

# Nicholas Light

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/10396346/publications.pdf>

Version: 2024-02-01

11  
papers

1,017  
citations

1040056

9  
h-index

1372567

10  
g-index

12  
all docs

12  
docs citations

12  
times ranked

3184  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pathogenic Germline Variants in Cancer Susceptibility Genes in Children and Young Adults With Rhabdomyosarcoma. <i>JCO Precision Oncology</i> , 2021, 5, 75-87.	3.0	27
2	Lineage-defined leiomyosarcoma subtypes emerge years before diagnosis and determine patient survival. <i>Nature Communications</i> , 2021, 12, 4496.	12.8	28
3	DNA Polymerase and Mismatch Repair Exert Distinct Microsatellite Instability Signatures in Normal and Malignant Human Cells. <i>Cancer Discovery</i> , 2021, 11, 1176-1191.	9.4	46
4	Myc and Loss of p53 Cooperate to Drive Formation of Choroid Plexus Carcinoma. <i>Cancer Research</i> , 2019, 79, 2208-2219.	0.9	15
5	Ewing-like sarcoma: An emerging family of round cell sarcomas. <i>Journal of Cellular Physiology</i> , 2019, 234, 7999-8007.	4.1	68
6	HGG-17. TUMOR MUTATIONAL BURDEN ANALYSIS OF PEDIATRIC TUMORS PROVIDES A DIAGNOSTIC TOOL FOR GERMLINE REPLICATION REPAIR DEFICIENCY AND PREDICT RESPONSE TO IMMUNE CHECKPOINT INHIBITION. <i>Neuro-Oncology</i> , 2018, 20, i92-i92.	1.2	0
7	Explosive mutation accumulation triggered by heterozygous human Pol $\mu$ proofreading-deficiency is driven by suppression of mismatch repair. <i>ELife</i> , 2018, 7, .	6.0	33
8	Comprehensive Analysis of Hypermutation in Human Cancer. <i>Cell</i> , 2017, 171, 1042-1056.e10.	28.9	596
9	Interrogation of allelic chromatin states in human cells by high-density ChIP-genotyping. <i>Epigenetics</i> , 2014, 9, 1238-1251.	2.7	9
10	Allelic expression mapping across cellular lineages to establish impact of non-coding $\langle \text{sc} \rangle \text{SNP} \langle / \text{sc} \rangle$ s. <i>Molecular Systems Biology</i> , 2014, 10, 754.	7.2	21
11	Analysis of expressed SNPs identifies variable extents of expression from the human inactive X chromosome. <i>Genome Biology</i> , 2013, 14, R122.	9.6	174