

# Judy Geissler

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3662184/publications.pdf>

Version: 2024-02-01

12  
papers

238  
citations

1163117

8  
h-index

1281871

11  
g-index

12  
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12  
docs citations

12  
times ranked

599  
citing authors

#	ARTICLE	IF	CITATIONS
1	Primary Immunodeficiency Caused by an Exonized Retroposed Gene Copy Inserted in the <i>CYBB</i> Gene. <i>Human Mutation</i> , 2014, 35, 486-496.	2.5	38
2	Factor H-Related (FHR)-1 and FHR-2 Form Homo- and Heterodimers, while FHR-5 Circulates Only As Homodimer in Human Plasma. <i>Frontiers in Immunology</i> , 2017, 8, 1328.	4.8	38
3	Neutrophil responsiveness to IgG, as determined by fixed ratios of mRNA levels for activating and inhibitory Fc $\gamma$ RII (CD32), is stable over time and unaffected by cytokines. <i>Blood</i> , 2006, 108, 584-590.	1.4	35
4	Complement Factor H-Related Protein 3 Serum Levels Are Low Compared to Factor H and Mainly Determined by Gene Copy Number Variation in CFHR3. <i>PLoS ONE</i> , 2016, 11, e0152164.	2.5	30
5	MKL1 deficiency results in a severe neutrophil motility defect due to impaired actin polymerization. <i>Blood</i> , 2020, 135, 2171-2181.	1.4	29
6	Genetic variation of human neutrophil Fc $\gamma$ receptors and SIRP $\alpha$ in antibody-dependent cellular cytotoxicity towards cancer cells. <i>European Journal of Immunology</i> , 2018, 48, 344-354.	2.9	28
7	Hermansky-Pudlak syndrome type 2: Aberrant pre-mRNA splicing and mislocalization of granule proteins in neutrophils. <i>Human Mutation</i> , 2017, 38, 1402-1411.	2.5	21
8	Mutation in an exonic splicing enhancer site causing chronic granulomatous disease. <i>Blood Cells, Molecules, and Diseases</i> , 2017, 66, 50-57.	1.4	13
9	Generation and characterization of a control and patient-derived human iPSC line containing the Hermansky Pudlak type 2 (HPS2) associated heterozygous compound mutation in AP3B1. <i>Stem Cell Research</i> , 2021, 54, 102444.	0.7	3
10	Generation and characterization of a human iPSC line SANi007-A from a patient with a heterozygous dominant mutation in ELANE. <i>Stem Cell Research</i> , 2021, 55, 102440.	0.7	1
11	Generation and characterization of a human iPSC line SANi008-A from a Ch $\alpha$ diak-Higashi Syndrome patient. <i>Stem Cell Research</i> , 2021, 55, 102442.	0.7	1
12	Generation and characterization of a human iPSC line SANi006-A from a Gray Platelet Syndrome patient. <i>Stem Cell Research</i> , 2021, 55, 102443.	0.7	1