CLINICAL STUDY

Associations of remission status and lanreotide treatment with quality of life in patients with treated acromegaly

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Abstract

Objective: Acromegaly has an important impact on quality of life (QOL). The aim of this study was to evaluate the associations of remission status and lanreotide treatment with QOL in patients with treated acromegaly, by the newly developed disease-generated acromegaly QOL questionnaire (ACROQOL).

Design: Cross-sectional study.

Methods: Fifty-two patients with treated acromegaly were recruited to complete the Chinese version of the ACROQOL translated and validated from the English version. These patients were divided into controlled and uncontrolled groups based on the latest remission criteria and further subdivided into four groups according to the present treatment with lanreotide or not. Comparisons between groups were analyzed.

Results: There was no difference between controlled and uncontrolled groups in the ACROQOL scores of total score, both scales and psychological subscales. However, in the controlled group, present treatment with lanreotide, in comparison with no treatment, showed worse ACROQOL scores in total score \( P = 0.021 \), psychological scale \( P = 0.011 \), psychological subscale ‘appearance’ \( P = 0.032 \) and ‘personal relations’ \( P = 0.010 \).

Conclusions: The lanreotide treatment was negatively associated with QOL in biochemically controlled acromegalic patients, especially in the psychological aspect.

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Introduction

Acromegaly is a rare disorder with an annual incidence of 3–4 cases per million and a prevalence of 50–70 cases per million population, being more common in women and mostly caused by a growth hormone (GH)-secreting pituitary adenoma (1–2). Mortality in acromegaly is at least twice that of the general population, mostly related to cardiovascular diseases and malignancies (3–8). One large retrospective cohort study demonstrated that post-treatment GH levels <2.5 μg/l (116.25 pmol/l) result in an overall mortality rate similar to that of the general population (4). The most recent broadly accepted definition of biochemical cure of acromegaly was based on the Internal Consensus Conference of Cortina d’Ampezzo in 1999 (9). Aggressive management is mandatory to achieve biochemical disease control and thereby restore life expectancy to normal. Present treatment modalities include transsphenoidal surgery, radiotherapy and medical therapy (dopamine agonists, somatostatin analogues and GH receptor antagonist).

Recently, besides the biochemical variables, clinicians have started to focus on the health-related quality of life (HR-QOL) as the outcome measurement of acromegaly and its treatment. The acromegaly quality of life questionnaire (ACROQOL), initially developed in Spain and first published in 2002, is the first acromegaly disease-generated quality of life questionnaires to specifically assess dimensions of HR-QOL most likely to be affected in patients with acromegaly (10–13). The original Spanish questionnaire was first translated into English following recommended standard methodology (10, 14).

Several studies have demonstrated that administration of somatostatin analogues ameliorates the abnormal biochemical profiles and the associated somatic impairments (15–19). However, whether these improvements translate into improved physical function remains largely unknown, and their impact on HR-QOL is still controversial (20). To date, there is paucity of data even attempting to correlate disease modification endpoints with quality of life in acromegaly.
The present study was aimed to evaluate the associations of remission status and slow-release somatostatin analogue (lanreotide) treatment with HR-QOL in patients with treated acromegaly using the Chinese version of ACROQOL.

Subjects and methods

Study protocol

Using a cross-sectional design, we prospectively recruited the patients with treated acromegaly between September 2005 and January 2006 at the Division of Endocrinology and Metabolism of the National Taiwan University Hospital, Taipei, Taiwan. At first, we translated the validated English version of ACROQOL into Chinese following recommended standard methodology (10, 14). The English version of ACROQOL is shown in Figure 1, comprising 22 questions covering physical (8 items) and psychological aspects (14 items), which is further divided into two subscales: one assesses appearance-related aspects (7 items) and the other measures the impact of disease on personal relationships (7 items).

The English version was translated into Chinese by two professional, bilingual translators with lots of experience in translating HR-QOL questionnaires. Both translations were compared with each other and with the original English version at a consensus meeting consisting of endocrinologists specifying in acromegaly. If the translation was clear and correct, no changes were made. If there were doubts or contrasting opinions with the project director, a consensus was reached after extensive discussion, to produce the first Chinese version of the questionnaire. This version was then independently translated back into English to ascertain equivalent significance in both the languages. After the second meeting, the second consensus Chinese version was produced, and presented to three Chinese-speaking Taiwanese patients with acromegaly to assess and correct for clarity, cultural relevance, comprehension and suitable wording, thus providing the final Chinese version of the ACROQOL questionnaire.

