Substrate reduction therapy as a new treatment option for patients with Gaucher disease type 1: A review of literatures

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Introduction

Gaucher disease type 1 (GD1) is an inherited lysosomal storage disorder caused by deficiency of acid β-glucosidase. The diminished enzyme activity leads to the accumulation of substrates and results in multi-systemic manifestations including hepatosplenomegaly, anemia, thrombocytopenia, and bone diseases. Enzyme replacement therapy (ERT) by infusion of recombinant protein has been the standard treatment for over 20 years. Despite the successful long-term treatment with ERT, several unmet needs remain in the treatment of GD1 such as severe pulmonary and skeletal manifestations. Substrate reduction therapy (SRT) reduces the accumulation of substrates by inhibiting their biosynthesis. Eliglustat, a new oral SRT, was approved in United States and Europe as a first-line therapy for treating adult patients with GD1 who have compatible CYP2D6 metabolism phenotypes. Although eliglustat is not yet available in Korea, introduction and summary of this new treatment modality are provided in this paper by review of literatures. Despite the fact that there are only limited studies to draw resolute conclusions, the current data demonstrated that eliglustat is not inferior to ERT in terms of its clinical efficacy. The approval of eliglustat enables eligible adult GD1 patients to have the option of oral therapy although it still needs further studies on long-term outcomes. The individual patient should be assessed carefully for the choice of treatment modality when eliglustat becomes available in Korea. Furthermore, the clinical guidelines for Korean patients with GD1 regarding the use of eliglustat needs to be developed in near future.

Key words: Gaucher disease, Substrate reduction therapy, Enzyme replacement therapy, Eliglustat.
GBA gene [2].

For treatment of GD1, enzyme replacement therapy (ERT) with intravenous infusion of recombinant acid β-glucosidase has been the standard treatment for more than two decades. ERT has proven to improve systemic manifestations of GD1 by degradation of accumulated glucosylceramide [2–4]. Despite the successful long-term treatment with ERT, there are still several unmet needs in the treatment of GD1 that continue to exist. Some patients experience skeletal symptoms or severe pulmonary symptoms which are refractory to ERT [5].

Substrate reduction therapy (SRT) reduces the production of the substrates by inhibiting glucosylceramide synthase, the rate-limiting step of glucosylceramides synthesis, thereby decreasing the over-production and accumulation of substrates [6]. The drugs for SRT are administered orally in contrast to intravenous ERT. The first developed oral SRT drug for treatment of GD1 was miglustat (Zavesca®, Actelion Pharmaceuticals, Allschwil, Switzerland), which was approved for mild to moderately affected GD1 patients who cannot receive ERT [7,8]. Unlike miglustat which is a second-line therapy to ERT, eliglustat (Cerdela®, Sanofi Genzyme, Cambridge, MA, USA) was approved by the Food and Drug Administration (FDA; 2014) and the European Medicines Agency (EMA; 2015) as a first-line treatment for adult patients with GD1 who are CYP2D6 extensive, intermediate, or poor metabolizers (>90% of patients) [9–11]. The recommendations of eliglustat treatment are also recently published by advisory council of experts in US and Europe [10,11].

Although ERT for patients with GD1 has been used successfully for more than 20 years in Korea, the physicians have no clinical experience with eliglustat as it is not yet available in Korea. The aim of this paper is to introduce the new treatment modality using eliglustat in managing patients with GD1 by review of literatures. This review could help to provide guidelines for the selection of treatment options of Korean GD1 patients in near future.

Enzyme Replacement Therapy

ERT has been treatment of choice for over 20 years since it has been available in early 1990s. Three recombinant acid β-glucosidase products are approved for the treatment of GD1; imiglucerase (Cerezyme®, Sanofi Genzyme), velaglucerase alpha (VPRIV®, Shire Human Genetic Therapies, Lexington, MA, USA), and taliglucerase alpha (ELELYSO®, Pfizer Labs, New York, NY, USA). In Korea, the non-comparable biologic imiglucerase (Abcertin®, ISU Abxis, Seoul, Korea) is also available since its approval by Korean Ministry of Food and Drug Safety in 2012 [12]. ERT is administered by intravenous infusion of 60 U/kg of drug every other week. ERT leads to breakdown of stored glucosylceramide and can achieve amelioration of hepatosplenomegaly, anemia, thrombocytopenia, and bone manifestations [2–4]. Longitudinal data from International Collaborative Gaucher Group (ICGG) Gaucher Registry demonstrated that most of the hematologic and visceral benefits occur during the first year of ERT, with maintained or sustained improvement in all parameters for at least 8 years of treatment [2,13]. Although approximately 7% of patients have experienced recurrent mild adverse events, the majority of adverse events were mild infusion-related reactions including itching and urticaria during infusions, which could be easily controlled by slowing the infusion rate and using antihistamine or corticosteroid [2]. Nonetheless, there are several limitations of ERT that entail a great burden for patients due to nature of life-long intravenous injections. First of all, the inconvenience of every other week infusion can interrupt social activities of the patients and reduce quality of life. The expensive medical costs also should be considered. Furthermore, ERT cannot completely prevent severe pulmonary or bone complications despite the long-term treatment [14].

