Christina Therkildsen

List of Publications by Year in descending order

36 papers	1,296 citations	623734 14 h-index	361022 35 g-index
41 all docs	41 docs citations	41 times ranked	2653 citing authors

#	Article	IF	Citations
1	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.	2.4	365
2	The predictive value ofKRAS, NRAS, BRAF, PIK3CAand PTEN for anti-EGFR treatment in metastatic colorectal cancer: A systematic review and meta-analysis. Acta OncolÃ³gica, 2014, 53, 852-864.	1.8	324
3	Urinary Tract Cancer in Lynch Syndrome; Increased Risk in Carriers of MSH2 Mutations. Urology, 2015, 86, 1212-1217.	1.0	74
4	Sarcomas associated with hereditary nonpolyposis colorectal cancer: broad anatomical and morphological spectrum. Familial Cancer, 2009, 8, 209-213.	1.9	58
5	Frequent mismatch-repair defects link prostate cancer to Lynch syndrome. BMC Urology, 2016, 16, 15.	1.4	52
6	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.	1.5	42
7	Familial colorectal cancer type X: genetic profiles and phenotypic features. Modern Pathology, 2015, 28, 30-36.	5.5	37
8	Towards gene- and gender-based risk estimates in Lynch syndrome; age-specific incidences for 13 extra-colorectal cancer types. British Journal of Cancer, 2017, 117, 1702-1710.	6.4	36
9	An Update on Immune Checkpoint Therapy for the Treatment of Lynch Syndrome. Clinical and Experimental Gastroenterology, 2021, Volume 14, 181-197.	2.3	36
10	Risk-reducing hysterectomy and bilateral salpingo-oophorectomy in female heterozygotes of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. Genetics in Medicine, 2021, 23, 705-712.	2.4	28
11	Distinct Gene Expression Signatures in Lynch Syndrome and Familial Colorectal Cancer Type X. PLoS ONE, 2013, 8, e71755.	2.5	28
12	Molecular subtype classification of urothelial carcinoma in Lynch syndrome. Molecular Oncology, 2018, 12, 1286-1295.	4.6	25
13	Gain of chromosomal region 20q and loss of 18 discriminates between Lynch syndrome and familial colorectal cancer. European Journal of Cancer, 2013, 49, 1226-1235.	2.8	23
14	New Pathogenic Germline Variants in Very Early Onset and Familial Colorectal Cancer Patients. Frontiers in Genetics, 2020, 11, 566266.	2.3	16
15	Immunoprofiles of colorectal cancer from Lynch syndrome. OncoImmunology, 2019, 8, e1515612.	4.6	14
16	The influence of marital status and partner concordance on participation in colorectal cancer screening. European Journal of Public Health, 2021, 31, 340-346.	0.3	14
17	Increased risk of male cancer and identification of a potential prostate cancer cluster region in BRCA2. Acta OncolÃ³gica, 2016, 55, 38-44.	1.8	13
18	An alternative approach to establishing unbiased colorectal cancer risk estimation in Lynch syndrome. Genetics in Medicine, 2019, 21, 2706-2712.	2.4	11

#	Article	IF	Citations
19	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.	2.8	11
20	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.	2.4	11
21	Lynch syndrome-associated epithelial ovarian cancer and its immunological profile. Gynecologic Oncology, 2021, 162, 686-693.	1.4	10
22	BRCA1/BRCA2 founder mutations and cancer risks: impact in the western Danish population. Familial Cancer, 2016, 15, 507-512.	1.9	9
23	Immunohistochemical Screening of Upper Tract Urothelial Carcinomas for Lynch Syndrome Diagnostics: A Systematic Review. Urology, 2022, 165, 44-53.	1.0	8
24	Cancer risks and immunohistochemical profiles linked to the Danish MLH1 Lynch syndrome founder mutation. Familial Cancer, 2012, 11, 579-585.	1.9	7
25	Renal cell cancer linked to Lynch syndrome: Increased incidence and loss of mismatch repair protein expression. International Journal of Urology, 2016, 23, 528-529.	1.0	7
26	Characterization of burning mouth syndrome profiles based on response to a local anaesthetic lozenge. Oral Diseases, 2020, 26, 656-669.	3.0	7
27	Broadening risk profile in familial colorectal cancer type X; increased risk for five cancer types in the national Danish cohort. BMC Cancer, 2020, 20, 345.	2.6	5
28	An effect from anticipation also in hereditary nonpolyposis colorectal cancer families without identified mutations. Cancer Epidemiology, 2009, 33, 231-234.	1.9	4
29	Deranged Wnt signaling is frequent in hereditary nonpolyposis colorectal cancer. Familial Cancer, 2011, 10, 239-243.	1.9	4
30	Histological and Molecular Adipose Tissue Changes Are Related to Metabolic Syndrome Rather Than Lipodystrophy in Human Immunodeficiency Virus-Infected Patients: A Cross-Sectional Study. Journal of Infectious Diseases, 2018, 218, 1090-1098.	4.0	4
31	Risk of multiple colorectal cancer development depends on age and subgroup in individuals with hereditary predisposition. Familial Cancer, 2019, 18, 183-191.	1.9	4
32	Use of primary health care and participation in colorectal cancer screening â€“ a Danish national register-based study. Acta OncolÃ³gica, 2020, 59, 1002-1006.	1.8	3
33	Functional implications of the p.Cys680Arg mutation in the MLH1 mismatch repair protein. Molecular Genetics & Genomic Medicine, 2014, 2, 352-355.	1.2	1
34	Abstract LB-439: Distinct tumorigenic pathways within the hereditary nonpolyposis colorectal cancer. , 2012, , .		0
35	Abstract 5216: Broadening tumor spectrum in Lynch syndrome: increased incidence for 15 distinct cancer types. , 2016, , .		0
36	Colorectal cancer in adolescents and young adults with Lynch syndrome: a Danish register-based study. BMJ Open, 2021, 11, e053538.	1.9	0