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

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*Źródło

orphanet