A Microdeletion at 12q24.31 Can Mimic Beckwith-Wiedemann Syndrome Neonatally

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Abstract

We report on a patient who was initially suspected to have Beckwith-Wiedemann syndrome because of recurrent neonatal hypoglycaemias, macroglossia and overgrowth, but in whom no 11p15 abnormality could be found. Follow-up showed continued overgrowth and disturbed glucose homeostasis, a marked developmental delay, and severe behavioural problems especially caused by anxieties. Array comparative genomic hybridization analysis showed a de novo 12q24.31 interstitial deletion, which was confirmed by fluorescence in situ hybridization. The deleted region contains amongst others: HNF1 homeobox A (HNF1A) which is important for the regulation of gene expression in the liver and involved in maturity-onset diabetes of the young type 3 and insulin resistance; acyl-CoA dehydrogenase short chain (ACADS) which encodes an enzyme important in mitochondrial fatty acid beta-oxidation and can cause short-chain acyl-CoA dehydrogenase (SCAD) deficiency, and purinergic receptor P2X7 (P2RX7) which encodes a ligand-gated ion channel, and of which polymorphisms are found with increased frequency in patients with psychiatric disorders, especially anxieties. We conclude the present patient has a hitherto undescribed contiguous gene syndrome, which can initially resemble Beckwith-Wiedemann syndrome.

Key Words

ACADS · Anxieties · Beckwith-Wiedemann syndrome · Contiguous gene syndrome · Deletion 12q24.31 · HNF1A · Macroglossia · Neonatal hyperinsulinism · Overgrowth · P2RX7

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Case Report

The proband was the first and only child of healthy, non-consanguineous parents. Family history was unremarkable. At 36 weeks of pregnancy the mother had a sudden unexplained weight gain. No clues for maternal diabetes were found. An emergency caesarean section was performed at term because of macrosomia. The girl weighed 4,094 g (90th–95th centile). At birth she had a large tongue which filled the mouth but did not cause respiratory problems. She did not have ear creases and her umbilicus was normal. She had immediate hypoglycaemias, which were found to be caused by hyperinsulinism. Management consisted of chlorthalidone, diazoxide and frequent nasogastric tube feedings, which remained needed during infancy to maintain a normoglycaemia. Weight gain was considerable: at 1 year her weight was 11.8 kg (98th centile). In early childhood gradually glucose control stabilized and she was successfully weaned off medication at the age of 4 years. The presence of a large tongue, recurrent neonatal hypoglycaemias, and rapid weight gain led to the suspicion of her having BWS (fig. 1a). Cytogenetic analysis showed a normal female karyotype, 46,XX, without evidence for an 11p15 duplication. A microsatellite marker mapping to the tyrosine hydroxylase locus was used to exclude mosaic 11p15 paternal isodisomy. A mutation screen of ABCC8 (OMIM 500509) gave normal results.

Developmental progress was very slow. She walked at 18 months and was able to use a few words at 2 years. She continued to be unsteady on her feet with a broad based gait. She developed marked anxieties, specifically about plants, small pieces of paper, loud noises and crowds which disturbed daily life significantly. A diagnosis of an autistic spectrum disorder was made at the age of 4 years. She continued to show overgrowth both in height and weight, despite dietary regimens. Urinary continence was achieved at 11 years. Except for significant constipation her general health has been good.

Re-examination at 11 years of age showed an overgrown girl, moderately to severely retarded, with a head circumference of 54.8 cm (75th centile), height of 157.5 cm (98th centile) and a weight of 91.5 kg (98th–99.6th centile). She had upslanted palpebral fissures, a broad nasal base, full cheeks, irregularly placed teeth, full and everted lower lip, and narrow but large ears with a thick helix (fig. 1b, c). She had inverted nipples, a single small truncal café-au-lait spot, large hands and feet, mild tapering of the fingers, and short toes. The left 4th finger was proximally implanted, and shortening of the 4th metacarpal bone was confirmed radiologically.

