Fukuyama Type Congenital Muscular Dystrophy in a Turkish Child

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ABSTRACT: Congenital muscular dystrophy (CMD) is a heterogeneous group of disorders which is associated with more or less degrees of cerebral involvement. There are four separate entities within CMD nosology. Among these Fukuyama’s CMD (FCMD) is highly prevalent in Japan, whereas the classic form with normal or subnormal intelligence, also known as the occidental type, covers the vast majority of cases in the West. We report a case of FCMD seen in a Turkish child.

RÉSUMÉ: Dystrophie musculaire congénitale de type Fukuyama chez un enfant Turque. La dystrophie musculaire congénitale (DMC) est un groupe hétérogène de maladies qui sont associées à différents degrés d’atteinte cérébrale. La DMC en tant qu’entité nosologique comprend quatre entités séparées. Parmi celles-ci, la DMC de Fukuyama (DMCF) a une prévalence élevée au Japon, alors que la forme classique avec une intelligence normale ou subnormale, aussi connue comme étant la forme occidentale, couvre la grande majorité des cas identifiés dans l’hémisphère ouest. Nous rapportons un cas de DMCF observé chez un enfant turque.


Congenital muscular dystrophy (CMD) is a unique type of muscular disorder characterized by probable autosomal recessive inheritance, early onset hypotonia, involvement of facial muscles, joint contracture and slowly progressive course. Mentality may or may not be involved. Clinical presentations are rather heterogeneous and several types of CMD have been described. Basically there are four major sub-groups within CMD nosology: classic occidental type CMD with normal or subnormal intelligence; Fukuyama’s CMD (FCMD); the cerebro-ocular dysplasia-muscular dystrophy syndrome (COD-MD); and some forms which do not fit in any category.1-5 Among these FCMD is distinguished by developmental central nervous system (CNS) abnormalities, severe mental retardation and related symptomatology.6 In Japan this seems to be one of the frequent causes of the floppy infant syndrome, whereas outside Japan the association of CMD with CNS involvement is very rare, and only a few cases have been reported.7,8 We report a case of FCMD observed in a 27 month old Turkish child.

CASE REPORT

A 27 month old female child coming from the northern part of Turkey was referred to our clinic for evaluation of floppiness and developmental delay since birth. She was born of non-consanguineous parents as the second child. An 8 year old male sibling is normal. There was no family history of muscle disease or mental retardation. Fetal movements were noted to be decreased in utero, but pregnancy was uneventful otherwise. Delivery was at term and was spontaneous vaginal. The birth weight was 3.1 kg. The neonatal period was complicated, also known as the occidental type, covers the vast majority of cases in the West. We report a case of FCMD seen in a Turkish child.

Up at 20 months, sat without support at 26 months, smiled and knew her mother at 10 months, and barely uttered single phrases at 26 months. She had experienced two generalized afebrile seizures at 8 and 10 months.

Physical examination on admission showed evidence of undernourishment and marked hypotonia. Her head circumference, length and weight were below the third percentile. There were no dysmorphic features. Her facial appearance was mask like. Ocular movements were full and fundi were normal. Findings from examination of the chest and abdomen were unremarkable. Her EKG and cardiac echogram were normal. An EEG recording had scattered sharp transients from both hemispheres and diffusely slow background activity. EMG showed generalized low voltages without denervation. A CT scan of the head revealed cortical atrophy and white matter hyperlucency around the frontal and occipital ventricular horns (Figure 1). Biopsy from the gastrocnemius showed variation in fibre size and moderate fibrous tissue infiltration along with mild fat replacement and scattered fibre necrosis (Figure 2). There was no evidence of type grouping. On the basis of these findings we made a diagnosis of FCMD.

DISCUSSION

The etiology of FCMD remains unclear. A genetic factor may be important,9 but some authors have suggested an infectious etiology.10,11 FCMD has even been reported in Japanese infants conceived and born outside Japan.12 The major changes in the CNS represent an arrest in the migration and differentiation of neurons early in the course of fetal development. This defect is expressed as microcephaly, polymicrogyria, pachygyria and heterotopias.13 Other abnormalities include fusion of the frontal lobes, hydrocephalus, periventricular cysts, optic nerve atrophy, hypoplasia of pyramidal tracts, reduction in anterior...
of CMD in different geographic areas, or merely to authors’ interests. The existence of various forms of CMD is very likely associated with more or less marked disorders in cerebral organogenesis.\textsuperscript{15,16} Efforts should be made to routinely screen CMD patients regardless of mental status and autopsy studies wherever possible should be performed, to recognize more aspects of this heterogeneous disease.

REFERENCES