

Klinične smernice ERN GENTURIS

Smernice za spremljanje raka pri posameznikih s sindromom s PTEN povezanih hamartomov
[PHTS pocket guide SI \(genturis.eu\)](#)

Dedni sindrom raka, povezan s TP53

[hTP53rc quick guide SI \(genturis.eu\)](#)

https://www.genturis.eu/l=eng/Assets/ERN-GENTURIS_hTP53rc_Plain-Language-Summary_SI-Version9321.pdf

Švanomatoza

https://www.genturis.eu/l=eng/Assets/SWN-pocket-guide_SI-Version7211.pdf

https://www.genturis.eu/l=eng/Assets/ERN-GENTURIS_SWN_Plain-Language-Summary_SI-Version7192.pdf

Nevrofibromatoza tipa 1

https://www.genturis.eu/l=eng/Assets/Slovenian_NF1-Guideline-summary_v2-Version11614.pdf

[NF1-Guideline---ERN-GENTURIS_v1.16-Version9330.pdf](#)

Smernice v soavtorstvu z ERN GENTURIS, ERN EURACAN in ESMO

Gastrointestinalni stromalni tumorji

[Gastrointestinal stromal tumours: ESMO–EURACAN–GENTURIS Clinical Practice Guidelines for diagnosis, treatment and follow-up - ScienceDirect](#)

Kostni sarkomi

[Bone sarcomas: ESMO–EURACAN–GENTURIS–ERN PaedCan Clinical Practice Guideline for diagnosis, treatment and follow-up - ScienceDirect](#)

Mehkotkivni in visceralni sarkomi

[Soft tissue and visceral sarcomas: ESMO–EURACAN–GENTURIS Clinical Practice Guidelines for diagnosis, treatment and follow-up☆ - ScienceDirect](#)

Priporočila mreže ERN GENTURIS (smernice za diagnosticiranje in spremljanje)

Nevrofibromatoza tipa 2 in švanomatoza

[Updated diagnostic criteria and nomenclature for neurofibromatosis type 2 and schwannomatosis: An international consensus recommendation - ScienceDirect](#)

[Revised diagnostic criteria for neurofibromatosis type 1 and Legius syndrome: an international consensus recommendation - ScienceDirect](#)

Sindrom Lynch

[The Manchester International Consensus Group recommendations for the management of gynecological cancers in Lynch syndrome - ScienceDirect](#)

[Thieme E-Journals - Endoscopy / Abstract \(thieme-connect.de\)](#)

[European guidelines from the EHTG and ESCP for Lynch syndrome: an updated third edition of the Mallorca guidelines based on gene and gender | British Journal of Surgery | Oxford Academic \(oup.com\)](#)

Polipoze in Peutz-Jeghersov sindrom

[Thieme E-Journals - Endoscopy / Abstract \(thieme-connect.de\)](#)

[JCM | Free Full-Text | The Management of Peutz–Jeghers Syndrome: European Hereditary Tumour Group \(EHTG\) Guideline \(mdpi.com\)](#)

Dedni kolorektalni rak

[Guidelines for the management of hereditary colorectal cancer from the British Society of Gastroenterology \(BSG\)/Association of Coloproctology of Great Britain and Ireland \(ACPGBI\)/United Kingdom Cancer Genetics Group \(UKCGG\) | Gut \(bmj.com\)](#)

Drugi dedni sindromi

Sindrom von Hippel-Lindau (VHL)

[von Hippel-Lindau disease: Updated guideline for diagnosis and surveillance - ScienceDirect](#)

Sindrom Gorlin

[guideline for the clinical management of basal cell naevus syndrome \(Gorlin–Goltz syndrome\)* | British Journal of Dermatology | Oxford Academic \(oup.com\)](#)

Nosilci patogenih/verjetno patogenih različic v genu DICER1

[Surveillance recommendations for DICER1 pathogenic variant carriers: a report from the SIOPE Host Genome Working Group and CanGene-CanVar Clinical Guideline Working Group | SpringerLink](#)

Dedni difuzni rak želodca

[Hereditary diffuse gastric cancer: updated clinical practice guidelines - ScienceDirect](#)

Drobnocelični rak jajčnikov

[Small-Cell Carcinoma of the Ovary, Hypercalcemic Type–Genetics, New Treatment Targets, and Current Management Guidelines | Clinical Cancer Research | American Association for Cancer Research \(aacrjournals.org\)](#)