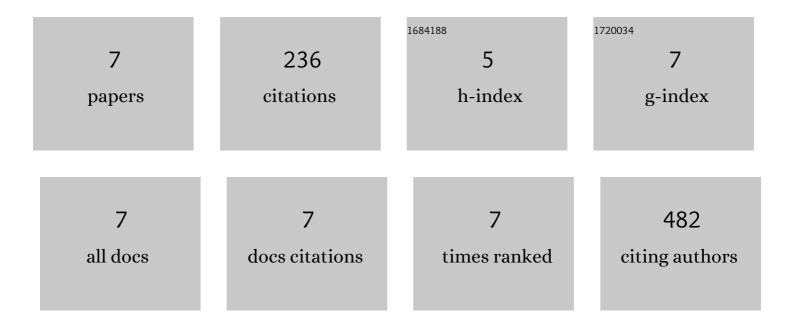
## C Anthony Scott

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/12195513/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Mutations in <i>C8ORF37</i> cause Bardet Biedl syndrome (BBS21). Human Molecular Genetics, 2016, 25, 2283-2294.	2.9	91
2	Hypomorphic mutations in <i>TRNT1</i> cause retinitis pigmentosa with erythrocytic microcytosis. Human Molecular Genetics, 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. Human Molecular Genetics, 2020, 29, 591-604.	2.9	34
4	Automated, highâ€ŧhroughput, in vivo analysis of visual function using the zebrafish. Developmental Dynamics, 2016, 245, 605-613.	1.8	27
5	A High-Throughput Assay for Congenital and Age-Related Eye Diseases in Zebrafish. Biomedicines, 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. Gene Therapy, 2021, , .	4.5	5
7	Functional Role of the RNA-Binding Protein Rbm24a and Its Target sox2 in Microphthalmia. Biomedicines, 2021, 9, 100.	3.2	3