Optic neuritis in a child with biotinidase deficiency: case report and literature review

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Abstract: Optic atrophy has often been reported in children with biotinidase deficiency. The visual prognosis is usually poor. This report is of a 6-year-old boy with an early onset of biotinidase deficiency who presented with acute profound visual loss in both eyes. Fundoscopy revealed swollen discs in both eyes, and the imaging was consistent with bilateral optic neuritis. He was treated with systemic corticosteroid, and commenced on oral biotin. The final visual outcome was promising.

Keywords: optic neuritis, children, biotinidase deficiency

Introduction

Acute optic neuropathy is an inflammatory disorder of the optic nerve. Childhood optic neuritis is an uncommon condition which differs from adult onset optic neuritis. It is frequently associated with systemic infections such as measles, mumps, chicken pox, pertussis, infectious mononucleosis, and immunizations. Less common causes include multiple sclerosis or part of a more diffused demyelinating disorder such as acute disseminated encephalomyelitis or neuromyelitis optica.

Biotinidase deficiency is an autosomal recessive inborn error of biotin metabolism. Poor vision secondary to optic atrophy in patients with biotinidase deficiency has been described in the literature. 1–7 This report is on a young boy with biotinidase deficiency who suffered an acute attack of optic neuritis in both eyes. Rapid clinical diagnosis and prompt management are essential in preventing devastating visual loss in this rare entity.

Case report

A 6-year-old Malay boy who had been diagnosed with biotinidase deficiency endured sudden reduced vision in both eyes within a period of 5 days. His teacher noted that he had difficulty with colors during an art lesson. His mother observed that he bumped into objects at home.

Based on further questioning, it was discovered that he had retro-orbital pain and a headache. There was no similar history in the past. He had no history of fever, symptoms of upper respiratory tract infection, recent vaccination, bleeding tendencies, or trauma.

He was diagnosed with biotinidase deficiency at 18 months of age when he presented with generalized impetigo, delayed developmental milestone, and recurrent episodes of seizure. His plasma biotin concentration was confirmed as low. His elder brother

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slowly. He was also recommenced on oral biotin 10 mg daily.

1 mg/kg daily for 11 days. The dosage was tapered down
remarkable improvement with the above treatment.
The optic nerve functions were monitored closely. He showed
1 month. Color vision and red desaturation were subsequently
evacuation, and was prescribed a metered-dose-inhaler of
salbutamol. There had been no seizure attacks for the past
3 years and his anticonvulsant treatment was discontinued.

On examination, visual acuity was 1/160 in both eyes. The
anterior segment examination was unremarkable in both eyes
except for the presence of a mild relative afferent pupillary
defect in the right eye. Funduscopy revealed swollen and
hyperemic optic discs in both eyes (Figure 1A and B). The
retina was normal in both eyes. Color vision and red desatura-
tion were generally reduced in both eyes. Visual field assess-
ment was not performed as the patient was uncooperative.

The patient was well oriented and afebrile. He was intel-
lectually normal. He had a steady gait. The other cranial nerve
examinations were normal, and there was no lymphadenopathy
elicited. Both lungs were clear, with no sign of exacerbation of
bronchial asthma. His hair and hearing assessments were normal.
Examination of the lower limbs revealed numerous scars from
previous impetigo, healed eczema, and dry scaly skin.

Blood investigations showed normal white cell count.
There was no evidence of leukocytosis or eosinophilia.
Erythrocyte sedimentation rate was raised to 30 mm during
the first hour. Serology screening for toxoplasmosis and
herpes infection was negative. Antinuclear antibody screen-
ing and skin tuberculin tests were also negative.

The biotinidase activity in dried blood spots was less
than 0.1 pmol/min/µL (normal range: 6.3–9.3 pmol/min/µL).
Plasma levels of 3-hydroxyisovalerate and 3-hydroxypropio-
plate were increased. Similarly, the urinary levels of organic
acids 3-hydroxyisovalerate and 3-hydroxypropionate were
also raised.

Visual evoke potential was delayed bilaterally. Magnetic
resonance (MR) imaging of orbit was consistent with optic
neuritis. The brain imaging was normal (Figure 2A and B).
Cerebrospinal fluid analysis was within normal range for cell
counts, protein, sugar, and gamma globulin levels.

He was started on intravenous methylprednisolone
125 mg 6-hourly for 3 days, followed by oral prednisolone
1 mg/kg daily for 11 days. The dosage was tapered down
slowly. He was also recommenced on oral biotin 10 mg daily.
The optic nerve functions were monitored closely. He showed
remarkable improvement with the above treatment.

His visual acuity improved to 6/7.5 in both eyes after
1 month. Color vision and red desaturation were subsequently
improved. Both fundi showed normal and a well-defined margin
of the optic discs (Figure 1C and D). The urinary organic acid
profile returned to normal 1 month after oral biotin therapy.
His case was followed up for 1 year and visual acuity in both
eyes remained good with no sign of recurrence.

