

A pediatric case of anaphylaxis due to octreotide

Dilek Azkur,¹ Tamer Yoldas,² Muge Toyran¹ and Can Naci Kocabas¹

Summary

Octreotide is an octapeptide that mimics natural somatostatin pharmacologically. It is a potent inhibitor of growth hormone, glucagon and insulin, which is used for treatment of acromegaly, symptomatic treatment of carcinoid tumours, and vasoactive intestinal peptide secreting tumors. It is also used for chylothorax, chemotherapy induced diarrhea and, as it inhibits the exocrine production of pancreatic enzymes, for acute and chronic pancreatitis. Gallbladder stones, diarrhea, nausea, vomiting, hypoglycemia/hyperglycemia, headache, and abdominal discomfort are some of the common adverse effects of octreotide and it may rarely cause anaphylaxis. We present here a child who had chronic pancreatitis and had an anaphylactic reaction to octreotide. To our knowledge this is the first pediatric case of anaphylaxis with octreotide who was successfully desensitized. (*Asian Pac J Allergy Immunol* 2011;29:361-3)

Key words: child, octreotide, anaphylaxis, desensitization, pancreatitis

Introduction

Octreotide is an octapeptide that mimics natural somatostatin pharmacologically. It is a potent inhibitor of growth hormone, glucagon and insulin, which is used for treatment of acromegaly,¹ symptomatic treatment of carcinoid tumours,² and vasoactive intestinal peptide secreting tumors.³ It is also used for chylothorax⁴, chemotherapy induced diarrhea^{5,6} and, as it inhibits the exocrine production of pancreatic enzymes, for acute and chronic pancreatitis.⁷⁻⁹ Gallbladder stones, diarrhea, nausea, vomiting, hypoglycemia /hyperglycemia, headache, and abdominal discomfort are some of the common

adverse effects of octreotide¹⁰ and it may rarely cause anaphylaxis.^{10,11} We present here a child who had chronic pancreatitis and had an anaphylactic reaction to octreotide. To our knowledge this is the first pediatric case of anaphylaxis with octreotide who was successfully desensitized.

Case report

A 12-yr-old white boy was admitted to our hospital with recurrent abdominal pain which was diagnosed as chronic pancreatitis. Besides other treatment strategies, octreotide was one of the drugs used. First octreotide treatment was given intravenously (iv) at 6 mcg/kg/dose once a day for 15 days without any adverse reaction. The patient was discharged from the hospital as he was clinically stable. However, two weeks after the first pancreatitis episode, the patient was readmitted with recurring symptoms and octreotide treatment was started again along with other treatment strategies. During the intravenous administration of the tenth dose of octreotide (1.3 mcg/kg/dose, iv, every 8 hours), he immediately experienced flushing of his face, erythema over the arms, periorbital and perioral swelling, upper airway breathing difficulty, cough and abdominal pain. Following the development of these symptoms, the octreotide infusion was stopped immediately. He did not have hypotension or wheezing but he had flushing and perioral and periorbital angioedema on physical examination. Pheniramine maleate was given by the iv route and his complaints began to subside within 5 minutes and the physical findings returned to normal completely within an hour. Aside from abdominal pain, none of these symptoms were present before the drug was given and the abdominal pain was dramatically augmented during the infusion. The patient was not taking any medication during the octreotide infusion. Treatment with other drugs (antibiotics and proton pump inhibitors) was continued and there was no reaction with these medications. The patient did not have pyrexia, infection, other than pancreatitis, peptic ulcer or malnutrition. Liver and kidney function tests were within normal limits but the serum lipase and pancreatic

From the ¹Department of Pediatric Allergy and Immunology
²Department of Pediatrics Ankara Children's Hematology
 Oncology Training and Research Hospital, Ankara, Turkey
 Corresponding author: Can Naci Kocabas
 E-mail: cankocabas@yahoo.com
 Submitted date: 25/7/2011
 Accepted date: 31/10/2011



Table 1. Desensitization Protocol for Octreotide

	Volume, mL	Infusion Time, min	Time Accumulated	Dose Administered, mcg	Cumulative Dose, mcg
Solution A	100	30	30	1	1
Solution B	100	30	60	5	6
Solution C	100	30	90	10	16
Solution D	100	30	120	20	36
Solution E	100	30	150	40	76

