Cornelia de Lange syndrome: A rare genetic disorder

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Abstract

Cornelia de Lange Syndrome (CdLS) was first reported by Vrolik in 1849 and Brachmann in 1916, followed by Cornelia de Lange in 1933, after whom the syndrome is named. This disorder has a varied presentation but is mainly characterized by distinctive facial features, growth retardation, microcephaly, hirsutism, psychomotor delay, intellectual disability, and malformations of the upper limbs. Initial diagnosis is usually based on clinical features following specific diagnostic scoring systems. The precise prevalence of the disease is unknown but is estimated to be 1–10:100,000. Depending on the mutated gene, Cornelia de Lange syndrome (CdLS) can be inherited in an autosomal dominant manner, when it is caused by variations in the NIPBL, SMC2, or RAD21 genes, or it can have an X-linked inheritance when it is caused by variations in the SMC1A or HDAC8 genes. However, most cases (more than 99%) result from new (de novo) mutations, which means that are not inherited from the parents and occur in people with no family history of the condition about 30% of the people affected by the syndrome do not have any known cause. Many studies focused on the importance of neurologic findings and reported an incidence of epilepsy in CdLS ranging from 14% to 25%, especially in the classic and more severe form of the syndrome, but there is no data about its electroclinical features and long-term outcome. Life expectancy is relatively normal for people with CdLS and most affected children live well into adulthood. However, certain features of this condition, particularly severe malformations of the heart or throat, may decrease life expectancy in some affected people. The diagnosis is suspected clinically and later confirmed by clinical exome sequencing.

Keywords: Cornelia de Lange Syndrome (CdLS), Epilepsy, Developmental Delay, Clinical Exome Sequencing.

Introduction

Cornelia de-Lange syndrome (CdLS), also known as Brachmann de-Lange syndrome, comprises of congenital malformations, growth retardation, and neurodevelopment delay [1-8]. The diagnosis of CdLS is usually done after birth, but the syndrome could be suspected after the first trimester of pregnancy by ultrasonography [9-16]. Avagliano et al [17] Proposed that following a sequence of detailed scans and examinations, CdLS-affected fetuses could be diagnosed in utero, when one or more characteristics, such as fetal growth rate, limb defects, facial abnormalities, diaphragmatic hernia, and heart diseases. A combination of signs and symptoms defines the classic CdLS phenotype [18]. The cardinal features of CdLS are synophrys, short nose, concave nasal ridge and/or upturned nasal tip, long and/or smooth philtrum, thin upper lip vermilion, downturned corners of the mouth, hand oligodactyly and/or adactyly, congenital diaphragmatic hernia. CdLS maybe associated with global developmental delay and/or intellectual disability, prenatal growth retardation, postnatal growth retardation, microcephaly (prenatally and/or postnatally), small hands and/or feet, short fifth finger and hirsutism [18]. A gene responsible for CdLS–NIPBL on chromosome 5–was discovered in 2004 by researchers at Children's Hospital of Philadelphia. In 2006, a second gene–SMC1A on the X chromosome–was found by Italian scientists. A third gene discovery was announced in 2007 [19]. The gene SMC3 is on chromosome 10 and was also discovered by the research team in Philadelphia. The latter two genes seem to correlate with a milder form of the syndrome. The vast majority of cases are due to spontaneous mutations, although the defective gene can be inherited from either parent, making it autosomal dominant. The types of mutations seen in CdLS rarely include large deletions and 50% have detectable point mutations (frame shift, splice site, nonsense, and missense) [20].
Case Report

We report a case of a 5-year-old boy, born of a non-consanguineous marriage, second by birth order, presented with generalized tonic-clonic epilepsy since 3 years of age. There was a history of developmental delay and poor performance at school. His birth history was uneventful with a low birth weight of 2.2 kg. On examination, microcephaly along with a flat occiput was present. The other anthropometric parameters were normal. The patient had a low pitched, husky voice. His eyebrows were thick, bushy, joined in the midline (synophrys), and with long curled eyelashes [Figure 1]. Right eye divergent squint was present. The mouth was fish-like with long philtrum [Figure 2] and micrognathia was also present. The upper lip was thin and long [Figure 2] and facial hypertrichosis was present. Clinodactylly was present [Figure 3]. The scalp hairline was low set [Figure 4]. He had large ears [Figure 5]. He also had behavioral disorders in the form of hyperactivity. In early life, the child had feeding difficulties due to regurgitation of feeds and also there was history suggestive of hypertonia as the mother had difficulty in changing his diapers. His IQ testing was suggestive of a mild social maturation deficit.

![Fig-1: Synophrys with increased facial hair.](image1)

![Fig-2: Thin upper lip with long philtrum.](image2)

![Fig-3: Clinodactylly](image3)
Blood investigations for convulsion profile and thyroid studies were WNL. EEG and MRI were absolutely normal. Antiepileptic medication, physiotherapy, and speech therapy were started. Parents were counseled regarding the possibility of the syndrome. Clinical Exome Sequencing was suggestive of NIBPL mutation and Genetic counseling was advised.

**Discussion**

CdLS is a very characteristic syndrome that can be identified immediately because of the salient clinical features. A differential diagnosis like Fryns syndrome and fetal alcohol syndrome should be ruled out. Our case had all the typical facial features, microcephaly, generalized hypertrichosis, developmental delay, and clinodactyly.

Life expectancy is normal unless major malformations like apnea, cardiac, and gastrointestinal complications occur. Seizures are common in CdLS (individuals with SMC1A variants: 45%; with NIPBL variants: 15%; without molecular confirmation: 20–26%) [21-23]. In a series of 14 clinically diagnosed patients [24] and an overview of individuals with mostly clinically diagnosed CdLS [23], partial epilepsy was the most common type; the age of onset was typically before 2 years of age, and 35 of 39 individuals for whom full data were available reacted well to standard therapy. Specifically, sodium valproate was found to be effective [24]. However, our patient had generalized tonic-clonic epilepsy, which responded well to valparin therapy. Structural brain abnormalities can occur, especially in individuals with NIPBL variants. NIPBL is known to regulate cortical neuron migration in mice [25] and, indeed, cortical malformations have been described in individuals with CdLS caused by NIPBL variants [26], as have small callosal bodies, white matter abnormalities, cerebellar anomalies, and brainstem abnormalities. Individuals with CdLS may have specific deficits in executive function beyond what is expected given the level of intellectual disability, especially in mental flexibility and visual short-term memory [27]. Hence, evaluating their cognitive strengths and weaknesses and
structuring their environment accordingly is of utmost importance. In a study of 42 individuals with NIPBL variants, a (fairly weak) positive correlation was observed between chronological age and behavioral difficulties [28], and statistically significant correlations were found between chronological age and measures of interest and pleasure, and insistence on sameness [29], indicating older individuals exhibit more difficulties. Parents should be counseled about prenatal diagnosis in the subsequent pregnancy.

Prenatal molecular testing can be performed on samples obtained from chorionic villous sampling or amniocentesis or by testing embryonic cells obtained through in vitro fertilization [18].

The major indications for prenatal diagnostics are an earlier child with CdLS, a new pregnancy in a family with a known genetic alteration in a CdLS gene, or, as occurs most frequently, no family history but features suggestive of CdLS on fetal ultrasonography [18]. They usually have a vast range of health problems, making it important for the pediatrician to be aware of the child's special needs. The multidisciplinary treatment approach is the key to success in managing children with CdLS.

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References


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