Discussion

Deficiency of biotinidase enzyme causes loss of biocytin
in urine and progressive depletion of biotin which results
in multiple carboxylase deficiency. It is characterized by
neurological and cutaneous manifestations, and metabolic
abnormalities. Classically, it is presented with hypotonia,
seizures, ataxia, dermatitis, alopecia, and recurrent infections
during the first year of life.8 Progressive neurological damage
such as sensorineural hearing loss, optic atrophy, ataxia, and
mental retardation has been described in older children.8–10

Salbert et al described 51% ophthalmic abnormalities
in 78 symptomatic children with biotinidase deficiency.1
This included 30% infections, 13% optic neuropathies, 13%
motility disturbances, 4% retinal pigment changes, and 1%
pupillary findings, while the most common features were
optic atrophy and keratoconjunctivitis.1

Table 1 summarizes cases of optic neuropathy in children
with biotinidase deficiency published in PubMed12 from
1997 to 2011. Our search was made on keywords of “optic
neuropathy,” “child,” and “biotinidase deficiency”. We
documented seven patients, and reviewed their age, gender,
ocular features, funduscopy, systemic features and onset of
biotinidase deficiency, treatment received, and their final
visual outcome. They presented with ocular signs at ages
ranging between 5 and 15 years old. Six of seven patients
(85.7%) presented with optic atrophy in both eyes, while the
remaining one patient (14.3%) displayed normal looking
optic discs bilaterally.2–5,9–10 Unfortunately, the final visual
outcome was poor in 71.4% (five patients).2–5

In contrast, Puertas et al reported a 12-year-old boy with
biotinidase deficiency and presented symptoms of acute
retrobulbar neuritis in both eyes.9 Similarly, our patient also
exhibited acute papillitis in both eyes. These two patients were
identified during the acute attack, and they were successfully
treated with a combination of intravenous corticosteroid and
oral biotin. Their final visual acuity was satisfactory.

Our patient had poor compliance to biotin therapy.
His biotinidase activity was confirmed low (less than
0.1 pmol/min/µL), and this supports the diagnosis of
biotinidase deficiency. Raised urinary levels of organic
acids 3-hydroxyisovalerate and 3-hydroxypropionate suggest
multiple carboxylase deficiencies in our patient.

and two paternal uncles were also diagnosed with biotinidase
deficiencies. Both parents had a consanguinity marriage.

He was started on a 6 mg daily dose of oral biotin,
however the compliance was poor. He was also suffering
from bronchial asthma with a history of recurrent acute
exacerbation, and was prescribed a metered-dose-inhaler of
salbutamol. There had been no seizure attacks for the past
3 years and his anticonvulsant treatment was discontinued.

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Figure 1 (A and B) fundus photographs show bilateral swollen disc on presentation. (C and D) resolved bilateral disc swelling at one month after treatment.

Figure 2 (A and B) MRI of the optic nerve shows expansion with slight increase signal intensity on T2-weighted image involving both optic nerves (left) and enhancement in T1-weighted image post IV contrast. (C–F) MRI of the optic nerve (white arrow) and both optic nerves (white double arrowheads) shows expansion and enhancement on T1-weighted image and post gadolinium images.

