Case Report

Meckel-Gruber syndrome: A rare and lethal anomaly

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ABSTRACT
Meckel-Gruber syndrome is a condition that belongs to the ciliopathies, a category of diseases thought to be caused by dysfunction of cilia and flagella. Polycystic liver and kidney disease, Bardet-Biedl syndrome, Alstrom syndrome, and Joubert syndrome also belong to the same group [5].

The pathophysiology of MGS is complex. One study screened patients with Meckel syndrome for mutations in B9D1 and B9D2 and identified homozygous c.301A>C (p.Ser101Arg) B9D2 mutation. The data showed that the domain-containing proteins Mks1, B9d1, and B9d2 interact physically; the p.Ser101Arg mutation abrogates the ability of B9d2 to interact with Mks1, which suggests that this mutation compromised B9d2 function. The data further showed that B9d1 is required for normal Hedgehog (Hh) signaling, ciliogenesis, and ciliary protein localization. B9d1 and B9d2 are essential components of a B9 protein complex and, when this is disrupted, Meckel syndrome results [6].

The triad of occipital encephalocele, large polycystic kidneys, and postaxial polydactyly characterizes Meckel-Gruber syndrome.

KEY words: Meckel-Gruber syndrome, anomaly, child, Sudan.

INTRODUCTION
Meckel-Gruber syndrome (MGS) is a rare, lethal autosomal recessive condition mapped to 6 different loci in different chromosomes 17q21-24 (MKS1), 11q13 (MKS2), 8q21.3-q22.1 (MKS3),[1] 12q21.31-q21.33 (MKS4),[2] 16q12.2 (MKS5),[3] and 4p15.3 (MKS6).[4] This mapping suggests genetic heterogeneity in Meckel-Gruber syndrome. More than 200 cases have been reported.

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How to cite this article:
Meckel-Gruber syndrome. Cystic dysplasia of the kidneys is the most constant and characteristic feature of Meckel-Gruber syndrome. Associated abnormalities include oral clefting, genital anomalies; CNS malformations, including Dandy-Walker, Arnold-Chiari malformation, and liver fibrosis. Cardiac lesions like atrial septal defect, coarctation of the aorta and pulmonary stenosis may be present [7,8]. Worldwide, the incidence of Meckel-Gruber syndrome is 1 per 13,250-140,000 live births. Individuals of Finnish descent have a higher incidence (1 per 9000 live births, one person in 50 is a carrier). The incidence is also higher among Belgians and Bedouins in Kuwait, with 1 affected birth in 3,500 (carrier rate 1 in 30). The highest incidence is reported in the Gujarati Indians, with 1 affected birth per 1,300 (carrier rate, 1 in 18) [9].

Prenatal ultrasound is currently the best method available to diagnose Meckel-Gruber syndrome and is available in 2-dimensional (2D), 3-dimensional (3D), and 4-dimensional (4D) modalities. The latter is particularly useful in assessing facial features and deformities, musculoskeletal malformations, and limitation of movement. MRI is a valuable complement to ultrasonography in assessing fetal anomalies in the presence of severe oligohydramnios. It is mainly used when ultrasonography findings are inconclusive or are insufficient to guide treatment choices [10].

Chromosome analysis is essential to exclude trisomy 13, which mimics Meckel-Gruber syndrome. Trisomy 13 carries a 1% recurrence risk, as opposed to the 25% recurrence rate for Meckel-Gruber syndrome. Linkage or mutation analysis is not yet available. The mortality is 100% and most babies die in utero or shortly after birth. Pulmonary hypoplasia is the leading cause of death. Other causes include liver and renal failure [11].

We report a Sudanese female baby with a typical triad of Meckel-Gruber syndrome in whom the diagnosis was not made antenatally but immediately after birth, she died on day three of life. To our knowledge this is the first case report of Meckel-Gruber syndrome from Sudan.

CASE REPORT

A female Sudanese baby born at term by emergency Caesarian section due to failure of progress, she was born with low Apgar score requiring admission to the neonatal unit because of respiratory distress and multiple congenital anomalies. A routine U/S during the 2nd trimester showed hydrocephalus, occipital encephalocele, bilateral hydronephrosis and cleft lip and palate. Folic acid was started during the 2nd trimester. The parents are first degree relatives with no family history of similar condition and no history of miscarriage or stillbirth.

Examination of the baby revealed a sick baby with respiratory distress (RR 80/min), weight was 4Kg, head circumference was 38 cm. There was occipital encephalocele (figure 1). Unilateral complete cleft lip and palate, low set ears and hypertelorism. Also there was post axial polydactyly in both hands and feet (figure 2). The rest of the examination revealed bimanually palpable kidneys, normal female genitalia, normal back and patent anus.

Routine investigations were normal including CBC, renal and liver function tests. Cranial U/S showed dilated ventricles and abdominal U/S showed findings consistent with infantile polycystic kidneys and normal liver. Of note, chromosomal analysis and DNA studies were not available. Based on the above a clinical diagnosis of Meckel-Gruber syndrome was made and the parents were fully counseled regarding the grave prognosis of this condition and the risk of recurrence.

Management of the baby involved oxygen administration, intravenous fluid fluids and antibiotics. The baby remained sick and distressed and on day three developed convulsions requiring phenobarbitone, phenytoin and calcium supplements,
after which she developed a prolonged apnoea and
died on the same day.

**DISCUSSION**

The first reports of Meckel-Gruber syndrome (MKS) was published in 1822 by Johann Friedrich Meckel. G.B. Gruber also published reports of patients with Meckel-Gruber syndrome in 1934 and gave it the name dysencephalia splanchnocystica. Meckel-Gruber syndrome is also known as Meckel syndrome and Gruber syndrome [12,13]. Our case had the typical triad of occipital encephalocele, polycystic kidneys and polydactyly and the diagnosis was made after birth. Unfortunately,
REFERENCES


ACKNOWLEDGEMENT

We would like to thank the family of the baby mentioned in this report for permitting the use of the case details and photographs. Thanks are also extended to the staff of the neonatal unit at Soba university hospital. Special thanks to Sr. Ragya and Sr. Raja, senior nurses at the neonatal unit, Soba university hospital.