

Silvia Martin-Almedina

List of Publications by Year in descending order

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7
papers

447
citations

1478505

6
h-index

1720034

7
g-index

8
all docs

8
docs citations

8
times ranked

876
citing authors

#	ARTICLE	IF	CITATIONS
1	Janus-faced EPHB4-associated disorders: novel pathogenic variants and unreported intrafamilial overlapping phenotypes. <i>Genetics in Medicine</i> , 2021, 23, 1315-1324.	2.4	6
2	Development and physiological functions of the lymphatic system: insights from human genetic studies of primary lymphedema. <i>Physiological Reviews</i> , 2021, 101, 1809-1871.	28.8	32
3	Human phenotypes caused by <i>PIEZO1</i> mutations; one gene, two overlapping phenotypes?. <i>Journal of Physiology</i> , 2018, 596, 985-992.	2.9	36
4	A Novel Splice-Site Mutation in VEGFC Is Associated with Congenital Primary Lymphoedema of Gordon. <i>International Journal of Molecular Sciences</i> , 2018, 19, 2259.	4.1	10
5	VIPAR, a quantitative approach to 3D histopathology applied to lymphatic malformations. <i>JCI Insight</i> , 2017, 2, .	5.0	33
6	EPHB4 kinase-inactivating mutations cause autosomal dominant lymphatic-related hydrops fetalis. <i>Journal of Clinical Investigation</i> , 2016, 126, 3080-3088.	8.2	83
7	Novel mutations in PIEZO1 cause an autosomal recessive generalized lymphatic dysplasia with non-immune hydrops fetalis. <i>Nature Communications</i> , 2015, 6, 8085.	12.8	247