

# CITATION REPORT

List of articles citing

## Recurrent Coding Sequence Variation Explains Only A Small Fraction of the Genetic Architecture of Colorectal Cancer

DOI: 10.1038/srep16286  
Scientific Reports, 2015, 5, 16286.

**Source:** <https://exaly.com/paper-pdf/60304637/citation-report.pdf>

**Version:** 2024-04-28

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
23	Correspondence: SEMA4A variation and risk of colorectal cancer. <i>Nature Communications</i> , <b>2016</b> , 7, 10611	17.4	6
22	Somatic POLE proofreading domain mutation, immune response, and prognosis in colorectal cancer: a retrospective, pooled biomarker study. <i>The Lancet Gastroenterology and Hepatology</i> , <b>2016</b> , 1, 207-216	18.8	160
21	Rare disruptive mutations and their contribution to the heritable risk of colorectal cancer. <i>Nature Communications</i> , <b>2016</b> , 7, 11883	17.4	86
20	Evidence for genetic association between chromosome 1q loci and predisposition to colorectal neoplasia. <i>British Journal of Cancer</i> , <b>2017</b> , 117, 1215-1223	8.7	8
19	Validation of Recently Proposed Colorectal Cancer Susceptibility Gene Variants in an Analysis of Families and Patients-a Systematic Review. <i>Gastroenterology</i> , <b>2017</b> , 152, 75-77.e4	13.3	62
18	Coding variants in NOD-like receptors: An association study on risk and survival of colorectal cancer. <i>PLoS ONE</i> , <b>2018</b> , 13, e0199350	3.7	5
17	A Rare Missense Variant in TCF7L2 Associates with Colorectal Cancer Risk by Interacting with a GWAS-Identified Regulatory Variant in the MYC Enhancer. <i>Cancer Research</i> , <b>2018</b> , 78, 5164-5172	10.1	46
16	Genome-wide scan of the effect of common nSNPs on colorectal cancer survival outcome. <i>British Journal of Cancer</i> , <b>2018</b> , 119, 988-993	8.7	4
15	Exploring causality in the association between circulating 25-hydroxyvitamin D and colorectal cancer risk: a large Mendelian randomisation study. <i>BMC Medicine</i> , <b>2018</b> , 16, 142	11.4	40
14	Personalized prediction of genes with tumor-causing somatic mutations based on multi-modal deep Boltzmann machine. <i>Neurocomputing</i> , <b>2019</b> , 324, 51-62	5.4	5
13	Novel significant stage-specific differentially expressed genes in hepatocellular carcinoma. <i>BMC Cancer</i> , <b>2019</b> , 19, 663	4.8	31
12	Effects of common genetic variants associated with colorectal cancer risk on survival outcomes after diagnosis: A large population-based cohort study. <i>International Journal of Cancer</i> , <b>2019</b> , 145, 2427-2432	7.5	8
11	MicroRNA-binding site polymorphisms and risk of colorectal cancer: A systematic review and meta-analysis. <i>Cancer Medicine</i> , <b>2019</b> , 8, 7477-7499	4.8	13
10	Head-to-Head Comparison of Family History of Colorectal Cancer and a Genetic Risk Score for Colorectal Cancer Risk Stratification. <i>Clinical and Translational Gastroenterology</i> , <b>2019</b> , 10, e00106	4.2	3
9	Screening of molecular targets and construction of a ceRNA network for oxaliplatin resistance in colorectal cancer.. <i>RSC Advances</i> , <b>2019</b> , 9, 31413-31424	3.7	1
8	LNK protein: Low expression in human colorectal carcinoma and relationship with tumor invasion. <i>Biomedicine and Pharmacotherapy</i> , <b>2020</b> , 121, 109467	7.5	2
7	A Systematic Analysis of Interactions between Environmental Risk Factors and Genetic Variation in Susceptibility to Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , <b>2020</b> , 29, 1145-1153 <sup>4</sup>		9

6	Rare Variants in the DNA Repair Pathway and the Risk of Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , <b>2021</b> , 30, 895-903	4	1
5	Differential genetic influences over colorectal cancer risk and gene expression in large bowel mucosa. <i>International Journal of Cancer</i> , <b>2021</b> , 149, 1100-1108	7.5	2
4	Colorectal cancer risk variant rs7017386 modulates two oncogenic lncRNAs expression via ATF1-mediated long-range chromatin loop. <i>Cancer Letters</i> , <b>2021</b> , 518, 140-151	9.9	1
3	Novel Significant Stage-Specific Differentially Expressed Genes in Liver Hepatocellular Carcinoma.		0
2	Recurrent, low-frequency coding variants contributing to colorectal cancer in the Swedish population. <i>PLoS ONE</i> , <b>2018</b> , 13, e0193547	3.7	3
1	SNP-Target Genes Interaction Perturbing the Cancer Risk in the Post-GWAS. <b>2022</b> , 14, 5636		1