CMA analysis identifies homozygous deletion of $MCPH1$ in 2 brothers with primary Microcephaly-1

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Abstract

**Background:** Homozygous mutations and deletions of the microcephalin gene ($MCPH1$; OMIM *607117) have been identified as a cause of autosomal recessive primary microcephaly and intellectual disability (MIM #251200). Previous studies in families of Asian descent suggest that the severity of the phenotype may vary based on the extent of the genomic alteration. We report chromosome microarray (CMA) findings and the first described family study of a patient with primary microcephaly in a consanguineous Hispanic family.

**Case presentation:** The proband, a boy born at full-term to consanguineous parents from Mexico, presented at 35 months of age with microcephaly, abnormal brain MRI findings, underdeveloped right lung, almond-shaped eyes, epicanthal folds, low hairline, large ears, smooth philtrum, thin upper lip, and developmental delay. MRI of the brain showed a small dermoid or lipoma (without mass effect) within the interpeduncular cistern and prominent arachnoid granulation. The underdeveloped right lung was managed with long-acting inhaled corticosteroids. Otherwise the proband did not have any other significant medical history. The proband had 2 older brothers, ages 14 and 16, from the same consanguineous parents. The 14-year-old brother had a phenotype similar to that of the proband, while both parents and the oldest brother did not have the same phenotypic findings as the proband. The SNP-based CMA analysis of the proband detected a homozygous 250-kb microdeletion at 8p23.2p23.1, extending from 6,061,169 to 6,310,738 bp [hg19]. This genomic alteration encompasses the first 8 exons of $MCPH1$. Follow-up studies detected the same homozygous deletion in the affected brother, segregating with microcephaly and intellectual disability. Regions of homozygosity (ROHs) were also observed in the affected brother. Since ROHs are associated with an increased risk for recessive disorders, presence of ROH may also contribute to the phenotype of the affected brothers. The parents were both hemizygous for the deletion.

**Conclusion:** Here we report a homozygous deletion of multiple exons of the $MCPH1$ gene that was associated with primary microcephaly and intellectual disability in a Hispanic family. In the context of previous studies, our results support the idea that deletions involving multiple exons cause a more severe phenotype than point mutations.

**Keywords:** Primary microcephaly, $MCPH1$ gene, Microdeletion
Background

Primary microcephaly (MCPH) is a genetically heterogeneous disorder, predominately showing an autosomal recessive mode of inheritance. MCPH is a developmental defect of the brain characterized by a congenital small cranium with a reduced occipito-frontal head circumference (OFC) of more than 2 standard deviations (SD) below the mean for age, sex, and ethnicity (severe microcephaly OFC < 3 SD). MCPH is also characterized by mild to moderate intellectual disability and mild seizures, simplified gyral pattern, periventricular neuronal heterotopias, polymicrogyria, speech delay, hyperactivity and attention deficit, aggressiveness, focal or generalized seizures, and delay of developmental milestones and pyramidal signs [1, 2].

Mutations in 18 genes are known to cause MCPH; MCPH1 (*607117) was the first gene to be identified [3, 4]. Whole exome (WES) or the whole genome (WGS) studies revealed an additional 17 OMIM genes associated with MCPH including WDR62 (*613583), CDK5RAP2 (*608201), CASCS (*609173), ASPM (*605481), CENPJ (*609279), STIL (*181590), CEP135 (*611423), CEP152 (*613529), ZNF335 (*610827), PHC1 (*609728), CDK6 (*603368), CENPE (*117143), CENPF (*60235), PLK4 (*605031), TUBGCP6 (*610053), CEP63 (*614724), NDE1 (*609449) [5].

MCPH caused by MCPH1 mutation presents with congenital microcephaly, intellectual disability, and a head circumference that is reduced to a greater degree than height [6].

