Home Treatment With Intravenous Enzyme Replacement Therapy for Gaucher Disease: An International Collaborative Study of 33 Patients

By A. Zimran, C.E.M. Hollak, A. Abrahamov, M.H.J. van Oers, M. Kelly, and E. Beutler

Intravenous enzyme replacement therapy (Alglucerase; Ceredase; Genzyme Corp, Boston, MA) is an effective and safe treatment for patients with type 1 Gaucher disease. In an attempt to reduce its high cost, a “low-dose high-frequency” protocol (30 U/kg/mo, 3 times a week) was introduced and found to be as effective as the original high-dose protocol (60 U/kg every 2 weeks). Because receiving frequent infusions creates a burden for many patients, we have implemented a program of home treatment for our patients. We now report the safety and feasibility of low-dose/high-frequency home intravenous enzyme-replacement therapy in 33 patients with Gaucher disease. The chronic nature of the treatment, its safety, lack of adverse effects, the stable condition of most patients, and the need to reduce the high cost make enzyme replacement for Gaucher disease a good candidate for intravenous home therapy.

Instructions and performance of home treatment. Instructions for the safe and sterile administration of the drug were given to all patients during the initial in-hospital therapy. After this period, they were usually able to perform an infusion either into a peripheral vein or into the Port-a-cath system by themselves or with the help of another person. Although most of the Dutch patients were trained to perform the insertion of the needles by themselves, many of the Israeli and American patients received help from either a family member or a visiting nurse (Table 1). Most of the patients received Ceredase vials containing 50 U of the enzyme (10 U/mL). In other cases, when 50 U vials were not available, strict guidelines for the preservation of undiluted enzyme (in the standard 400 U/mL vials) were given. The appropriate quantity of the enzyme was diluted aseptically in 100 mL of saline (0.9%), followed by administration by gravity or infusion-pumps either immediately or within 1 week. The enzyme, diluted in 100 mL of physiologic saline for administration, was stable for several weeks (Fig 1).

RESULTS

Our patient population included 17 women and 16 men, ranging in age from 4 to 53 years old, with a mean age of 29.5. Their mean severity score index was 15.7 (range, 5 to 30), indicating moderate-to-severe phenotypic expression of the disease. They received a total of approximately 4,500 infusions during a period that ranged between 13 and 82 weeks per patient.

Side effects. One patient complained of some abdominal discomfort after Ceredase infusion. Another patient reported 2 episodes of a visual disturbance occurring during...

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The stability of Alglucerase diluted to a concentration of approximately 1 U/mL in preservative-free isotonic sodium chloride solution for infusion into a patient with Gaucher disease. The residual enzyme from the IV tubing was stored at 4°C and assayed for acid β-glucosidase activity using 4-methyl umbelliferyl β-glucoside as substrate according to the method of Raghaven et al. Even after 60 days of storage, over 70% of the enzyme activity was still present.

**DISCUSSION**

Increasing numbers of patients suffering from different major diseases are receiving intravenous therapy at home. These home treatments include hemodialysis, intravenous antibiotics, γ-globulin, chemotherapy, blood, parenteral nutrition, heparin, and factor VIII. Although these treatments are associated with some risk of complications, they were still found suitable for home therapy. Among the major advantages of home therapy are the ability to maintain a near-normal lifestyle, including work or school attendance, in a comfortable and private environment; the patient’s choice of the most convenient time for therapy; and cost-effectiveness. With a proper selection of patients, basic education, and periodic monitoring and with the advent of various access devices that provide an easy way to insert a needle, intravenous home treatment has become feasible for very sick patients who otherwise would be hospitalized for long periods of time. Although originally used mainly by adults, recent reports have shown the applicability of this approach to children.

Low-dose enzyme-replacement therapy for Gaucher disease seems to be an ideal candidate for intravenous home therapy. The treatment, administered over a long period of time with high frequency, is very safe and practically free of significant complications. Most of the patients are clinically stable and, therefore, do not require frequent examinations by a physician. In some of the patients, especially children, contact with other patients with various hematologic and malignant disorders may create additional fears. The lowering of the cost of the treatment is particularly important, because alglucerase is presently one of the most expensive drugs in the world.

Because, safety is a major consideration, after a single report of a putative anaphylactic reaction in a patient treated by the high-dose protocol in hospital (G. Pastore, Mount Sinai Medical Center, New York, NY, personal communication), patients were treated as outpatients in the United States dollars, $1,700, $4,810, and $2,750, respectively, per month. The costs of home treatment plus a monthly visit to the hospital for follow-up are estimated to be $240, $500, and $320, respectively, representing approximately a 90% reduction in the costs (excluding the cost of the enzyme).

**Table 1. Person Performing the Infusion**

<table>
<thead>
<tr>
<th>Person Performing Infusion</th>
<th>No. of Infusions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient</td>
<td>19</td>
</tr>
<tr>
<td>Parent</td>
<td>8</td>
</tr>
<tr>
<td>Brother/Sister</td>
<td>1</td>
</tr>
<tr>
<td>Son/Daughter</td>
<td>1</td>
</tr>
<tr>
<td>Spouse</td>
<td>2</td>
</tr>
<tr>
<td>Nurse</td>
<td>4</td>
</tr>
</tbody>
</table>

In 3 patients the infusions were performed by more than one person.
munication), we reviewed the course of our population of 61 Gaucher patients treated with Ceredase (both in the hospital and at home) and found 1 patient with a single minor episode of hives and pruritus that did not reoccur during later infusions. Combining the minimal adverse effects with the large number of infusions (~9,000 to date), we suggest that the safety profile of low-dose Ceredase compares very favorably with that of other medications that are administered at home. The absence of severe infectious complications is also remarkable in this population of Gaucher patients, which includes some who are relatively immunocompromised because of splenectomy or defective neutrophil function. Thus, we find that home intravenous enzyme-replacement therapy for Gaucher disease is safe, feasible, and well accepted by the patients and their families.

REFERENCES

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