

References for *ACADVL*

October 29, 2009

- Andresen, B. S., P. Bross, et al. (1996). "Cloning and characterization of human very-long-chain acyl-CoA dehydrogenase cDNA, chromosomal assignment of the gene and identification in four patients of nine different mutations within the VLCAD gene." Hum Mol Genet **5**(4): 461-72.
- Andresen, B. S., S. Olpin, et al. (1999). "Clear correlation of genotype with disease phenotype in very-long-chain acyl-CoA dehydrogenase deficiency." Am J Hum Genet **64**(2): 479-94.
- Boneh, A., B. S. Andresen, et al. (2006). "VLCAD deficiency: pitfalls in newborn screening and confirmation of diagnosis by mutation analysis." Mol Genet Metab **88**(2): 166-70.
- Goetzman, E. S., Y. Wang, et al. (2007). "Expression and characterization of mutations in human very long-chain acyl-CoA dehydrogenase using a prokaryotic system." Mol Genet Metab **91**(2): 138-47.
- Gregersen, N., B. S. Andresen, et al. (2001). "Mutation analysis in mitochondrial fatty acid oxidation defects: Exemplified by acyl-CoA dehydrogenase deficiencies, with special focus on genotype-phenotype relationship." Hum Mutat **18**(3): 169-89.
- Mathur, A., H. F. Sims, et al. (1999). "Molecular heterogeneity in very-long-chain acyl-CoA dehydrogenase deficiency causing pediatric cardiomyopathy and sudden death." Circulation **99**(10): 1337-43.
- McKusick, V. A. (2007). "201475 ACYL-CoA DEHYDROGENASE, VERY LONG-CHAIN, DEFICIENCY OF." OMIM, created 1993.
- Rinaldo, P., D. Matern, et al. (2002). "Fatty acid oxidation disorders." Annu Rev Physiol **64**: 477-502.
- Straussberg, R. and A. W. Strauss (2002). "A novel mutation of late-onset very-long-chain acyl-CoA dehydrogenase deficiency." Pediatr Neurol **27**(2): 136-7.