Occurrence of hemolytic anemia in patients with GBS treated with high-dose IVIg

ABSTRACT

Objective: We describe an underrecognized side effect of high-dose IV immunoglobulin (IVIg), hemolytic anemia.

Background: There are no established guidelines on treating patients with Guillain-Barré syndrome (GBS) who relapse or do not improve after a standard course of treatment (IVIg or plasma exchange). Some centers will opt for a second course of the initial treatment. There is an ongoing trial of a second course of IVIg in patients with severe GBS.

Methods: We retrospectively reviewed 4 patients with severe GBS who received high-dose IVIg. One patient inadvertently received a high dose of IVIg for Miller Fisher syndrome. All patients received a total of at least 2 courses of the standard dose of IVIg (total >4 g/kg). We review their clinical course and side effects.

Results: All patients with non-O blood types developed clinically significant hemolytic anemia requiring blood transfusion.

Conclusion: Hemolytic anemia may limit doses of IVIg for treatment of severe GBS in patients with non-O blood types. Neurology Neuroimmunol Neuroinflammation 2014;1:e50; doi: 10.1212/NXI.0000000000000050

GLOSSARY

GBS = Guillain-Barré syndrome; HA = hemolytic anemia; IVIg = IV immunoglobulin; MFS = Miller Fisher syndrome.

Guillain-Barré syndrome (GBS) is the most common cause of acute flaccid neuromuscular paralysis, resulting in significant morbidity and cost. There is a proposed model to identify patients with poor prognosis. However, treatment options beyond IV immunoglobulin (IVIg) and plasma exchange are limited. An ongoing trial is evaluating a second course of IVIg for severe GBS. This trial has prompted treating neurologists to consider an additional course of IVIg if patients do not respond initially. We alert neurologists to significant hemolytic anemia (HA) in patients with non-O blood types treated with high-dose IVIg within a short time interval.

METHODS This is a retrospective case series. We review 4 patients with severe GBS and one patient with Miller Fisher syndrome (MFS) admitted to our institution from 2012 to 2013. All patients received very high-dose IVIg (total dose of at least 4 g/kg, equivalent to 2 standard treatment courses [2 g/kg]) administered over fewer than 14 days. All patients with GBS did not respond initially, prompting a second course of treatment (table). The patient with MFS received a high dose inadvertently. We gathered data regarding the incidence of HA in these patients. As this was a retrospective review of a small case series, institutional review board approval was not required.

RESULTS All of the patients with non-O blood types (3/5) treated with very high-dose IVIg developed clinically significant anemia requiring blood transfusion. When tested (2/3 cases), there was clear evidence of hemolysis (positive direct antiglobulin test, low haptoglobin, high lactate dehydrogenase/bilirubin). In addition, there were no other medical comorbidities or medications commonly implicated in causing autoimmune hemolytic anemia (table). Patient 4 developed hematuria prior to discharge with normal

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Haptoglobin. Note patient 3 was not tested for these parameters.

Abbreviations: AMAN = acute motor axonal neuropathy; GBS = Guillain-Barré syndrome; IVIg = IV immunoglobulin; MFS = Miller Fisher syndrome; MRC = Medical Research Council.

Hemolysis testing indicates the patient’s serum was consistent with hemolysis: high lactate dehydrogenase, positive direct antiglobulin test, and low haptoglobin. Note patient 3 was not tested for these parameters.

We propose that risk of IVIg-related HA is increased with high doses, significant in non-O blood types, and more likely to occur when administered over a short interval (i.e., <2 weeks). HA has not been frequently observed in patients receiving maintenance IVIg, likely due to the lower dose, longer time intervals between infusions (typically 3–4 weeks), and 2–3 week half-life of anti-A/B IgG. Although all of our patients had GBS/MFS, HA has been reported in many other conditions treated with high-dose IVIg such as postpolio syndrome, chronic inflammatory demyelinating polyneuropathy, myasthenia gravis, renal transplantation, and Kawasaki disease.3–7

The applicability of our findings is limited by a small number of cases, retrospective analysis, and hemolysis testing in only 2 of the 3 patients. However, our findings raise relevant, novel considerations such as a possible maximum tolerated dose of IVIg in a short time frame. In addition, practitioners should exercise caution if considering escalating IVIg dosages to >2 g/kg in <14 days. Hemoglobin and creatinine levels should be monitored closely during and after treatment with high-dose IVIg (up to 1–2 weeks) to detect this untoward complication. In the literature, nadir hemoglobin level seemed to occur within 2 weeks of treatment, commonly within 4–5 days.3–7 If anemia occurs, testing should include direct antiglobulin test, peripheral blood smear, bilirubin, haptoglobin, lactate dehydrogenase, and reticulocyte count. IVIg lots can be tested for anti-A and anti-B titers. In addition, a crossmatch could be performed with serum and IVIg. Manufacturers should label anti-A and anti-B titers on their product and consider more stringent industry standards. We propose that there is a maximal tolerated dose of IVIg over a 2-week period, which is dependent on patient ABO blood group and IVIg’s anti-A/anti-B titers.

DISCUSSION

We propose that risk of IVIg-related HA is increased with high doses, significant in non-O blood types, and more likely to occur when administered over a short interval (i.e., <2 weeks). HA has not been frequently observed in patients receiving maintenance IVIg, likely due to the lower dose, longer time intervals between infusions (typically 3–4 weeks), and 2–3 week half-life of anti-A/B IgG. Although all of our patients had GBS/MFS, HA has been reported in many other conditions treated with high-dose IVIg such as postpolio syndrome, chronic inflammatory demyelinating polyneuropathy, myasthenia gravis, renal transplantation, and Kawasaki disease.3–7

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AUTHOR CONTRIBUTIONS

Dr. Nguyen: study concept, design, acquisition of data, analysis, interpretation. Dr. Biliciler: critical revision of the manuscript for important intellectual content. Dr. Wahed: critical revision of the manuscript for important intellectual content. Dr. Sheikh: critical revision of the manuscript for important intellectual content, study supervision.

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DISCLOSURE

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