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Obesity-colitis-hypothyroidism-cardiac hypertrophy-developmental delay syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Obesity-colitis-hypothyroidism-cardiac hypertrophy-developmental delay syndrome</u>.

ORPHA:88643

Obesity-colitis-hypothyroidism-cardiac hypertrophy-developmental delay syndrome is characterised by precocious obesity, congenital hypothyroidism, neonatal colitis, cardiac hypertrophy, craniosynostosis and developmental delay. It has been described in two brothers, one of whom died within the first month of life. The parents of the two children were nonconsanguineous and in good health, however, the pregnancies were complicated by a maternal HELLP syndrome (Haemolysis, Elevated Liver enzymes and Low Platelets). The mode of inheritance has not yet been clearly established.

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