Substrate Reduction Therapy

SRT aims to reduce accumulating glucosylceramide and related materials by inhibiting their synthesis. Comparing with recombinant proteins used in ERT, inhibitors of glucosylceramide synthesis used in SRT are small molecules that can be taken orally and potentially diffuse into various tissues, including the central nervous system and bones [15].

Miglustat, the first developed SRT drug, has been available in both Europe and US since its approval in 2002 and 2003, respectively [11]. Significant improvement in hepatosplenomegaly and biochemical markers has been demonstrated with miglustat treatment [7,8,16]. However, miglustat, iminosugar of a synthetic analog of D-glucose, is a non-selective inhibitor of glucosidases as well as an inhibitor of glucosylceramide synthase and causes considerable side effects including gastrointestinal complaints (abdominal cramping and diarrhea) by inhibiting intestinal glycosidases. These complications have led many patients to discontinue the treatment [7,8], designating miglustat as a second-line therapeutic option for limited cases who cannot receive ERT [11].
Although miglustat can cross the blood brain barrier in murine models, improvements of neurological manifestations in type III GD are controversial [15,16]. It remains as the first-line drug for Niemann-Pick type C disease.

Eliglustat, a new selective glycosylceramide synthase inhibitor, was recently approved as a first-line therapy for adult patients with GD1 by FDA and EMA [10,11]. Eliglustat has stronger inhibitory potency than miglustat [15]. In contrast to miglustat, eliglustat does not cause gastrointestinal side effect because it is not a potent inhibitor of intestinal glycosidases that prevents off-target actions.

Phase II and III clinical trials demonstrated that eliglustat significantly reduces spleen and liver volumes, increases levels of hemoglobin and platelet counts compared with placebo in treatment-naïve patients with GD1 [17,18] and maintains long-term stability [17,19]. In patients whose disease status had been stabilized with ERT, a switch to eliglustat treatment showed non-inferiority compared with ERT [20]. Although eliglustat has been well-tolerated in clinical trials, the long-term observation for monitoring of adverse event is necessary for this newly approved drug. Commonly reported adverse events include headache, migraine, arthralgia, nausea, abdominal pain, diarrhea, and dizziness which were mild to moderate in severity [10]. As the effect of eliglustat on possible long-term complications of GD such as multiple myeloma, hematologic malignancies, Parkinsonism and peripheral neuropathy is currently undetermined [10], the long-term follow up is needed.

**Eliglustat Dosing and Drug Interactions**

Dosing of eliglustat is determined based on the patient's genetic CYP2D6 metabolizer status because eliglustat is extensively metabolized by CYP2D6, and less extensively, CYP3A [11]. There are four main phenotypes of CYP2D6 metabolizers poor, intermediate, extensive, and ultra-rapid. A recommended dose has not been determined for ultra-rapid metabolizers and eliglustat therapy is not recommended in these patients currently [11]. Therefore, CYP2D6 genotyping is necessary to determine the patient's eligibility and dosing. More than 90% of Caucasian population is known to be extensive or intermediate metabolizer. However, the distribution of CYP2D6 polymorphism is not well known in Korean population. Lee et al. [21,22] reported that approximately 98% of Koreans are extensive or intermediate metabolizer among 400 Koreans assessed in the study. Furthermore, if the patient uses concomitant drugs that can affect CYP2D6 or CYP3A activity, the dose of eliglustat is recommended to be adjusted. The recommendations for eliglustat standard dosing and recommendations with concomitant drugs are summarized in Table 1.

The use of eliglustat in patients with renal insufficiency or cardiac conditions has not been studied yet and eliglustat is not recommended in patients with such underlying diseases [11]. Also, the use in a pregnant or lactating woman with GD is not recommended either [11].

