

# Raquel Ruiz-García

## List of Publications by Year in descending order

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Version: 2024-02-01

26  
papers

807  
citations

516710

16  
h-index

642732

23  
g-index

27  
all docs

27  
docs citations

27  
times ranked

1127  
citing authors

#	ARTICLE	IF	CITATIONS
1	Associations of paediatric demyelinating and encephalitic syndromes with myelin oligodendrocyte glycoprotein antibodies: a multicentre observational study. <i>Lancet Neurology</i> , The, 2020, 19, 234-246.	10.2	207
2	Isolated IgA Anti- $\alpha$ 2 Glycoprotein I Antibodies in Patients with Clinical Criteria for Antiphospholipid Syndrome. <i>Journal of Immunology Research</i> , 2014, 2014, 1-8.	2.2	68
3	The Diagnostic Value of Onconeural Antibodies Depends on How They Are Tested. <i>Frontiers in Immunology</i> , 2020, 11, 1482.	4.8	60
4	Limitations of a Commercial Assay as Diagnostic Test of Autoimmune Encephalitis. <i>Frontiers in Immunology</i> , 2021, 12, 691536.	4.8	46
5	Late-onset neuromyelitis optica spectrum disorder. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2019, 6, .	6.0	44
6	Paraneoplastic cerebellar ataxia and antibodies to metabotropic glutamate receptor 2. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020, 7, .	6.0	39
7	Mutations in PI3K110 $\beta$ cause impaired natural killer cell function partially rescued by rapamycin treatment. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 605-617.e7.	2.9	36
8	Seizure-related 6 homolog like 2 autoimmunity. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	6.0	36
9	Primary Immune Regulatory Disorders With an Autoimmune Lymphoproliferative Syndrome-Like Phenotype: Immunologic Evaluation, Early Diagnosis and Management. <i>Frontiers in Immunology</i> , 2021, 12, 671755.	4.8	35
10	Harmful Effect of Preformed Anti-MICA Antibodies on Renal Allograft Evolution in Early Posttransplantation Period. <i>Transplantation</i> , 2013, 96, 70-78.	1.0	28
11	Thymoma and Autoimmune Encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2021, 8, .	6.0	28
12	Caveats and Pitfalls of SOX1 Autoantibody Testing With a Commercial Line Blot Assay in Paraneoplastic Neurological Investigations. <i>Frontiers in Immunology</i> , 2019, 10, 769.	4.8	26
13	Autoimmune lymphoproliferative syndrome due to somatic FAS mutation (ALPS-sFAS) combined with a germline caspase-10 (CASP10) variation. <i>Immunobiology</i> , 2016, 221, 40-47.	1.9	25
14	Decreased activation-induced cell death by EBV-transformed B-cells from a patient with autoimmune lymphoproliferative syndrome caused by a novel FASLG mutation. <i>Pediatric Research</i> , 2015, 78, 603-608.	2.3	21
15	Acquired Senescent T-Cell Phenotype Correlates with Clinical Severity in GATA Binding Protein 2-Deficient Patients. <i>Frontiers in Immunology</i> , 2017, 8, 802.	4.8	18
16	Toll-like receptor 3 deficiency in autoimmune encephalitis post-herpes simplex encephalitis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2019, 6, e611.	6.0	18
17	Frequency and relevance of IgM, and IgA antibodies against MOG in MOG-IgG-associated disease. <i>Multiple Sclerosis and Related Disorders</i> , 2019, 28, 230-234.	2.0	18
18	Acquired and Innate Immunity Impairment and Severe Disseminated Mycobacterium genavense Infection in a Patient With a NF- $\kappa$ B1 Deficiency. <i>Frontiers in Immunology</i> , 2018, 9, 3148.	4.8	14

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19	A case of partial dedicator of cytokinesis 8 deficiency with altered effector phenotype and impaired CD8+ and natural killer cell cytotoxicity. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 134, 218-221.e7.	2.9	12
20	Extreme Phenotypes With Identical Mutations: Two Patients With Same Non-sense NHEJ1 Homozygous Mutation. <i>Frontiers in Immunology</i> , 2019, 9, 2959.	4.8	12
21	Fatal <i>Pneumocystis jirovecii</i> and Cytomegalovirus Infections in an Infant With Normal TRECs Count. <i>Pediatric Infectious Disease Journal</i> , 2019, 38, 157-160.	2.0	10
22	Autoimmune Septin-5 Disease Presenting as Spinocerebellar Ataxia and Nystagmus. <i>Neurology</i> , 2021, 97, 291-292.	1.1	5
23	Homozygous R136S mutation in PRNP gene causes inherited early onset prion disease. <i>Alzheimer's Research and Therapy</i> , 2021, 13, 176.	6.2	1
24	Allodynia, rigidity and gait disturbance. <i>Practical Neurology</i> , 2020, , practneurol-2020-002669.	1.1	0
25	Autoimmune Lymphoproliferative Syndrome in Children with Nonmalignant Organomegaly, Chronic Immune Cytopenia, and Newly Diagnosed Lymphoma. <i>Turkish Journal of Haematology</i> , 2021, 38, 145-150.	0.5	0
26	Pseudoathetosis and pseudodystonia in sensory ganglionopathy due to anti- $\epsilon$ -Amphiphysin and anti- $\epsilon$ -Hu antibodies: a rare presentation underlying a neoplastic etiology. <i>Movement Disorders Clinical Practice</i> , 0, , .	1.5	0