Short Communication

Long-term outcomes with agalsidase alfa enzyme replacement therapy: Analysis using deconstructed composite events

Michael Becka,⁎, Derralynn Hughesb, Christoph Kampmannc, Guillem Pintos-Morelc, Uma Ramaswami, Michael L. Westd, Roberto Giugliane

a University Medical Center, Mainz, Germany
b Royal Free London NHS Foundation Trust, University College London, London, UK
c University Hospital and Research Institute " Germans Trias i Pujol", Badalona, Universitat Autònoma de Barcelona, Barcelona, Spain
d Department of Medicine, Dalhousie University, Halifax, NS, Canada
e Medical Genetics Service HCPA/Department of Genetics UFRGS and INAGEMP, Porto Alegre, Brazil

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ABSTRACT

This is a retrospective analysis of Fabry Outcome Survey data from children/adults (n = 677) receiving agalsidase alfa enzyme replacement therapy for a median of 3 years, examining cerebrovascular, cardiac, and renal morbidity endpoints separately. Cardiac events occurred at younger ages than cerebrovascular or renal events, cerebrovascular events were more frequent in females than males, and males were more likely to experience cardiac and renal events at a younger age than females.

1. Introduction

Fabry disease (OMIM 301500) is a rare inherited lysosomal storage disorder associated with reduced activity of α-galactosidase A (EC 3.2.1.22) [1] and characterized by significant morbidity, including renal impairment, cardiomyopathy, and cerebrovascular events [2–5].

In a recent analysis, we retrospectively compared long-term composite morbidity outcomes in patients from the Fabry Outcome Survey (FOS; sponsored by Shire) who were receiving agalsidase alfa enzyme replacement therapy (ERT) with published outcomes in untreated patients with Fabry disease [6]. Long-term ERT appeared to slow the progression of renal impairment and cardiomyopathy, and composite morbidity events occurred later in treated patients. Our current objective was to further examine and describe long-term morbidity rates separately by cerebrovascular, cardiac, and renal event origin in patients from FOS receiving agalsidase alfa ERT.

2. Materials and methods

Data were extracted from FOS for the period from database inception in 2001 to November 2012. Patient disposition is shown in Fig. A.1. The FOS registry collects standardized information from patients who are managed at participating centers and provide signed informed consent. FOS has been approved by the ethics committee/institutional review board of all participating centers and all procedures were in accordance with the Declaration of Helsinki of 1975, revised in 2013.

Members of the FOS evaluable treated cohort (n = 677) had all received agalsidase alfa ERT, including 86 children and 591 adults who initiated treatment before or after FOS entry. Patients who had received a kidney transplant or initiated dialysis before starting ERT were excluded, as were those who had received another ERT before FOS entry; 66 patients who left FOS were censored from the date they left.

Estimated glomerular filtration rate (eGFR) was calculated in adults (patients ≥ 18 years of age) only, using a four-variable Modification of Diet in Renal Disease equation [7]. Baseline left ventricular hypertrophy (LVH) was defined by left ventricular mass index (LVMI) > 48 g/m².7 in females or > 50 g/m².7 in males. Baseline data regarding proteinuria and Fabry phenotype were not available.

Morbidity as age at first event was examined separately for cerebrovascular, cardiac, and renal events. Cerebrovascular events included stroke, transient ischemic attack, or prolonged reversible ischemic neurologic deficit. Cardiac events included myocardial infarction, cardiac surgery, arrhythmia, angina, or heart failure. Renal events included renal transplant, dialysis, or ≥ 33% increase (in two consecutive values) from baseline in serum creatinine. Kaplan-Meier analyses were conducted for age at morbidity (from birth to first event) with censoring at last visit.

Abbreviations: eGFR, estimated glomerular filtration rate; ERT, enzyme replacement therapy; FOS, Fabry Outcome Survey; LVH, left ventricular hypertrophy; NE, not estimable

⁎ Corresponding author at: Children’s Hospital, University Medical Center, J.G. University Mainz, Langenbeckstrasse 1, 55101 Mainz, Germany.

E-mail address: Michael.Beck@unimedizin-mainz.de (M. Beck).

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3. Results

Baseline demographic and clinical characteristics are shown in Table A.1. Among the 677 patients (360 male, 317 female) in the overall FOS evaluable treated cohort, median (range) age at first Fabry disease symptom was 12 (0–72) years and median (range) age at diagnosis was 31 (0–79) years. Median (range) time on treatment (time from treatment initiation to data extraction) was 3 (0–14) years. At baseline, 63/447 (14.1%) of adult patients with a baseline eGFR value available had eGFR < 60 mL/min/1.73 m² and 143/330 (43.3% of patients with a baseline LVMi value available) had LVH.

