Leupold, D., Bojasch, M. and Jakobs, C. 3-Hydroxy-3-methylglutaryl-CoA lyase deficiency in an infant with macrocephaly and mild metabolic acidosis. *Eur. J. Pediatr.* 138 (1982) 73–76

Lowry, O. H., Roseborough, N. J., Farr, A. L. and Randall, R. J. Protein measurement with folin-phenol reagent. *J. Biol. Chem.* 193 (1951) 265–275

Moses, S. W., Aviram, M., Geiger, R., Berger, R. and Smit, P. C. 3-Hydroxy-3-methylglutaryl-coenzyme A lyase deficiency. (Case report). *J. Inher. Metab. Dis.* 12 (1989) 341–342

Srere, P. A. Citrate synthase, EC 4.1.3.7, citrate-oxaloacetate-lyase. In Lowenstein, J. M. (ed.), *Methods in Enzymology*, vol. 13, Academic Press, London, 1969, pp. 3–11

Stacey, T. E., de Sousa, C., Tracey, B. M., Whitelaw, A., Mistry, J., Timbrell, P. and Chalmers, R. A. Dizygotic twins with 3-hydroxy-3-methylglutaric aciduria; unusual presentation, family studies and dietary management. *Eur. J. Pediatr.* 144 (1985) 177–181

Tanaka, K. and Rosenberg, L. E. Disorders of branched chain amino acid and organic acid metabolism. In Stanbury, J. B., Wyngaarden, J. B., Fredrickson, D. S., Goldstein, J. L. and Brown, M. S. (eds.), *The Metabolic Basis of Inherited Disease*, (5th edn.), McGraw-Hill, New York, 1983, pp. 440–473

Wanders, R. J. A., Schutgens, R. B. H. and Zoeters, P. H. M. 3-Hydroxy-3-methylglutaryl CoA lyase in human skin fibroblasts: study of its properties and deficient activity in 3-hydroxy 3-methylglutaric aciduria patients using a simple spectrophotometric method. *Clin. Chim. Acta* 171 (1988a) 95–102

Wanders, R. J. A., Schutgens, R. B. H. and Zoeters, P. H. M. Prenatal diagnosis of 3-hydroxy-3-methylglutaric aciduria via enzyme activity measurements in chorionic villi, chorionic villous fibroblasts or amniocytes using a simple spectrophotometric method. *J. Inher. Metab. Dis.* 11 (1988b) 430

Wilson, W. G., Cass, M. B., Sovik, O., Gibson, K. M. and Sweetman, L. A child with acute pancreatitis and recurrent hypoglycemia due to 3-hydroxy-3-methylglutaryl-CoA lyase deficiency. *Eur. J. Pediatr.* 142 (1984) 289–291

Wysocki, S. J. and Haehnel, R. 3-Hydroxy-3-methylglutaric aciduria: 3-hydroxy-3-methyl-glutaryl coenzyme A lyase level in leukocytes. *Clin. Chim. Acta* 73 (1976) 373–375

Wysocki, S. J. and Haehnel, R. 3-Hydroxy-3-methylglutaryl CoA lyase deficiency: a review. *J. Inher. Metab. Dis.* 9 (1986) 225–233

Zoghbi, H. Y., Spence, J. E., Beaudet, A. L., O'Brien, W. E., Goodman, C. J. and Gibson, K. M. Atypical presentation and neuropathological studies in 3-hydroxy-3-methylglutaryl-CoA lyase deficiency. *Ann. Neurol.* 20 (1986) 367–369

J. Inher. Metab. Dis. 13 (1990) 164

ERRATUM

Villasana *et al.* Neurological deterioration in adult phenylketonuria. Vol. 12 No. 4 (1989) 451–457

In Table 1, page 454, the serum phenylalanine concentrations before diet for Subjects A and B should read 27.8 and 27.1 mg/100 ml respectively, rather than 17.8 and 27.4.