

# Loïc Garçon

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2934136/publications.pdf>

Version: 2024-02-01

13  
papers

672  
citations

1163117

8  
h-index

1199594

12  
g-index

13  
all docs

13  
docs citations

13  
times ranked

818  
citing authors

#	ARTICLE	IF	CITATIONS
1	Dehydrated hereditary stomatocytosis linked to gain-of-function mutations in mechanically activated PIEZO1 ion channels. <i>Nature Communications</i> , 2013, 4, 1884.	12.8	282
2	Recommendations regarding splenectomy in hereditary hemolytic anemias. <i>Haematologica</i> , 2017, 102, 1304-1313.	3.5	138
3	Clinical and biological features in <i>PIEZO1</i>-hereditary xerocytosis and Gardos channelopathy: a retrospective series of 126 patients. <i>Haematologica</i> , 2019, 104, 1554-1564.	3.5	76
4	PIEZO1 activation delays erythroid differentiation of normal and hereditary xerocytosis-derived human progenitor cells. <i>Haematologica</i> , 2020, 105, 610-622.	3.5	47
5	Red blood cell Gardos channel (KCNN4): the essential determinant of erythrocyte dehydration in hereditary xerocytosis. <i>Haematologica</i> , 2017, 102, e415-e418.	3.5	42
6	Long-term follow-up of subtotal splenectomy for hereditary spherocytosis: a single-center study. <i>Blood</i> , 2016, 127, 1616-1618.	1.4	31
7	Recent advances in the pathophysiology of <sc>PIEZO1</sc>-related hereditary xerocytosis. <i>American Journal of Hematology</i> , 2021, 96, 1017-1026.	4.1	28
8	Piezo1-xerocytosis red cell metabolome shows impaired glycolysis and increased hemoglobin oxygen affinity. <i>Blood Advances</i> , 2021, 5, 84-88.	5.2	10
9	Multiple thrombosis in a patient with <sc>Gardos</sc> channelopathy and a new <sc><i>KCNN4</i></sc> mutation. <i>American Journal of Hematology</i> , 2021, 96, E318-E321.	4.1	6
10	PIEZO1, sensing the touch during erythropoiesis. <i>Current Opinion in Hematology</i> , 2022, 29, 112-118.	2.5	6
11	Subtotal and total splenectomy for hereditary pyropoikilocytosis: Benefits and outcomes. <i>American Journal of Hematology</i> , 2018, 93, E340-E342.	4.1	4
12	Acquired spherocytosis due to somatic <sc><i>ANK1</i></sc> mutations as a manifestation of clonal hematopoiesis in elderly patients. <i>American Journal of Hematology</i> , 2022, 97, .	4.1	2
13	Hereditary Spherocytosis: Indication of splenectomy and its long-term complications. <i>Hematologie</i> , 2018, 24, 126-133.	0.0	0