# Christina Therkildsen

## List of Publications by Year in descending order



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|---|---|---|---|
| 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 | Immunohistochemical Screening of Upper Tract Urothelial Carcinomas for Lynch Syndrome Diagnostics: A Systematic Review. Urology, 2022, 165, 44-53. | 1.0 | 8 |
| 2 | 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 |
| 3 | 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 |
| 4 | An Update on Immune Checkpoint Therapy for the Treatment of Lynch Syndrome. Clinical and Experimental Gastroenterology, 2021, Volume 14, 181-197. | 2.3 | 36 |
| 5 | 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 |
| 6 | Lynch syndrome-associated epithelial ovarian cancer and its immunological profile. Gynecologic Oncology, 2021, 162, 686-693. | 1.4 | 10 |
| 7 | 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 |
| 8 | Colorectal cancer in adolescents and young adults with Lynch syndrome: a Danish register-based study. BMJ Open, 2021, 11, e053538. | 1.9 | 0 |
| 9 | 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 |
| 10 | Characterization of burning mouth syndrome profiles based on response to a local anaesthetic lozenge. Oral Diseases, 2020, 26, 656-669. | 3.0 | 7 |
| 11 | New Pathogenic Germline Variants in Very Early Onset and Familial Colorectal Cancer Patients. Frontiers in Genetics, 2020, 11, 566266. | 2.3 | 16 |
| 12 | 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 |
| 13 | 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 |
| 14 | An alternative approach to establishing unbiased colorectal cancer risk estimation in Lynch syndrome. Genetics in Medicine, 2019, 21, 2706-2712. | 2.4 | 11 |
| 15 | 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 |
| 16 | Immunoprofiles of colorectal cancer from Lynch syndrome. OncoImmunology, 2019, 8, e1515612. | 4.6 | 14 |
| 17 | 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 |
| 18 | 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 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Molecular subtype classification of urothelial carcinoma in Lynch syndrome. Molecular Oncology, 2018, 12, 1286-1295. | 4.6 | 25 |
| 20 | 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 |
| 21 | 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 |
| 22 | Frequent mismatch-repair defects link prostate cancer to Lynch syndrome. BMC Urology, 2016, 16, 15. | 1.4 | 52 |
| 23 | BRCA1/BRCA2 founder mutations and cancer risks: impact in the western Danish population. Familial Cancer, 2016, 15, 507-512. | 1.9 | 9 |
| 24 | 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 |
| 25 | Abstract 5216: Broadening tumor spectrum in Lynch syndrome: increased incidence for 15 distinct cancer types. , 2016, , . | | 0 |
| 26 | Urinary Tract Cancer in Lynch Syndrome; Increased Risk in Carriers of MSH2 Mutations. Urology, 2015, 86, 1212-1217. | 1.0 | 74 |
| 27 | Familial colorectal cancer type X: genetic profiles and phenotypic features. Modern Pathology, 2015, 28, 30-36. | 5.5 | 37 |
| 28 | Functional implications of the p.Cys680Arg mutation in the MLH1 mismatch repair protein. Molecular Genetics & Genomic Medicine, 2014, 2, 352-355. | 1.2 | 1 |
| 29 | 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 |
| 30 | 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 |
| 31 | Distinct Gene Expression Signatures in Lynch Syndrome and Familial Colorectal Cancer Type X. PLoS ONE, 2013, 8, e71755. | 2.5 | 28 |
| 32 | Cancer risks and immunohistochemical profiles linked to the Danish MLH1 Lynch syndrome founder mutation. Familial Cancer, 2012, 11, 579-585. | 1.9 | 7 |
| 33 | Abstract LB-439: Distinct tumorigenic pathways within the hereditary nonpolyposis colorectal cancer. , 2012, , . | | 0 |
| 34 | Deranged Wnt signaling is frequent in hereditary nonpolyposis colorectal cancer. Familial Cancer, 2011, 10, 239-243. | 1.9 | 4 |
| 35 | An effect from anticipation also in hereditary nonpolyposis colorectal cancer families without identified mutations. Cancer Epidemiology, 2009, 33, 231-234. | 1.9 | 4 |
| 36 | Sarcomas associated with hereditary nonpolyposis colorectal cancer: broad anatomical and morphological spectrum. Familial Cancer, 2009, 8, 209-213. | 1.9 | 58 |