Opis choroby *

Definicja

A rare neoplastic disease characterized by the occurrence of a hormonal syndrome resulting from secretion of humoral factors (including polypeptides, vasoactive amines, and prostaglandins) from a functional neuroendocrine tumor (particularly from the midgut), typically manifesting with increased bowel movements and diarrhea, episodic vasoactive flushes (particularly of the face), hypotension, tachycardia, venous telangiectasia, dyspnea, and bronchospasms, as well as long-term fibrotic changes in the mesentery, retroperitoneum, and of the cardiac valves.

Dane

Klasyfikacja
Zespół kliniczny

Synonimy
Malignant carcinoid syndrome
Złośliwy zespół rakowiaka

Kod ORPHA
100093

Kod OMIM
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Kod ICD10
E34.0

Kod ICD11
5B10

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*Źródło

orphanet