Correspondence

To the editor:

Severe cold hemagglutinin disease (CHD) successfully treated with rituximab

Autoimmune hemolytic anemia results from an IgM antibody (Ab)–red cell interaction and is referred to as cold hemagglutinin disease (CHD). CHD is most commonly associated with infectious or lymphoproliferative diseases but may also be coupled with malignancies or autoimmune or immunodeficiency syndromes. Cold agglutinin–mediated hemolysis occurs at low temperatures, may be severe, and is notoriously difficult to treat. CHD patients with infections often have a short clinical course, whereas those with lymphoma require therapy. Therapeutic maneuvers successful in patients with warm Ab–associated autoimmune hemolytic anemia such as corticosteroids, intravenous immunoglobulin G (IgG), and splenectomy are usually ineffective in CHD.1,2 Novel treatment approaches of CHD are based on a better understanding of the immunologic abnormalities associated with this disorder, as well as the availability of sophisticated biotechnology products.1-5 Rituximab is a genetically engineered chimeric monoclonal antibody that targets the CD20 antigen on B cells currently used for the treatment of non-Hodgkin lymphoma. In vitro studies have demonstrated that the antibody binds human C1q and induces complement-dependent cytotoxicity, antibody-dependent, cell-mediated cytotoxicity, and apoptosis. In clinical trials of patients with malignant lymphoma, rituximab depleted circulating B cells with the first doses, leading to ongoing remissions and remaining effective for several months.3-5 In these diseases, as in autoimmune thrombocytopenia (ITP), rituximab presumably acts by elimination of CD20+ clonotypic precursor B cells and/or CD20+ plasma cells, applying to immunoglobulin-mediated diseases of B lymphocytes.6 Here we report on 2 patients with severe CHD, who were both successfully treated with the chimeric anti-CD20 monoclonal Ab rituximab. The patients’ disease course and response to rituximab are displayed in Figure 1. Both patients were seronegative for HIV-1 and HIV-2, and other infections were ruled out by appropriate diagnostic procedures.

Figure 1. Changes in biologic variables before and after rituximab therapy. (A) First patient; (B) second patient. Pred indicates prednisone; PP, plasmapheresis; Cy, cyclophosphamide; and MMF, mycophenolate mofetil.

The first patient, a 50-year-old female, was admitted to our hospital in February 2001 because of severe anemia (hemoglobin level, 7.3 g/dL), jaundice, weakness, and dyspnea. Respiratory sounds were normal, and the spleen and liver were not enlarged. Laboratory data showed marked reticulocytosis (7.54%) and bilirubinemia (7.8 mg/dL), positive direct Coombs test, a high cold agglutinin titer of 1:256 (normal, 1:32), and low haptoglobin level (21 mg/dL). A bone marrow (BM) biopsy revealed a markedly hyperplastic erythropoiesis and agglutination of erythroid precursors. Immune phenotyping showed only polyclonal B cells and no cytogenetic abnormalities. Initially, prednisone (2 mg/kg/d) was administered and slowly tapered (Figure 1A). But 5 months later the direct Coombs test became positive again, with a concomitant increase in cold agglutinins (1:1024), impressive decrease of hemoglobin level to 5.9 g/dL, elevated reticulocyte percentage (10.1%), elevated lactate dehydrogenase (LDH) level (458 U/L), elevated bilirubin level (6.8 mg/dL), and haptoglobin level below 5 mg/dL. Cyclophosphamide (1000 mg/m²) was given but did not improve the clinical course. Therefore, mycophenolate mofetil (MMF; 1 g/d, then 2 g/d) was started, which has been described as beneficial in autoimmune hemolytic anemia.2 Nevertheless, the above parameters did not improve; on the contrary, severe illness of the patient persisted with daily worsening of the clinical condition. Emergency plasmapheresis was carried out, resulting in only brief clearance of cold agglutinins. Since the effect of plasmapheresis did not lead to a marked improvement in the clinical course, rituximab was given (375 mg/m², intravenously, weekly for 4 courses). Already with the first infusion, this led to a marked improvement of the patient’s clinical condition and to a continuously rising hemoglobin level and marked decrease of reticulocyte percentage and bilirubin and LDH levels (Figure 1A). MMF was discontinued with the first rituximab infusion, and only the latter continued for 3 additional weeks. Tolerance to treatment was excellent, with no side effects. At last follow-up, 9 months after rituximab therapy, the hemoglobin level is 12.6 g/dL, bilirubin level, 1.6 mg/dL, and LDH level, 188 U/L, indicating continuing remission of CHD.
To the editor:

Pharmacology of PEG-asparaginase in childhood acute lymphoblastic leukemia (ALL)

Asparaginase (ASNase) has long been considered to be an important element in the management of childhood ALL. Its antileukemic effect is thought to be related to a metabolic deficiency reflected by the blasts’ incapability to synthesize asparagine (ASN) from aspartic acid. Treatment with ASNase aims therefore at depleting the blood of ASN in order to exhaust the substrate supply that became selectively essential to the malignant cells. Highly interesting findings on PEG-ASNase, which is a polyethylene

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References

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