The MCPH1/microcephalin gene (*607117) is located at chromosome 8p23 and has a genomic size of 241,905 bp. The open reading frame is 8032 bp and consists of 14 exons that encode 835 amino acids; 3 isoforms have been reported so far [4]. Microcephalin, the encoded protein, is implicated in chromosome condensation and cellular responses that are induced by DNA damage [7]. This protein is thought to have a role in G2/M checkpoint arrest via maintenance of inhibitory phosphorylation of cyclin-dependent kinase 1 [7–9]. Loss of the microcephalin protein thus triggers early mitotic entry of neuroprogenitor cells leading to the inability to maintain brain size [7–9].

Two types of mutations in MCPH1 have been reported in the literature. A missense mutation was identified in patients with a less severe cellular phenotype and mild microcephaly [6, 10, 11]. Deletions in MCPH1 have also been reported. A deletion of the first 6 exons of the gene was identified in an Iranian family with intellectual disability and mild microcephaly; premature chromosome condensation in at least 10% to 15% of cells was also reported for this family [11]. In addition, a deletion of the first 11 exons was identified in an Asian-Indian patient [12].
Here we present a consanguineous Hispanic family with primary microcephaly and intellectual disabilities associated with \( \text{MCPH1} \) deletions.

**Case presentation**

The proband presented at 35 months of age with microcephaly (44 cm head circumference, which is more than 2 SD below the third percentile for age), dysmorphic facial features (almond shaped eyes, epicanthal folds, bilateral esotropia, low hairline, large ears, smooth philtrum, and thin upper lip), and intellectual disability or developmental delay.

The proband’s MRI of the brain showed a small dermoid or lipoma (without mass effect) within the interpeduncular cistern and prominent (1.2 × 0.4 cm) arachnoid granulation (Fig. 1). The proband also had an underdeveloped right lung, which was managed with long-acting inhaled corticosteroids. Otherwise the proband did not have any other significant medical history. The proband was born via full-term vaginal delivery at 2720 g. The 34-year old mother was G4P3A1L3 and had no known maternal complications or infections. The proband had no prenatal, perinatal, or postnatal complications, and he was discharged on day 2 of life. He did not walk until he was 13 months and did not speak words until 2 years old, so he received physical and speech therapy for diagnosis of developmental delay. At 48 months, <25% of his speech was intelligible to strangers, and he did not form sentences.

The proband had two brothers who were 14 and 16 years old and from the same consanguineous parents (Fig. 2). One of the brothers (V.6) presented with a similar phenotype as the proband (V.8), while both parents were phenotypically normal (IV.6 and IV.7). The pedigree indicated consanguinity in offspring who had microcephaly, intellectual disabilities, ptosis, hearing loss, and short stature. Hypertension, diabetes, high cholesterol, arthritis, stroke, and osteoporosis were also present in these offspring.

The consanguineous parents of the proband had normal head circumferences for their ages (57 cm for the
father, 57.5 cm for the mother). They did not finish high school and stated it was due to marriage at young age. Their country of origin was Mexico. There was no history of seizures or cardiac, renal, skeletal, or metabolic disease noted in the proband or any of his first degree relatives. In order to determine the cause of the patient’s abnormality, CMA was ordered for both parents.

**Methods**

Genomic DNA was extracted from whole blood using the Gentra Puregene kit (Qiagen-Sciences, Maryland, USA). Microdeletion/microduplication screening was performed for the proband, his parents, and available brothers using a SNP-array platform (CytoScan HD; Affymetrix, Santa Clara, CA), following the manufacturer’s instruction. The CytoScan HD array has 2.67 million probes, including 1.9 million copy number probes and 0.75 million SNP probes. Array data were analyzed using the Chromosome Analysis Suite (ChAS) (Affymetrix, Inc.) software v2.0. FISH analysis using BlueGnome probes RP11-1111 (Illumina, San Diego, CA, USA) for the deleted region 8p23.2p23.1 was performed on metaphase cells according to the manufacturer’s protocol. Subsequently, 50 cells were examined carefully.

