Experience with cinacalcet in familial hypocalciuric hypercalcemia resulting from inactivating mutations in calcium sensing receptor (CaSR)

Abstract

Background: Familial hypocalciuric hypercalcemia (FHH) is usually characterized by asymptomatic hypercalcemia, hypocalciuria and normal PTH. The diagnosis of FHH is often incidental and seldom require any medical intervention.

Patient description: We are reporting Case 1 an 8 years old female incidentally diagnosed with hypercalcemia at the age of 3 years. She had complaints of constipation, polyuria and nocturia. Urinary calcium excretion was extremely low. A trial of Cinacalcet was initiated at 7 years of age, which provided complete symptomatic relief, increased urinary calcium excretion and decreased serum calcium levels. Case 2 is a 7 year old male was evaluated for leg cramps and high serum calcium level (11.3 mg/dL). The patient complained of occasional leg pains. Case 3 is a 6 year old boy the younger brother of Case 2 was referred for high serum calcium level (11.1 mg/dL). Family history is significant for hypercalcemia in their father. After confirmation of a CaSR mutation, both were started on Cinacalcet 30mg twice a day. Over the course of treatment the calcium levels were normalized and leg pain resolved in case 2.

Result: Mutation analysis of the CASR gene in case 1 revealed a misssense mutation (CGA-CAA, R185Q). In case 2 and case 3 a heterozygous mutation (GCC->ACC, A194T) was found in the extracellular domain in the CaSR encoded by exon 4.

Conclusion: We are reporting a novel mutation of CaSR in a severe presentation in case 1 and lack of symptoms in parents would suggest a de-novo mutation. In cases 2 and 3 the mutation was probably inherited from the father, consistent with autosomal dominant pattern of inheritance of FHH. The use of Cinacalcet successfully improved calcium levels and alleviated all the associated symptoms.

Abbreviations: FHH, Familial hypercalcemia hypocalciuria; CaSR, Calcium Sensing 20 Receptor; PTH, Parathyroid Hormone

Introduction

Familial hypocalciuric hypercalcemia (FHH) is a metabolic abnormality that is usually asymptomatic. Occasionally, patients may complain of weakness, fatigue, polydipsia, and psychiatric disorders. It is present in about 2% of patients and the survival rates of affected patients are no different than those in the general population. FHH is an autosomal dominant disorder with high penetrance. Most people with FHH have heterozygous loss of function mutations in the Calcium Sensing Receptor (CaSR) gene. Common mutations locus associated with FHH occur on chromosomes 3 and 19 (19p13.3 and 19q13). The CaSR is expressed in the parathyroid, thyroid, kidney, gut, pancreas, and bone. In a CaSR with normal function, high Ca concentrations stimulate the CASRs, which signal calcitonin production and urinary Ca excretion, while suppressing the synthesis of parathyroid hormone (PTH). The CaSR mutation in FHH results in reduced sensitivity of CaSRs to circulating calcium. The decreased receptor sensitivity in the distal nephron of the kidney results in hypocalciuria. As a result of the decreased sensitivity to calcium, a normal level of calcium is perceived as low by the parathyroid and PTH is released to bring up the calcium concentration. Thus the kidney responds by reducing calcium excretion in the urine as the CaSR has reduced sensitivity to high circulation calcium levels.

