

# Emmanuelle Szenker-Ravi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9498293/publications.pdf>

Version: 2024-02-01

11  
papers

782  
citations

933410

10  
h-index

1199563

12  
g-index

14  
all docs

14  
docs citations

14  
times ranked

1631  
citing authors

#	ARTICLE	IF	CITATIONS
1	The double face of the histone variant H3.3. <i>Cell Research</i> , 2011, 21, 421-434.	12.0	324
2	RSPO2 inhibition of RNF43 and ZNRF3 governs limb development independently of LGR4/5/6. <i>Nature</i> , 2018, 557, 564-569.	27.8	141
3	Developmental roles of histone H3 variants and their chaperones. <i>Trends in Genetics</i> , 2013, 29, 630-640.	6.7	104
4	<i>CC2D2A</i> mutations in Meckel and Joubert syndromes indicate a genotype-phenotype correlation. <i>Human Mutation</i> , 2009, 30, 1574-1582.	2.5	80
5	A Human Pleiotropic Multiorgan Condition Caused by Deficient Wnt Secretion. <i>New England Journal of Medicine</i> , 2021, 385, 1292-1301.	27.0	23
6	Next-generation sequencing in a series of 80 fetuses with complex cardiac malformations and/or heterotaxy. <i>Human Mutation</i> , 2020, 41, 2167-2178.	2.5	21
7	R-spondin signalling is essential for the maintenance and differentiation of mouse nephron progenitors. <i>ELife</i> , 2020, 9, .	6.0	20
8	R-SPONDIN2 mesenchymal cells form the bud tip progenitor niche during human lung development. <i>Developmental Cell</i> , 2022, 57, 1598-1614.e8.	7.0	19
9	Homozygous Null TBX4 Mutations Lead to Posterior Amelia with Pelvic and Pulmonary Hypoplasia. <i>American Journal of Human Genetics</i> , 2019, 105, 1294-1301.	6.2	17
10	Discovery of a genetic module essential for assigning left-right asymmetry in humans and ancestral vertebrates. <i>Nature Genetics</i> , 2022, 54, 62-72.	21.4	16
11	A <i>GLI3</i> variant leading to polydactyly in heterozygotes and Pallister-Hall-like syndrome in a homozygote. <i>Clinical Genetics</i> , 2020, 97, 915-919.	2.0	3