

# Good Syndrome, a Rare Cause of Refractory Chronic Diarrhea and Recurrent Pneumonia in a Chinese Patient after Thymectomy

Junyan Qu,<sup>a</sup> Xiaoju Lü,<sup>a</sup> Qin Gao,<sup>a</sup> Yan Zhang<sup>b</sup>

Center of Infectious Disease, West China Hospital, Sichuan University, Chengdu, China<sup>a</sup>; Department of Gastroenterology, West China Hospital, Sichuan University, Chengdu, China<sup>b</sup>

**The diagnosis of Good syndrome is very difficult. It has various symptoms, and these symptoms can be present at different periods. In this report we present a patient with refractory chronic diarrhea, recurrent pneumonia, and dysgammaglobulinemia after thymectomy, who was finally then diagnosed with Good syndrome.**

## CASE REPORT

In June 2006, a 41-year-old man was admitted to a local hospital for watery diarrhea (about 10 bowel movements per day). A thymoma with a size of 10 × 8 cm was identified by computed tomography (CT), and then a thymectomy was performed. His bowel movement returned to normal after the surgery. Two months later, he was admitted to the hospital again with more-severe watery diarrhea (10 to 15 bowel movements per day). *Escherichia coli* was identified by stool culture, levofloxacin (0.4 g intravenously daily) was given, and he recovered after 5 days. From December 2006 to January 2011, he was repeatedly admitted to hospitals for recurrent diarrhea, chronic cough, and fever. Colonoscopy was normal. His condition improved after antibiotic therapy each time. In January 2011, he was admitted to our hospital for severe watery diarrhea (15 to 20 bowel movements per day, accompanied by urgency and incontinence), productive cough, and weakness.

On physical examination, the patient was malnourished. Diffuse rales were heard in the bilateral lower lung. A CT scan of the chest revealed several infiltrates on the lungs. Gastroscopy and capsule endoscopy showed chronic superficial gastritis and scattered congestion as well as swelling in the small intestine, respectively. Colonoscopy revealed focal hemorrhage in the transverse colon. Blood chemistry was normal except for a decreased serum albumin level of 31.8 g/liter (reference range, 35 to 55 g/liter). His hemoglobin level was 97 g/liter (reference range, 120 to 160 g/liter). Dysgammaglobulinemia was noted with an IgG concentration of 6.93 g/liter (reference range, 8 to 15.5 g/liter), but the IgM and IgA concentrations were normal. Flow cytometry showed a CD4<sup>+</sup> lymphocyte percentage of 13% and an inverted CD4<sup>+</sup>/CD8<sup>+</sup> ratio of 0.16. Tests for IgM antibodies to cytomegalovirus (CMV) and herpes simplex virus were positive. Quantitative PCR testing of CMV DNA was also positive at 18,200 copies per milliliter. Sputum culture showed *Escherichia coli* and *Haemophilus influenzae* infection. Multiple stool cultures were negative. Repeated HIV serology was negative. Bone marrow aspiration and biopsy showed no obvious abnormalities. Good syndrome (GS) was diagnosed.

Piperacillin sodium-sulbactam sodium (3 g every 12 h) and acyclovir (5 mg/kg of body weight every 8 h) were given. Monthly intravenous immunoglobulin (IVIG) was scheduled. Two weeks later, his gastrointestinal (GI) and respiratory conditions improved. His bowel movement was one to two times per day, and

daily stool weight was about 300 g. He recovered with a good appetite and gained weight. A CT scan showed that the infiltrates in the lung were significantly reduced. He continued with IVIG monthly after being discharged. To date, the patient's bowel movement has remained normal.

Good syndrome was first described in 1954; it is a rare association of thymoma and adult-onset immunodeficiency (1). The incidence is not clear yet. It has a worldwide distribution, but most cases have been described in Europe. To date, only 17 cases have been reported in China. The immunodeficiency in GS is characterized by reduced or absent B cells in peripheral blood, dysgammaglobulinemia, and T-cell immunodeficiency. The clinical manifestations of GS include thymoma, opportunistic infections, diarrhea, and autoimmune manifestations, such as myasthenia gravis, pure red cell aplasia (PRCA), and aplastic anemia (2).

