Carrier testing in adults with a family history of mental retardation, stereotypic movements, epilepsy, and/or cerebral malformations in whom sequence analysis was negative.

### Methodology
DNA isolated from peripheral blood is hybridized to a CGH array to detect deletions and duplications. The targeted CGH array has overlapping probes which cover the entire genomic region.

Please note that a "backbone" of probes across the entire genome are included on the array for analytical and quality control purposes. Rarely, off-target copy number variants causative of disease may be identified that may or may not be related to the patient's phenotype. Only known pathogenic off-target copy number variants will be reported. Off-target copy number variants of unknown clinical significance will not be reported.

### Detection
Detection is limited to duplications and deletions. The CGH array will not detect point or intronic mutations. Results of molecular analysis must be interpreted in the context of the patient's clinical and/or biochemical phenotype.

### Specimen Requirements
Submit only 1 of the following specimen types

* Preferred specimen type: Whole Blood

**Type: Whole Blood**

Specimen Requirements:

- In EDTA (purple top) or ACD (yellow top) tube:
  - Infants (2 years): 3-5 ml
  - Older Children & Adults: 5-10 ml

Specimen Collection and Shipping: Refrigerate until time of shipment. Ship sample within 5 days of collection at room temperature with overnight delivery.

**Type: Saliva**

Specimen Requirements:
Oragene™ Saliva Collection kit (available through EGL) used according to manufacturer instructions.

Specimen Collection and Shipping: Store sample at room temperature. Ship sample within 5 days of collection at room temperature with overnight delivery.

**Special Instructions**

Sequence analysis is required before deletion/duplication analysis by targeted CGH array. If sequencing is performed outside of EGL Genetics, please submit a copy of the sequencing report with the test requisition.

**Related Tests**

- Sequence analysis of the *MEF2C* gene is available and is required before deletion/duplication analysis.
- Custom diagnostic mutation analysis (KM) is available to family members if mutations are identified by targeted mutation testing or sequencing analysis.
- Prenatal testing is available only for known familial mutations to individuals who are confirmed carriers of mutations. Please contact the laboratory genetic counselor to discuss appropriate testing prior to collecting a prenatal specimen.