Oculocutaneous albinism in a patient with 17p13.2-pter duplication – a review on the molecular syndromology of 17p13 duplication

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**Background.** Chromosomal duplications involving 17p13.3 have recently been defined as a new distinctive syndrome with several diagnosed patients. Some variation is known to occur in the breakpoints of the duplicated region and, consequently, in the phenotype as well.

**Aims.** We report on a patient, the fifth to our knowledge, a 4-year-old girl with a pure *de novo* subtelomeric 17p13.2-pter duplication. She presents all of the facial features described so far for this duplication and in addition, a unilateral palmar transversal crease and oculocutaneous albinism which has not been reported previously.

**Methods.** A detailed molecular description of the reported aberration and correlation with the observed phenotypical features based on a literature review. We discuss the possible molecular etiology of albinism in regard to the mode of inheritance.

**Conclusion.** The new data provided here may be useful for further genotype correlations in syndromes with oculocutaneous albinism, especially of autosomal dominant inheritance.

**Key words:** chromosomal duplication, 17p13.3, albinism

**INTRODUCTION**

Chromosomal duplications involving 17p13.3 have recently been defined as a new distinctive syndrome (MIM#613215) with more than 40 diagnosed patients. This disorder represents a contiguous gene duplication syndrome involving the genes *PAFAH1B1* (*LIS1*) and/or *YWHAE*. The same region on chromosome 17p13.3 is deleted in Miller-Dieker lissencephaly syndrome (MDLS). The phenotypic features of 17p13.3 duplication may include: intrauterine growth retardation (IUGR), mild to moderate psychomotor delay, hypotonia, and craniofacial dysmorphism such as: high forehead with frontal bossing, small nose and mouth, subtle hand/foot malformations. Nevertheless, the phenotype varies depending on the breakpoints of the duplicated region and, consequently, on the gene content. Bruno et al. suggested that there are 2 classes of microduplications of 17p13.3. Class I involves *YWHAE*, but not *PAFAH1B1*, whereas class II duplications involve *PAFAH1B1* and may also include *CRK* and *YWHAE* (ref.4).

Here, we report on another patient, a 4-year-old girl, with a pure *de novo* subtelomeric class II duplication of 17p13.2-pter identified by MLPA (Fig. 1). This finding was confirmed and further characterized by FISH and array CGH studies (Fig. 2, 3, respectively). It was found that a 5.77 Mb duplicated region of 17p13.2-pter resided on the non-deleted terminal region of chromosome 14p.

**CLINICAL SUMMARY**

The girl is the second child of healthy nonconsanguineous parents, aged 27 years. She was born by vaginal delivery in the 39th week of gestation after an uneventful pregnancy. Her birth weight was 2680 g (5th-10th centile), birth length, 55 cm, and head circumference, 32 cm. Her Apgar score was 10 points. Clinical examination of the newborn showed capillary haemangioma on the eyelids, forehead and nape, which disappeared by the time of genetic evaluation.

The girl’s development was delayed: she began to sit at 12 months and walk at 24 months; her gait at 4 years was unstable. At the age of 2 years she spoke a few words but then stopped speaking. Her anthropometric results at 1 year 5 months were: height 77.7 cm (25th-50th centile), weight 9 kg (25th-50th centile), OFC (occipital frontal circumference) 44 cm (3rd-10th centile), and then at 5 years, during follow-up at our genetic department: height 108 cm (25th-50th centile), weight 18.5 kg (25th-50th centile), and OFC 49 cm (3rd-10th centile).

At the age of 7 months she developed epilepsy. Echocardiography and ultrasonography of the brain and abdomen were normal. Investigation for biotinidase deficiency and organic acidurias revealed no abnormalities, and plasma amino acids were normal. Ophthalmological examination showed myopia: (R: - 3.5 Dsf, L: - 2.25 Dsf), and nystagmus. Moreover, lack of retinal pigment with
Fig. 1.  MLPA analysis presenting a terminal duplication of the short arm of chromosome 17. In our case, the fluorescence signal (peak height) for the 17pter region is increased (blue peak) with respect to the control sample (red peak).

visualization of the choroidal blood vessels, and altered visual evoked potentials (VEP) were noted. This picture, together with other features – hypopigmentation of the skin and hair – for someone could be suggestive with the diagnosis of oculocutaneous albinism type 2 (OCA2) or type 4 (OCA4), which are very similar.

