Effects of enzyme replacement therapy on pain and health related quality of life in patients with Fabry disease: data from FOS (Fabry Outcome Survey)

B Hoffmann, A Garcia de Lorenzo, A Mehta, M Beck, U Widmer, R Ricci, on behalf of the FOS European Investigators

Background: Fabry disease is an X linked lysosomal storage disease caused by deficiency of the lysosomal enzyme α-galactosidase A. This leads to accumulation of globotriaosylceramide in nearly all tissues, including the blood vessels, kidney, myocardium, and nervous system. Symptoms often begin in childhood and include acropaesthesia, with burning or tingling pain that spreads from the extremities to more proximal sites.

Aims: This study set out to evaluate pain and its influence on quality of life in patients with Fabry disease receiving enzyme replacement therapy (ERT) with agalsidase alfa.

Methods: Data were obtained from the Fabry Outcome Survey. Pain was measured using the Brief Pain Inventory (BPI), and health-related quality of life (HRQoL) was documented with the European Quality of Life Questionnaire (EQ-SD).

Results: The mean (SD) score for “pain at its worst” on the BPI prior to ERT was 5.1 (2.7). One year after commencement of ERT, this had improved by 0.5, and improved by a further 0.6 after 2 years (p<0.05). Similar statistically significant improvements were seen for “pain on average” and “pain now” after 2 years of ERT. The mean HRQoL utility score prior to ERT was 0.66 (0.32). After 12 months of treatment with agalsidase alfa, this had improved to 0.74 (0.26; p<0.05); this improvement was maintained after 2 years.

Conclusions: ERT with agalsidase alfa significantly reduces pain and improves quality of life in patients with Fabry disease.
patients and, importantly, the effect of ERT on QoL has not been evaluated.

This paper aims to describe QoL and pain related QoL in patients enrolled in the Fabry Outcome Survey (FOS), the most comprehensive database of patients with Fabry disease, both before and during ERT with agalsidase alfa.

METHODS
FOS is a European database for patients with Fabry disease. At the time of this analysis (March 2004), 545 patients (264 female and 281 male) were registered on the database. Of these, 314 patients (58%) were receiving intravenous ERT with agalsidase alfa every 2 weeks at a dose of 0.2 mg/kg. Data are collected anonymously and entered by physicians or specialist nurses into the database. The FOS database has been approved by the ethics review boards of all participating institutions, and all patients gave written informed consent. The FOS database and its properties have been described previously.\(^\text{25}\)

Evaluating quality of life in FOS
According to the World Health Organization (WHO) Quality of Life Working Group, QoL is defined as an individual’s perception of their position in life. This has to be seen in the context of the culture and value system in which the person lives. Furthermore, it must be related to the individual’s goals, expectations, standards, and concerns.\(^\text{34}\) To evaluate an item of such complexity would appear unfeasible. The WHO definition of health as “a state of complete physical, mental, and social wellbeing and not merely the absence of disease” has similar practical limitations.\(^\text{35}\) The concept of HRQoL has therefore been established. Instruments that aim to measure HRQoL are designed solely to measure QoL with respect to an individual’s health state.

There are several instruments available to measure HRQoL. One of these is the European Quality of Life questionnaire (Euro-QOL; EQ-5D), which covers five dimensions: mobility, pain/discomfort, self care, anxiety/depression, and usual activities. Each dimension comprises three levels (no problems, some/moderate problems, extreme problems). The questionnaire is designed for self completion and validated for different languages. It was developed to measure pain and its influence on life.\(^\text{41}\) An EQ-5D score of 1 means “full health” and a score of 0 is equivalent to “death”. By calculation, it is also possible to get a health state “worse than death” (<0).\(^\text{39}\) Representative EQ-5D data from a “normal” population are available for comparison.\(^\text{36}\)

