CLINICAL STUDY

Soluble α -Klotho: a novel serum biomarker for the activity of GH-producing pituitary adenomas

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Abstract

Objective: Klotho is a lifespan-influencing gene expressed mainly in the kidneys. Soluble α -Klotho (α KL) is released into the circulation. In this study, we present baseline α KL serum levels of patients with acromegaly compared with controls with other pituitary adenomas and assess changes following transsphenoidal surgery.

Design: Prospective controlled study.

Methods: We measured soluble α KL (sandwich ELISA) and IGF1 (RIA) in sera of 14 patients (eight females and six males) with active acromegaly and in 22 control patients (13 females and nine males) operated for non-GH-producing pituitary adenomas. Immunohistochemical staining for Klotho was performed in resected adenomas and in normal pituitary tissue samples.

Results: Soluble α KL was high in the acromegaly group preoperatively (median 4217 pg/ml, interquartile range (IQR) 1812–6623 pg/ml) and declined after surgery during early follow-up (2–6 days; median 645 pg/ml, IQR 550–1303 pg/ml) ($P < 0.001$) and during late follow-up (2–3 months post-operatively; median 902 pg/ml, IQR 497–1340 pg/ml; $P < 0.001$). In controls, preoperative soluble α KL was significantly lower than in acromegalics, 532 pg/ml (400–677 pg/ml; $P < 0.001$). Following surgery, soluble α KL remained low during early and late follow-up – changes over time within the control group were not statistically significant. These results were independent of age, sex and kidney function. Klotho staining was equal or slightly decreased in GH-positive adenomas compared with controls.

Conclusion: High soluble α KL serum levels were specific to GH-producing adenomas and decreased rapidly following adenoma removal. Thus, soluble α KL appears to be a new specific and sensitive biomarker reflecting disease activity in acromegaly. Similar Klotho staining patterns in controls and acromegalics suggest that the rise in serum α KL is caused by systemic actions of pituitary GH rather than due to increased expression of Klotho by the pituitary (adenoma).

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Introduction

The α -klotho gene (α KL) was serendipitously discovered as a lifespan-influencing gene in mice after recognition that its disruption caused accelerated ageing (1). Lifespan extension was achieved by overexpression (2). In humans, variants of α KL were associated with ageing (3) and the phenotype of a rare homozygous missense mutation was described (4). The gene and protein were named after the Greek goddess Klotho who spins the thread of life. α KL encodes α KL protein, a 130 kDa type I membrane protein (1014 amino acids long) considered the founder of the Klotho family that is predominantly expressed in the kidneys, choroid plexus and several endocrine organs including the pituitary, parathyroid,

testis, ovary, placenta and pancreas (1). The α KL protein exists in two forms with distinct functions. Membrane-bound α KL is a co-receptor for fibroblast growth factor 23, a bone-derived phosphaturic hormone that inhibits renal phosphate reabsorption and calcitriol production (5, 6). Soluble α KL attenuates insulin/insulin-like growth factor 1 (IGF1) signalling and regulates calcium homeostasis (7, 8). The extracellular domain of the membrane-bound form can be enzymatically cleaved (ectodomain shedding) and released as soluble α KL into blood, urine and cerebrospinal fluid (9, 10). Recently, a sandwich ELISA for the measurement of soluble α KL was established and is now commercially available (11). Soluble α KL was inversely related to age in healthy subjects (11) and

with mortality in the elderly (12). High levels of soluble α KL were found in human umbilical cord blood (13), and based on a positive correlation between plasma levels of soluble α KL and growth and metabolic parameters in premature and term neonates, it has been speculated that Klotho may play a role in the stimulation of growth (14).

GH excess due to benign adenomas of the pituitary gland is the major cause for acromegaly with an incidence of approximately four cases per 1 million persons per year (15). Acromegaly is usually diagnosed with a considerable delay and therefore possibly associated with increased mortality even after curative transsphenoidal surgery (16). Clinical features develop slowly over many years and comprise metabolic derangements and pathognomonic changes in the patient's appearance, mainly soft tissue swelling and skeletal bone growth, resulting in typical acral enlargement and coarse facial features. Metabolic changes include increased plasma glucose in the wake of insulin resistance despite reduced visceral fat and high levels of serum phosphate accompanied by higher than normal renal glomerular filtration (17, 18, 19). Both insulin resistance and elevated serum phosphate are associated with increased mortality in the general population (20, 21). In the setting of reduced visceral fat and increased glomerular filtration, insulin resistance and serum phosphate elevation are unusual, and the underlying mechanisms are unclear. Currently, GH (the hormone directly produced by the adenoma) and IGF1 (a GH-dependent, predominantly liver-derived hormone) are the classical biochemical markers of disease activity in acromegaly.

