Case Report

Two Cases of Primary Cold Agglutinin Disease Associated with Megaloblastic Anemia

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1. Introduction

Cold agglutinin disease (CAD), or cold antibody autoimmune hemolytic anemia, is characterized by mild anemia with reticulocytosis, positive direct Coombs test, elevated levels of lactate dehydrogenase (LDH), low levels of serum haptoglobin, and high titers of cold agglutinin [1–3]. CAD manifests as either a primary disease, that is, chronic CAD, or secondary to Waldenström's macroglobulinemia (WM) or B-cell type malignant lymphoma [4, 5]. Secondary CAD also occurs in association with systemic lupus erythematosus [6] or transiently upon Epstein-Barr virus or mycoplasma pneumoniae infection [7]. Cold agglutinins, which are specific for the I-antigen expressed on the surface of red blood cells, belong to the IgM subclass and, in the majority of patients with primary CAD, are monoclonal IgM-kappa antibodies [1–3]. Primary CAD is most often seen in elderly patients (median age at onset is 67 years (range 30–92 years)) and the incidence rate is 1 per 1 million people per year [2]. Primary CAD may develop in association with various hematological/immunological diseases, including pernicious anemia [8] and common variable immunodeficiency (CVID) [9]. Here, we report the cases of two elderly Japanese patients with primary CAD who showed clinical features of megaloblastic anemia due to decreased vitamin 12 levels. In addition, one of these patients also showed probable CVID in addition to typical CAD symptoms.

2. Case Presentation

2.1. Case 1. A 67-year-old male was diagnosed with CAD in 2009. Since then, over the past 3 years, he had maintained Hb levels at 15.0 to 16.5 g/dL but complained of peripheral coldness and cyanosis of the limbs in association with Raynaud’s phenomenon, particularly in cold seasons; however, he did not receive any specific therapy. The patient was hospitalized due to progression of anemia and hemoglobinuria in December 2012. In the summer of that year he had Hb level...
at 16.2 g/dL and then became anemic over the fall-to-winter period. His prior medical history revealed alcoholic liver dysfunction, mild diabetes mellitus, and hypertension. There was no history of inappropriate dietary intake or drug use and no recent ongoing excess alcohol use. On admission, the patient (height 167 cm and body weight 73.4 kg) was anemic (Hb 8.1 g/dL) and slightly icteric, with total bilirubin levels of 2.5 mg/dL. He also had macrocytic anemia. A peripheral blood film revealed marked red blood cell agglutination (Figure 1). At Cs can show no lymphadenopathy or splenomegaly. The laboratory data are summarized in Table 1.

During the 3 years prior to hospitalization, his cold agglutinin titer remained high (1:2,048); however, upon hospitalization it was 1: >8,192. He also had monoclonal M-proteins (IgM-kappa) but normal IgG, IgA, and IgM; however, complement levels were low (Table 1). In this case, no bone marrow analyses were performed; however, during the entire course of CAD, he did not show any signs of lymphoproliferative diseases (serum sIL-2R remained within normal range and there were negative CT findings). In addition, the patient had low vitamin 12 levels, confirming megaloblastic anemia, with positive anti-intrinsic factor as well as antiparietal cell antibodies. Gastrointestinal endoscopy revealed atrophic gastritis. In addition to vitamin B12 supplementation (mecobalamin 500 μg × 3/day), he was treated with four doses of weekly rituximab (375 mg/m²/dose), which increased the Hb levels from 8.1 g/dL to 14.7 g/dL and reduced serum LDH levels from 1,119 IU/L to 201 IU/L 2 months later. MCV was normalized in 2 months following vitamin B12 administration. For the last 2 years, he has been doing well without rituximab maintenance therapy, with Hb levels >15.0 g/dL, LDH levels around 160 IU/L, a cold agglutinin titer of 1:2,048, and no episodes of acute hemolysis.

