Case Report
Update Review and Clinical Presentation in Congenital Insensitivity to Pain and Anhidrosis

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Introduction. Congenital insensitivity to pain and anhidrosis (CIPA) or hereditary sensory and autonomic neuropathy type IV is an extremely rare syndrome. Three clinical findings define the syndrome: insensitivity to pain, impossibility to sweat, and mental retardation [3, 4]. Only a few hundreds of cases of CIPA have been recently published worldwide [5, 6]. This condition occurs with an incidence of 1 in 125 million newborns [7].

The pathogenesis of CIPA is characterized by a genetic loss-of-function mutation of the NTKRI gene ( locus 1q 21-22) [8, 9]. Multiple new mutations have been progressively described [10–16]. NTRK1 mutations imply an alteration in TrkA, a NGF receptor. NGF is involved in surveillance of nociceptive sensory neurons and sympathetic autonomic neurons and collaborates in the activation and homeostasis of other cellular types so that a NTRK1 mutation will cause deficient development of [17–20]

(1) the afferent somatic sensory system for pain and temperature, located in the dorsal root ganglion sensory neurons,

(2) the autonomic sympathetic neuronal system, which implies loss of the innervation of eccrine sweat glands by sympathetic neurons,

(3) the central nervous system,

(4) the bidirectional communication between the immune system and the nervous system (NGF has a relevant role in the signal pathway of B lymphocytes through three processes: Trk A phosphorylation, cytoskeleton assemblage, and MAP kinase activation).

The molecular alteration in the function of NGF in turn also alters the normal process of fracture consolidation [21].
Normal osteoblast/osteoprogenitor differentiation and proliferation are hindered, tending to result in fibroblast differentiation of multipotent stromal mesenchymal cells and periosteal cells.

Bone metabolism is also affected by the lack of nociceptive fibers, present not only in the skin but also in the skeletal system [22]. Due to the trophic role that nociceptive fibers may play in the skeletal system, bone fractures are very common [23].

2. Case Presentation

Medical record and radiographic data of the present case were reviewed and reported in a study approved by the department of documentation of our hospital. The patient's parents also gave their consent. A thorough review of the PubMed literature on CIPA and associated medical conditions mentioned in this paper was performed (Table 1). This case report is an illustrative example of a patient affected by CIPA.

We present a case involving a seven-year-old, female child of Spanish nationality. She had been evaluated in another center for episodes of recurrent fever. After a long diagnostic process including a pertinent genetic study which detected two mutations in the NTRK1 gene responsible for CIPA, she was diagnosed with the syndrome [8]. Her parents were healthy, and no consanguinity was present.

Clinical exploration revealed absence of a pain response, recurrent episodes of fever, sweating deregulation, mental retardation, cutaneous autolesions, fracture without consolidation, avascular necrosis (Figure 1), demineralized bones, generalized osseous destruction (Figure 2), warm and dry skin with thickening of the soles and palms, and lower limb edema (Figures 3(a), 3(b), and 3(c)) [5, 24–26].

The patient was referred to our center four months after fracture of the middle shaft of the right tibia. Radiologic signs of hypertrophic pseudoarthrosis were present (Figure 4). An elastic intramedullary nailing was carried out [27]. Complete radiological consolidation of the fracture was achieved five months after the surgery (Figure 5).

In the following months, several fractures occurred, including a fifth metatarsal fracture in the right foot (Figure 1) and a fourth metatarsal fracture in the left foot, a right femoral middle shaft fracture that was surgically treated with good results (Figures 6(a), 6(b), and 6(c)), and an epiphysiolyis at the distal shaft of the right tibia. In CIPA, due to the alteration of the bone fracture metabolism, hypertrophic bone callus (Figures 1, 4, and 6(c)) and pseudoarthrosis (Figure 4) are very common. In the present patient, bone consolidation was only achieved when a surgical technique was applied.

During this period of time with recurrent fractures, treatment with bisphosphonates was started. A dose of 1 mg/Kg/day during 3 consecutive days of intravenous pamidronate was administered every four months, for one year. We obtained good results in preventing new fractures at upper and lower limbs, skull, and spine bones at 5 years of follow-up. No adverse effects were seen regarding pamidronate infusion or during the follow-up.

