Verena Steinke-Lange

List of Publications by Year
in descending order

26 papers	1,380 citations	623188 14 h-index	525886 27 g-index
29 all docs	29 docs citations	29 times ranked	1555 citing authors

#	Article	IF	Citations
1	Adenoma and colorectal cancer risks in Lynch syndrome, Lynchâ€like syndrome and familial colorectal cancer type X. International Journal of Cancer, 2022, 150, 56-66.	2.3	2
2	Clinical Validity of Circulating Tumor DNA as Prognostic and Predictive Marker for Personalized Colorectal Cancer Patient Management. Cancers, 2022, 14, 851.	1.7	5
3	Long-term chemoprevention in patients with adenomatous polyposis coli: an observational study. Familial Cancer, 2022, 21, 463-472.	0.9	4
4	Splicing analyses for variants in MMR genes: best practice recommendations from the European Mismatch Repair Working Group. European Journal of Human Genetics, 2022, 30, 1051-1059.	1.4	7
5	The â€œunnaturalâ€•history of colorectal cancer in Lynch syndrome: Lessons from colonoscopy surveillance. International Journal of Cancer, 2021, 148, 800-811.	2.3	55
6	Value of upper <scp>gastrointestinal</scp> endoscopy for gastric cancer surveillance in patients with Lynch syndrome. International Journal of Cancer, 2021, 148, 106-114.	2.3	28
7	Uptake of hysterectomy and bilateral salpingo-oophorectomy in carriers of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. European Journal of Cancer, 2021, 148, 124-133.	1.3	11
8	Revised diagnostic criteria for neurofibromatosis type 1 and Legius syndrome: an international consensus recommendation. Genetics in Medicine, 2021, 23, 1506-1513.	1.1	290
9	No Difference in Penetrance between Truncating and Missense/Aberrant Splicing Pathogenic Variants in MLH1 and MSH2: A Prospective Lynch Syndrome Database Study. Journal of Clinical Medicine, 2021, 10, 2856.	1.0	11
10	Early detection of duodenal cancer by upper <scp>gastrointestinal</scp>â€endoscopy in Lynch syndrome. International Journal of Cancer, 2021, 149, 2052-2062.	2.3	4
11	Variation in the risk of colorectal cancer in families with Lynch syndrome: a retrospective cohort study. Lancet Oncology, The, 2021, 22, 1014-1022.	5.1	58
12	Liquid Biopsy Hotspot Variant Assays: Analytical Validation for Application in Residual Disease Detection and Treatment Monitoring. Clinical Chemistry, 2021, 67, 1483-1491.	1.5	7
13	Somatic mosaics in hereditary tumor predisposition syndromes. European Journal of Medical Genetics, 2021, 64, 104360.	0.7	8
14	Overview of hereditary breast and ovarian cancer (HBOC) guidelines across Europe. European Journal of Medical Genetics, 2021, 64, 104350.	0.7	22
15	Cancer risks by gene, age, and gender in 6350 carriers of pathogenic mismatch repair variants: findings from the Prospective Lynch Syndrome Database. Genetics in Medicine, 2020, 22, 15-25.	1.1	365
16	Risk-Reducing Gynecological Surgery in Lynch Syndrome: Results of an International Survey from the Prospective Lynch Syndrome Database. Journal of Clinical Medicine, 2020, 9, 2290.	1.0	12
17	Ageâ€dependent performance of <scp>BRAF</scp> mutation testing in Lynch syndrome diagnostics. International Journal of Cancer, 2020, 147, 2801-2810.	2.3	17
18	Associations of Pathogenic Variants in MLH1, MSH2, and MSH6 With Risk of Colorectal Adenomas and Tumors and With Somatic Mutations in Patients With Lynch Syndrome. Gastroenterology, 2020, 158, 1326-1333.	0.6	60

#	Article	IF	Citations
19	Prevalence of CNV-neutral structural genomic rearrangements in MLH1, MSH2, and PMS2 not detectable in routine NGS diagnostics. Familial Cancer, 2020, 19, 161-167.	0.9	11
20	Cancer risks in Lynch syndrome, Lynch-like syndrome, and familial colorectal cancer type X: a prospective cohort study. BMC Cancer, 2020, 20, 460.	1.1	32
21	Full-length transcript amplification and sequencing as universal method to test mRNA integrity and biallelic expression in mismatch repair genes. European Journal of Human Genetics, 2019, 27, 1808-1820.	1.4	16
22	Survival by colon cancer stage and screening interval in Lynch syndrome: a prospective Lynch syndrome database report. Hereditary Cancer in Clinical Practice, 2019, 17, 28.	0.6	27
23	The Apparent Genetic Anticipation in PMS2-Associated Lynch Syndrome Families Is Explained by Birth-cohort Effect. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 1010-1014.	1.1	6
24	Lack of association between screening interval and cancer stage in Lynch syndrome may be accounted for by over-diagnosis; a prospective Lynch syndrome database report. Hereditary Cancer in Clinical Practice, 2019, 17, 8.	0.6	42
25	Cancer Risks for PMS2-Associated Lynch Syndrome. Journal of Clinical Oncology, 2018, 36, 2961-2968.	0.8	147
26	No Difference in Colorectal Cancer Incidence or Stage at Detection by Colonoscopy Among 3 Countries With Different Lynch Syndrome Surveillance Policies. Gastroenterology, 2018, 155, 1400-1409.e2.	0.6	112