A case report: Pfeiffer Syndrome Type 2

*Gwang-jun Kim, Sang-hoon Lee, Hyoung-moo Park

Chung-ang University Hospital, Department of Obstetrics and Gynecology,

Seoul, Korea

Abstract

Pfeiffer syndrome is an autosomal dominant syndrome consisting of craniosynostosis, a midfacial hypoplasia, ocular proptosis, broad thumbs and medially deviated great toes. The prenatal diagnosis can be possible with ultrasonography after second trimester. We present one case of Pfeiffer syndrome type 2 boy with his prenatal ultrasound findings.