



Name of Journal: *World Journal of Gastrointestinal Oncology*

Manuscript NO: 53940

Manuscript Type: REVIEW

Gastrointestinal neuroendocrine tumors in 2020

Ahmed M. Gastrointestinal neuroendocrine tumors

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Abstract

Gastrointestinal neuroendocrine tumors are rare slow-growing tumors with distinct histological, biological, and clinical characteristics that have increased in incidence and prevalence within the last few decades. They contain chromogranin A, synaptophysin and neuron-specific enolase which are necessary for making a diagnosis of neuroendocrine tumor. Ki-67 index and mitotic index correlate with cellular proliferation. Serum chromogranin A is the most commonly used biomarker to assess the bulk of disease and monitor treatment and is raised in

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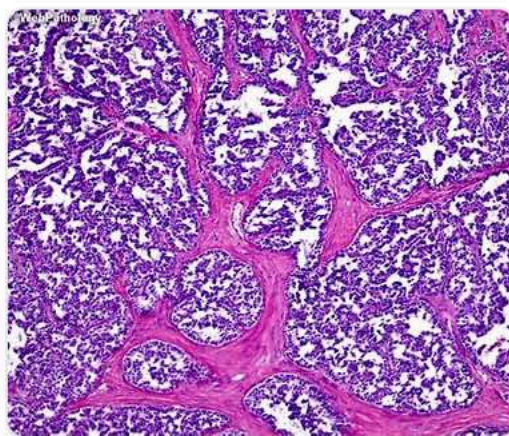
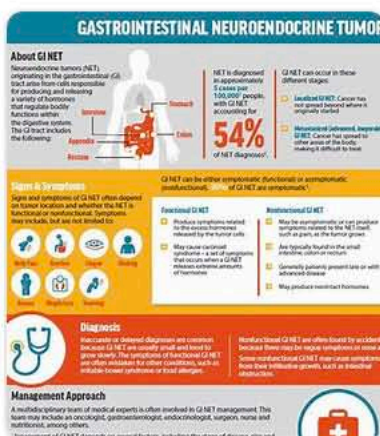
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Neuroendocrine Tumor of the Gastrointestinal Tract: Statistics Approved by the **Cancer.Net** Editorial Board , 01/2020 ON THIS PAGE: You will find information about the number of people who are diagnosed with a GI tract NET each year.

Images of Gastrointestinal Neuroendocrine Tumors in 2020

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Published online on February 24, 2020 Abstract **Neuroendocrine tumors** (NET) of thegastrointestinal tract and pancreas are extremely rare in pediatric population and limited data is available. In most cases, NET of the **gastrointestinal** tract in children are located in the appendix.

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