Discrepant GH and IGF-I values in the evaluation of treated acromegalic patients; an ongoing challenge. A meta-analysis.

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INTRODUCTION - OBJECTIVES

Growth Hormone (GH) and Insulin-like Growth Factor 1 (IGF-I) are currently the principal biomarkers used to assess disease activity in acromegaly and any discrepancy between them, renders interpretation of results inconclusive. The purpose of this study was to assess the frequency of discrepant results and identify parameters that might affect the emergence of this phenomenon.

METHODS

A systematic review of Medline and Scopus was performed (1987 – 2013) followed by a meta-analysis to address the frequency of discrepant results between GH and IGF-I levels. Meta-regression was performed using the year of publication as a continuous variable and subgroup analyses included the different types of GH testing and GH assays used, as well as the results of patients treated with Somatostatin Analogues (SSAs) compared to those treated with non-pharmaceutical modalities.

RESULTS

The analysis retrieved 39 eligible studies totaling 7071 patients. The pooled discordance rate between GH and IGF-I was 25.7% (95% CI: 22.3-29.4) and the predominant format was that of elevated IGF-I with normal GH levels [15.3% (95% CI: 12.5-18.7)].

No significant correlation between the discordance rate and the year of publication was shown  Coeff: +0.003 95% CI: -0.008 to +0.134 (p=0.589)

The use of ultrasensitive GH assays resulted in higher discordance rates (30.7%, 95% CI: 25.9-35.9) vs. (19.8%, 95% CI: 14.1-27.2, p=0.04). On the contrary, GH nadir values obtained during an OGTT (GHn) yielded the lowest discordance rates over a random GH value (GHr) and over the mean value (GHm) of a day curve (22.0%, vs. 29.1 vs. 33.5%, p=0.041).

CONCLUSIONS

Discrepancy between GH and IGF-I results is encountered in almost a quarter of treated patients with acromegaly, especially when using ultrasensitive GH assays or in patients receiving SSAs, a fact that the clinician should take into consideration when making clinical decisions.

REFERENCES


The Greek School of Endocrinology Initiative was organized and funded by Ipsen. Statistical support and publication costs associated with this work were also funded by Ipsen. The development at all stages of the concept and content was the sole responsibility of the authors.

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