Reduction in Cutis Verticis Gyrata after Transsphenoidal Pituitary Resection of Growth Hormone–Secreting Adenoma

Daniel Pinggera1 Delia Anna Marsoner1 Marcel Seiz-Rosenhagen2 Claudius Thomé1 Christian Franz Freyschlag1

1 Department of Neurosurgery, Medical University Innsbruck, Innsbruck, Austria
2 Department of Neurosurgery, University Medical Center Mannheim, University of Heidelberg, Mannheim, Germany

Abstract

Aim Cutis verticis gyrata (CVG) describes a hypertrophy and folding of the skin that appears as gyration of the scalp. Secondary CVG is a common symptom of growth hormone (GH)–producing adenoma of the pituitary and an early sign of acromegaly. We hypothesized that surgical removal of a pituitary adenoma with a consecutive decrease in hormone levels will cause a reduction in gyration.

Methods Imaging and laboratory examination of 10 patients (median age: 44 years) with GH-producing adenoma have been evaluated retrospectively. Surgical treatment consisted of endonasal transsphenoidal resection of the adenoma, followed by endocrinologic and radiographic follow-up. CVG was calculated as area under the curve of the scalp on preoperative coronal T1-weighted MRI, compared with postoperative follow-up imaging after 12 months (AUC, measured in cm²). In laboratory examination, the levels of insulin-like growth factor-1 (IGF-1) were analyzed accordingly as its levels are constant throughout the day.

Results After 12 months, we found a statistically significant \( p = 0.008 \) correlation between the decrease in IGF-1 levels (median: 219.00 ng/mL) and the reduction in CVG (median: 0.12 cm²).

Conclusion Resection of GH-producing pituitary adenoma leads to a decrease in CVG in patients suffering from acromegaly, which is strongly related to the levels of IGF-1.

Keywords
- acromegaly
- cutis verticis gyrata
- pituitary adenoma

Introduction

Pituitary adenomas account for 10 to 20% of all intracranial tumors and growth hormone (GH)–producing adenoma may occur in up to 10% of this population.1 GH-producing adenomas cause an aberrant secretion of GH and, consecutively, insulin-like growth factor-1 (IGF-1), leading to the distinctive features of acromegaly.2 The prevalence of acromegaly is 60/1,000,000 with an incidence of 3 to 4 per million per year.3

Acromegaly may present with a variety of different symptoms, ranging from subtle signs of acral overgrowth, jaw prognathism, fasting hyperglycemia, and hyperhidrosis to more severe diseases such as diabetes mellitus, hypertension, and cardiac failure. Besides medical history, physical findings, and imaging, diagnosis is made by measurement of high-serum IGF-1 and high-glucose-suppressed GH levels.4 Transsphenoidal surgery is the treatment of choice according to current guidelines, whereas control rates are strongly dependent on the neurosurgeons experience and the size of the adenoma.5,6 Therapeutic success is defined by normalization of IGF-1 and...
GH serum levels (<1 µg/L and a GH nadir of <0.4 µg/L in oral glucose tolerance test [OGTT]).

Skin changes have been recognized an important component of acromegaly. IGF-1 is stimulated by GH and produced primarily by the liver. It activates the IGF-1 receptor, a tyrosine kinase receptor, located in the cell membrane. There are no major differences in relation to sex or circadian rhythm. IGF-1 receptor signaling regulates skin development, growth and differentiation of fibroblasts, and keratinocytes. This fact leads to severe cutaneous manifestations such as cutis verticis gyrata (CVG), acanthosis nigricans, or hyperhidrosis.

In our study we wanted to focus on CVG, a disorder characterized by thickening of the scalp due to hypertrophy of the skin, leading to skin folds and grooves that resemble the gyri of the brain surface. It has an estimated prevalence of 1 in 100,000 in males and 0.026 in 100,000 in females. The term was first used in 1901 by Unna. It may be classified into a primary and a secondary form. Primary CVG has no obvious cause, usually appears after puberty, and is often associated with neurologic conditions, such as epilepsy or mental retardation. Secondary CVG is frequently associated with endocrine or neoplastic conditions such as acromegaly, myxedema, or amyloidosis. The number of folds varies from 2 to 12, and the direction is mostly anterior to posterior. They cannot be corrected by pressure or traction and the skin color is normally unaffected.

These skin changes are well known to impair the quality of life in patients with acromegaly. Therefore, we want to investigate whether transsphenoidal surgery of GH-producing adenoma leads to a decrease in CVG in patients suffering from acromegaly related to the levels of IGF-1.

