

Opis choroby *

Definicja

A rare inherited cancer syndrome, characterized by the development of multiple neuroendocrine tumors of the parathyroids, gastro-entero-pancreatic tract, and anterior pituitary gland, and less commonly the adrenal cortical gland, thymus and bronchi, with other non-endocrine tumors in some patients.

Dane

Klasyfikacja

Choroba

Synonimy

MEN1

MEN 1

Zespół Wermera

Wermer syndrome

Kod ORPHA

652

Kod OMIM

131100

Kod ICD10

D44.8

Kod ICD11

2F7A.0

*Źródło

orphanet