We recently found that serum-soluble α KL is markedly elevated in patients with acromegaly and that this α KL excess reversed following adenoma removal (22). In order to assess whether high serum levels of soluble α KL are specific to patients with GH-producing pituitary adenomas, we now present a prospective controlled study that documents the baseline serum levels of α KL in patients with acromegaly compared with a control group of patients with other pituitary adenomas. Additionally, we monitor the temporal changes in α KL levels following transsphenoidal surgery, and we report detailed immunohistochemical analysis of the adenomas removed.

Materials and methods

Patient characteristics

We included 14 consecutive patients with active acromegaly (eight females and six males) with a mean age of 48 years (range 29–84 years) who underwent transsphenoidal surgery at the University Hospital Zurich. The preoperative diagnosis of acromegaly was based on pathognomonic clinical findings and biochemical markers (excess IGF1 and GH, nonsuppressible

during a 75 g oral glucose tolerance test (oGTT)). Patients were excluded when histopathological examination failed to identify a GH-producing adenoma. As a control group, we included 22 patients (13 females and nine males) with a mean age of 48 years (range 14–81 years) operated for pituitary adenomas not producing GH (13 non-functioning adenomas (NFAs) and nine prolactinomas). All patients provided written informed consent and the study was approved by the Local Ethics Committee. Detailed patient characteristics are summarized in Tables 1 and 2. Tumour volume calculation was based on preoperative magnetic resonance imaging (MRI) and the diameter method (tumour volume = $\frac{4}{3} \times \pi \times \frac{1}{2}x \times \frac{1}{2}y \times \frac{1}{2}z$), where x , y and z are the maximum diameters within the three axes. Surgical strategy was transnasal transsphenoidal using microsurgical technique and intraoperative MRI (PoleStar N20, 0.15T, Medtronic Navigation, Minneapolis, MN, USA). All procedures were performed by the senior author (R-L Bernays) – details on the surgical strategy have been described previously (23).

Histopathology

All pathology materials, consisting of H&E and reticulin-stained sections, a full panel of anterior pituitary hormone immunohistochemistry, and the MIB-1 (Ki-67) proliferation marker were reviewed to confirm the diagnosis. Pituitary adenomas were then categorized as either GH-positive adenomas, adenomas without hormone expression (NFAs) or prolactinomas.

GH antibodies were obtained from Thermo scientific (Waltham, MA, USA; MA5-11926, 1:3000) and Klotho antibodies were purchased from Abcam (Cambridge, UK; ab68208, 1:40). Immunophenotypic analysis was performed using a Leica Bond-Max automated immunostainer employing 2 μ m-thick, formalin-fixed, paraffin-embedded sections. GH signal detection was performed using 3,3'-diaminobenzidine (brown) and 3-amino-9-ethylcarbazole (red) for Klotho detection respectively. Sections were counterstained with haematoxylin. For co-localization analysis, the sections were treated with citric acid for 15 min at 95 °C and immunofluorescent co-stainings were performed according to the Bond staining protocol. GH-positive areas were detected using the secondary antibody Alexa 488 anti-mouse (1:1000) and Alexa 594 anti-rabbit antibody (1:1000) for Klotho signals respectively.

Assays

All blood samples were drawn around the same time in the morning after overnight fasting. Soluble α KL was determined using a sandwich ELISA described by Yamazaki *et al.* (11) (Kyowa Hakko Kirin Co. Ltd., Tokyo, Japan) according to the manufacturer's instructions. IGF1 was measured by RIA after the removal of carrier proteins as described elsewhere (22) and

Table 1 (A) Patient characteristics of the acromegaly group ($n=14$); immunohistological reactivity to other hormones than GH is shown in parentheses. (B) Patient characteristics of the control group ($n=22$); minor immunohistological hormonal reactivity is shown in parentheses. Preoperative GH, α -Klotho and IGF1 serum levels in controls with minor GH positivity were within the normal range (see [Supplementary Table 1](#), see section on supplementary data given at the end of this article) and the latter two markers did not show the post-operative decrease characteristic for the acromegaly group.

