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Novel clinical finding in *MECP2* Duplication Syndrome: a case report

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MECP2 Duplication Syndrome

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Sir,

MECP2 Duplication Syndrome (*MECP2* DS) is a newly described genetic condition, with approximately 120 affected males being reported insofar [1-11]. The clinical features of this syndrome are hypotonia with feeding difficulties in the first months of life, followed by spasticity in childhood; delayed motor developmental milestones; severe speech delay; dysmorphic features (brachycephaly, midfacial hypoplasia, large ears, and flat nasal bridge); moderate to severe intellectual disability; autistic features; seizures; recurrent respiratory infections; hypoplasia of the corpus callosum; severe constipation; less frequently, bladder dysfunction (urinary retention; bladder dilatation) [8]. Movement disorders (*i.e.* choreiform movements, stereotypies and repetitive behaviors - especially midline hand movements), were reported in various percentages of patients; rocking, spinning, and self-injurious behavior were also reported [11]. Most females heterozygous for *MECP2* duplication are asymptomatic, but can exhibit features of the broad autism phenotype or other symptoms like anxiety, depression, compulsions [12]. These patients management is multidisciplinary, including the treatment of feeding difficulties and of respiratory infection, physical therapy, speech therapy, ergotherapy, and specific behavioural therapy.

We report on a 6 year-old boy, the first child of healthy nonconsanguineous parents, born after an uneventful pregnancy, with a birth weight 2500 g, Apgar score 10. No movement disorders or attention disorders in the family history were reported. Shortly after birth, the boy presented hypotonia, with delayed psychomotor development: he held his head at 10 months, sat at 15 months, walked alone at 36 months, and spoke first words at 40 months. In the first year of life the patient had feeding difficulties and recurrent respiratory infections. The clinical examination showed: weight 23 kg (Pc 75), height 115 cm (Pc 50), occipitofrontal circumference 48 cm (< 2 DS); facial dysmorphism (hypertelorism; partial palpebral ptosis; open, carp-shaped

mouth; micrognathia) (figure 1); cryptorchidism; bilateral pyramidal syndrome; severe intellectual disability; severe speech delay; autistic features; drooling. A prominent feature of the clinical picture was severe hyperkinesia, manifested as random non-purposeful movements, occurring throughout the day, in any environment (at home, in kindergarten, in other public spaces), more prominent during the morning and in the afternoon, impairing daily activities (the child was not able to play or to stay at the table to do a task for more than 5 minutes, thus not being able to participate in different activities together with other children). The patient presented also stereotyped movements (hand-flapping movements) and agitation during sleep. Hyperkinesia was, actually, the main reason for referral to our Department. The patient had not chorea, tremor, or epileptic seizures. To our knowledge, this is the first case with confirmed *MECP2* duplication and hyperkinesia, the *MECP2* duplication syndrome patients being considered rather hypoactive [13]. Biological tests, EEG and cerebral MRI were normal.

Genetic tests included: GTG-banded karyotyping by standard procedures, which was normal, and aCGH on 44K Agilent platform which revealed a 1.43 Mb duplication of Xq28 (152824755-153262507, NCBI Build 36) (figure 2). This region contains the following genes *AVPR2*, *ARHGAP4*, *ARD1A*, *RENBP*, *HCFC1*, *TMEM187*, *IRAK1*, *MECP2*, *OPN1LW*, *OPN1MW*, *OPN1MW2*, *TEX28*, *TKTL1*, *FLNA*, *EMD* [14]

Our patient management included educational and rehabilitation support: speech therapy, cognitive stimulation, behavioural therapy, with slight improvement of autistic features, but with no progress in speech development. Association of behavioural therapy with neuroleptic (risperidone) improved the hyperkinesia.

Discussions

MECP2 duplications often involve several genes besides *MECP2*, including genes known to act in central nervous system development and/or reported in association with MR: *LICAM*, *FLNA*, *EMD* [3]. Our case presents a duplication of Xq28, including *MECP2* gene, and *IRAK1* but, notably, not *LICAM*. Comparisons with two CNV public databases [15] found part of the duplicated loci reported as CNVs detected in healthy subjects, both as gains and losses (*IRAK1*, *OPN1LW*, *OPN1MW*, *OPN1MW2*, *TEX28*, *TKTL1*, *FLNA*, *EMD*); therefore, the role of these genes in the pathologic condition of the patient is unclear.

The clinical features of our patient largely overlap with those described by other authors, namely infantile hypotonia, followed by spasticity, feeding difficulties, severe speech delay, dysmorphic features, intellectual disability, and recurrent respiratory infections. A novel clinical feature, not reported by date in *MECP2* DS, was severe hyperkinesia. Clinical characterization of larger patient groups is required in order to establish if hyperkinesia might stand as a major sign or just add to the clinical variability of *MECP2* DS.

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Figures legend.

Figure. 1. Frontal view of the patient showing dysmorphic facial features.

Figure 2. aCGH profile of chromosome X showing a 1.43 Mb duplication of Xq28.