At 5 years of follow-up, patient has progressed.
Table 1: Thorough review of the PubMed literature on CIPA and associated medical conditions mentioned in this paper was performed.

<table>
<thead>
<tr>
<th>References</th>
<th>Year of publication</th>
<th>Particularity of the observation and remarks for each reading</th>
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<tbody>
<tr>
<td>Dearborn [1]</td>
<td>1932</td>
<td>First reference, in literature, to a similar disease</td>
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<tr>
<td>Nishida [3]</td>
<td>1951</td>
<td>Three clinical representative findings: insensitivity to pain, inability to sweat, and mental retardation</td>
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<tr>
<td>Tunçbilek et al. [4]</td>
<td>2005</td>
<td>Only 32 cases have been published worldwide</td>
</tr>
<tr>
<td>Rosenberg et al. [5]</td>
<td>1994</td>
<td>Only some hundreds of cases have been published worldwide</td>
</tr>
<tr>
<td>Daneshjou et al. [7]</td>
<td>2012</td>
<td>Incidence 1 in 125 million newborns</td>
</tr>
<tr>
<td>Indo et al. [9]</td>
<td>1997</td>
<td>Autosomal recessive disorder</td>
</tr>
<tr>
<td>Indo et al. [8]</td>
<td>1996</td>
<td>Not only autosomal recessive inheritance, but also uniparental disomy (non-Mendelian inheritance of autosomal recessive disease from a single carrier parent, as the exposed case)</td>
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<tr>
<td>Indo [25]</td>
<td>2002</td>
<td>A very profuse resume of clinical and genetic characteristics of CIPA</td>
</tr>
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<td>Indo [18]</td>
<td>2010</td>
<td>NGF receptor failure causes a deficient development of dorsal root neurons (pain and temperature sensory system)</td>
</tr>
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<td>Tanaka et al. [20]</td>
<td>1990</td>
<td>Central nervous system</td>
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<td>Schwarzkopf et al. [17]</td>
<td>2005</td>
<td>The signal pathway of B lymphocytes</td>
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<td>Melamed et al. [21]</td>
<td>2004</td>
<td></td>
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<tr>
<td>Grills and Schuijers [24]</td>
<td>1998</td>
<td>NGF function disruption also causes an altered process of fracture consolidation</td>
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<tr>
<td>Fruchtman et al. [26]</td>
<td>2013</td>
<td>Descriptive clinical presentation including morbidity conditions (some of these clinical facts are also present in the case reported)</td>
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<tr>
<td>Yang et al. [27]</td>
<td>2013</td>
<td></td>
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<tr>
<td>Jarade et al. [35]</td>
<td>2002</td>
<td>Ocular manifestations</td>
</tr>
<tr>
<td>Oliveira et al. [40]</td>
<td>2009</td>
<td>Anaesthetic considerations</td>
</tr>
<tr>
<td>Abdulla et al. [33]</td>
<td>2014</td>
<td>Heterotopic ossification and callus formation following fractures, eventually Charcot's joint</td>
</tr>
<tr>
<td>Schreiber et al. [41]</td>
<td>2005</td>
<td>Insulin-related difficulties</td>
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demineralization. Surgical fracture repair allows for early weight bearing, diminishing the risk of further osteopenia, which is also usually present in these patients as a part of their associated neurogenic arthropathy (Figure 3) [21].

For all of these reasons, we recommend early surgical treatment of fractures. It allows for more rapid functional recovery, reducing the risk of accelerated osteopenia due to immobilization.
The use of bisphosphonates in patients affected by CIPA had never been mentioned before in literature. Due to our previous good experience with pamidronate in treating osteoporotic fractures for disuse in children with different medical conditions [29, 30], we made a therapeutic approach with pamidronate as a compassionate use in this child. We obtained good results in preventing new fractures.

These two therapeutic observations might be relevant in the absence of specific treatment for CIPA. However, we may not forget that further studies addressing CIPA management are needed to provide more rigorous and scientific conclusions.

CIPA may present various signs and symptoms that can be misleading. The differential diagnoses of this pathology include radicular hereditary sensory neuropathy (HSN I); hereditary sensory and autonomic neuropathy (HSN II); familial dysautonomia or Riley-Day syndrome (HSN III) [31]; congenital indifference to pain (HSN V) [32]; and Lesch-Nyhan syndrome. Corneal ulcers are also relatively frequent in patients with CIPA. A differential diagnosis of neurotrophic keratitis may be taken into consideration [33, 34]. Among all these diagnostic possibilities and according to Raspall-Chaure [29], CIPA must be the first diagnostic hypothesis when assessing a patient with insensitivity to pain, anhidrosis, and self-mutilation.