Case	Age	Sex	Staining	Tumour volume (mm ³)	BMI (kg/m ²)	Preoperative GFR (ml/min per 1.73 m ²)
(A) Patient characteristics of the acromegaly group						
1	43	M	GH++	880	29.2	106
2	30	F	GH++ (PRL+)	2492	25.7	120
3	54	M	GH++	576	27.8	97
4	84	M	GH++ (PRL+)	4775	28.6	59
5	40	F	GH++ (LH+++ , HCG+)	110	38.6	99
6	41	F	GH++	3541	24.6	118
7	43	F	GH++ (PRL+)	7948	23	113
8	42	F	GH++	1026	33.7	118
9	53	F	GH++ (PRL+)	286	32	104
10	64	F	GH++	3519	20.7	104
11	52	M	GH++	1334	38.7	101
12	42	F	GH++	4524	24.5	97
13	29	M	GH++	1634	33.7	127
14	45	M	GH++	50	37.6	85
(B) Patient characteristics of the control group						
1	75	F	NFA	5864	25	83
2	29	F	PRL++ (GH+)	280	20.4	129
3	75	F	NFA (FSH+)	5483	22.9	89
4	81	M	NFA	1021	31.7	82
5	32	F	PRL++	339	20.2	100
6	49	M	NFA	11 310	29.3	88
7	39	M	NFA (FSH++ , TSH+)	2714	23.2	96
8	47	M	NFA	1407	22.2	100
9	30	M	NFA	16 085	27.5	123
10	65	M	PRL++	8181	29.4	75
11	14	F	PRL++	2246	17.9	133
12	61	F	NFA (GH+ , TSH+)	4516	27.3	94
13	63	F	PRL++ (GH+)	16 118	28.7	79
14	36	F	PRL++ (ACTH+ , GH+)	403	26.6	76
15	68	M	NFA	2368	27.5	72
16	69	M	NFA	2356	34	48
17	48	F	NFA	628	20.2	109
18	17	M	PRL++	377	21.9	129
19	18	F	PRL++ (GH+)	622	15.8	128
20	57	F	NFA (LH+)	15 683	19.7	101
21	32	F	PRL++	452	18.1	119
22	54	F	NFA (FSH+)	674	27.3	100

GH by IRMA (hGH-RIATC; CIS Bio International, Oris Industries, Gif-Sur-Yvette, France). Creatinine was measured using the kinetic Jaffé method on a Roche COBAS 8000 analyzer (Roche Diagnostics) and glomerular filtration rate (GFR) was estimated using the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) formula. These values were recorded as soluble α KL is associated with renal function (11). Both soluble α KL and IGF1 were measured preoperatively, at least once shortly after surgery (2–6 days post-operatively, before discharge) and again at the first outpatient follow-up (2–3 months after surgery). In all control patients and in a subgroup of seven acromegaly patients, multiple short-term post-operative measurements were performed. GH was measured preoperatively in all patients.

Statistical analyses

Statistical analyses were performed using commercially available software IBM SPSS Statistics 20 (SPSS, Inc.) and Matlab (www.mathworks.com). Continuous variables are presented as median with interquartile range (IQR). Preoperative variables were compared between groups using the Mann–Whitney U test or the χ^2 test as appropriate. The Wilcoxon signed-rank test was used for comparing pre- and post-operative values within one group. To corroborate the results of bivariate testing, we constructed a general linear model (GLM) for repeated measures. The dependent variable was soluble α KL after log-transformation. The within-subjects factor was time with the levels preoperative, early post-operative and late post-operative. Between-subjects

Table 2 Group characteristics are presented as median and interquartile range (IQR).

	Acromegaly (<i>n</i> =14)	Control (<i>n</i> =22)	<i>P</i> value
Sex (M/F)	6/8	9/13	0.91 ^a
Age (years)	43 (41–53)	49 (31–66)	0.79 ^b
BMI (kg/m ²)	28.9 (24.6–34.7)	24.1 (20.2–27.8)	0.010 ^b
Preoperative GFR (ml/min per 1.73 m ²)	104 (97–118)	98 (81.25–120)	0.35 ^b
Preoperative GH (ng/ml)	10.3 (7.01–43.10)	0.29 (0.13–0.74)	<0.001 ^b
Tumour volume (mm ³)	1483 (503–3786)	2301 (579–6443)	0.45 ^b
Hospital stay (days)	6 (5–7)	6 (5–7)	0.60 ^b

^a χ^2 Test.^bMann–Whitney *U* test.

factors were group and sex, and covariates were age and preoperative GFR. Two-tailed *P* values <0.05 were considered statistically significant.