The urinary calcium: creatinine ratio is important to distinguish FHH from primary hyperparathyroidism, both of them present with hypercalcemia, and hence the urinary calcium: creatinine ratio should be calculated. Hypocalciuria is an important distinguishing feature of FHH but not primary hyperparathyroidism. If the ratio is less than or equal to 0.01, then the result is suggestive of FHH. However, 20% of FHH patients have been reported to have a ratio greater than 0.01. So, urine concentrations of calcium can be normal in FHH, while the urine is diluted in primary hyperparathyroidism. Whereas the urinary calcium: creatinine ratio in primary hyperparathyroidism is higher A CaSR mutation analysis can be used as an essential tool to arrive at a diagnosis of FHH. A homozygous loss of function mutation of CaSR results in neonatal severe primary hyperparathyroidism (NS-PHT), which has a more severe presentation than FHH. This condition presents in the neonatal period with hypercalcemia, parathyroid gland hyperplasia, hypotonia, dehydration, bone demineralization, fractures, polyuria, failure to thrive, and respiratory distress. There currently is no FDA (Food and Drug Administration) approved treatment for FHH or NSPHT. Parathyroidectomy is an effective treatment option in NSPHT, but with significant risk for complications. FHH is often left untreated and patients are counseled regarding the benign
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Case 2: A 7 year old male was sent to the Pediatric Endocrine clinic by pediatrician for the evaluation of the continuous high serum calcium level (11.3 mg/dL). The patient complained of occasional leg pains. He was diagnosed with bipolar disorder prior to the visit, which was well controlled on medications. Patient had a DEXA scan which did not show any osteoporosis or osteopenia (Bone mineral density Z score: +1.6 to +1.8). Renal Sonogram was also normal. Laboratory evaluation showed serum phosphorus of 4.2 mg/dL, a high intact PTH (69 mg/dL), and normal vitamin D-25 and vitamin D-1,25 levels (see Table). He was started on Cinacalcet 30mg twice a day. Over the course of treatment his calcium levels were normalized (9.5 mg/dL). Case 2 was followed for 4 years with normocalcemia.

Case 3: A 6 year old boy younger brother of Case 2 was referred to our clinic by pediatrician for the evaluation of the high serum calcium level (11.1 mg/dL). At the time of presentation he had no acute complaints and past medical history was non-contributory. Family history is significant for hypercalcemia in the father who appeared asymptomatic. Physical exam was normal. Renal and Thyroid sonogram were normal. Patient had a DEXA scan on which did not indicate any osteoporosis or osteopenia. MRI of head and neck were also performed and did not show anything suggestive of a parathyroid adenoma. The laboratory evaluation showed serum phosphorus of 3.7 mg/dL, normal PTH (44pg/mL) and normal vitamin-D 25 and vitamin-D 1, 25 (Table 1). Patient was started on Cinacalcet 30mg twice a day, and his calcium concentrations decreased to normal ranges (9.8 mg/dL). Serum Phosphorus increased (4.5 mg/dL) and PTH decreased slightly to 40 pg/mL. Case 3 was followed for 4 years as well and has been normocalcemic.

Table 1 The table changes in the biochemical and hormonal profile of the 3 patients with FHH, who were treated by Cinacalcet

<table>
<thead>
<tr>
<th>Norms</th>
<th>Prior cinacalcet</th>
<th>After cinacalcet</th>
<th>Prior</th>
<th>After</th>
<th>Prior</th>
<th>After</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calcium</td>
<td>8.9-10.4 mg/dL</td>
<td>14.9</td>
<td>10.3</td>
<td>11.3</td>
<td>9.5</td>
<td>11.1</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>4.5-8 mg/dL</td>
<td>2.9</td>
<td>3.5</td>
<td>4.2</td>
<td>5.1</td>
<td>3.7</td>
</tr>
<tr>
<td>Magnesium</td>
<td>2.3-5 mg/dL</td>
<td>2.4</td>
<td>-</td>
<td>2.5</td>
<td>-</td>
<td>2.3</td>
</tr>
<tr>
<td>Intact-PTH</td>
<td>9.59 pg/mL</td>
<td>121-169</td>
<td>93.7</td>
<td>69</td>
<td>120.6</td>
<td>40</td>
</tr>
<tr>
<td>Urinary calcium/creatinine ratio</td>
<td>0.001</td>
<td>0.02</td>
<td>1.1</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Vitamin D 25-Hydroxy</td>
<td>32-100 ng/mL</td>
<td>18</td>
<td>27</td>
<td>31.9</td>
<td>45</td>
<td>40</td>
</tr>
<tr>
<td>Vitamin D 1, 25 Dihydroxy</td>
<td>27-71 pg/mL</td>
<td>51</td>
<td>-</td>
<td>65</td>
<td>57</td>
<td>65</td>
</tr>
<tr>
<td>Alkaline Phosphatase</td>
<td>93-309 ng/mL</td>
<td>-</td>
<td>374</td>
<td>42</td>
<td>-</td>
<td>201</td>
</tr>
</tbody>
</table>

Figure 1 Shows the decrease in serum calcium levels of the 3 cases with CaSR mutations which were treated with Cinacalcet. Case was treated with the high dose upto 240 mg/day of Cinacalcet divided twice a day. The other 2 case were treated with 60 mg/day divided twice a day.

