

Opis choroby *

Definicja

A rare subtype of renal cell carcinoma, occurring in the context of end-stage kidney disease and acquired cystic kidney disease, characterized by a usually well circumscribed, solid, multifocal, bilateral tumor with inter- or intracellular microlumen formation (leading to cribriform architecture). Tumors are often diagnosed incidentally in early stages, although complications caused by renal cysts (dull flank or abdominal pain, fever) or renal parenchymal bleeding may mask the underlying neoplastic process. Most have an indolent behavior.

Dane

Klasyfikacja

Choroba

Kod ORPHA

404514

Kod OMIM

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Kod ICD10

C64

Kod ICD11

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*Źródło

orphanet