**Results and discussion**

The SNP-microarray analysis of the proband’s DNA identified a homozygous microdeletion of 250 kb at

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**Table 1** Clinical and Molecular Presentation of Proband and Comparison to Patients Reported in the Literature to have MCPH1 Deletions

| Features                        | Proband, 1 Patient | Garshasbi 2006 [11], 6 Patients | Pfau 2013 [12], 1 Patient |
|---------------------------------|--------------------|---------------------------------|--------------------------|
| Age                             | 35 months          | 18-32 years                     | 10 months                |
| Gender                          | Male               | 4 Males, 2 females              | Male                     |
| Race                            | Hispanic           | Iranian                         | Asian Indian             |
| Microcephaly (Head circumference)| 44 cm (<−2 SD)     | 49-50 cm (~3 SD), borderline to mild microcephaly | 38 cm (~−5 SD) |
| Chromosome deletion             | 8p23.2p23.1 (hg19: 6,061,169-6,310,738) | 8p22.2p23.2 (hg19: ~6239 kb-6300 kb) | 8p23.1(hg19: 6164,466 × 2,6197,889 × 1-6295,040 × 1,6399,527 × 2) |
| Deletion description            | 250 kb (203 kb of upstream sequence and exons 1-8 of 14 (NM_024596) | 150-200 kb that covered 25 kb of upstream sequence and exons 1-6 of 11 | 97-175 kb, promoter region and exons 1-11 of 14 |
| Facial features                 | Almond shaped eyes, epicanthal folds, bilateral esotropia, low hairline, large ears, smooth philtrum, and thin upper lip | Not described | Mild facial dysmorphism |
| Intellectual disability         | Physical and speech delay. At 48 months, <25% of his speech was intelligible to strangers, and he did not form sentences. | Mild to moderate mental retardation | Developmentally, patient had a social smile, stranger recognition, verbalized with coos and babbling, clapped when asked, and was able to progress from lying recumbent to rolling over, sitting and crawling without assistance. His progress had resulted in graduation from state developmental services. |
The deletion extended from 6,061,169 to 6,310,738 bp (UCSC genome Browser; http://genome.ucsc.edu/; hg19 release). This genomic alteration indicated a homozygous loss (zero copy) of the first 8 exons of the microcephalin gene (MCPH1; OMIM #607117).

The homozygous deletion detected in our proband was also observed in one of his brothers and segregated with microcephaly and intellectual disability (Fig. 3). The parents were both found to be hemizygous for this deletion (Fig. 3) and admitted to consanguinity, which supported the autosomal recessive nature of this pathogenic deletion for microcephaly and intellectual disability. FISH testing confirmed homozygous deletion of the MCPH1 gene in the proband and hemizygous deletion of the same gene in his mother and father (Fig. 4a, b). The older, unaffected brother was not available for testing.

The clinical and molecular features of the affected siblings presented herein are similar to those of patients reported in the literature to also have MCPH1 deletions (Table 1) [11, 12]. Garshasbi et al. reported 6 individuals (4 males, 2 females) of an Iranian family who had primary microcephaly and intellectual disability. FISH testing confirmed homozygous deletion of the MCPH1 gene in the proband and hemizygous deletion of the same gene in his mother and father (Fig. 4a, b). The older, unaffected brother was not available for testing.

Conclusions
Here we report a homozygous deletion of multiple exons of the MCPH1 gene that was associated with primary microcephaly and intellectual disability; to our knowledge, this is the first such report in a Hispanic family. In the context of previous studies, our results support the idea that deletions involving multiple exons cause a more severe phenotype than point mutations.

Abbreviations
CMA: Chromosomal microarray analysis; MCPH1: Microcephalin gene; Oligo-SNP array: Oligonucleotide-single nucleotide polymorphism array

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Authors’ contributions
MH designed and conducted the project, drafted and finalized the manuscript. The rest of the authors provided input to the project, made critical comments on the drafted manuscript, gathered clinical information. All authors read and approved the final manuscript.
Authors' information
Morteza Hemmat - the first and corresponding author. The clinical information was provided by Melissa J Rumple. The rest of the authors helped to draft the manuscript All authors read and approved the final manuscript.

Ethics approval and consent to participate
Consent form was signed by the patient's parent and it is available upon request.

Consent for publication
Consent forms for publication and pictures was signed by the patient's parent and it is available upon request.

Competing interests
The authors declare that they have no competing interests.

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