

**NEUROENDOCRINE TUMORS (CURRENT CLINICAL  
ONCOLOGY)**

**Lorraine Kathrine Ekern**

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Neuroendocrine Tumors Editors James C. Yao, MD Department of Gastrointestinal Medical Oncology The University of Texas M.D. Anderson Cancer Center.

Neuroendocrine Tumors--Current and Future Clinical Advances. (1) Department of Medical Oncology, Dana-Farber Cancer Institute, Harvard.

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Oberndorfer S, Frankfurt Z. Prolactinomas are initially treated with cabergoline or bromocriptine, which are long acting dopamine agonists [20].

The youngest reported MEN1 related tumor was a pituitary adenoma in a child 5y. High blood alcohol levels in women. Often, the ambiguity of the symptoms from excess hormones produced by PNETs results in a delay in diagnosis [25]. However, familial clusterings of NETs are rarely reported.

The mean age at diagnosis of pheochromocytoma in patients with NF1, 42 years, diagnosis and treatment of adrenal disease is the same as the sporadic counterparts.