Barth syndrome, originally described in 1979 by Neustein et al. and later by Barth et al. as an X-linked cardiосkeletal myopathy with abnormal mitochondria and neutropenia, is a severe disorder that classically presents in infancy with heart failure and sepsis and may be associated with 3-methylglutaconic aciduria. Growth retardation, abnormalities of carnitine and cholesterol and mitochondrial respiratory chain deficiency, particularly cytochrome C, are also notable. A high percentage of affected children reportedly succumb early in life from this disorder.

Bione et al. identified the disease-causing gene as TAZ (G4.5), a novel gene containing 11 alternatively spliced exons. The gene, which is located on the X chromosome at Xq28, encodes a protein family called tafazzins, the function of which remains unknown. Multiple mutations in TAZ including missense, nonsense and splicing mutations, as well as small deletions and insertions, have been identified. However, phenotype-genotype correlations have not been identified. Further, it has been shown that mutations in TAZ result not only in classic Barth syndrome but also in left ventricular noncompaction (LVNC), X-linked infantile cardiomyopathy, X-linked endocardial fibroelastosis (EFE) and dilated cardiomyopathy (DCM). Thus, mutations in TAZ can result in a broad spectrum of clinical diseases including, but not limited to classical Barth syndrome.

The John Welsh Cardiovascular Diagnostic Laboratory offers molecular genetic testing for TAZ mutations. Symptomatic males is tested by DNA sequencing of all 11 exons of the TAZ gene. We strongly recommend initial testing of an affected male, if available, in order to provide the greatest test sensitivity and clearest interpretation of results for subsequent carrier female testing. If an affected male is unavailable for testing, testing of females at high risk is offered. Genetic counseling is recommended for all individuals in order to identify additional at-risk family members and to discuss reproductive issues.

### REASONS FOR REFERRAL
- Molecular confirmation of the diagnosis of Barth syndrome, X-linked infantile cardiomyopathy, left ventricular noncompaction, and X-linked endocardial fibroelastosis (EFE) in affected males
- Carrier testing in females with a family history of Barth syndrome, X-linked infantile cardiomyopathy, left ventricular noncompaction, and X-linked endocardial fibroelastosis (EFE)
- Carrier testing is not offered for asymptomatic minor females. Please call for additional information.

### METHODOLOGY
Genomic DNA is analyzed for TAZ mutations by DNA sequencing of all 11 exons of the TAZ gene, as well as the exon/intron junctions and a portion of the 5' and 3' untranslated regions. Patient DNA is sequenced in both the forward and reverse orientations. If a mutation is identified, additional family members are analyzed only for the familial mutation by automatic fluorescent DNA sequencing.

### SERVICE FEES
- **Index Case (Male or Female)**: $600 per sample
- **Additional Family Members**: $300 per sample; known familial mutation only
- **CPT Codes**: 81406, 81403

### SPECIMEN REQUIREMENTS
- **Blood (preferred)**: EDTA (purple-top) tubes: **Adult**: 5 cc **Child**: 5 cc **Infant**: 2-3 cc
- **Tissue**: Frozen (preferred), RNA later
- **Other Body Fluids and Formalin-fixed, Paraffin-embedded Tissue**: Call to inquire