

UW Cytogenetic and Molecular Genetic Laboratories

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Test Menu Updates

The UW Cytogenetic and Molecular Genetic Services is consolidating our microarray testing menu. Effectively immediately, we will offer the following microarray testing options:

Germline Microarray Analysis

SLH Test Code 890: Provides high resolution, genome-wide assessment of copy number variants (CNVs) and absence of heterozygosity (AOH). Postnatal (whole blood) and prenatal (chorionic villi, amniotic fluid, products of conception) specimen types accepted.

Targeted Microarray Analysis (Family studies)

SLH Test Code 890FAM: Provides analysis of previously characterized familial CNVs and/or regions of homozygosity (ROH). This test may also be used to confirm CNVs or ROH identified in a research laboratory or with another methodology. Postnatal (whole blood) and prenatal (chorionic villi, amniotic fluid) specimen types accepted.

Oncology Microarray Analysis

SLH Test Code 8900NC: Provides high resolution, genome-wide assessment of copy number variants (CNVs) and copy neutral loss of heterozygosity (cn-LOH). Oncology (bone marrow or whole blood) specimen types accepted.

The following test codes will be retired: 890PREC and 890PRET

Test descriptions, sample requirements, and CPT codes are available on our website: http://www.slh.wisc.edu/clinical/cytogenetics/

Additional Discontinued Test Offerings

Effective immediately, we will no longer offer the following FISH analysis testing options:

SLH Test Code 871F25: Angelman syndrome (deletion 15q11.2) SLH Test Code 871F26: Cri du chat syndrome (deletion 5p15.2) SLH Test Code 871F27: Deletion 1p36 deletion SLH Test Code 871F30: Prader-Willi syndrome (deletion 15q11.2) SLH Test Code 871F33: Wolf-Hirschhorn syndrome (deletion 4p16.3)

Please call our laboratories at 608-262-0402 with any questions.