Nikolai Klena

List of Publications by Year in Descending Order

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

17	989	11	2 O
papers	citations	h-index	g-index
20 ext. papers	1,308 ext. citations	21 avg, IF	3.28 L-index

#	Paper	IF	Citations
17	Visualizing the native cellular organization by coupling cryofixation with expansion microscopy (Cryo-ExM) <i>Nature Methods</i> , 2022 ,	21.6	5
16	Kinesin-1 activity recorded in living cells with a precipitating dye. <i>Nature Communications</i> , 2021 , 12, 146	317.4	6
15	Overview of the centriole architecture. Current Opinion in Structural Biology, 2021, 66, 58-65	8.1	19
14	A helical inner scaffold provides a structural basis for centriole cohesion. <i>Science Advances</i> , 2020 , 6, eaa	z 4 4.37	54
13	Architecture of the centriole cartwheel-containing region revealed by cryo-electron tomography. <i>EMBO Journal</i> , 2020 , 39, e106246	13	22
12	Biliary-Atresia-Associated Mannosidase-1-Alpha-2 Gene Regulates Biliary and Ciliary Morphogenesis and Laterality. <i>Frontiers in Physiology</i> , 2020 , 11, 538701	4.6	4
11	Isolation and Fluorescence Imaging for Single-particle Reconstruction of Chlamydomonas Centrioles. <i>Journal of Visualized Experiments</i> , 2018 ,	1.6	4
10	The complex genetics of hypoplastic left heart syndrome. <i>Nature Genetics</i> , 2017 , 49, 1152-1159	36.3	107
9	Role of Cilia and Left-Right Patterning in Congenital Heart Disease 2016 , 67-79		2
8	Genetic link between renal birth defects and congenital heart disease. <i>Nature Communications</i> , 2016 , 7, 11103	17.4	32
7	Global genetic analysis in mice unveils central role for cilia in congenital heart disease. <i>Nature</i> , 2015 , 521, 520-4	50.4	256
6	MMP21 is mutated in human heterotaxy and is required for normal left-right asymmetry in vertebrates. <i>Nature Genetics</i> , 2015 , 47, 1260-3	36.3	52
5	Discovery of four recessive developmental disorders using probabilistic genotype and phenotype matching among 4,125 families. <i>Nature Genetics</i> , 2015 , 47, 1363-9	36.3	91
4	CCDC151 mutations cause primary ciliary dyskinesia by disruption of the outer dynein arm docking complex formation. <i>American Journal of Human Genetics</i> , 2014 , 95, 257-74	11	113
3	Role of cilia in structural birth defects: insights from ciliopathy mutant mouse models. <i>Birth Defects Research Part C: Embryo Today Reviews</i> , 2014 , 102, 115-25		21
2	DYX1C1 is required for axonemal dynein assembly and ciliary motility. <i>Nature Genetics</i> , 2013 , 45, 995-10	- 0 33 .3	197
1	In situ architecture of the ciliary base reveals the stepwise assembly of IFT trains		2