

Precision Medicine Testing Recommendations For: Renal Cell Carcinoma (RCC)

Renal Cell Carcinoma (RCC) – Genetic Risk

Who should be tested?

- Genetic testing is recommended for those with a hereditary renal cell carcinoma syndrome based on any of the following:
- 1st or 2nd degree relative with RCC diagnosis
- Diagnosed ≤ 46 years old
- Bilateral or multifocal tumors
- Multifocal papillary histology
- HLRSS-associated RCC, RCC with Fumarate hydratase (FH) deficiency
- BHDS related histology (multiple chromophobe, oncocytoma, oncocytic hybrid)
- Angiomyolipomas of the kidney and one TSC criterion in same person
- Succinate dehydrogenase (SDH) deficient RCC histology

When and what should be tested?

- Upon or before presentation
- Peripheral blood, saliva or buccal mucosa swab

What tests should be considered?

- NCCN/FDA Approved
 - BAP1
 - FLCN
 - FH
 - MET
 - SDHA/B/C/D
 - TSC1, TSC2
 - VHL

*Ensure inclusion into the molecularly targeted set of testing recommendations