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Nota di contenuto	Intro -- Contents -- Introduction -- References -- Epidemiology and Screening in RCC -- Epidemiology -- Risk Factors -- Rationale for Screening -- Screening Test -- The Screening Population -- Screening Implementation and Public Acceptability -- Current Nuances -- References -- Hereditary Renal Cancer Predisposition Syndromes -- Von Hippel Lindau (VHL) Disease -- Clinicopathological Hallmarks -- Genetics and Molecular Pathogenesis -- Clinical Management and Therapeutic Approaches -- Germline MET Variants: Hereditary Papillary Renal Cell Carcinoma (HPRC) Syndrome -- Clinicopathological Hallmarks -- Genetics and Molecular Pathogenesis -- Clinical Management and Therapeutic Approaches -- Herediatry Leiomyomatosis and Renal Cell Carcinoma (HLRCC) Syndrome -- Clinicopathological Hallmarks -- Germline Genetics and Molecular Pathogenesis -- Clinical Management and Therapeutic Considerations -- Succinate Dehydrogenase Deficient RCC -- Clinicopathological Hallmarks -- Genetics and Molecular Pathogenesis -- Clinical Management and Therapeutic Considerations -- Disruption of the TCA Cycle: Oncometabolites in RCC -- Birt-Hogg-Dube (BHD) Syndrome -- Clinicopathological Hallmarks -- Genetics and Molecular Pathogenesis -- Clinical Management and Therapeutic Approaches -- BRCA1-Associated Protein (BAP1) Tumour Predisposition Syndrome -- Clinicopathological Hallmarks -- Genomics and Molecular Pathogenesis -- Clinical Management and Therapeutic Approaches -- Tuberous

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Chromophobe Renal Cell Carcinoma (ChRCC) -- Collecting Duct Carcinoma (CDC) -- Renal Medullary Carcinoma (RMC) -- MiT (Microphthalmia-Associated Transcription Factor) Family Translocation Renal Cell Carcinomas -- Succinate Dehydrogenase-Deficient Renal Cell Carcinoma -- Mucinous Tubular and Spindle Cell Carcinoma (MTSRCC) -- Tubulocystic Renal Cell Carcinoma (TCRCC) -- Acquired Cystic Disease-Associated Renal Cell Carcinoma (ACD-RCC) -- Clear Cell Papillary Renal Cell Carcinoma (CCP-RCC).
Renal Cell Carcinoma, Unclassified.