According to literature, the first step in the diagnosis of CIPA syndrome is consideration of the clinical presentation based on the combination of three basic signs: insensitivity to pain, anhidrosis, and mental retardation [3, 4]. Other possible signs may be associated: impaired temperature sensation [5], facial alterations [6], mandibular osteolysis [7], dental caries [6], and premature tooth loss [6]; repetitive soft tissue and osseous infections of hematogenous origin [33], mainly caused by S. aureus [25]; self-mutilating behavior [7]; occasional microcephaly [5, 24]; urine and fecal incontinence [11]; growth disturbances; and heterotopic ossification [7, 35, 36].

Neurological laboratory tests may provide additional information. Short-latency somatosensory evoked potentials show marked prolongation of the central conduction time [19] and microneurography reveals abnormal activity of somatic A-delta and C fibers in the nerves of the skin [6, 37, 38]. A negative sympathetic skin response may also be helpful in the diagnosis due to the lack of sudomotor nerves in skin biopsy [38].

Pharmacologic tests that evaluate autonomic function are also useful. The Mecholyl test produces prompt pupillary miosis [24], pain test results abnormal [6, 13, 24], there is an absence of a flare reaction to the histamine test [24] (although we may find some normal responses to subdermal histamine injection) [11], and the sweat test using pilocarpine reveals a disruption of sweat gland function. Histopathologic evaluation shows a hyperplastic epidermis with acanthosis and hyperkeratosis and a decreased amount of sweat and sebaceous glands [6].

Finally, molecular evaluation that reveals mutations of the NTKRI gene provides a definitive diagnosis [19, 24].

About the anesthetic considerations [39, 40], although pain stimuli are absent, anxiety associated with surgical procedures may generate stress and consequent hemodynamic instability. It is necessary to minimize preoperative apprehension and anxiety with the use of sedatives. Also the autonomic response to surgery is inconsistent and erratic, which results in difficulty determining the necessary anesthetic doses in advance. Finally, temperature control is crucial. Malignant hyperthermia or hypothermia may be lethal.

NGF-TrkA pathway has a role in the morphogenesis of the endocrine pancreas, in insulin secretion in vitro, and in insulin secretion in response to glucose. Patients with CIPA present with alterations of the first phase of insulin secretion [41].

The similarities between CIPA and reflex sympathetic dystrophy are very interesting. Both are characterized by neurogenic inflammation, skin alterations with vasomotor disruption, and osteopenia.
Some authors have focused on establishing a specific treatment for complex regional pain syndrome by studying the role of receptor tyrosine kinase for NGF in patients with CIPA [29]. The high incidence of infections in patients with CIPA is also problematic. Skin and deep bone infections are the most common types, and *Staphylococcus aureus* is the most commonly involved pathogen.

Resistance to antibiotics is a frequently occurring limitation in the treatment of these patients [25]. Temperature deregulation may cause recurrent fever, which may lead to death if not recognized early.

Other complications such as trauma or soft tissue/bone infection may decrease condition of the survival rate, although all are treatable conditions if diagnosed in a timely manner [7].

The best therapeutic approach to patients with CIPA appears to be based on prophylactic measures such as braces for early weight bearing in nonsurgical fractures and accurate follow-up to avoid missing complications. We propose an unusual treatment challenge, with an early surgical treatment for long bone fractures and early use of bisphosphonates as follows.

Therapeutic proposals are as follows:

1. Surgical fracture repair to achieve an early functional recovery that avoids a final destructive situation.

2. Bisphosphonates use to manage osteoporosis.

Addressing the cause of CIPA as opposed to solely symptomatic treatment seems to be the optimal therapeutic approach. If CIPA results from loss-of-function mutations in the NTRK1 gene encoding TrkA, then molecular treatment involving a receptor tyrosine kinase for NGF would be the most effective therapeutic technique.

**Disclosure**

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**Conflict of Interests**

None of the authors has directly received research funding and/or has potential conflict of interests.

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**References**


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