Psychological examination at 6 years assessed her development at the level of a 2 year old. Autism was excluded.

At 1 year 4 months she was referred for genetic counseling because of developmental delay and albinism. Physical examination revealed facial dysmorphe (Fig. 4): impression of hypertelorism, downslanting palpebral fissures with ptosis, slightly broad nasal bridge with bulbous nasal tip, small mouth, and open-mouth appearance.

RESULTS

Conventional cytogenetic analysis performed on G-banded metaphase chromosomes in the proband showed a normal karyotype. Laboratory tests toward Rett and Angelman syndromes (DNA methylation testing of the SNRPN region) revealed no aberration.

MLPA (multiplex ligation-dependent probe amplification) analysis for subtelomeric regions carried out using commercially available SALSA MLPA P036 and P070 human telomere kits (MRC-Holland, Amsterdam, the Netherlands) revealed the presence of a terminal duplication of the short arm of chromosome 17 (Fig. 1).

FISH (fluorescence in situ hybridization) analysis with a subtelomeric probe for chromosome 17p (Cytocell Ltd., Cambridge, UK) confirmed the duplication (three signals for the 17p region noted) showing that an extra copy of the 17pter fragment resided on the non-deleted terminal region of chromosome 14p (FISH with a whole-chromosome painting probe for chromosome 14 was also performed) (Fig. 2). Parental chromosome investigations for the 17pter duplication (karyotyping and FISH) gave normal results, proving the de novo occurrence of this aberration in the patient.

To precisely characterize this chromosomal aberration, a whole-genome oligonucleotide microarray [NimbleGen Human CGH 3x720K Whole-Genome Tiling v3.0 array which contains 720 000 oligonucleotide probes (probe length: 60-mer) with a median probe spacing of 2 509 bp] was subsequently performed. It showed the terminal duplication of chromosome 17 with the prox-
Fig. 3. Array CGH analysis showing a copy number gain (5.77 Mb in size) of chromosome region 17p13.2-pter with the proximal breakpoint at 5,770,000 bp in band p13.2. Data visualization with the use of Roche NimbleGen’s SignalMapTM v1.9 software.

The following eight genes from the MDLS critical region: PRPF8, RILP, SCARF1, PITPNA, SKIP, MYO1C, CRK, and YWHAE (Fig. 5). All had hypotonia, mild to moderate psychomotor delay, and dysmorphic features (the most common: frontal bossing, low-set ears, small nose with broad nasal bridge, hypertelorism and downslanting palpebral fissures, triangularly-shaped chin).

DISCUSSION

The patient presented herein is the next reported patient with a pure de novo 17p13.3 duplication. Interestingly, aberrations consisting solely of a unidirectional translocation of a segment of one chromosome (17p in our case) into another chromosome (14p in the presented patient) are rare. Usually, an accompanying deletion occurs and chromosome is duplicated when their sticky ends remain connected to one another. Nonetheless, in our patient, similarly as in the publications of Avela et al. and Kiiski et al., no reciprocal event could be detected.

The dysmorphic features observed in our proband are in agreement with the findings in the few probands described to date. These comprise a marked hypotonic and long face, downsloping palpebral fissures, low-set ears, small nose with round tip and small mouth. The only sign that has not been mentioned in the other patients and is observed in ours is hypopigmentation, albeit in the figures presented by Roos et al., blond hair is noted (especially in Patient 1 at 14 years). Three patients described by Roos et al. had aberrations ranging in size from 1.8 to 4.0 Mb, with a 1.8 Mb region of overlap, which included the following eight genes from the MDLS critical region: PRPF8, RILP, SCARF1, PITPNA, SKIP, MYO1C, CRK, and YWHAE.

In one patient described in the literature, failure to thrive and poor growth were noted, while another presented overgrowth, with a marfanoid habitus. In a third patient, brain MRI revealed hypoplasia of the corpus callosum and dilated lateral ventricles. Generally, the anthropometric parameters vary among the described patients from a tendency towards tall stature, normal growth, to even growth retardation.

