virus diarrhea noticed among 23 children out of 92 (25%) from rural area in the present study.

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Niemann-Pick Disease Type IS in Sibs with 20 Years Follow Up

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In 1983, we had reported two sibs with Niemann-Pick Disease (NPD) type B(I). Type B is now reclassified as type IS(2). Now we are reporting further follow up of these sibs. No such follow up has been reported so far in the Indian literature.

Case Reports

Case 1: The subject, bora of non consanguinous marriage from Indian Christain family was admitted in KEM Hospital,

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Bombay in 1975 (age 9 years), with a history of progressive distention of abdomen noticed since infancy; jaundice at 1, 5 and 9 years of age, recurrent lower respiratory tract infections; frequent episodes of hemoptysis and epistaxis. His early developmental milestones were normal and had no neurological manifestations at that time. His younger sister expired at 7 years of age, she had hepatosplenomegaly, but no neurological manifestations at that time. She was not completely investigated.

On examination, he had massive hepatosplenomegaly (spleen 10 cm and liver 7 cm). His fundus examination revealed increased sphingomyelin (20 times the normal control). The enzyme sphingomyelinase was undetectable in cultured skin fibroblasts, whereas other lysosomal enzymes showed normal activity.

During a follow up period from 1975 to 1990 he developed myoclonic jerks in 1983, requiring treatment with nitrazepam; deposits of sphingomyelin around the cornea starting from superior aspect and completing the circle over the next five years; intermittent cerebellar signs in 1985 and frank convulsions in 1988. In 1988, he also developed frequent loose stools with blood and mucus, not associated with evidence of infection. By 1989, he developed diffuse pulmonary infiltrates (Fig. 1), generalized lymphadenopathy and clubbing. He also developed marked skin pigmentation on lower extremities. In December 1989, he developed congestive cardiac failure, requiring hospitalization. His ECG revealed poor left ventricular function. He expired in June 1990, at the age of 24.

![Image of chest X-ray showing bilateral diffuse pulmonary infiltrates](image-url)
years. He was a fully grown adult with normally developed, secondary sexual characteristics.

Case 2: The elder sibling of Case 1 was noticed to have hepatosplenomegaly at 9 years of age. He was diagnosed to have Niemann Pick disease on bone marrow and liver biopsies at the age of 14 years, during family screening. His fundus examination revealed grayish white areola at the macula and had no neurological manifestation at that time. He studied up to 8th standard. However, he developed deterioration of mental milestones at 15 years, cerebellar signs and corneal deposits at 23 years, and one episode of diarrhea with blood and mucus at 29 years. At present he is 30 years, has jerky movements of hands and feet, incoherent speech, poor concentration and memory, but no frank convulsions at yet. He has no respiratory or cardiac problems.

Discussion

NPD has recently been reclassified into two broad Groups-I and II. Type I includes patients who are sphingomyelinase deficient (with activity less than 10% of normal) and predominant sphingomyelin and cholesterol deposits (as in our case); irrespective of neurological manifestations. Included in type H are a wide spectrum of patients with visceral or neurovisceral lipidosis with sphingomyelin deposition being less than other lipids and in whom exact nature of defect is not known. Each group is further subdivided into Acute (A), Subacute (S) and Chronic (C). Acute forms present in early infancy with rapid deterioration to death by 6 years of age. Chronic forms are usually first detected in adulthood(2).

Subacute forms develop visceral changes in early infancy or childhood, usually splenomegaly followed by hepatomegaly, but progress slowly as in our case. In these early onset cases, the lung fields show diffuse infiltrates contributing to respiratory problems such as asthma and infections as in first sib. These patients show marked phenotypic heterogeneity particularly with respect to the nature, extent and temporal sequence of nervous system involvement. Various manifestations reported include mental retardation, cerebellar ataxia or extrapyramidal manifestations, or more than one manifestation. Various ocular changes such as macular halos, cherry red spot and corneal deposits may be the only evidence of nervous system involvement(2).

Management of these patients is mainly symptomatic. Replacement of enzyme sphingomyelinase by repeated implantation of human amnion sheets has been tried with promising results. Implantation of pure epithelial cells separated from other cell types of the amnion may avoid the risk of host versus graft rejection(3). Antenatal diagnosis is possible by enzyme estimation(2).

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REFERENCES


Sirenomelia with Spinabifida

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Sirenomelia is an exceptionally rare congenital malformation characterized by a single fused lower limb. This deformity is variably known as symmelia, sympodia monopodia, sympus and due to its resemblance to the mermaid of Greek and Roman mythology popularly also known as mermaid baby. Sirenomelia has prevalence of 1.5 - 4.2 cases per 100,000 births(1). About 300 cases of this lethal anomaly have been reported in the world literature(2). In India over the last 15 years, 8 reports have appeared describing this condition(3). Most of these reports have only given the description of the dysmorphism but not have made any attempt to study the possible pathogenesis of the syndrome. In the present report we have done a postmortem arteriogram study to find out the possible vascular abnormalities which are supposed to be one of the mechanisms in the causation of this disorder(4). We also describe and discuss the association of neural tube defects along with siernomelia.

Case Report

A 23-year-old, gravida two, with 8 months amenorrhea was admitted with premature rupture of membranes and onset of labor pain. Her antenatal period was uneventful and there was no history of any drug intake. There was neither a past or a family history of malformations.

A 1.6 kg infant was born by breech delivery. The baby had severe birth asphyxia (Apgar score was 2 at 1 minute and 4 at 5 and 10 minutes). All attempts to resuscitate the baby failed and the baby died after 40 minutes. Physical examination of the infant showed soft tissue fusion in lower extremities except the feet (Fig. 1). The feet were normally formed with toes, normal in number and appearance. Genitalia was represented by a small bud like projection. The anal and urethral openings were absent. Umbilical cord showed a single umbilical artery. The child had normal upper half of the body. The placenta was normal.

Radiological studies demonstrated normal bony structures in the lower extremities and feet. There was bilateral dislocation of hip and crowding of ribs. All the cervical vertebrae and thoracic vertebrae T1 to T4 showed spinabifida defect. The heart was of normal size and lungs were hypoplastic (Fig. 2).