Although up to 31.8% of patients with GS can have diarrhea, the cause of diarrhea is still not clear. Many factors may be related to the cause of diarrhea, such as infections and malabsorption. Immunodeficiency can increase the risk of opportunistic infections in the GI tract. Many pathogens have been identified, including CMV, *Campylobacter* spp., and *Giardia lamblia*, etc. (3–6), among which *Salmonella* spp. were the most common one. Hypogammaglobulinemia and mucosal lesions resembling villous atrophy can also lead to malabsorption (7, 8). Other causes of diarrhea may include bacterial overgrowth (9) and immune-mediated colitis (10). Ulcerative colitis also has been reported and may be an explanation for the diarrhea in many patients. Although we could not identify definite pathogens in this patient's stool, his diarrhea still was alleviated by antibacterial treatment, which suggested that his diarrhea may be caused by infection by an unusual pathogen.

Sinopulmonary infection is the most commonly described infection in GS (2). The pathogens identified included *Haemophilus*

Received 15 March 2013 Returned for modification 1 April 2013

Accepted 9 May 2013

Published ahead of print 22 May 2013

Address correspondence to Yan Zhang, zhangyan201302@126.com.

Copyright © 2013, American Society for Microbiology. All Rights Reserved.

doi:10.1128/CVI.00141-13

*influenza* and *Streptococcus pneumoniae*, etc. Tuberculosis also has been reported. CMV is the most commonly described viral infection in GS (2). About 21.1% of patients with GS presented with CMV infection. The clinical manifestations of CMV infection in GS include pneumonia and colitis, etc. (2). In our patient, *Escherichia coli*, *Haemophilus influenzae*, CMV, and herpes simplex virus were identified at the same time. To our knowledge, this is the first report of two kinds of bacterial and viral infections in one patient with GS. We believe that the recurrent infection was related to his humoral immunity deficiency and his low CD4<sup>+</sup>/CD8<sup>+</sup> ratio.

The pathogenesis of immunodeficiency in GS remains unknown. The thymus is an organ that mediates important immune functions. In thymoma, derangement of the microenvironment of the medullary and cortical compartments in the thymus may lead to loss of self-tolerance (11), impaired immune surveillance, and various autoimmune phenomena (12).

The standard treatment of thymoma is surgical removal or debulking of the tumor (13); however, the clinical outcomes of thymectomy differ. For some patients, symptoms improved (12), for some symptoms became aggravated (14), and for the others there was no change (15, 16). Our patient's symptoms became worse after the surgery. The mechanism is not fully understood. In almost all cases, thymectomy does not restore immune function. The thymus plays a central role in the generation and maintenance of peripheral T-cell populations. Thymectomy was not able to change the number of peripheral T cells, which may be related to thymectomy extending the half-life of immature T cells; in fact, thymectomy can reduce the number of T<sub>H</sub> cells (17), which may lead to a reduction in the number of activated B cells. The mechanism of chronic diarrhea in GS is a complex mechanism, with multiple factors involved and thymoma itself may be one of the factors.

In conclusion, in a patient with thymoma and recurrent opportunistic infection, Good syndrome should be considered, and the immunological parameters should be checked. Currently, there is no satisfactory treatment. Thymectomy can prevent locally invasive growth and metastasis of thymoma, but it does not reverse dysimmunity (18). Gammaglobulin replacement was recommended to suppress infections associated with GS. A recent report showed that the use of IL-7 or IL-5 cytokines might be able to rebuild immunity in patients with GS (19).

