

# Mary Risinger

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5227845/publications.pdf>

Version: 2024-02-01

14  
papers

258  
citations

1307594

7  
h-index

1058476

14  
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15  
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15  
docs citations

15  
times ranked

352  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genotype-phenotype correlations in hereditary elliptocytosis and hereditary pyropoikilocytosis. <i>Blood Cells, Molecules, and Diseases</i> , 2016, 61, 4-9.	1.4	60
2	Red cell membrane disorders: structure meets function. <i>Blood</i> , 2020, 136, 1250-1261.	1.4	47
3	Inhibition of Band 3 tyrosine phosphorylation: a new mechanism for treatment of sickle cell disease. <i>British Journal of Haematology</i> , 2020, 190, 599-609.	2.5	46
4	The Spectrum of SPTA1-Associated Hereditary Spherocytosis. <i>Frontiers in Physiology</i> , 2019, 10, 815.	2.8	32
5	Cytokinesis failure in RhoA-deficient mouse erythroblasts involves actomyosin and midbody dysregulation and triggers p53 activation. <i>Blood</i> , 2015, 126, 1473-1482.	1.4	26
6	Rare Hereditary Hemolytic Anemias. <i>Hematology/Oncology Clinics of North America</i> , 2019, 33, 373-392.	2.2	22
7	Hereditary xerocytosis: Diagnostic considerations. <i>American Journal of Hematology</i> , 2018, 93, E67-E69.	4.1	7
8	Peroxiredoxin II (PRDX2) Is a Novel Candidate Gene for Congenital Dyserythropoietic Anemia. <i>Blood</i> , 2018, 132, 3605-3605.	1.4	4
9	Hereditary elliptocytosis-associated alpha-spectrin mutation p.L155dup as a modifier of sickle cell disease severity. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27531.	1.5	2
10	The Spectrum of Alpha-Spectrin Associated Hereditary Spherocytosis. <i>Blood</i> , 2015, 126, 941-941.	1.4	2
11	Clinical Application of Massively Parallel Sequencing in the Diagnosis of Hereditary Hemolytic and Dyserythropoietic Anemias. <i>Blood</i> , 2016, 128, 4746-4746.	1.4	2
12	Evaluation of Phenotype-Genotype Correlation in Two Common PIEZO1 Mutations p.R2456H and p.L2495_E2495dup. <i>Blood</i> , 2018, 132, 1040-1040.	1.4	1
13	<i>VPS4A</i> : A Novel Candidate Gene for Congenital Dyserythropoietic Anemia. <i>Blood</i> , 2017, 130, 923-923.	1.4	1
14	RGL2 Deficiency Impairs Human Erythropoiesis By Altering Terminal Erythroid Differentiation and Apoptosis. <i>Blood</i> , 2017, 130, 8-8.	1.4	0