



Personalized Medicine in Benign and Malignant Adrenal Tumors

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Message from the Guest Editors

Dear Colleagues,

Adrenocortical carcinoma is a rare endocrine tumor characterized by a grim prognosis, a strong propensity to recur after surgery, and a limited range of available therapeutic options. Therefore, great efforts are required to identify reliable prognostic factors, novel methods to secure early diagnosis and new markers for a timely detection of tumor recurrence following adrenalectomy. Given that the available medical therapies have limited efficacy and are quite toxic, the identification of new molecular targets that can be druggable and the development of innovative therapeutic agents is a definitive priority to improve patient care. In the meanwhile, studies focusing on the efficacy and safety of existing therapies, as well as on quality of life, should be promoted.

This Special Issue will focus on all of these aspects, in particular on genetic, new “omics” markers, differential diagnosis, prognostic factors, as well as therapeutic approaches and their effects. Both original research and review articles are welcome.

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Guest Editors





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Message from the Editor-in-Chief

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