Emmanuelle Szenker-Ravi

List of Publications by Year
in descending order

11	782	933447 10	1199594 12
papers	citations	h-index	g-index

14	14	14	1631
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Discovery of a genetic module essential for assigning leftâ€“right asymmetry in humans and ancestral vertebrates. Nature Genetics, 2022, 54, 62-72.	21.4	16
2	R-SPONDIN2 mesenchymal cells form the bud tip progenitor niche during human lung development. Developmental Cell, 2022, 57, 1598-1614.e8.	7.0	19
3	A Human Pleiotropic Multiorgan Condition Caused by Deficient Wnt Secretion. New England Journal of Medicine, 2021, 385, 1292-1301.	27.0	23
4	Nextâ€generation sequencing in a series of 80 fetuses with complex cardiac malformations and/or heterotaxy. Human Mutation, 2020, 41, 2167-2178.	2.5	21
5	A <scp>GLI3</scp> variant leading to polydactyly in heterozygotes and Pallisterâ€Hallâ€like syndrome in a homozygote. Clinical Genetics, 2020, 97, 915-919.	2.0	3
6	R-spondin signalling is essential for the maintenance and differentiation of mouse nephron progenitors. ELife, 2020, 9, .	6.0	20
7	Homozygous Null TBX4 Mutations Lead to Posterior Amelia with Pelvic and Pulmonary Hypoplasia. American Journal of Human Genetics, 2019, 105, 1294-1301.	6.2	17
8	RSPO2 inhibition of RNF43 and ZNRF3 governs limb development independently of LGR4/5/6. Nature, 2018, 557, 564-569.	27.8	141
9	Developmental roles of histone H3 variants and their chaperones. Trends in Genetics, 2013, 29, 630-640.	6.7	104
10	The double face of the histone variant H3.3. Cell Research, 2011, 21, 421-434.	12.0	324
11	CC2D2Amutations in Meckel and Joubert syndromes indicate a genotype-phenotype correlation. Human Mutation, 2009, 30, 1574-1582.	2.5	80