The study project was approved by the Institutional Review Board of the National Taiwan University Hospital. Patients were asked to answer the 22 items

Scale 1 (Physical, 8 items)

- Item 1 My legs feel weak
- Item 3 I get depressed
- Item 9 I have problems carrying out my usual activities
- Item 13 The illness affects my performance at work or in my usual tasks
- Item 14 My joints ache
- Item 15 I am usually tired
- Item 19 I feel like a sick person
- Item 22 I feel weak

Scale 2 (Psychological, 14 items)

subscale 2–1 (Appearance, 7 items)

- Item 2 I feel ugly
- Item 4 I look awful in photographs
- Item 7 I look different in the mirror
- Item 11 Some parts of my body (nose, feet, hands, ...) are too big
- Item 12 I have problems doing things with my hands, e.g. sewing or handling tools
- Item 16 I snore at night
- Item 17 It is hard for me to articulate words due to the size of my tongue

subscale 2–2 (Personal relationships, 7 items)

- Item 5 I avoid going out very much with friends because of my appearance
- Item 6 I try to avoid socializing
- Item 8 I feel rejected by people because of my illness
- Item 10 People stare at me because of my appearance
- Item 18 I have problems with sexual relationships
- Item 20 The physical changes produced by my illness govern my life
- Item 21 I have little sexual appetite

Figure 1 The acromegaly quality of life questionnaire (ACROQOL).
of the Chinese version of the ACROQOL either by telephone visit or during their regular clinical visit. Out of 52 patients, 47 completed the questionnaire in our outpatient clinic and 4 by phone because they were treated in another hospital. All were conducted by the same researcher (S-C Hua) and in the same fashion. The individual clinical variables, including age, gender, age of definite diagnosis, macro- or microadenoma, operation history, repeat operation history, radiotherapy history, the previous and the present treatment modalities, latest GH/insulin-like growth factor-I (IGF-I) levels, whether on hormone replacement therapy, and morbidities related to acromegaly were recorded by chart review. Those treated by slow-release somatostatin analogues were all using lanreotide every 14 days. For the four patients treated in another hospital, we obtained the clinical variables provided by the patients themselves. Two of them had markedly elevated GH/IGF-I levels and another two had oral glucose tolerance test (OGTT) in our outpatient clinic for confirmation of suppressible GH level.

The study was performed according to the declaration of Helsinki. Each patient gave informed consent personally or by mail.

Assessment of biochemically controlled (remission) status

The remission criteria were defined according to the Internal Consensus Conference of Cortina d’Ampezzo in 1999 (9) as follows: normal age- and gender-matched IGF-I concentration, basal GH level less than 2.5 µg/l (116.25 pmol/l) and suppression of GH to <1 µg/l (46.5 pmol/l) during OGTT.

Grouping

The enrolled patients were divided into controlled and uncontrolled groups based on the remission criteria defined by the Internal Consensus Conference of Cortina d’Ampezzo in 1999. The controlled and the uncontrolled groups were further subdivided into with or without present lanreotide treatment groups.

Tumour classification

Pituitary adenomas were classified by the size on the magnetic resonance imaging, computed tomography or intra-operative findings: tumour diameter of <1 cm was defined as microadenoma. Those ≥1 cm were defined as macroadenoma.

Hormone replacement

Pituitary function of each patient was periodically evaluated. Thyroid deficiency was defined as low free thyroxine (T4) level. Adrenal insufficiency was defined as low cortisol level with requirement of glucocorticoids supplementation and failure to be tapered off. Gonadal insufficiency was defined as low testosterone (male) or oestrogen/progesterone (female) levels.

Scoring of the ACROQOL

Each of the 22 items was answered in a 1 (worst) to 5 (best) Likert scale, measuring either the frequency of occurrence (always, most of the time, sometimes, rarely and never) or the degree of agreement with the item (completely agree, moderately agree, neither agree nor disagree, moderately disagree and completely disagree). As suggested by the authors of the questionnaire (11), a global score and scores of the respective scales and subscales were obtained by adding the results using the following formula:

\[
(X - Y) ÷ (5Y - Y)100
\]

where \(X\) is the sum of the answers (between 1 and 5 for each item) of the respective scale and \(Y\) is the number of items of the respective scale (22 for the total ACROQOL-score, 8 for the physical scale, 14 for the psychological scale, 7 for the subscales ‘appearance’ and ‘personal relations’). The formula converts the different scores to a scale between a minimum of 0 (worst) and a maximum of 100 (best).

