



MICROGNATHIA WITH GLOSSOPTOSIS (PIERRE - ROBIN SYNDROME) - Report of a case

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A case of micrognathia with glossoptosis (Pierre - Robin Syndrome) is presented. This syndrome is characterised by micrognathia, glossoptosis and cleft palate. Clinical features and skull radiographs revealed typical abnormalities associated with the syndrome. No haematological, biochemical or immunological abnormalities could be seen in this patient. Pedigree analysis showed an autosomal dominant mode of transmission of the disease. Chromosomal studies did not reveal any structural or numerical discrepancies.

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