

References for *ARG1*

November 2, 2009

- Bernar, J., R. A. Hanson, et al. (1986). "Arginase deficiency in a 12-year-old boy with mild impairment of intellectual function." J Pediatr **108**(3): 432-5.
- Cederbaum, S. D., K. N. Shaw, et al. (1977). "Hyperargininemia." J Pediatr **90**(4): 569-73.
- Cederbaum, S. and E. A. Crombez (2009). "Arginase Deficiency." GeneReviews, created 2004.
<http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=gene&part=arg1>.
- Crombez, E. A. and S. D. Cederbaum (2005). "Hyperargininemia due to liver arginase deficiency." Mol Genet Metab **84**(3): 243-51.
- Iyer, R., C. P. Jenkinson, et al. (1998). "The human arginases and arginase deficiency." J Inherit Metab Dis **21 Suppl 1**: 86-100.
- McKusick, V. A. (2007). "237300 CARBAMOYL PHOSPHATE SYNTHETASE I DEFICIENCY, HYPERAMMONEMIA DUE TO." OMIM, created 1986.
- McKusick, V. A. (2009). "207800 ARGININEMIA." OMIM, created 1986.
- McKusick, V. A. (2009). "207900 ARGININOSUCCINIC ACIDURIA." OMIM, created 1986.
- McKusick, V. A. (2009). "215700 CITRULLINEMIA, CLASSIC." OMIM, created 1986.
- McKusick, V. A. (2009). "311250 ORNITHINE TRANSCARBAMYLASE DEFICIENCY, HYPERAMMONEMIA DUE TO." OMIM, created 1986.
- Scaglia, F. and B. Lee (2006). "Clinical, biochemical, and molecular spectrum of hyperargininemia due to arginase I deficiency." Am J Med Genet C Semin Med Genet **142C**(2): 113-20.
- Summar, M. L. (2005). "Urea Cycle Disorders Overview." GeneReviews, created 2003.
<http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=gene&part=uca-overview>.
- Uchino, T., S. E. Snyderman, et al. (1995). "Molecular basis of phenotypic variation in patients with argininemia." Hum Genet **96**(3): 255-60.
- Vockley, J. G., B. K. Goodman, et al. (1996). "Loss of function mutations in conserved regions of the human arginase I gene." Biochem Mol Med **59**(1): 44-51.