Results

Patient characteristics

Both study groups, acromegalics (eight females and six males; mean age 48 years) and controls (13 females and nine males; mean age 49 years), had successful transsphenoidal removal (at least debulking) of their adenomas – detailed histopathological results are shown in Fig. 1. Within the group of patients with active acromegaly, median estimated tumour volume was 1484 mm³ (IQR 503–3786 mm³). Of all 14 acromegalic patients, two (14%) patients presented with microadenomas, whereas 12 (86%) presented with macroadenomas. In the control group (*n*=22; 13 females and nine males; 13 NFAs and nine prolactinomas; mean age of 49 years), median preoperative tumour volume was 2301 mm³ (580–6444 mm³). Microadenomas were seen in two cases (9%). When preoperative tumour volume was compared between both groups, no statistical difference was identified (*P*=0.45, Mann–Whitney *U* test).

Soluble serum α KL

Soluble α KL was high in the acromegaly group before surgery (Figs 1 and 2) with a soluble α KL median of 4217 pg/ml (1813–6624 pg/ml), then levels declined after removal of the GH-producing adenoma to a median of 646 pg/ml (550–1303 pg/ml) (*P*<0.001, Wilcoxon signed-rank test) during early follow-up (2–6 days post-operatively), then to a median of 902 pg/ml (498–1341 pg/ml; *P*<0.001) during late follow-up (2–3 months post-operatively) – soluble α KL kinetics are shown in Fig. 1. Compared with acromegalics, the preoperative median of soluble α KL in controls was significantly lower, 532 pg/ml (400–678 pg/ml; *P*<0.001). Following surgery, soluble α KL stayed low with 404 pg/ml (320–635 pg/ml) during early follow-up and 524 pg/ml (359–621 pg/ml) during

late follow-up – changes over time within the control group were not statistically significant. The relative drop of soluble α KL (early follow-up/preoperative) was more pronounced in the acromegaly group, 0.25 (range 0.1–0.5) compared with controls, 0.95 (range 0.5–1.6) (*P*<0.001). Short-term kinetics of α KL levels of individual acromegalic patients compared with controls is plotted in Fig. 2A. To corroborate the results of bivariate testing, we constructed a GLM for repeated measures after log-transformation of our data. The only significant interaction was found between TIME and GROUP (*F*=33, hypothesis degrees of freedom (df) 2, error df 17, *P*<0.001) – the temporal changes of the two groups on a log-scale are not parallel. Group differences and significant post-operative decrease in the acromegaly group concerning soluble α KL levels were independent of age, sex and kidney function (GFR).

IGF1

As expected, preoperative median IGF1 levels were higher in the acromegaly group, 483 ng/ml (367–640 ng/ml) compared with the control group, 86 ng/ml (53–136 ng/ml) (*P*<0.001). Within the acromegaly group, median preoperative IGF1 levels of 483 ng/ml returned to median early post-operative (2–6 days post-operatively) IGF1 levels of 182 ng/ml (144–229 ng/ml; *P*<0.001), whereas no significant difference was found between preoperative and post-operative IGF1 levels within the control group. The long-term IGF1 time course within the acromegaly group (Fig. 1A) and controls (Fig. 2A) is illustrated in Fig. 1. The short-term time course of IGF1 levels of individual acromegalic patients compared with controls are plotted in Fig. 2B.

GH

Preoperative GH levels were significantly higher in the acromegaly group as expected (*P*<0.001). In all acromegalic patients, surgery resulted in a significant decrease in GH with median preoperative levels of 10 ng/ml (7–43 ng/ml) and median post-operative values of 1.9 ng/ml (0.6–2.5 ng/ml) (*P*<0.001) and

Figure 1 Time course of serum-soluble α -Klotho in acromegalics (A, red boxes ($n=12$)) and controls (B, blue boxes ($n=12$)) and IGF1 (white boxes) in both groups. Preoperative, early (2–6 days post-operatively, at discharge from hospital) and late (2–3 months post-operatively) follow-up values are plotted. Note the different scales in (A) and (B).

disease features improved as judged by all patients and treating physicians. Median preoperative GH levels in the control group were 0.3 ng/ml (0.1–0.7 ng/ml) – post-operative measurements of GH were only performed in the acromegaly group. The covariation of soluble α KL, IGF1 and GH is shown in [Supplementary Figure 1](#), see section on [supplementary data](#) given at the end of this article.