Statistical analysis

Quantitative data are presented as mean ± S.D. and qualitative data as percentages. Comparisons among groups were performed using Student’s \(t\)-test for quantitative data and \(\chi^2\)-test for qualitative variables. Both Student’s \(t\)-test and multivariate linear regression analyses were performed to assess independent association of whether current lanreotide treatment or not with the ACROQOL scores of total score, respective scales, and subscales. All statistical methods were carried out with SPSS 12.0 for Windows software (SPSS Inc., Chicago, IL, USA). A \(P\) value <0.05 was considered statistically significant.

Results

The cohort consisted of 52 patients with treated acromegaly. Half of the patients were male. The mean age was 51.9 ± 10.1 (S.D.) years, range 20–76 years. Mean time interval between diagnosis and study assessment was 12.6 ± 7.1 years, range 1–30 years. At diagnosis, 35 patients (67.3%) had a macroadenoma and 17 (32.7%) had a microadenoma. Out of these 52 patients, 16 (six male, ten female) were on hormone replacement because of surgery-related hypopituitarism. Of these 16 patients, 11 were on thyroxine replacement, 15 on glucocorticoids replacement, and 2...
males and 4 females were on sex steroid replacement. Out of ten females, two were menopausal and free of postmenopausal symptoms, who did not accept sex steroid replacement.

The clinical characteristics of the biochemically controlled and uncontrolled groups are shown in Table 1. Out of all 52 patients, 30 (57.7%) were biochemically categorized as the controlled group and 22 (42.3%) were categorized as the uncontrolled group. There was significant difference between groups in operation history, latest GH and IGF-I level ($P=0.042$, $<0.001$, $<0.001$ respectively). There was no significant difference between groups in age, gender, age of definite diagnosis, repeated operation history, radiotherapy history, percentage of macroadenoma, percentage of hormone replacement, latest GH/IGF-I level and rate of morbidity related to acromegaly.

Distribution of the ACROQOL scores of total score, scales and subscales in the four groups was shown in Table 4. There was a significant difference between the ‘controlled with present lanreotide treatment’ and ‘controlled without present lanreotide treatment’ groups in total score ($P=0.021$), psychological scale ($P=0.011$), psychological subscale ‘appearance’ ($P=0.032$) and ‘personal relationships’ ($P=0.010$).

**Discussion**

Our study demonstrated that there was no significant difference between biochemically controlled and uncontrolled acromegaly in the ACROQOL scores of total score, physical scale, psychological scale and psychological subscales. In the biochemically controlled group, those presently treated with slow-release somatostatin analogues (lanreotide), in comparison with without, had significantly worse quality of life, including the total score, psychological scale, and psychological subscales ‘appearance’ and ‘personal relationships’. On the contrary, in the biochemically uncontrolled group, there was no difference between whether current treatment with lanreotide or not in the ACROQOL scores of total score, both scales, and subscales.

Although the abundance of evidences has demonstrated that administration of somatostatin analogues ameliorates the abnormal biochemical profiles and the associated somatic impairments (15–19), whether these improvements translate into improved physical function remains largely unknown, and their impact on HR-QOL is still controversial (20). Before the development of disease-generated ACROQOL, only two studies assessed the impact of slow-release somatostatin analogues on quality of life in patients with acromegaly.