Molecular Analyses

Microarray analysis using the BlueGnome CytoChip 2.01 (Genome Assembly NCBI 36) showed a cryptic interstitial deletion of the long arm of chromosome 12. The breakpoints of the deletion were within band 12q24.31. The deletion was between 1.58 Mb and 2.53 Mb in length and included the BACs RP11-18C24, RP11-44F24 and RP11-87C12 (fig. 2). The deletion was confirmed by fluorescent in situ hybridisation (FISH) using probes RP11-18C24 and RP11-87C12. Both parents gave a normal result by FISH using probe RP11-87C12, indicating that the deletion had arisen de novo.

Annotated genes in the deleted region are shown in figure 2 and include HNF1 homeobox A (HNF1A), acyl-coenzyme A dehydrogenase, C2 to C3 short chain (ACADS), purinergic receptor P2X, ligand-gated ion
channel 7 (P2RX7), 4-hydroxyphenylpyruvate dioxygenase (HPD) and B-cell CLL/lymphoma 7A (BCL7A). The HNF1A gene has been shown to be important in glucose metabolism. Mutations in HNF1A (OMIM 142410) are associated with maturity-onset diabetes of the young type 3 (MODY3), susceptibility to insulin dependent diabetes mellitus, insulin resistance and hepatic adenomas.

ACADS (OMIM 606885) is a tetrameric mitochondrial flavoprotein similar to MCAD. Patients with homozygous mutations in ACADS present with variable features including hypoglycaemia, developmental delay, seizures, and behavioural difficulties. In some patients these symptoms are transient [Van Maldegem et al., 2006]. Polymorphisms in P2RX7 (OMIM 602566) are found with increased frequency in patients with chronic lymphatic leukaemia and psychiatric disorders such anxiety, phobias, depression and bipolar disorder [Kapoor, 2009]. HPD (OMIM 609695) is involved in the tyrosine catabolic pathway; heterozygous mutations are associated with Hawkinsinuria. Affected patients have failure to thrive and transient metabolic acidosis. Finally, variants in BCL7A (OMIM 601406) have been shown to be a risk factor for the development of non-Hodgkin lymphoma [Morton et al., 2009].

**Discussion**

There have been only limited reports of patients with interstitial deletions involving band 12q24.31 and to our knowledge none with the same breakpoints or clinical features as our case. A male with a 12q24.31q24.33 deletion, developmental delay, tracheomalacia, genital abnormalities and minor cardiac defects was reported by Sathya and co-workers [1999]. Plotner et al. [2003] described a second cytogenetically visible case with microcephaly, moderate developmental delay, cardiac defects, growth failure, non-specific dysmorphic features and a 12q24.31q24.32 deletion. A review of terminal 12q deletions can be found elsewhere [Van Karnebeek, 2002].

The function of several of the genes within the deleted region seems to explain most of the features in the present proband well. The heterozygous deficiency of HNF1A may be responsible for the abnormal insulin levels in the neonatal period, and the unstable glucose homeostasis thereafter. It has been reported for patients with HNF4A mutations to have macrosomia at birth and hyperinsulinaemic hypoglycaemias in the neonatal period, evolving in decreased insulin secretion and diabetes later in life [Pearson et al., 2007]. This has not been reported for HNF1A though. The P2RX7 deletion might explain her unusual behavioural and social problems, and it is possible that the ACADS deficiency plays a role in her developmental delay, although this is much less certain. As
such the present patient may be characterized as having a true contiguous gene syndrome. The deletion of several genes known to be involved in cancer development urged us to install a regular follow-up of the patient.

The initial resemblance of the proband to patients with Beckwith-Wiedemann syndrome is important for etiologic investigations in similar patients, and stresses the need to search for submicroscopic chromosome imbalances. The psychiatric profile in the proband and the deletion of P2RX7 deserves further attention. Studies of series of patients with similar psychiatric illnesses for mutations in the receptor P2RX7 are warranted. Possibly this may lead to better understanding of the aetiology and pathogenesis of such psychiatric disorders, and ultimately may lead to new therapeutic avenues.

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References