Evaluating pain in FOS
The Brief Pain Inventory (BPI) contains a series of questions relating to pain and its interference with life. Each question can be answered by circling a number between 0 and 10. According to the scoring of “pain at its worst” the patients can be assigned to one of three groups: a pain score of 1–4 is ascribed to mild pain, a score of 5–6 is defined as moderate pain, and a score of 7–10 represents severe pain.\(^\text{40}\) Like the EQ-5D, the BPI is validated for different languages. It was developed to measure pain and its influence on life.\(^\text{41}\) Although the BPI was primarily designed to measure pain in patients with cancer, it has become widely accepted in different healthcare settings where the impact of pain has to be recorded.\(^\text{42}\)–\(^\text{44}\) Finally, the BPI has been shown to be responsive to interventional treatments.\(^\text{45}\)

Data sample and statistics
The data presented here result from evaluations at baseline and after a mean (SD) of 12 (3) months and 24 (3) months of treatment with agalsidase alfa. Baseline values are taken as the value reported up to 6 months preceding or up to 3 months after the onset of ERT with agalsidase alfa.

In FOS, patients are asked to fill out both the EQ-5D and BPI questionnaires regularly. For this evaluation, we excluded children enrolled in FOS, as reliable reference data for the EQ-5D and BPI are only available for adults.

Wilcoxon’s rank sum test, Student’s t test and Spearman’s rank correlation were used where appropriate. Numbers are presented as mean (SD).

RESULTS
HRQoL
Prior to ERT, the mean EQ-5D utility score was 0.66 (0.32) \((n = 120; 47 \text{ women}, 73 \text{ men})\), with no difference observed in utility score between sexes. The EQ-5D utility scores were significantly lower than those reported for the normal UK population, matched for age and sex \((p<0.05; \text{table 1})\).\(^\text{38}\)

Information was available from 59 patients \((20 \text{ women}, 39 \text{ men})\) prior to ERT and after 1 year of treatment. These patients had a mean (SD) EQ-5D score of 0.64 (0.32) at baseline. After 12 months of treatment with agalsidase alfa,

<table>
<thead>
<tr>
<th>EQ-5D utility score</th>
<th>Difference from normal UK population data</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median (10th to 90th percentile)</td>
<td>Mean (SD)</td>
</tr>
<tr>
<td>All patients ((n = 120))</td>
<td>0.76 (0.13 to 1.00)</td>
</tr>
<tr>
<td>Women ((n = 47))</td>
<td>0.80 (0.12 to 1.00)</td>
</tr>
<tr>
<td>Men ((n = 73))</td>
<td>0.76 (0.14 to 1.00)</td>
</tr>
</tbody>
</table>

*p<0.05 difference between patients with Fabry disease and UK reference data.

Figure 1 Improvement in health related quality of life after 1 year of treatment with agalsidase alfa in patients with Fabry disease \((n = 59; 20 \text{ women}, 39 \text{ men})\) measured using the European Quality of Life Questionnaire (EQ-5D).

Table 1 Median and mean European Quality of Life Questionnaire (EQ-5D) utility scores at baseline in 120 patients with Fabry disease compared with data from the normal UK population\(^\text{38}\)
this had improved significantly (0.74 (0.26), p<0.05; fig 1) and no longer differed from the reference data for the normal UK population. No differences were found between men and women (data not shown). On the pain/discomfort dimension of the EQ-5D, 14 patients reported no problems, 37 reported moderate problems, and 8 reported extreme problems at baseline. An improvement on this scale occurred after 1 year of treatment with agalsidase alfa, with 21 patients reporting no problems, 36 reporting moderate problems, and 2 reporting extreme problems.

For 28 patients (four women, 24 men), longitudinal data were available for two consecutive years of ERT. The mean (SD) EQ-5D score prior to ERT in these patients was 0.50 (0.32). The significant improvement seen after 1 year was maintained after 2 years of treatment with agalsidase alfa in this group (fig 2), and, likewise, the EQ-5D score after 2 years did not differ significantly from the scores for the UK reference population.

On the pain/discomfort dimension of the EQ-5D, 1 patient had reported no problems at baseline, 21 had reported moderate problems, and 6 had reported extreme problems. Following 2 years of treatment with agalsidase alfa, 9 of these patients now reported no problems, 18 reported moderate problems, and 2 reporting extreme problems.