2.2. Case 2. A 55-year-old male (height 170 cm and body weight 65.4 kg) was diagnosed with CAD in 2012 when he did not show anemia, apart from a slight increase in total bilirubin levels (2.2–3.0 mg/dL). One year later, after exposure to cold, he again visited a clinic and was found to be mildly anemic (Hb 8.0 g/dL); however, he was not systematically tested or treated. In his history he had inappropriate dietary intake but no drug use. In February 2014, he was hospitalized via emergency medical transport due to the fact that he was severely anemic (Hb 4.3 g/dL) as well as icteric, with a peripheral blood smear showing significant red blood cell agglutination (data not shown). A CT scan revealed no lymphadenopathy, but pulmonary edema, bilateral pleural fluid, and mild splenomegaly were noted. The laboratory data are summarized in Table 1. His cold agglutinin level was extremely high (1:16,834) and he had monoclonal M-proteins (IgM-kappa) in association with hypogammaglobulinemia (IgG, 584 mg/dL) and hypocomplementemia. In addition, he had megaloblastic anemia associated with low serum levels of vitamin B12 and folate. A bone marrow smear showed an increase in megaloblasts associated with dyserythropoietic features of polychromatophilic erythroblasts (Figure 2), with an M/E ratio of 0.92; these data were compatible with a diagnosis of both hemolytic and megaloblastic anemia. No increase in the number of myeloma cells or lymphoma cells was noted. Bone marrow cells showed a normal karyotype (46 XY; 20/20). He was treated with a heated RBC transfusion (a total of 10 U) and received oral folate (5 mg/day × 3/day) and vitamin 12 (mecobalamin 500 μg × 3/day) but declined intravenous rituximab therapy. Unfortunately, this patient was discharged not for medical recovery but for his personal reasons and became lost to follow-up.

3. Discussion

Both patients showed clinical features typical of chronic CAD with high titers of cold agglutinin. Direct Coombs test (DCT) was negative, as shown in Table 1, for both patients with use of wide spectrum antiserum. It was reasoned that our patients were not in a hemolytic phase when DCT was tested. In fact, in patients with mycoplasma pneumoniae-related CAD, the incidence of patients with a positive DCT was shown to fall with time [10]. After the interval of 2-3 years from the first recognition of CAD symptoms, both of our patients required intensive therapy for hemolytic anemia upon hospitalization. CAD may develop in association with WM, B-cell type lymphoma, SLE, or upon Epstein-Barr virus
Table 1: Laboratory data of 2 CAD cases.

|                     | Case 1       | Case 2       |
|---------------------|--------------|--------------|
| Age (years)/sex     | 67/M         | 55/M         |
| WBC (3000–8500)/μL  | 9900         | 6300         |
| Hb (12.5–17.5) g/dL | 8.1          | 4.3          |
| MCV (84.6–100.6) fL| 115          | 110          |
| Reticulocytes (0.3–1.1) % | 11.5 | 8.2          |
| PLTs (115,000–305,000)/μL | 198,000 | 147,000     |
| Haptoglobin (19–170) mg/dL | 67 | 2            |
| AST (13–37) IU/L    | 45           | 25           |
| ALT (8–45) IU/L     | 37           | 14           |
| LDH (122–228) IU/L  | 1119         | 1021         |
| Total bilirubin (0.3–1.3) mg/dL | 2.50 | 7.94         |
| Direct bilirubin (0.1–0.3) mg/dL | 1.10 | 1.30         |
| Total protein (6.7–8.3) g/dL | 7.0 | 5.9          |
| Albumin (4.1–5.2) g/dL | 4.0 | 4.3          |
| BUN (7.8–18.9) mg/dL | 19.2 | 8.3          |
| Creatinine (0.64–1.11) mg/dL | 0.88 | 0.86         |
| CRP (0–0.29) mg/dL  | 0.60         | 0.92         |
| Direct Coombs test* | Negative     | Negative     |
| Cold agglutinin titer | 1: >8,192  | 1:16,384     |
| M-protein           | Positive (IgM-kappa) | Positive (IgM-kappa) |
| Cryoglobulin        | Negative     | NT           |
| HBV/HCV             | Negative/negative | Negative/NT |
| ANA                 | <<40         | <<40         |
| Soluble IL-2R (122–496) U/mL | 437 | NT          |
| IgG/IgA/IgM (820–1740/90–400/31–200) | 1065/554/217 | 584/NT/164 |
| C3/C4/CH50 (80–140/11–34/30–45) | 60/5.0/10.8 | 56/6.3/7.0 |
| Folate (3.6–12.9) ng/mL | 10.9 | 3.2          |
| Vitamin B12 (233–914) pg/mL | 85 | 173          |
| Anti-IF-Ab          | Positive     | NT           |
| Anti-PC-Ab          | Positive     | NT           |

