Special Issue

Current Updates and Future Directions in Neuroendocrine Neoplams (NENs): Understanding Biology, Diagnosis, Management and Research Efforts in NENs

Message from the Guest Editors

Neuroendocrine neoplasms (NENs) are categorized as neuroendocrine tumors (NETs) or neuroendocrine carcinomas (NECs). These are rare malignancies which occur, for example, in the gastrointestinal tract, islets of the pancreas, lung, adrenal medulla, thyroid C-cells, etc., and are heterogeneous groups of neoplasms with unique tumor biology, natural history, and clinical management issues. Most NETs are sporadic, but they can be part of familial cancer syndromes such as multiple endocrine neoplasia type 1 (MEN1), multiple endocrine neoplasia type 2A/B neurofibromatosis type 1 (NF1) or Von Hippel-Lindau (VHL) syndrome. The standard treatment for localized NETs is surgical resection; however, a variety of therapeutic options are available for patients with advanced NETs. When to apply a given option, what combination therapeutic approach should be used, how long treatment should be continued remain unclear and controversial topics. Moreover, patients with these endocrine cancers seek expert advice in the management of their care. This Special Issue will is a collaboration of research which will help us to further understand the biology of endocrine cancers.

Guest Editors

Dr. Jaydira del Rivero

Developmental Therapeutics Branch, Center for Cancer Research, National Cancer Institute, National Institutes of Health, Bethesda, MD 20892, USA

Dr. Aman Chauhan

Department of Medicine, Division of Medical Oncology, University of Miami, Miami, FL 33136, USA

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Cancers
Editorial Office
MDPI, Grosspeteranlage 5
4052 Basel, Switzerland
Tel: +41 61 683 77 34
cancers@mdpi.com

mdpi.com/journal/cancers





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Message from the Editor-in-Chief

Cancers is an international online journal addressing both clinical and basic science issues related to cancer research. The journal is publishing in Open Access format, which will certainly evolve to ensure that the journal takes full advantage of the rapidly changing world of information and knowledge dissemination. It publishes high-quality clinical, translational, and basic science research on cancer prevention, initiation, progression, and treatment, as well as other related topics, particularly to capture the most seminal studies in the rapidly growing area of immunology, immunotherapy, and tumor microenvironment.

Editor-in-Chief

Prof. Dr. Samuel C. Mok.

Department of Gynecologic Oncology and Reproductive Medicine, The University of Texas MD Anderson Cancer Center, Houston, TX 77030, USA

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