## REFERENCES

1. Good, RA. 1954. Agammaglobulinemia—a provocative experiment of nature. *Bull. Univ. Minn.* 26:1–19.
2. Kelesidis T, Yang O. 2010. Good's syndrome remains a mystery after 55 years: a systematic review of the scientific evidence. *Clin. Immunol.* 135:347–363.
3. Tarr PE, Sneller MC, Mechanic LJ, Economides A, Eger CM, Strober W, Cunningham-Rundles C, Lucey DR. 2001. Infections in patients with immunodeficiency with thymoma (Good syndrome). Report of 5 cases and review of the literature. *Medicine (Baltimore, MD)* 80:123–133.
4. Agarwal S, Cunningham-Rundles C. 2007. Thymoma and immunodeficiency (Good syndrome): a report of 2 unusual cases and review of the literature. *Ann. Allergy Asthma Immunol.* 98:185–190.
5. Khanna S, Kumar A, Tandon R. 2004. Good's syndrome: an unusual cause of chronic diarrhea. *Indian J. Gastroenterol.* 23:152–153.
6. Koriyama N, Fukumoto O, Fukudome M, Aso K, Hagiwara T, Arimura K, Naka-zaki M, Arima N, Eizuru Y, Tei C. 2004. Successful treatment of Good syndrome with cytomegalovirus duodenenteritis using a combination of ganciclovir and immunoglobulin with high anticytomegalovirus antibody titer. *Am. J. Med. Sci.* 327:49–54.
7. Verne GN, Amann ST, Cosgrove C, Cerda JJ. 1997. Chronic diarrhea associated with thymoma and hypogammaglobulinemia (Good's syndrome). *South. Med. J.* 90:444–446.
8. Ghoshal UC, Goel A, Ghoshal U, Jain M, Misra A, Choudhuri G. 2011. Chronic diarrhea and malabsorption due to hypogammaglobulinemia: a report on twelve patients. *Indian J. Gastroenterol.* 30:170–174.
9. Puebla-Maestu A, Martín-Lorente JL, Arias-García L, Sáez-Royuela F, Gento-Peña E, Pérez-Alvarez JC, Ojeda-Giménez C. 2003. Good's syndrome and chronic diarrhea. *Gastroenterol. Hepatol.* 26:245–247.
10. Kornacki S, Hansen FC, III, Lazenby A. 1995. Graft-versus-host-like colitis associated with malignant thymoma. *Am. J. Surg. Pathol.* 19:224–228.
11. Salaün J, Corbel C, Le Douarin NM. 2005. Regulatory T cells in the establishment and maintenance of self-tolerance: role of the thymic epithelium. *Int. J. Dev. Biol.* 49:137–142.
12. Yildiz O, Ozguroglu M, Turna H, Yanmaz T, Kaynak K, Akman C, Cetin SE, Oz B, Celik A. 2007. Thymoma with chronic diarrhea: report of two cases and review of the literature. *Med. Oncol.* 24:119–123.
13. Johnson SB, Eng TY, Giaccone G, Thomas CR, Jr. 2001. Thymoma: update for the new millennium. *Oncologist* 6:239–246.
14. Ho JK, Wong MM, Tai TK, Tse DM. 2010. A rare combination of recurrent pneumonia, diarrhoea, and visual loss in a patient after thymectomy: Good syndrome. *Hong Kong Med. J.* 16:493–496.
15. de Jesus NP, Carvalho PM, Dias FM, Gaspar EM, de Moura JJ. 2008. Dementia in a patient with thymoma and hypogammaglobulinaemia (Good's syndrome). *Cases J.* 13:90.
16. Ohuchi M, Inoue S, Hanaoka J, Igarashi T, Tezuka N, Ozaki Y, Teramoto K. 2007. Good syndrome coexisting with leukopenia. *Ann. Thorac. Surg.* 84:2095–2097.
17. Tanchot C, Rocha B. 1995. The peripheral T cell repertoire: independent homeostatic regulation of virgin and activated CD8<sup>+</sup> T cell pools. *Eur. J. Immunol.* 25:2127–2136.
18. Souadjian JV, Enriquez P, Silverstein MN, Pépin JM. 1974. The spectrum of diseases associated with thymoma. Coincidence or syndrome? *Arch. Intern. Med.* 134:374–379.
19. Ternavasio-de la Vega HG, Velasco-Tirado V, Pozo-Rosado L, Soler-Fernández MC, Pérez-Andres M, Orfao A, Sánchez-Sánchez R, González-Villaron L. 2011. Persistence of immunological alteration after thymectomy in Good's syndrome: a clue to its pathogenesis. *Cytometry B Clin. Cytom.* 80:339–342.