amylase levels were high. The patient developed the same reaction to the subsequent dose of octreotide. Octreotide treatment was stopped again and he was referred to our department of Pediatric Allergy and Immunology for evaluation of an allergic reaction to octreotide. With his symptoms of flushing, angioedema, upper airway obstruction and abdominal pain recurring during octreotide infusions, the patient was suspected of having anaphylaxis to octreotide. The patient had no personal or family history of atopic disease or of hypersensitivity to any drug. Skin prick tests with inhalants and food allergens all showed negative results. Skin prick test at 1/1 concentration and intradermal tests at of 1/1000, 1/100, 1/10, and 1/1 with octreotide were performed five weeks after the anaphylactic reaction. Intradermal testing at concentrations of 1/10 and 1/1 were positive (3 mm and 4 mm respectively, with a negative saline control and positive histamine control of 7x7 mm). The patient was diagnosed as anaphylaxis caused by octreotide hypersensitivity. As the patient had an exacerbation of pancreatitis at the time of the diagnosis, he needed octreotide treatment and desensitization to the drug was considered. The desensitization protocol was carried out with continuous monitoring (pulse-oximetry, non-invasive blood pressure, and heart rate) and appropriate resuscitation equipment and medications at the bedside (Table 1). Premedication with corticosteroids and/or antihistaminics was not given before desensitization. Increasing doses of the drug were given by iv infusion, with 30 minutes intervals between doses, until a cumulative dose of 76 mcg (1.3 mcg/kg/dose) was reached. The patient did not experience any reaction during desensitization for octreotide. Treatment was continued for five days without any reaction at 1.3 mcg/kg/dose every 8 hours.

Discussion

Anaphylaxis is a severe, potentially fatal, systemic allergic reaction that occurs suddenly after contact with an allergy-causing substance.¹² Foods and drugs are the most common causes of anaphylaxis.^{13,14} When there is no alternative drug that can be used instead of the agent causing anaphylaxis, desensitization of the patient can be considered.

To our knowledge, no pediatric case of anaphylaxis to octreotide has been reported to date. The diagnosis of octreotide hypersensitivity was based on the clinical picture and skin testing of the patient. He was successfully desensitized and octreotide treatment was completed without any allergic reaction. In conclusion, although rare, anaphylaxis due to octreotide can occur in children and desensitization can be considered when treatment is needed to be continued.

References

1. Fleseriu M. Clinical efficacy and safety results for dose escalation of somatostatin receptor ligands in patients with acromegaly: a literature review. *Pituitary*. 2011;14:184-93.
2. Khuroo MS, Khuroo MS, Khuroo NS. Treatment of type I gastric neuroendocrine tumors with somatostatin analogs. *J Gastroenterol Hepatol*. 2010;25:548-54.
3. Song S, Shi R, Li B, Liu Y. Diagnosis and treatment of pancreatic vasoactive intestinal peptide endocrine tumors. *Pancreas*. 2009;38:811-4.
4. Tatar T, Kilic D, Ozkan M, Hatipoglu A, Aslamaci S. Management of Chylothorax with Octreotide after Congenital Heart Surgery. *Thorac Cardiovasc Surg*. 2011;59:298-301.
5. Benson AB 3rd, Ajani JA, Catalano RB, Engelking C, Kornblau SM, Martenson JA Jr, et al. Recommended guidelines for the treatment of cancer treatment-induced diarrhea. *J Clin Oncol*. 2004;22:2918-26.



6. Barbounis V, Koumakis G, Vassilomanolakis M, Demiri M, Efremidis AP. Control of irinotecan-induced diarrhea by octreotide after loperamide failure. *Support Care Cancer*. 2001;9:258-60.
7. Paran H, Mayo A, Paran D, Neufeld D, Shwartz I, Zissin R, et al. Octreotide treatment in patients with severe acute pancreatitis. *Dig Dis Sci*. 2000;45:2247-51.
8. Fiedler F, Jauernig G, Keim V, Richter A, Bender HJ. Octreotide treatment in patients with necrotizing pancreatitis and pulmonary failure. *Intensive Care Med*. 1996;22:909-15.
9. Lieb JG 2nd, Shuster JJ, Theriaque D, Curington C, Cintrón M, Toskes PP. A pilot study of Octreotide LAR vs. octreotide tid for pain and quality of life in chronic pancreatitis. *JOP*. 2009;10:518-22.
10. Novartis. Sandostatín (octreotide acetate) package insert, East Hanover, New Jersey, January 2010.
11. eHealthMe: Real world drug outcomes [Internet]. [place unknown]: eHealthMe; c 2011. Octreotide acetate side effect: Anaphylaxis; [cited 2011 July 10]; [about 1 screens]. Available from: <http://www.ehealthme.com/ds/octreotide+acetate/anaphylaxis>
12. Sampson HA, Muñoz-Furlong A, Campbell RL, Adkinson NF Jr, Bock SA, Branum A, et al. Second symposium on the definition and management of anaphylaxis: summary report--second National Institute of Allergy and Infectious Disease/Food Allergy and Anaphylaxis Network symposium. *Ann Emerg Med*. 2006;47:373-80.
13. Lieberman P. Epidemiology of anaphylaxis. *Curr Opin Allergy Clin Immunol*. 2008;8:316-20.
14. Moro Moro M, Tejedor Alonso MA, Esteban Hernández J, Múgica García MV, Rosado Ingelmo A, Vila Albelda C. Incidence of anaphylaxis and subtypes of anaphylaxis in a general hospital emergency department. *J Investig Allergol Clin Immunol*. 2011;21:142-9.