Materials and Methods

Data of all patients with surgically treated GH-producing adenoma and the signs of acromegaly between 2006 and 2012 were retrospectively evaluated for presence of CVG. As institutional review board approval is not needed for retrospective cohorts in Austria, informed consent was not required. We identified 12 patients with CVG who underwent endonasal transsphenoidal surgery for GH-producing adenoma; 2 patients had to be excluded due to a lack of imaging data or serum levels of IGF-1, resulting in 10 eligible patients. Serum levels of IGF-1 were recorded using identical IGH arrays (standard value: 56–201 ng/mL). Magnetic resonance imaging (MRI) was performed preoperatively and 12 months after surgery.

Defined areas under the curve (AUC) of preoperative coronal T1-weighted MRI were compared with postoperative follow-up imaging after 12 months (AUC, measured in cm²). A standardized measurement protocol was applied to assess CVG (Fig. 1). The red and blue area were measured together to minimize errors in measurement. All measurement were done pre- and postoperatively with Agfa HealthCare IMPAX 6 (Agfa HealthCare GmbH, Bonn, Germany).

All statistical analyses were conducted using IBM SPSS Statistics 21 (IBM Corp., Armonk, New York, United States). Differences with an error probability of less than 0.05 were
considered statistically significant. A t-test was used to evaluate the reduction in the AUC related to the IGF-1 serum levels as well as to determine the difference between the AUC pre- and postoperatively.

**Results**

In 10 patients we found a 4:6 male/female ratio with a mean age at surgery of 44 years (range: 31–74 years). The mean total AUC (red and blue area together, Fig. 1) at surgery was 13.72 cm² (range: 12.82–14.66 cm²) and 13.64 cm² (range: 12.75–14.34 cm²) 12 months after surgery. In nine cases a decrease in the AUC was seen, with a slight increase in one patient (►Fig. 2). This patient was afterward assigned to further drug therapy. The mean reduction in the AUC was 0.12 cm² (range: −0.8–0.52 cm²).

The mean IGF-1 levels at time of surgery was 469 ng/mL (range: 190–1,155) and 205 ng/mL (range: 114–490) at the 12-month follow-up evaluation. In all 10 cases the serum levels of IGF-1 decreased (►Fig. 3). We were able to demonstrate that both the decreases in the total AUC correlate to the decreasing serum levels of IGF-1 according to the t-test ($p = 0.008$).

**Discussion**

To the best of our knowledge, this is the first study dealing with changes in CVG related to postoperative reduction in GH levels. It could be demonstrated that a postoperative reduction in IGF1 will lead to a reduction in CVG.

Although the diagnosis of CVG is primarily clinical, imaging plays a role in differentiating primary CVG from...
secondary CVG. The appearance of CVG on computed tomography (CT) and MRI is characteristic, and because of the anterior-posterior direction of the folds, the coronal plane is regarded best to visualize CVG. In four of our patients, imaging data were available only of a pituitary MRI protocol, including coronal plane only around sella, but CVG was clearly visible. Contrary to the available literature, we only reviewed cases of secondary CVG alone, leading to a small sample size.

Despite the small number of patients, our epidemiologic data are in agreement with the literature. The mean patient age was over 40 years with a female predominance. Similar to the study by Schöfl et al, we only evaluated the levels of IGF-1, as the GH assay used at our center changed during the observed period, and in light of the enormous interassay variability, data could be found misleading. Furthermore, recent data confirm discordance between GH and IGF-1 after surgery, so that we decided to focus on one marker. The increase in total CVG in one patient cannot be explained definitively, since levels of IGF-1 were decreasing, but the AUC of the skin was decreasing too. Overall, the correlation was better seen in total AUC. We hypothesize that the measurement is more accurate when done completely.

All patients showed postoperative normalized levels of IGF-1, and we believe that this decrease in IGF-1 stops and reverses the hypertrophy of the skin, as described for other cutaneous manifestations as acanthosis nigricans and oily skin. This may contribute a better health-related quality of life as skin manifestations are often complained by patients; nevertheless, the normal levels of IGF-1 alone improve health-related quality of life. Reduction in CVG is probably only a side effect of surgically treated acromegaly, but perhaps this could be a decision-making aid to undergo surgery, considering the “cosmetic” disturbances mentioned by patients. The limitations of our study include its retrospective nature and its limited follow-up of 1 year.

Conclusion

Transsphenoidal resection of GH-producing adenoma of the pituitary leads to a decrease in CVG in patients suffering from acromegaly related to the decreasing levels of IGF-1.

Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper.

Congress

Portions of this work were presented in poster and oral form at the EANS Annual Meeting 2013, Tel Aviv, Israel, and at the 49th Annual Meeting of the Austrian Neurosurgical Society 2013, Innsbruck, Austria.

References


