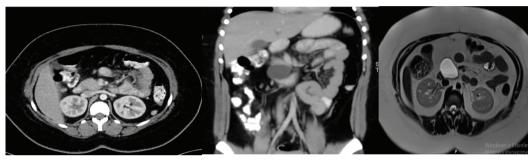
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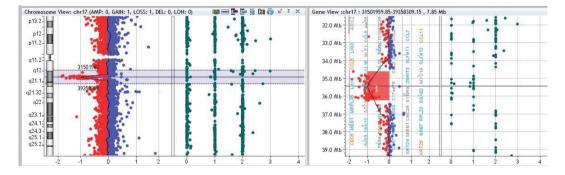
Journal of Clinical Research in Pediatric Endocrinology

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Abdominal computed tomography and magnetic resonance images demonstrating a cystic lesion in the head of the pancreas, and agenesis of the pancreatic neck, trunk and tail



Microarray analysis demonstrated a prominent, heterozygous deletion of a 1.63 Mb-spanning DNA sequence at chromosomal location 17q12, which included the HNF1B gene

A Case of Diabetes Mellitus Type MODY5 as a Feature of 17q12 Deletion Syndrome
Yaşar Köstek H et al.

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What is already known on this topic? What this study adds?

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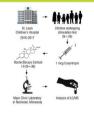
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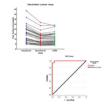
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Book Chapters: Darendeliler F. Growth Hormone Treatment in Rare Disorders: The KIGS Experience. In: Ranke MB, Price DA, Reiter EO (eds). Growth Hormone Therapy in Pediatrics: 20 Years of KIGS. Basel, Karger, 2007;213-239.

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Review

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Tom Mazur, Jennifer O'Donnell, Peter A. Lee

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 - Enver Şimşek, Oğuz Çilingir, Tülay Şimşek, Sinem Kocagil, Ebru Erzurumluoğlu Gökalp, Meliha Demiral, Çiğdem Binay
- 146 Serum Ghrelin and Glucagon-like Peptide 1 Levels in Children with Prader-Willi and Bardet-Biedl Syndromes Doğa Türkkahraman, Suat Tekin, Merve Güllü, Güzin Aykal
- Associations of Adipocyte-derived Versican and Macrophage-derived Biglycan with Body Adipose Tissue and Hepatosteatosis in Obese Children

 Reyhan Deveci Sevim, Mustafa Gök, Özge Çevik, Ömer Erdoğan, Sebla Güneş, Tolga Ünüvar, Ahmet Anık
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