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Epigenetics of gastroenteropancreatic neuroendocrine tumors: A clinicopathologic perspective

Brendan M Finnerty, Katherine D Gray, Maureen D Moore, Rasa Zarnegar, Thomas J Fahey III

Abstract

30 Gastroenteropancreatic neuroendocrine tumors (GEP-NETs) are a heterogeneous group of rare tumors whose site-specific tumor incidence and clinical behavior vary widely. Genetic alterations associated with familial inherited syndromes have been well defined; however, the genetic profile of sporadic tumors is less clear as their tumorigenesis does not appear to be controlled by classic oncogenes such as P53, RB, or

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