Special Issue

Clinical Updates on Acromegaly

Message from the Guest Editors

This Special Issue aims to compile a collection of articles on the latest and most relevant research directions in the field of acromegaly. Acromegaly is an endocrine disease mainly caused by growth hormone-secreting pituitary adenoma. Persistent excessive secretion of growth hormone (GH) and its peripheral mediator, insulin-like growth factor-1 (IGF-1), results in numerous complications, including those related to the cardiovascular, respiratory, and musculoskeletal systems. Personalized diagnostic and therapeutic management is essential in providing care to each patient with acromegaly. We invite specialists in the field of neuroendocrinology to share their experiences and observations. Original papers and review articles are welcome.

Guest Editors

Prof. Dr. Marek Bolanowski

Dr. Jowita Halupczok-Żyła

Prof. Dr. Aleksandra Gilis-Januszewska

Prof. Dr. Przemysław Witek

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Journal of Clinical Medicine
Editorial Office
MDPI, Grosspeteranlage 5
4052 Basel, Switzerland
Tel: +41 61 683 77 34
jcm@mdpi.com

mdpi.com/journal/jcm

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There has been an explosion of gene and target based research and therapeutics in the multitude of fields that compose clinical medicine. The *Journal of Clinical Medicine*'s (*JCM*) staff and editorial board are dedicated to providing cutting edge, timely, and peer-reviewed articles covering the diverse subspecialties of clinical medicine. The journal publishes concise, innovative, and exciting research articles as well as clinically significant articles and reviews that are pertinent to the myriad of disciplines within medicine. The articles published are relevant to both primary care physicians and specialists. The journal's full-texts are archived in PubMed Central and indexed in PubMed. Please consider submitting your manuscripts for publication to our journal and check us out on-line!

Editors-in-Chief

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Internal Medicine Department, University Hospital Strasbourg, 67000 Strasbourg, France

Prof. Dr. Kent Doi
Department of Acute Care Medicine, University of Tokyo, Tokyo, Japan

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