*Employed with wide spectrum antiserum, not with monospecific anti-C3 antibody, WBC: white blood cell counts, PLTs: platelet counts, AST: aspartate aminotransferase, ALT: alanine aminotransferase, LDH: lactate dehydrogenase, BUN: blood urea nitrogen, CRP: C-reactive protein, and ANA: antinuclear antibody; units for IgG, IgA, IgM, C3, and C4 are mg/dL, and unit for CH50 is U/mL; IF: intrinsic factor, PC: parietal cell, Ab: antibody, and NT: not tested.

or mycoplasma pneumoniae infection [4–7]; however, these two patients had no such underlying diseases. In addition, both patients showed monoclonal IgM-kappa antibodies, compatible with primary CAD or WM, differentiated from other types of CAD (Figure 3). However, WM was ruled out; thus primary CAD was diagnosed. In addition to clinical symptoms and laboratory data typical of CAD, both patients had megaloblastic anemia due to vitamin B12 deficiency. Macrocytic anemia in our cases was not false elevations of the MCV due to cold agglutinins [11]. The reduced complement levels (C3, C4, and CH50 were all reduced) observed in these patients were similar to those reported in previous studies [12, 13]. Ulvestad et al. reported that, of 15 primary CAD patients, nine showed decreased serum C3, 11 showed C4, and six showed reduced CH50, indicating that patients with CAD experience continuous complement consumption [13]. Furthermore, Case 2 had mild hypogammaglobulinemia.

In primary CAD, both the cold-reactive antibodies and the IgG-kappa antibodies on circulating red cells play a role in the various hematological/immunological abnormalities [12]. However, few studies have reported cooccurrence of CAD and megaloblastic anemia. Such an association in our cases might be incidental; however, given that CAD is an autoimmune disease caused by multiple autoantibodies, it may not be a coincidence, particularly as one study reported a case of CAD with pernicious anemia [8]. It is possible that the megaloblastic anemia in our cases was actually pernicious anemia, a complex disorder consisting of hematological, gastric, and immunological alterations, as we confirmed the evidence for antibodies against intrinsic factor or gastric parietal cells in Case 1, although antibodies against the intrinsic factor are not specific pernicious anemia. Other factors such as chronic use of alcohol in Case 1 and inappropriate dietary...
Figure 2: Bone marrow smear from Case 2 showing megaloblastic changes associated with the dyserythropoietic features of polychromatric erythroblasts.

Figure 3: The majority of primary CAD and WM-associated CAD cases (>70%) show cold agglutinins of monoclonal IgM-kappa antibodies. On the other hand, lymphoma-associated CAD is associated with monoclonal IgM-lambda antibodies [2]. Cold agglutinins associated with postinfectious diseases such as Epstein-Barr virus or mycoplasma pneumoniae infection-related CAD can be polyclonal [1]. CAD: cold agglutinin disease; ML: malignant lymphoma; WM: Waldenström's macroglobulinemia; p.i.: postinfection; k: kappa; L: lambda; p: polyclonal.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