### Pain

BPI scores prior to the initiation of ERT with agalsidase alfa are shown in table 2. The mean BPI score for ‘‘pain at its worst’’ was 5.1 (2.7) and the mean score for ‘‘pain at its least’’ was 2.1 (2.5). Before beginning ERT, patients reported that in the previous 24 hours pain interfered most with general activity (3.3 (3.1)) and mood (3.3 (3.2)), and least with walking ability (2.5 (3.1)) and sleep (2.5 (3.2)).

After a mean duration of 12 months of ERT with agalsidase alfa, 41 patients completed the questionnaire. ‘‘Pain on average’’, ‘‘pain now’’ and ‘‘pain at its worst’’ had improved (mean scores had decreased by 0.6, 0.4, and 0.5, respectively); however, this was not statistically significant. The mean value for ‘‘pain at its least’’ did not change (fig 3). At baseline, ‘‘pain on average’’ had been reported as none in 4 patients, mild in 21, moderate in 9, and severe in 7. Following 1 year of treatment with agalsidase alfa, ‘‘pain on average’’ was reported as none in 8 patients, mild in 20, moderate in 10 and severe in 3.

The effects of 2 years of ERT with agalsidase alfa on mean BPI scores are shown in fig 4 for the 20 patients followed longitudinally over this period. Values for ‘‘pain on average’’ decreased by 0.7 after 1 year and by a further 0.9 after 2 years of treatment. ‘‘Pain now’’ decreased by 0.9 after 1 year and by a further 0.4 after 2 years of treatment. For ‘‘pain at its worst’’ the decrease was 0.5 during the first year and 0.6 during the second year of treatment. All these changes between baseline and 2 years were statistically significant (p<0.05; fig 4). Although the mean score for ‘‘pain at its least’’ decreased by 0.5 and 0.3 after 1 and 2 years, respectively, these changes were not statistically significant. At baseline, ‘‘pain on average’’ had been reported as mild in 10 patients, moderate in 5, and severe in 5. Following 2 years of treatment with agalsidase alfa, ‘‘pain on average’’ was

### Table 2 Brief Pain Inventory scores before enzyme replacement therapy in patients with Fabry disease

<table>
<thead>
<tr>
<th>BPI dimension</th>
<th>n</th>
<th>Mean (SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pain at its worst</td>
<td>90</td>
<td>5.1 (2.7)</td>
</tr>
<tr>
<td>Pain at its least</td>
<td>89</td>
<td>2.1 (2.5)</td>
</tr>
<tr>
<td>Pain on average</td>
<td>90</td>
<td>4.1 (2.5)</td>
</tr>
<tr>
<td>Pain now</td>
<td>91</td>
<td>2.9 (2.8)</td>
</tr>
</tbody>
</table>

Interference of pain with:
- General activity  102 3.3 (3.1)
- Mood              103 3.3 (3.2)
- Walking ability   102 2.5 (3.1)
- Normal work       101 3.4 (3.4)
- Social relations  103 2.7 (3.0)
- Sleep             102 2.5 (3.2)
- Enjoyment of life 103 3.0 (3.2)

In total, 114 patients completed the questionnaire, although not all patients answered all questions.
reported as none in 4 patients, mild in 12, moderate in 3, and severe in 1.

**Relationship between HRQoL and pain**

At baseline, there was a statistically significant negative correlation between the EQ-5D and all dimensions of the BPI (fig 5). There was no correlation between pain or EQ-5D score and age.

**DISCUSSION**

QoL is impaired in patients with Fabry disease prior to treatment with ERT. In the current study, the EQ-5D scores of patients with Fabry disease at baseline were found to be significantly decreased compared with age and sex matched control data from the UK population. Compared with other patient groups who have completed this questionnaire, patients with Fabry disease were found to score slightly worse on the EQ-5D than those who have undergone liver transplantation and women with breast cancer; however, worse on the EQ-5D than those who have undergone liver transplantation and women with breast cancer; however, the EQ-5D score illustrates that this moderate pain affects QoL.

After 12 months of treatment with agalsidase alfa, QoL improved significantly in patients with Fabry disease, with no differences observed between men and women. Importantly, the improvement was maintained after 24 months of treatment. In contrast, EQ-5D scores decrease with age in the normal population and in untreated patients with Fabry disease.