BMI and kidney function

BMI was significantly higher in the acromegaly group ($P=0.010$) with a median BMI of 28.9 kg/m² (24.6–34.7 kg/m²) compared with a median BMI of 24.1 kg/m² (20.2–27.8 kg/m²) in the control group. In terms of preoperative kidney function, there was no significant difference regarding preoperative GFR ($P=0.35$) with a median preoperative GFR of 104 ml/min per 1.73 m² (97–118 ml/min per 1.73 m²) and 98 ml/min per 1.73 m² (81.25–120 ml/min per 1.73 m²) in acromegalics and in controls respectively. However, distinct changes were observed in response to surgery: post-operative GFR decreased in patients undergoing transsphenoidal surgery for GH-producing adenomas, median Δ GFR (post-operative GFR–preoperative GFR) was -4.50 ml/min per 1.73 m² (-9.00 to -0.75 ml/min per 1.73 m²), whereas GFR increased in controls with a median Δ GFR of

6.50 ml/min per 1.73 m² (3.50–11.50 ml/min per 1.73 m²) ($P<0.001$). The difference between preoperative and post-operative GFR was statistically significant in both acromegalics ($P=0.006$) and controls ($P=0.001$).

Immunohistochemistry

GH-producing adenomas (Fig. 3A, B and C) showed variable GH expression within the tumour (brown) from samples with strong, diffuse immunoreactivity to those with weaker and/or focal paranuclear staining. These staining patterns corresponded to either densely or sparsely granulated subtypes. In cases with focal or weak GH staining, effacement of the normal lobular architecture confirmed the presence of tumour. The Klotho expression pattern (red) in corresponding areas was more diffuse and independent of GH-positive cells. The cells in the GH-negative control group, consisting of hormone-inactive pituitary adenomas (Fig. 3D and E), and a prolactinoma (Fig. 3F) showed diffuse and strong positivity for Klotho. There is no stringent

Figure 2 Individual short-term time course of soluble α -Klotho (A) and IGF1 (B) in acromegaly patients ($n=7$) (coloured lines) compared with box-plots of all control patients.

Figure 3 Klotho expression in GH-producing adenomas vs the control group. Three examples of GH-producing adenomas (A, case 11; B, case 7 and C, case 8) with GH expression (brown) and diffuse Klotho expression (red) are depicted in corresponding areas. In the control group, hormone-inactive pituitary adenomas (D, case 1 and E, case 4) and a prolactinoma (F, case 10) lack GH positivity. Klotho expression seems to be equal or slightly increased in the control compared with GH-positive adenomas. Original magnification 400 $\times$.

co-localization of these two markers. Immunofluorescence analysis (Fig. 4) emphasizes that Klotho (Alexa 594, red) is independently expressed from GH-positive cells (Alexa 488, green). Original magnification 400 $\times$. Klotho and GH staining in normal pituitary is shown in Fig. 5.

Discussion

This is the first prospective controlled study that documents serum levels of α KL in patients with active acromegaly compared with a control group of patients with other pituitary adenomas. Moreover, we monitored α KL levels over time (short- and long-term follow-up) following transsphenoidal surgery. Our results show a highly significant difference in preoperative α KL levels between acromegalics and controls, indicating that α KL excess is specific for GH-positive adenomas. Reversal of α KL occurs rapidly after surgery and there are no significant differences between short- and long-term follow-up, suggesting that α KL is a very sensitive marker for disease activity of acromegaly. Both the preoperative group difference and the rapid post-operative decrease in α KL levels in acromegalics remained significant after adjusting our results for age, sex and kidney function. The mechanisms leading to soluble α KL excess in the serum in active acromegaly remain unclear. Soluble α KL could arise either from a distinct transcript (24) or from

ectodomain shedding of membrane Klotho (9, 10, 25). It remains unclear whether in acromegalics membrane-associated Klotho (mainly found in the kidneys) and soluble α KL (as detected in the serum) rise concurrently or whether elevated soluble α KL is unrelated or possibly inversely related to the abundance of plasma membrane Klotho in the kidneys – possibly resulting from enhanced enzymatic activity. Admittedly, we cannot provide our own experimental data to support this favoured hypothesis. α KL (130 kDa) appears to result from proteolytic ectodomain clipping (10). Two members of the 'A Disintegrin and Metalloproteinase' (ADAM) family, ADAM10 and ADAM17, have been suggested as the responsible enzymes (9), and the activity of secretases (25, 26) shedding the ectodomain from the integral membrane Klotho may be increased in acromegaly, either directly by GH or indirectly by factors or a proteolytic activity induced by GH.