Citation: Anthonypillai J, Sinha SK, Mamkin A, et al. Experience with cinacalcet in familial hypocalciuric hypercalcemia resulting from inactivating mutations in calcium sensing receptor (CaSR). Endocrinal Metab Int J. 2018;6(2):137–140. DOI: 10.15406/emij.2018.06.00167
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Materials and methods

Hormonal testing

Vitamin D, 25-Hydroxy and Vitamin D, 1, 25-Dihydroxy were measured by HPLC-MS mass spectrometry. The Intact Parathyroid Hormone (iPTH) was measured by immunoenzymetric assay. DNA isolation and sequencing DNA was extracted from peripheral leukocytes and all exons of the CaSR genes were amplified by PCR using primer pairs and standard conditions. Written informed consent and assent to do the mutational analysis was obtained from the parents and subjects respectively. The study was approved by the Institutional Review Board of University of California, San Francisco.

Results

Mutation analysis of the CASR gene from peripheral leukocyte from case 1 revealed a previously described heterozygous missense mutation (CGA→CAA, R185Q). In case 2 and case 3 a heterozygous mutation (GCC→ACC, A194T) was found in the extracellular domain in the CaSR encoded by exon 4 of the CASR gene and in the unaffected father. The mother’s genomic DNA is of normal sequence.

Discussion

Calcimimetics are allosteric modulators of CaSR that make the mutant receptor more sensitive to serum calcium levels. This increased sensitivity results in a reduction of PTH and serum calcium levels. Cinacalcet is currently the only drug in this class approved for human use and is currently indicated for treatment of secondary hyperparathyroidism in patients with chronic kidney disease and for the treatment of hypercalcemia in patients with primary hyperparathyroidism.9–18 Cinacalcet has been used in cases with FHH but currently is not FDA approved for treatment of FHH or NSPHT. Cinacalcet is usually well tolerated the most reported side effects are nausea and vomiting.12,17

FHH is considered a benign entity because it is often asymptomatic and cardiac abnormalities or cardiac dysrhythmias are not associated with the high serum calcium levels.1 However, in our cases, patients complained of polyuria, nocturia, enuresis, and constipation which were all relieved on administration of Cinacalcet and correction of hypercalcemia. These symptoms although not as serious as the cardiac manifestations, but were severe enough to the effect the normal functioning of daily activities. The symptoms reported by patients were often restricting and at times debilitating.

In all three of our cases, serum calcium levels were reduced to within normallimits and serum phosphorus was increased after administration of Cinacalcet. Serum PTH levels were decreased in both cases 1 and 3 after Cinacalcet administration. In addition to our cases, Cinacalcet has been used in both adult and pediatric patients to treat FHH and NSHPT.6–8,19 These results are similar to cases presented in a 37 year old woman with FHH suffering from recurrent pancreatitis due to hypercalcemia.7 Similarly Cinacalcet was used in a 26 year old male with psychosis, hypercalcemia and osteoporosis.6 In a pediatric case report a 6 year old boy with FHH was described with dense tye-panosclerosis involving multiple calcium deposits, whose condition improved with surgical revision and treatment with Cinacalcet which resulted in the surgical scar healing without excessive calcium depo-sits.3

Conclusion

In conclusion, our experience with Cinacalcet in treating pediatric patients with FHH, in addition to the few reports involving adults and a child, support the beneficial effects by alleviation of symptoms of polyuria, nocturia, enuresis and constipation after calcimimetic administration in patients with loss of function mutations of the CaSR. We reported a novel mutation of CaSR in a severe presentation in case 1 and lack of symptoms in parents would suggest a de-novo mutation. Whereas, in the two brothers was probably inherited from the father, consistent with autosomal dominant pattern of inheritance of FHH.

Acknowledgment

None.

Conflict of interest

None to declare.

References