Similar to other questionnaires that generate a generic index of health states, such as the Short Form-36, the EQ-5D has some limitations. It was established to detect and to measure changes in HRQoL reported by the patient and thus does not provide an objective measure with which to detect an improvement or deterioration in health state. Therefore, improvement of HRQoL does not imply improvement in an individual's physical health state, but an improvement in the patient's perception of their health state. Several reports have described improvements in different organ systems under ERT, but it is unclear whether these are reflected by improvements in the patient’s perception of health. Measures of HRQoL, or so called “patient reported outcomes”, have become useful and important tools in evaluating healthcare interventions. Such tools are likely to be important in evaluating therapy in Fabry disease, as there are no biomarkers of disease severity in this condition.

Before treatment with agalsidase alfa, the pain reported by patients with Fabry disease was, on average, categorised as moderate (mean (SD) BPI score, 5.1 (2.7)); however, the term “moderate pain” is unlikely to reflect the whole burden of the pain experienced by patients with Fabry disease. The statistically significant negative correlation between pain and the EQ-5D score illustrates that this moderate pain affects HRQoL and thus has a major impact in patients with Fabry disease.

Scores for “pain on average”, “pain now”, and “pain at its worst” on the BPI improved, but not significantly, following 1 year of treatment with agalsidase alfa. However, after 2 years of treatment, there was a significant improvement in all three scores.

Several questionnaires, in particular the BPI, have been accepted as useful tools for evaluating pain. Nevertheless, there may still be concerns about using such purely subjective instruments. A more objective measure of pain evaluation is quantitative sensory testing, as used by Dütsch et al to evaluate heat pain perception in patients with Fabry disease. This group presented 25 stimuli of different intensity up to 49°C to patients with Fabry disease and asked the patients to grade their response to the stimulus on a visual analogue scale ranging from 0 (no discomfort or pain) to 10 (most discomfort or pain). Their findings clearly indicated impaired Aδ and, to a lesser extent, C nerve fibre function in patients with Fabry disease. These findings could explain the lack of improvement of neuropathic pain in some patients in our cohort, as it is unlikely that pain perception via structurally damaged nerve fibres can be improved. Additionally, these findings provide support for early ERT in order to prevent irreversible structural damage to nerve tissue.
CONCLUSION

Fabry disease has a major effect on QoL. Patients with Fabry disease show significantly lower EQ-5D scores compared with normative population data. ERT with agalsidase alfa (Replagal) significantly improves QoL in patients after 1 year of treatment. These promising results were sustained after 2 years of ERT. Pain is a major contributor to the decreased QoL in Fabry disease, and “pain on average”, “pain now”, and “pain at its worst”, as measured by the BPI, were all significantly reduced after 2 years of ERT with agalsidase alfa.

ACKNOWLEDGEMENTS


Authors’ affiliations

B Hoffmann, University Children’s Hospital, Heinrich Heine University, Düsseldorf, Germany
A G de Lorenzo, Hospital Universitario, La Paz, Madrid, Spain
A Mehta, University College London, London, UK
M Beck, University of Mainz, Mainz, Germany
U Widmer, University of Zurich, Zurich, Switzerland
R Richfield, Institute of Clinical Pediatrics, UCSC, Rome, Italy

The FOS database is under the independent control of the FOS European board. Data analysis and conclusions in FOS are supported by TKT Europe–SS, Danderyd, Sweden. The sponsor had no role in the interpretation of data or writing of this report.

Competing interests: none declared

REFERENCES


Effects of enzyme replacement therapy on pain and health related quality of life in patients with Fabry disease: data from FOS (Fabry Outcome Survey)
B Hoffmann, A Garcia de Lorenzo, A Mehta, M Beck, U Widmer and R Ricci

doi: 10.1136/jmg.2004.025791

Updated information and services can be found at:
http://jmg.bmj.com/content/42/3/247

These include:

**References**
This article cites 42 articles, 5 of which you can access for free at:
http://jmg.bmj.com/content/42/3/247#BIBL

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Topic Collections**
Articles on similar topics can be found in the following collections

- Metabolic disorders (329)
- Immunology (including allergy) (604)

**Notes**

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/