In the overall FOS evaluable treated cohort, the number and percentage of patients who did not experience an event during follow-up was: cerebrovascular, 578 (85.4%); cardiac, 378 (55.8%); renal, 614 (90.7%); any morbidity event, 326 (48.2%; Table A.2). The Kaplan-Meier point estimate (95% CI) for median age in years at first event was: cerebrovascular, 70.9 (68.1, not estimable [NE] because of the small number of events) in females and NE (64.9, NE) in males (Fig. 1A); cardiac, 59.9 (56.8–61.9) in females and 50.6 (48.4–52.7) in males (Fig. 1B); renal, NE (NE, NE) in females and 78.4 (70.3, NE) in males (Fig. 1C).

Among female patients, the probability of not having a renal event was relatively high even at advanced ages; for example, at age 75 years, the probability of not having a renal event was ~90% in women and ~55% in men.

4. Discussion

The current analysis is an extension of our retrospective analysis [6], which found that agalsidase alfa ERT appeared to slow the progression of renal impairment and LVH, and composite morbidity events (including death) occurred later, in FOS patients receiving agalsidase alfa ERT compared with published findings in untreated patients with Fabry disease. The current work represents a purely descriptive analysis of the status of treated patients; because of the lack of external comparator groups, no comparison of agalsidase alfa treated and untreated patients was possible. Our findings are consistent with other published analyses showing stable or improved cardiac or renal parameters in patients receiving ERT with agalsidase alfa [3,8–12]. Compared with similar analyses in the published literature [8,13–15], our analysis is based on the largest dataset of clinical events in the long-term treatment of Fabry disease to date. However, the number of renal events recorded in the FOS database was still relatively small, limiting the analysis of renal events and indicating that renal function is well maintained over years of ERT.

Although the current analysis showed a somewhat higher proportion of patients with an event than that reported by Germain et al. [8], the definition of events was not comparable between these two analyses. For consistency with our previous analysis [6], which compared treated patients from FOS with untreated patients in the published literature [16], the definition of events was broader in our FOS analysis than in the analysis of Germain et al. [8], and therefore some of these events were less severe (e.g., transient ischemic attack, prolonged reversible ischemic neurologic deficit). In contrast, Germain et al. focused on severe clinical events, which were defined as "chronic dialysis, kidney transplant, myocardial infarction, congestive heart failure, major cardiac procedures (i.e., implantation of a balloon pump, cardioverter-defibrillator or first pacemaker; or bypass surgery), stroke and death" [8].

Several limitations of our analysis should be noted. This was a retrospective analysis of FOS registry data and the FOS patients were not randomly selected, which may contribute to selection and/or ascertainment bias. In general, registry analyses have a higher probability of patient loss from the analysis because of the relative lack of data completeness compared with randomized controlled trials. Comparisons were not made with untreated patients from this registry; consequently, no definitive conclusions can be made regarding effectiveness of ERT. FOS lacks a robust cohort of untreated patients for comparison with treated patients. In general, untreated patients in FOS have less severe disease and undergo fewer follow-up assessments; thus, fewer outcome data are available [17]. Although LVMi is not itself a clinical endpoint, LVH has been associated with both cardiac and renal clinical outcomes in patients with Fabry disease [17,18]. A potential source of bias is the censoring of patients who discontinued agalsidase alfa ERT.

Over a median period of 3 years of agalsidase alfa ERT in this registry population of patients with long-standing Fabry disease, cardiac events occurred more frequently and at younger ages than cerebrovascular or renal events. Cerebrovascular events were more frequent in females than in males. Males were more likely to experience cardiac and renal events at a younger age than females. It is possible that the presence of cardiac and renal events in males, and their subsequent treatment, may decrease the incidence of later cerebrovascular events relative to the incidence in females. Although we analyzed a cohort of FOS patients with a median age of 46.2 years, one must also consider that, from a US population perspective, the 10-year risk of stroke is higher in middle-aged women aged 55 years than middle-aged men aged 55 years [19]. The main determinant of stroke risk other than advancing age is blood pressure [19], which an earlier analysis showed to be uncontrolled in over half of untreated FOS patients at baseline before declining after 2 years of agalsidase alfa ERT [20]. Analysis of 10-year outcomes in patients receiving agalsidase alfa ERT using deconstructed composite events will shed more light on these observations.

Role of the funding source

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Therapies, Inc. Shire International GmbH provided funding for data collection and analysis, and provided funding to Excel Scientific Solutions for support in writing and editing this manuscript. Under the direction of the authors, Margit Rezabek, DVM, PhD, employee of Excel Scientific Solutions, provided writing assistance for this publication. Editorial assistance in formatting, proofreading, copyediting, and fact checking was also provided by Excel Scientific Solutions. Jaco Botha, MSc, of Shire International GmbH, and Svetlana Bizjajeva, PhD, formerly of Shire International GmbH, provided statistical support and reviewed and edited the manuscript for scientific accuracy. Sylvain Larroque, MSc, formerly of Shire International GmbH, planned and conducted statistical analyses.

