



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

Guest Editors:

Dr. Jaydira del Rivero

National Cancer Institute (NCI),
Bethesda, MD, USA

Dr. Aman Chauhan

Sylvester Comprehensive Cancer
Center, University of Miami, Coral
Gables, FL, USA

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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 be a collaboration of research which will help us to further understand the biology of endocrine cancers.





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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

Message from the Editor-in-Chief

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Cancers Editorial Office
MDPI, Grosspeteranlage 5
4052 Basel, Switzerland

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