References

[1] M. A. Gertz, "Cold agglutinin disease," *Haematologica*, vol. 91, no. 4, pp. 439–441, 2006.
[2] S. Berentsen, E. Ulvestad, R. Langholm et al., "Primary chronic cold agglutinin disease: a population based clinical study of 86 patients," *Haematologica*, vol. 91, no. 4, pp. 460–466, 2006.
[3] D. Crisp and W. Pruzanski, "B-cell neoplasms with homogeneous cold-reacting antibodies (cold agglutinins)," *The American Journal of Medicine*, vol. 72, no. 6, pp. 915–922, 1982.
[4] M. Zago-Novaretti, F. Khuri, K. B. Miller, and E. M. Berkman, "Waldenström's macroglobulinemia with an IgM paraprotein that is both a cold agglutinin and a cryoglobulin and has a suppressive effect on progenitor cell growth," *Transfusion*, vol. 34, no. 10, pp. 910–914, 1994.
[5] A. E. Eskazan, H. Akmurad, S. Ongoren, O. Ozer, and B. Ferhanoglu, "Primary gastrointestinal diffuse large B cell lymphoma presenting with cold agglutinin disease," *Case Reports in Gastroenterology*, vol. 5, no. 2, pp. 262–266, 2011.
[6] T. Kotani, T. Takeuchi, Y. Kawasaki et al., "Successful treatment of cold agglutinin disease with anti-CD20 antibody (rituximab) in a patient with systemic lupus erythematosus," *Lupus*, vol. 15, no. 10, pp. 683–685, 2006.
[7] S. Basu, A. Saifudeen, and P. Kaur, "Transient cold agglutinin disease with mycoplasma infection," *Journal of Association of Physicians of India*, vol. 57, no. 9, 2009.
[8] P. Narasimhan, T. Bharathan, and W. J. Watson-Williams, "Symmetrical gangrene due to cold hemagglutinin disease with pernicious anemia," *New York State Journal of Medicine*, vol. 74, no. 6, pp. 1046–1048, 1974.
[9] J. Wang and C. Cunningham-Rundles, "Treatment and outcome of autoimmune hematologic disease in common variable immunodeficiency (CVID)," *Journal of Autoimmunity*, vol. 25, no. 1, pp. 57–62, 2005.
[10] T. Feizi, "Cold agglutinins, the direct coombs' test and serum immunoglobulins in *Mycoplasma pneumoniae* infection," *Annals of the New York Academy of Sciences*, vol. 143, no. 1, pp. 801–812, 1967.
[11] P. G. Hattersley, P. W. Gerard, V. Caggiano, and D. R. Nash, "Erroneous values on the Model S Coulter Counter due to high titer cold autoagglutinins," *American Journal of Clinical Pathology*, vol. 55, no. 4, pp. 442–446, 1971.
[12] J. L. Moake and D. R. Schultz, "Hemolytic anemia associated with multiple autoantibodies and low serum complement," *The American Journal of Medicine*, vol. 38, no. 3, pp. 431–437, 1975.
[13] E. Ulvestad, S. Berentsen, K. Bø, and F. V. Shammas, “Clinical immunology of chronic cold agglutinin disease,” *European Journal of Haematology*, vol. 63, no. 4, pp. 259–266, 1999.

[14] M. Micheau, C. de Caqueray, D. Fontan, J. Bentegeat, J. M. Guillard, and P. Verger, “Proceedings: autoimmune hemolytic anemia with megaloblast formation in agammaglobulinemia,” *Archives Françaises de Pédiatrie*, vol. 32, no. 6, article 586, 1975 (French).

[15] P. Sève, L. Bourdillon, F. Sarrot-Reynaud et al., “Autoimmune hemolytic anemia and common variable immunodeficiency: a case-control study of 18 patients,” *Medicine*, vol. 87, no. 3, pp. 177–184, 2008.

[16] M. Pavic, P. Sève, C. Malcus et al., “Common variable immunodeficiency with autoimmune manifestations: study of nine cases; Interest of a peripheral B-cell compartment analysis in seven patients,” *Revue de Médecine Interne*, vol. 26, no. 2, pp. 95–102, 2005 (French).

[17] S. Berentsen, E. Ulvestad, B. T. Gjertsen et al., “Rituximab for primary chronic cold agglutinin disease: a prospective study of 37 courses of therapy in 27 patients,” *Blood*, vol. 103, no. 8, pp. 2925–2928, 2004.

[18] A. Vassou, V. Alymara, A. Chaidos, and K. L. Bourantas, “Beneficial effect of rituximab in combination with oral cyclophosphamide in primary chronic cold agglutinin disease,” *International Journal of Hematology*, vol. 81, no. 5, pp. 421–423, 2005.

[19] S. Berentsen, U. Randen, A. M. Vågan et al., “High response rate and durable remissions following fludarabine and rituximab combination therapy for chronic cold agglutinin disease,” *Blood*, vol. 116, no. 17, pp. 3180–3184, 2010.

[20] A. Gueli, D. Gottardi, H. Hu, I. Ricca, A. De Crescenzo, and C. Tarella, “Efficacy of rituximab-bendamustine in cold agglutinin haemolytic anaemia refractory to previous chemoimmunotherapy: a case report,” *Blood Transfusion*, vol. 11, no. 2, pp. 311–314, 2013.