To date, duplication of YWHAE gene is associated with overgrowth or relatively higher body weight and/or length,
but only if the duplicated region also included the CRK gene which is involved in growth regulation and cell differentiation.

The phenotype observed in 17p13.3 duplication varies, depending on YWHAE or PAFAH1B1 involvement. This has suggested the need to classify the patients into two groups according to which gene is involved in the chromosomal duplication (Fig. 5) (ref.4).

Bi et al. concluded that in the individuals with YWHAE duplication macrosomia, mild neurocognitive and pervasive developmental disorder, as well as subtle dysmorphic craniofacial features were observed, whereas the duplication including PAFAH1B1 resulted in a severe total body growth restriction and moderate to severe developmental delay2. Moreover, the patients with duplication of PAFAH1B1 but not YWHAE or CRK showed relative microcephaly or mild brain structural anomalies2,6.

In Table 1 we have summarized the major clinical features of 17p13 duplication in relation to gene content.

In our proband, apart from features observed in other cases and noted in Table 1, oculocutaneous albinism is present. We have no molecular confirmation of suggested by referring physician oculocutaneous albinism type 2, hence the occurrence of hypopigmentation in duplication 17p13 as a feature or coincidence cannot be proved. We were enabled to molecular test for OCA2 gene. However, the patient’s father also has blond hair and myopia (we do not know his other ocular findings) what makes the diagnosis of oculocutaneous albinism type 2 in our proband very unlikely. It is a disease of highly variable phenotype but of autosomal recessive inheritance in which, as a rule, parents are asymptomatic. We suppose that this type of albinism was suspected mainly because it is most common forms of albinism in the world and without taking into account the family history.

More than 30 genes are localized in the region duplicated in our patient (17p13.2-pter) (Fig. 5). Unfortunately, specific correlations of individual genes with the clinical symptoms of the patient could not be delineated. None of the analyzed genes within 17p13.2-pter are known to influence pigmentation. Taking into account the fact that patient’s father also has blond hair and myopia but no chromosomal aberration, the relationship of albinism with identified duplication is doubtful. Rather, it seems reasonable to suspect monogenic aberration of autosomal dominant inheritance, caused by a point mutation in another gene(s) outside of the duplicated chromosomal region.

In the mouse, for example, more than 150 genes are known to affect pigmentation9,10. In humans, the most studied genes resulting in isolated oculocutaneous albinism are TYR (on chromosome 11q14.3), OCA2 (on
chromosome 15q11.2-q12), TYRPI (on 9p23, structurally similar to TYR), and SLC45A2 (on 5p13.2). In our proband, also OCA other than type 2, as an autosomal recessive disorder, unlikely to be inherited from affected father, i.e. the more common forms: OCA1, OCA3 and OCA6 (ref.11), OCA5 (ref.10) and OCA7 (ref. 13). It can be expected that with the further advent of next-generation sequencing (whole exome sequencing: WES) we will find more molecular evidence for new genes contributing to the albino phenotype.

In this paper we delineated the molecular syndromology of duplication within 17p12.3-pter, which seems to manifest with a quite specific and recognizable phenotype. Moreover, we hope we have provided new data for further genotype correlation in syndromes with oculocutaneous albinism.

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REFERENCES


Table 1. Proposed genotype-phenotype correlation in duplication 17p13 (based on Bruno et al.).

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<tr>
<th>Duplicated gene</th>
<th>Major clinical features</th>
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<tr>
<td><strong>YWHAE</strong> (not <strong>PAFAH1B1</strong>)</td>
<td>autistic manifestations/behavioural problems, psychomotor delay, hand/foot malformations, tendency to overgrowth, craniofacial dysmorphism*</td>
</tr>
<tr>
<td><strong>PAFAH1B1</strong> (with/without <strong>CRK</strong> and <strong>YWHAE</strong>)</td>
<td>hypotonia, microcephaly/brain structural abnormalities, developmental delay, severe growth restriction, craniofacial dysmorphism*</td>
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*Prominent forehead and pointed chin are shared by both the class I and class II duplications.