Contributions

All authors reviewed and revised early and final drafts of the manuscript and were fully responsible for editorial decisions related to this manuscript.

Conflict of interest

Michael Beck has received unrestricted grants, travel support, and honoraria from Actelion, BioMarin, Genzyme, Shire, and Synageva. Derralynn Hughes has received travel/research grants and honoraria for speaking/consulting from Amicus, Protalix, Sanofi-Genzyme, and Shire. Christoph Kampmann has received research support from BioMarin and Shire; consulting/speaker fees from Amicus, BioMarin, Gore, and Shire; participated in clinical studies sponsored by Actelion, BioMarin, Genzyme, Gore, and Shire; and served on advisory committees for BioMarin and Shire. Guillo Pintos-Morell has received travel grants/honoraria for consulting/lectures from Alexion, BioMarin, and Shire. Uma Ramaswami has received travel/research grants and honoraria for lectures/consulting from Amicus, Alexion, Genzyme, Protalix, and Shire. Michael L. West has received research grants, consulting/speaker fees, and/or travel support from Actelion, Alexion, Amicus, AvroBio, Excelsior, GlaxoSmithKline, Sanofi-Genzyme, Shire, and Sumitomo. Roberto Giugliani has received consulting fees from Amicus, BioMarin, Genzyme, Shire, and Synageva/Alexion; participated in clinical trials sponsored by Amicus, BioMarin, Genzyme, GlaxoSmithKline, Shire, and Synageva/Alexion; and assisted in the design of and/or participated in clinical studies using products manufactured by Actelion, BioMarin, Genzyme, and Shire.

Funding

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Appendix A

Table A.1
Demographic and clinical characteristics of patients in the FOS evaluable treated cohort.

<table>
<thead>
<tr>
<th>Race, n (%)</th>
<th>Male, n = 360</th>
<th>Female, n = 317</th>
<th>Overall FOS evaluable treated cohort, n = 677</th>
</tr>
</thead>
<tbody>
<tr>
<td>White</td>
<td>294 (82.1)</td>
<td>261 (83.9)</td>
<td>555 (83.0)</td>
</tr>
<tr>
<td>Black</td>
<td>0 (0)</td>
<td>1 (0.3)</td>
<td>1 (0.1)</td>
</tr>
<tr>
<td>Asian</td>
<td>58 (16.2)</td>
<td>40 (12.9)</td>
<td>98 (14.6)</td>
</tr>
<tr>
<td>Other</td>
<td>6 (1.7)</td>
<td>9 (2.9)</td>
<td>15 (2.2)</td>
</tr>
<tr>
<td>Age at data extract (years)</td>
<td>n = 360</td>
<td>n = 317</td>
<td>n = 677</td>
</tr>
</tbody>
</table>

Fig. A.1. Disposition of patients. ERT, enzyme replacement therapy; FOS, Fabry Outcome Survey.
Table A.2
Number (percentage) [95% CI] of patients without a morbidity event.

<table>
<thead>
<tr>
<th></th>
<th>Male</th>
<th>Female</th>
<th>Overall FOS evaluable treated cohort</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n = 360</td>
<td>n = 317</td>
<td>n = 677</td>
</tr>
<tr>
<td>Cerebrovascular</td>
<td>320 (88.9)</td>
<td>258 (81.4)</td>
<td>578 (85.4)</td>
</tr>
<tr>
<td></td>
<td>[85.2–91.9]</td>
<td>[77.1–85.7]</td>
<td>[82.7–88.0]</td>
</tr>
<tr>
<td>Cardiac</td>
<td>209 (58.1)</td>
<td>169 (53.3)</td>
<td>378 (55.8)</td>
</tr>
<tr>
<td></td>
<td>[52.9–63.2]</td>
<td>[47.6–58.9]</td>
<td>[52.0–59.6]</td>
</tr>
<tr>
<td>Renal</td>
<td>311 (86.4)</td>
<td>303 (95.6)</td>
<td>614 (90.7)</td>
</tr>
<tr>
<td></td>
<td>[82.9–89.9]</td>
<td>[93.3–97.9]</td>
<td>[88.5–92.9]</td>
</tr>
<tr>
<td>Any event</td>
<td>185 (51.4)</td>
<td>141 (44.5)</td>
<td>326 (48.2)</td>
</tr>
<tr>
<td></td>
<td>[46.2–56.6]</td>
<td>[39.0–50.0]</td>
<td>[44.4–51.9]</td>
</tr>
</tbody>
</table>

BP, blood pressure; eGFR, estimated glomerular filtration rate; FOS, Fabry Outcome Survey; FOS-MSSI, Fabry Outcome Survey Mainz Severity Score Index; IQR, interquartile range; LVH, left ventricular hypertrophy.

References


