

# C Anthony Scott

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/12195513/publications.pdf>

Version: 2024-02-01

7  
papers

236  
citations

1684188

5  
h-index

1720034

7  
g-index

7  
all docs

7  
docs citations

7  
times ranked

482  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in <i>C8ORF37</i> cause Bardet Biedl syndrome (BBS21). <i>Human Molecular Genetics</i> , 2016, 25, 2283-2294.	2.9	91
2	Hypomorphic mutations in <i>TRNT1</i> cause retinitis pigmentosa with erythrocytic microcytosis. <i>Human Molecular Genetics</i> , 2016, 25, 44-56.	2.9	64
3	The master transcription factor SOX2, mutated in anophthalmia/microphthalmia, is post-transcriptionally regulated by the conserved RNA-binding protein RBM24 in vertebrate eye development. <i>Human Molecular Genetics</i> , 2020, 29, 591-604.	2.9	34
4	Automated, high-throughput, in vivo analysis of visual function using the zebrafish. <i>Developmental Dynamics</i> , 2016, 245, 605-613.	1.8	27
5	A High-Throughput Assay for Congenital and Age-Related Eye Diseases in Zebrafish. <i>Biomedicines</i> , 2019, 7, 28.	3.2	12
6	Development and biological characterization of a clinical gene transfer vector for the treatment of MAK-associated retinitis pigmentosa. <i>Gene Therapy</i> , 2021, , .	4.5	5
7	Functional Role of the RNA-Binding Protein Rbm24a and Its Target sox2 in Microphthalmia. <i>Biomedicines</i> , 2021, 9, 100.	3.2	3