For obvious reasons, renal biopsies to check for changes in membrane-bound Klotho abundance were not feasible in our study population. However, we analysed Klotho staining in GH-producing adenomas, in controls (NFAs and prolactinomas) and in normal pituitary tissue samples. The immunohistochemical staining pattern presented suggests that the rise in serum α KL is not explained by increased pituitary (adenoma) Klotho expression but rather due to an increase in pituitary GH secretion.

Figure 4 Immunofluorescence of a GH-positive adenoma showing GH-positive cells (Alexa 488, green) and diffuse positivity for Klotho (Alexa 594, red). Nuclear counterstain with Dapi (blue). Original magnification 100 $\times$.

Until now, GH (a hormone produced by the adenoma itself) and especially IGF1 (a GH-stimulated, predominantly liver-derived peptide) have been the 'classical' biomarkers for diagnosing and monitoring disease activity during the treatment of patients with acromegaly. In fact, their normalization has been linked to decreased mortality (27, 28, 29). However, it has been recognized that both parameters entail various shortcomings.

Serum levels of IGF1 are influenced not only by GH status but also by age, moreover, by gender (estrogens), race, genetic makeup, liver function, nutritional status, portal insulin, thyroid hormones and by concomitant inflammatory disease. Some of these influences,

particularly the former, may also have an impact on α KL (to an as yet unknown extent). Serum IGF1 is mainly derived from the liver and tightly bound to IGF binding proteins (IGFBPs) (30, 31). Changes in IGFBP concentrations contribute to the limitations known for a variety of IGF1 assays (32). To circumvent these problems, we used a classical and time-consuming assay in which carrier proteins are removed before the samples are incubated with the antibodies (33, 34).

Dynamic testing using oGTT to suppress GH is widely used; however, patients with acromegaly can demonstrate normal oGTT GH suppression despite elevated IGF1 levels (27, 35). In patients receiving non-surgical treatment of acromegaly, such as long-acting somatostatin analogues (LA-SRIFs) (36), pegvisomant (PEG-V) (37) or radiotherapy for GH-producing adenomas, GH values may be misleading due to highly irregular GH secretion pattern and flattened GH pulses (38).

The limitations (both biological and technical) of the assays used to measure GH (39) and IGF1 are well known (40), and it was stated that additional, possibly more specific and sensitive, biomarkers are desperately needed (41).

Our study has some notable limitations. The assay for α KL has been introduced only recently; therefore, knowledge of its shortcomings is limited. Additionally, the normal range for serum levels of soluble α KL has not been established. In our laboratory, we measured soluble α KL in the sera of 26 healthy volunteers (11 females and 15 males; mean age 39 years) as previously reported (22): α KL (median and IQR) was 596 (506–734) pg/ml. Similar to IGF1, serum-soluble α KL also decreases with increasing age; moreover, it may be low in patients with renal failure (11), which prompted us to routinely check creatinine. In the

Figure 5 Expression pattern in normal pituitary. Normal pituitary shows GH positivity (brown) in the GH-producing areas. Klotho positivity (red) is observed in some lobules of normal pituitary gland, whereas other parts are negative in IHC. Original magnification A, 100 $\times$ (upper panel) and B, 400 $\times$ (lower panel).

context of NFA, some of our patients may have been GH deficient, but the design of our study did not allow us to determine whether these patients had lower than normal IGF1 and α KL serum levels. Furthermore, the molecular mechanisms of α KL excess in active acromegaly and the functional impact of α KL in acromegaly disease biology remains unknown.

Conclusions

Acromegaly is (thus far, to our knowledge, up to this writing) the only acquired disease known to man with excessively elevated levels of soluble α KL. Highly elevated soluble α KL is specific to GH-producing adenomas of the pituitary gland and rapidly decreases following adenoma removal. Thus, soluble α KL appears to represent a new, quite specific and fairly sensitive biomarker reflecting disease activity in patients with acromegaly.

Supplementary data

This is linked to the online version of the paper at <http://dx.doi.org/10.1530/EJE-12-1045>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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