An open treatment study by Sonino et al. investigated the effects on quality of life of slow-release somatostatin analogue (lanreotide) in ten patients with active acromegaly. Hormone measurements and psychometric evaluation by means of self-rating scales were carried out at the baseline and after 1 and 2 months. Together with a significant decrease in GH and IGF-I, treatment with lanreotide significantly improved symptoms relating to psychological distress, well-being and social fears by employing the Symptom questionnaire, the Cognitive Scale of the screening List for Psychosocial Problems by Kellner and the Social Situation Questionnaire by Marks (21). Another small prospective study assessing various outcome measures including quality of life in nine patients with newly diagnosed acromegaly treated with...
Table 2 Distribution of the acromegaly quality of life questionnaire scores of total score, scales and subscales in biochemically controlled and uncontrolled groups.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Controlled (n=30)</th>
<th>Uncontrolled (n=22)</th>
<th>P</th>
<th>(P^*)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Scale 1</td>
<td>59.3±30.6</td>
<td>48.7±18.8</td>
<td>0.232</td>
<td>0.258</td>
</tr>
<tr>
<td>Scale 2</td>
<td>59.5±27.4</td>
<td>56.6±26.5</td>
<td>0.699</td>
<td>0.752</td>
</tr>
<tr>
<td>Subscale 1</td>
<td>48.5±27.8</td>
<td>45.6±29.9</td>
<td>0.727</td>
<td>0.740</td>
</tr>
<tr>
<td>Subscale 2</td>
<td>70.6±30.6</td>
<td>67.5±26.0</td>
<td>0.706</td>
<td>0.794</td>
</tr>
<tr>
<td>Total score</td>
<td>59.4±26.9</td>
<td>53.7±27.0</td>
<td>0.453</td>
<td>0.497</td>
</tr>
</tbody>
</table>

Comparisons between groups using univariate and multivariate regression analyses. \(P^*\) calculated by multivariate regression analysis after adjustment for age and gender. Scale 1, physical scale; scale 2, psychological scale; scale 2–1, psychological subscale ‘appearance’; scale 2–2, psychological subscale ‘personal relationships’.

octreotide-LAR for 6 months demonstrated significantly improvements in health perception and fatigue (22).

On the contrary, our study showed that treatment with slow-release somatostatin analogues (lanreotide) impairs quality of life, especially the psychological aspect. Furthermore, this association only occurred in the biochemical remission patients, not in the biochemically uncontrolled group. In contrast with the previous two studies, our study recruited a larger cohort to evaluate the effect in patients in remission status or not, and used a different measure of quality of life questionnaire, ACROQOL. ACROQOL is a disease-generated quality of life questionnaire and represents a reliable, construct valid and disease-specific tool for assessing HR-QOL in patients with acromegaly (23, 24). Besides the opposite results compared with the previous studies, the most interesting findings are that lanreotide treatment selectively impairs quality of life in patients with treated acromegaly in remission status, independent of age and gender. This may imply that the remission status achieved by lanreotide treatment would significantly worsen quality of life, especially the psychological aspect. This negative effect on quality of life in acromegalic patients may be expressed in patients in remission status. The acromegalic patients in biochemical remission may function relatively normal in daily life physically and their overall mortality rate may be similar to that of general population (4). However, we speculate that the requirement of having slow-release somatostatin analogue (lanreotide) injection every 2 weeks in hospital may give the patients more psychological distress, social fears and less sense of well-being. This kind of impairment may be exaggeratedly expressed in the patients in remission status.

There are two studies assessing quality of life in patients with controlled versus uncontrolled acromegaly using ACROQOL. Furthermore, both studies used the same remission criteria as ours. A prospective and cross-sectional study by Trepp et al. (23) shows the results on 33 patients with treated acromegaly completing the German version of the ACROQOL. Patients with biochemically uncontrolled acromegaly had significantly lower HR-QOL than patients with acromegaly in remission status.

Another cross-sectional study by Rowles et al. (13) on 80 patients with acromegaly demonstrated that there is no correlation between biochemical control and any measure of quality of life, including ACROQOL, Psychological General Well-Being Schedule, EuroQoL, and signs and symptoms score. Our study results also support the latter findings.

That acromegaly biochemically controlled by slow-release somatostatin analogues (lanreotide) leads to worse quality of life, and no difference between controlled and uncontrolled groups in the ACROQOL scores, in our study seems counterintuitive and runs contrary to the plethora of published data and vast personal experience of multiple researchers in this field: there is no doubt that controlled patients as a group individually feel better. Two potential explanations are possible. First, the magnitude of biochemical activity in