**Future Research Aspect of Substrate Reduction Therapy**

Additional clinical trials are warranted to characterize the efficacy of eliglustat in extended population cohort such as patients aged 65 years and older and children. As children were not included in the clinical trials, the additional studies are necessary to ensure its safety and efficacy in these populations. Further studies are also needed to evaluate the long-term efficacy and safety of eliglustat in these additional patient populations.
required to evaluate the efficacy, safety, and dosing strategies in pediatric population. Moreover, additional research is needed in CYP2D6 ultra–rapid metabolizers, who are currently not eligible for eliglustat therapy.

Potential advantages of SRT modalities over unmet needs of ERT are considered to be better outcomes of bone complications due to easier drug delivery to bones [15]. This issue needs to be clearly demonstrated by long-term clinical trial.

Although eliglustat crosses the blood–brain barrier, it is immediately transported back out of the brain by the multidrug transporter Pgp-1 [23]. This suggests that eliglustat would be ineffective for treating neuronopathic GD. Therefore, research for developing new small molecule which can be distributed into brain would be warranted.

Comparison of Substrate Reduction Therapy with Enzyme Replacement Therapy

Although the randomized controlled studies for direct comparison of eliglustat treatment with ERT has not been properly performed yet, two recent reports demonstrated the comparative outcomes of SRT and ERT. Ibrahim et al. [24] compared the clinical response to eliglustat in 46 treatment–naïve patients (26 patients’ data from phase II and 20 patients’ data from phase III ENGAGE clinical trials) with 75 imiglucerase–treated patients (data from ICGG Gaucher registry) by post-hoc comparison. The authors demonstrated that the degree of improvement in organ volumes and hematologic parameters from baseline was similar between eliglustat–treated patients and imiglucerase–treated patients during the initial 9 to 12 months of treatment [24]. Smid et al. [15] also demonstrated that biochemical markers including chitotriosidase, CCL18 and glucosylphingosine decreased comparably in small number of patients receiving eliglustat treatment (among six patients, four were treatment naïve and two were switched from ERT) and ERT (n=4). At this moment, the limited studies are insufficient to conclude resolutely whether ERT and SRT using eliglustat are equivalent in clinical efficacy and safety. Nonetheless, at least several data show that SRT using eliglustat is not inferior to ERT in terms of reduction of liver and spleen volume, and improvements in hematologic and biochemical parameters. The extended comparative studies on both SRT and ERT in larger

![Fig. 1. Suggested algorithm to determine eligibility of eliglustat therapy in adults with Gaucher disease type 1. Modified from the article of Balwani et al. (Mol Genet Metab 2016;117:95-103) [10]. *Long QT syndrome, use of Class IA, IC or III antiarrhythmic agents, congestive heart failure, recent acute myocardial infarction, bradycardia, heart block, ventricular arrhythmia.](image-url)
cohort are needed to determine optimal clinical guidelines for treatment of patients with GD1.

Deciding Choice of Therapy

SRT with eliglustat approved as a first-line therapy for eligible patients with GD1 provides a convenient daily oral therapy as an alternative to ERT in Europe and US [10,11]. Although eliglustat is not yet available in Korea, the physicians and patients would consider the eligibility of this new treatment modality as it can be available in near future. The decision to choose of therapy should be based on the patient characteristics, underlying conditions, individual patient’s needs and/or preferences for therapy, and access to each type of therapy [10]. For example, eliglustat is currently approved only for adult GD1 patients older than 18 years, whereas ERT is approved for children as well as adults. Therefore, symptomatic children should be started on ERT because early treatment of symptomatic patients can improve outcomes [2]. Balwani et al. [10] suggested algorithm for determining eligibility of eligulstat therapy in adults with GD1 (Fig. 1).

Conclusion

ERT has been the mainstay of treatment in GD1 patients for more than 20 years. However, there are unmet needs in ERT. The approval of eliglustat as a first-line therapy enables eligible adult GD1 patients to have the option of oral therapy although it still needs further studies on long-term outcome. Therefore, physicians should carefully assess individual patient to determine the choice of treatment modality and appropriateness of the therapy when eliglustat becomes available. Furthermore, the clinical guidelines for Korean patients with GD1 regarding the SRT using eliglustat, including starting therapy and monitoring patients, needs to be developed in near future.

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References