

# Jonathan Eintracht

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/10146752/publications.pdf>

Version: 2024-02-01

8  
papers

54  
citations

1937685

4  
h-index

1720034

7  
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8  
docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	The Use of Induced Pluripotent Stem Cells as a Model for Developmental Eye Disorders. <i>Frontiers in Cellular Neuroscience</i> , 2020, 14, 265.	3.7	15
2	Translational readthrough of ciliopathy genes BBS2 and ALMS1 restores protein, ciliogenesis and function in patient fibroblasts. <i>EBioMedicine</i> , 2021, 70, 103515.	6.1	14
3	CUGC for syndromic microphthalmia including next-generation sequencing-based approaches. <i>European Journal of Human Genetics</i> , 2020, 28, 679-690.	2.8	6
4	Generation of human iPSC line (UCLi013-A) from a patient with microphthalmia and aniridia, carrying a heterozygous missense mutation c.372C>A p.(Asn124Lys) in PAX6. <i>Stem Cell Research</i> , 2021, 51, 102184.	0.7	5
5	Generation of two human control iPS cell lines (UCLi016-A and UCLi017-A) from healthy donors with no known ocular conditions. <i>Stem Cell Research</i> , 2020, 49, 102113.	0.7	5
6	Identification of 4 novel human ocular coloboma genes ANK3, BMPR1B, PDGFRA, and CDH4 through evolutionary conserved vertebrate gene analysis. <i>Genetics in Medicine</i> , 2022, 24, 1073-1084.	2.4	4
7	Efficient embryoid-based method to improve generation of optic vesicles from human induced pluripotent stem cells. <i>F1000Research</i> , 0, 11, 324.	1.6	3
8	Generation of two human iPSC lines from patients with autosomal dominant retinitis pigmentosa (UCLi014-A) and autosomal recessive Leber congenital amaurosis (UCLi015-A), associated with RDH12 variants. <i>Stem Cell Research</i> , 2021, 54, 102449.	0.7	2