Abbreviation: MRI, magnetic resonance image.
<table>
<thead>
<tr>
<th>Authors</th>
<th>Age/gender</th>
<th>Ocular features</th>
<th>Funduscopic</th>
<th>Biotinidase deficiency</th>
<th>Treatment</th>
<th>Ocular outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rahman et al&lt;sup&gt;2&lt;/sup&gt;</td>
<td>5-year-old, female</td>
<td>VA reduced to 0.5/60 OU</td>
<td>Bilateral optic atrophy</td>
<td>5 years</td>
<td>Intractable seizures, hypotonia, ataxia, hearing loss, dermatitis, and alopecia</td>
<td>Oral biotin 10 mg daily</td>
</tr>
<tr>
<td>Lott et al&lt;sup&gt;3&lt;/sup&gt;, Wolf et al&lt;sup&gt;4&lt;/sup&gt;</td>
<td>10.5-year-old, male</td>
<td>Loss of visual acuity, scotoma, double vision</td>
<td>Bilateral optic atrophy</td>
<td>13.5 years</td>
<td>Limb weakness, spastic paraparesis and fatigue</td>
<td>Biotin treatment</td>
</tr>
<tr>
<td>Wolf et al&lt;sup&gt;4&lt;/sup&gt;, Ramaekers et al&lt;sup&gt;5&lt;/sup&gt;</td>
<td>10-year-old, male</td>
<td>VA reduced to 20/100 OU, cecocentral scotoma OU</td>
<td>Bilateral optic atrophy</td>
<td>15 years</td>
<td>Spastic paraparesis, limb weakness, ptosis and fatigue</td>
<td>Started oral biotin 10 mg daily at the age of 15-year-old</td>
</tr>
<tr>
<td>Wolf et al&lt;sup&gt;4&lt;/sup&gt;</td>
<td>15-year-old, female</td>
<td>Not tested</td>
<td>Bilateral optic atrophy</td>
<td>15 months</td>
<td>Seizure and rash</td>
<td>Biotin treatment</td>
</tr>
<tr>
<td>Wolf et al&lt;sup&gt;4&lt;/sup&gt;</td>
<td>8-year-old, male</td>
<td>Loss of visual acuity, scotoma, double vision</td>
<td>Bilateral optic atrophy</td>
<td>22–24 months</td>
<td>Seizure, alopecia and ataxia</td>
<td>Biotin treatment</td>
</tr>
<tr>
<td>Puertas et al&lt;sup&gt;9&lt;/sup&gt;</td>
<td>12-year-old, male</td>
<td>VA reduced 0.1 OU</td>
<td>Normal optic disc in both eyes</td>
<td>12 years</td>
<td>Sensorineural hearing loss, asthma, dermatitis, alopecia and retrolubular neuritis</td>
<td>IV corticosteroid, oral biotin 5 mg 12 hourly</td>
</tr>
<tr>
<td>Yang et al&lt;sup&gt;10&lt;/sup&gt;</td>
<td>5-year-old, male</td>
<td>VA reduced to 20/50 OU and bilateral blepharoconjunctivitis</td>
<td>Bilateral optic atrophy</td>
<td>7 years</td>
<td>Progressive movement disorders with spastic gait, intermittent ataxia, tremor, lehargy, anorexia, intermittent vomiting, and alopecia</td>
<td>Started oral biotin 20 mg/daily at the age of 14-year-old</td>
</tr>
<tr>
<td>Hayati et al&lt;sup&gt;11&lt;/sup&gt;</td>
<td>6-year-old, male</td>
<td>VA reduced to 1/160 OU</td>
<td>Bilateral swollen optic disc</td>
<td>18 months</td>
<td>Recurrent seizure</td>
<td>IV corticosteroid, restarted oral biotin 10 mg daily.</td>
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Abbreviations: OU, both eyes; VA, visual acuity.
Figure 3 (A) Possible mechanism explains retinal ganglion cell apoptosis and acidosis in biotinidase deficiency. (B) Possible mechanism describes neuronal damage.
**Figure 4 (A)** Possible mechanism improvement of energy production to retinal ganglion cell and resolved acidosis after biotin therapy in biotinidase deficiency. (B) Possible mechanism limits neuronal damage.
There are possible explanations regarding the occurrence of optic neuritis in children with biotinidase deficiency (Figure 3A and B). Decreased amount of biotin causes reduction of methylcrotonyl-CoA carboxylase activity. This leads to the accumulation of 3-methylcrotonyl-CoA, and subsequently the formation of 3-methylcrotonic acid. This will be modified later to 3-hydroxyisovaleric acid. Accumulation and storage of these two acids in the retinal ganglion cells directly affects the intracellular metabolism and leads to retinal ganglion cells apoptosis.

Due to high energy demand and the long course of their axons, retinal ganglion cells are very susceptible to intracellular metabolic defect. In biotinidase deficiency, the decreased amount of biotin causes a lower activity of propionyl-CoA carboxylase. This leads to the accumulation of propionyl-CoA which can be converted later into propionic acid resulting in acidosis.

In addition to that, propionyl-CoA can be processed into methylcitrate by an enzyme in the citric acid cycle. Methylcitrate blocks the normal function of citric acid. This causes limited energy production in the retinal ganglion cells and subsequently, results in cell apoptosis. The above factors may lead to optic neuropathy and consequently result in irreversible optic atrophy.

Corticosteroid is the standard modality for treatment of optic neuritis. It acts to decrease inflammation by suppressing migration of polymorphonuclear leukocytes and reversing increased capillary permeability. The Optic Neuritis Treatment Trial describes the recommended treatment for adult onset optic neuritis. Our patient tolerated the corticosteroid therapy well.

Children with profound biotinidase deficiency have less than 10% of mean serum biotinidase enzyme activity, while children with partial biotinidase deficiency have 10%–30% of mean serum biotinidase enzyme activity. Both profound and partial biotinidase deficiency are usually identified by newborn screening. Unfortunately, this test is not performed routinely in our country.

Optic neuritis is uncommon in children with biotinidase deficiency. It is important for ophthalmologists and pediatricians to be aware of this diagnosis when a child with biotinidase deficiency presents symptoms of acute visual loss. Prompt diagnosis and early treatment is mandatory to avoid irreversible visual loss in these children. The combination of systemic corticosteroid and biotin treatment seemed an effective treatment with our patient.

Disclosure

The authors report no conflict of interest in this work.

References