Recurrent Respiratory Tract Infection in an infant: Early Diagnosis by Clinical and Radiological Pearls

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ABSTRACT

Recurrent respiratory tract infections, a cause of concern for both parents and paediatricians, can have various etiologies entitled to different organ systems. Diagnosing the exact cause warrants both clinical acumen and timely investigations. Here, we are reporting an infant with recurrent respiratory tract infections, where adequate clinical examination prompted us to diagnose the extra-respiratory cause with simple investigations.

Keywords: Joubert Syndrome; Molar tooth sign; Recurrent RTI

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Recurrent respiratory tract infections (RTI) are one of the leading reasons for OPD consultation and hospitalisation in children, particularly the infants. More than six serious respiratory tract diseases a year are termed as recurrent RTI (RRI).\(^1\) Epidemiologists estimate that approximately 15% of children suffer from RRI.\(^1\) Though the disease mainly involves upper respiratory tract, involvement of lower respiratory tract (i.e. LRTI or pneumonia) is more serious and more often requires hospitalisation. Apart from diseases of the respiratory tract (e.g. congenital malformations, defective mucociliary clearance etc.), causes of RRI may also include cardiovascular diseases, aspiration, immunodeficiencies and neurological ailments. Here, we are presenting such a case of recurrent RTI in an infant, where the cause was lying elsewhere.

**CASE REPORT**

A 10 month old female child presented to our hospital OPD with complaints of repeated chest infections since birth. She had been hospitalised five times for breathlessness with or without fever since the age of one month. She was born at term out of non-consanguineous marriage with uneventful prenatal and natal periods. However, mother complained of feeding difficulties and occasional fast breathing from the immediate postnatal period. No convulsion was ever seen. There was obvious delay in achieving the motor milestones, though language and social milestones were almost normal. There was no history of similar illness in any of the two elder siblings.

On examination, her weight was below 3\(^{rd}\) percentile, but her length and head circumference were within normal limits. Facies of the child was insignificant, except the presence of prominent forehead. Respiratory system examination was essentially normal with presence of bilateral vesicular breath sounds without any adventitious sounds. But while examining the child for prolonged period, breathing pattern seemed abnormal with brief episodes of hyperpnoea interspersed with normal pattern. Neurologically, the child had generalised hypotonia, but deep tendon reflexes were normal and plantars were equivocal. Other systems including cardiovascular system didn’t reveal any significant abnormalities. Ocular and ophthalmoscopic examinations were also normal.

Routine investigations including CBC and chest X-ray (CXR) were within normal limits. All the previous CXRs available with the parents also didn’t show any feature of pneumonia. Keeping in mind the abnormal neurological findings on clinical examination, MRI of brain was ordered. Axial T1-weighted image showed the features of hypoplastic cerebellar vermis with elongated and thick superior cerebellar peduncles giving it the appearance of “molar tooth” in midbrain (Molar Tooth Sign or MTS) \([\text{Figure 1A]}\). T2-weighted image showed “bat-wing” shaped 4\(^{th}\) ventricle due to dilatation of the same \([\text{Figure 1B}]\). Echocardiography and abdominal ultrasonography were normal. Based on the clinical features and neuroimaging, a diagnosis of Joubert Syndrome (JS) was made.

**DISCUSSION**

JS is a rare congenital CNS malformation with autosomal recessive inheritance in most of them, though X-linked recessive cases has been reported.\(^2,3\) It was first described by French neurologist Marie Joubert in 1969 in four siblings with agenesis of cerebellar vermis presenting with episodic hyperpnea, abnormal eye movements, ataxia and intellectual disability.\(^4\) Several years later, the pathognomic midbrain-hindbrain malformation giving rise to the MTS in radiography was detected in JS.

Even in the absence of established diagnostic criteria, hypotonia, developmental delay, and one or both of the following – a) Irregular breathing pattern, b) Abnormal eye movements have been mentioned as essential for diagnosis of JS along with MRI signs.\(^2\) MRI plays a cornerstone in establishing the diagnosis of JS, because presence of MTS in cranial imaging is very much specific of this disease. Presence of abnormally deep interpeduncular fossa, elongated thick and maloriented superior cerebellar peduncles and absence or hypoplasia of cerebellar vermis gives the characteristic appearance of MTS.\(^5\) Subsequently, the fourth ventricle is moderately
dilated taking the appearance of a “bat wing” or an open “umbrella” on axial CT/MRI. These two signs are reported to be consistent and reliable imaging findings to distinguish JS from other posterior fossa abnormalities. The possibility of prenatal diagnosis of JS has been reported with the ultrasonographic findings showing increased nuchal translucency.\(^6\)

Joubert Syndrome Related Disorders (JSRD) is a group of disorders with developmental delay and multiple congenital anomalies involving mainly retina, kidney, liver and skeleton. The marked pleiotropism of JSRD can be explained by the 10 causative genes identified till date – JBTS1 to JBTS10. They all encode for proteins of primary cilium making JSRD part of an expanding group of diseases called “Ciliopathies.”\(^7\) These proteins play key role in the development and functioning of retinal photoreceptors, neurons, kidney tubules and bile ducts.\(^8\) In the developing cerebellum and brainstem these organelles regulating major signal transduction pathways have been implicated both in neuronal cell proliferation and axonal migration.\(^9\)

Clinically, JSRDs are classified into six phenotypic subgroups: Pure JS, JS with ocular defect, JS with renal defect, JS with oculorenal defect, JS with hepatic defect and JS with orofaciiodigital defect.\(^2\) Thus, diagnosis of JS by MTS in radiography should be followed by a diagnostic protocol to assess whether there is any other organ involvement. Prognosis also largely depends on the severity of the involvement of different organ systems, particularly retina, liver and kidney. There was no extraneuronal involvement in our case suggesting it as a case of Pure JS.

Recurrent RTI, as seen in our case, can be attributed to various factors, e.g. feeding difficulty causing micro-aspirations, hypotonia of respiratory muscles denying adequate cough response, or the ciliary abnormality also affecting the respiratory cilia. The breathing pattern in JS, an effortless hyperventilation in episodes, may have also contributed to overestimation of respiratory distress causing multiple hospitalisations in this case.

**CONCLUSIONS**

Considering the various etiologies of recurrent RTI involving multiple systems other than the respiratory system, careful clinical evaluation is of paramount importance before jumping into the battery of tests. Though there is no curative therapy for JS, early diagnosis dictates prognostic outcome and an opportunity for genetic counselling.
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