Table 3 Clinical characteristics of the four groups.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Controlled (n=30)</th>
<th>Uncontrolled (n=22)</th>
<th>P</th>
<th>(P^*)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>50.8±10.2</td>
<td>52.9±7.5</td>
<td>0.507</td>
<td></td>
</tr>
<tr>
<td>Male (%)</td>
<td>38.5</td>
<td>41.2</td>
<td>0.880</td>
<td></td>
</tr>
<tr>
<td>Age of diagnosis (years)</td>
<td>37.5±8.9</td>
<td>40.4±9.2</td>
<td>0.383</td>
<td></td>
</tr>
<tr>
<td>Disease duration (years)</td>
<td>13.3±5.7</td>
<td>12.5±7.2</td>
<td>0.751</td>
<td></td>
</tr>
<tr>
<td>Operated (%)</td>
<td>92.3</td>
<td>94.1</td>
<td>0.844</td>
<td></td>
</tr>
<tr>
<td>Re-operated (%)</td>
<td>7.7</td>
<td>5.9</td>
<td>0.844</td>
<td></td>
</tr>
<tr>
<td>Radiotherapy (%)</td>
<td>38.5</td>
<td>29.4</td>
<td>0.802</td>
<td></td>
</tr>
<tr>
<td>Macroadenoma (%)</td>
<td>69.2</td>
<td>58.8</td>
<td>0.558</td>
<td></td>
</tr>
<tr>
<td>Hormone replacement (%)</td>
<td>46.2</td>
<td>35.3</td>
<td>0.547</td>
<td></td>
</tr>
<tr>
<td>GH ((\mu)g/l), 1 (\mu)g/l=46.5 pmol/l</td>
<td>1.2±0.5</td>
<td>0.7±0.6</td>
<td>0.021*</td>
<td></td>
</tr>
<tr>
<td>IGF-I ((\mu)g/l), 1 (\mu)g/l=0.13 nmol/l</td>
<td>204.7±70.6</td>
<td>170.7±84.3</td>
<td>0.251</td>
<td></td>
</tr>
<tr>
<td>Morbidity (%)</td>
<td>38.5</td>
<td>47.1</td>
<td>0.638</td>
<td></td>
</tr>
</tbody>
</table>

GH, growth hormone; IGF-I, insulin-like growth factor-I. *\(P<0.05\) by Student’s \(t\)-test for quantitative data and \(\chi^2\)-test for qualitative variables.
uncontrolled patients was so mild (see Table 1), for example, a cross-sectional study by Damjanovic et al. (25) on 41 patients after transphenoidal pituitary surgery for somatotroph pituitary adenoma showed that those achieving complete biochemical cure and those with discordant GH and IGF-I levels had similar beneficial changes in cardiac performance, insulin resistance and body composition. It appears that even incomplete disease control may not necessarily result in a clinically relevant unfavourable outcome if the disease activity is mild. Second, the ACROQoL may not be a good or a sensitive enough instrument, or the Chinese version was not equivalent.

Until now there has been a paucity of data attempting to correlate biochemical control and slow-release somatostatin analogues with quality of life in acromegaly. Furthermore, our study and previously published studies showed inconsistent results and the impact remained controversial. As with previous studies, the major limitation is the limited sample size. Thus, further investigations on this issue are mandatory. The possibly unknown psychological side effects caused by slow-investigations on this issue are mandatory. The possibly major limitation is the limited sample size. Thus, further

galy. Furthermore, our study and previously published

somatostatin analogues with quality of life in acromegaly was not equivalent.

Table 4 Distribution of the acromegaly quality of life questionnaire scores of total score, scales and subscales in four groups.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Lanreotide (+) (n=13)</th>
<th>Lanreotide (−) (n=17)</th>
<th>P</th>
<th>P²</th>
</tr>
</thead>
<tbody>
<tr>
<td>Scale 1</td>
<td>49.5±34.4</td>
<td>66.7±25.9</td>
<td>0.129</td>
<td>0.118</td>
</tr>
<tr>
<td>Scale 2</td>
<td>45.5±29.7</td>
<td>70.3±20.3</td>
<td>0.011*</td>
<td>0.011*</td>
</tr>
<tr>
<td>Scale 2–1</td>
<td>36.3±29.5</td>
<td>57.8±23.2</td>
<td>0.033*</td>
<td>0.032*</td>
</tr>
<tr>
<td>Scale 2–2</td>
<td>54.7±34.1</td>
<td>82.8±21.4</td>
<td>0.010*</td>
<td>0.010*</td>
</tr>
<tr>
<td>Total score</td>
<td>46.9±29.8</td>
<td>69.0±20.5</td>
<td>0.023*</td>
<td>0.021*</td>
</tr>
</tbody>
</table>

Comparisons between groups using univariate and multivariate regression analyses. P² calculated by a multivariate regression analysis after adjustment for age and gender. *P value < 0.05. Scale 1, physical scale; scale 2, psychological scale; scale 2–1, psychological subscale ‘apparance’; scale 2–2, psychological subscale ‘personal relationships’.

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