Sophie Georgin-Lavialle

List of Publications by Year
in descending order

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Version: 2024-02-01

78	1,181	430874	501196
papers	citations	18 h-index	28 g-index

113	113	113	1198
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Correspondence on â€œSafety of vaccination against SARS-CoV-2 in people with rheumatic and musculoskeletal diseases: results from the EULAR Coronavirus Vaccine (COVAX) physician-reported registryâ€•by Machadoet al. Annals of the Rheumatic Diseases, 2023, 82, e228-e228.	0.9	0
2	Is neutrophilic dermatosis a manifestation of familial Mediterranean fever?. Scandinavian Journal of Rheumatology, 2022, 51, 42-49.	1.1	1
3	Azacitidine for patients with Vacuoles, E1 Enzyme, Xâ€linked, Autoinflammatory, Somatic syndrome (VEXAS) and myelodysplastic syndrome: data from the French VEXAS registry. British Journal of Haematology, 2022, 196, 969-974.	2.5	85
4	Tattooing and autoinflammatory diseases: a study among 197 French patients from the JIR cohort. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	1
5	UBA1 gene mutation in giant cell arteritis. Clinical Rheumatology, 2022, 41, 1257-1259.	2.2	6
6	Amyloidosis from the patient perspective: the French daily impact of amyloidosis study. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 165-174.	3.0	17
7	AA amyloidosis complicating cryopyrin-associated periodic syndrome: a study of 86 cases including 23 French patients and systematic review. Rheumatology, 2022, 61, 4827-4834.	1.9	8
8	Neurological manifestations in mevalonate kinase deficiency: A systematic review. Molecular Genetics and Metabolism, 2022, 136, 85-93.	1.1	7
9	Health Outcomes of 215 Mothers of Children With Autoimmune Congenital Heart Block: Analysis of the French Neonatal Lupus Syndrome Registry. Journal of Rheumatology, 2022, 49, 1124-1130.	2.0	3
10	On the Determinants of IDO Activity in Patients With Familial Mediterranean Fever. Modern Rheumatology, 2022, , .	1.8	0
11	COVID-19 infection among patients with autoinflammatory diseases: a study on 117 French patients compared with 1545 from the French RMD COVID-19 cohort: COVIMAI â€“ the French cohort study of SARS-CoV-2 infection in patient with systemic autoinflammatory diseases. RMD Open, 2022, 8, e002063.	3.8	7
12	Clinical course of COVID-19 in a cohort of 342 familial Mediterranean fever patients with a long-term treatment by colchicine in a French endemic area. Annals of the Rheumatic Diseases, 2021, 80, 539-540.	0.9	28
13	AA Amyloidosis Secondary to Primary Immune Deficiency: About 40 Cases Including 2 New French Cases and a Systematic Literature Review. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 745-752.e1.	3.8	3
14	Fast diagnostic test for familial Mediterranean fever based on a kinase inhibitor. Annals of the Rheumatic Diseases, 2021, 80, 128-132.	0.9	16
15	Prescription of interleukin-1 inhibitors in a French adult cohort of familial Mediterranean fever. European Journal of Internal Medicine, 2021, 84, 109-111.	2.2	2
16	Infections and AA amyloidosis: An overview. International Journal of Clinical Practice, 2021, 75, e13966.	1.7	12
17	LACC1 deficiency links juvenile arthritis with autophagy and metabolism in macrophages. Journal of Experimental Medicine, 2021, 218, .	8.5	17
18	â€œHelicobacter pylori in familial mediterranean fever: A series of 120 patients from literature and from franceâ€• Helicobacter, 2021, 26, e12789.	3.5	1

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19	Remdesivir for COVID-19 in Europe: will it provide value for money?. Lancet Respiratory Medicine,the, 2021, 9, 127-128.	10.7	14
20	Tumour necrosis factor receptor-1 associated periodic syndrome (TRAPS)-related AA amyloidosis: a national case series and systematic review. Rheumatology, 2021, 60, 5775-5784.	1.9	11
21	Amyloid Goiter in Familial Mediterranean Fever: Description of 42 Cases from a French Cohort and from Literature Review. Journal of Clinical Medicine, 2021, 10, 1983.	2.4	4
22	Association between familial Mediterranean fever and multiple sclerosis: A case series from the JIR cohort and systematic literature review. Multiple Sclerosis and Related Disorders, 2021, 50, 102834.	2.0	6
23	Daily multidisciplinary COVID-19 meeting: Experiences from a French university hospital. Respiratory Medicine and Research, 2021, 79, 100828.	0.6	0
24	Chronic hepatic involvement in the clinical spectrum of A20 haploinsufficiency. Liver International, 2021, 41, 1894-1900.	3.9	9
25	Abnormal electrochemical skin conductance values in patients with AA amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, , 1-2.	3.0	0
26	AA Amyloidosis in the Course of HIV Infection: A Report of 19 Cases Including 4 New French Cases and a Comprehensive Review of Literature. Nephron, 2021, 145, 675-683.	1.8	1
27	Thyroid disorders in familial Mediterranean fever: think about AA amyloidosis!. Clinical Rheumatology, 2021, 40, 3381-3382.	2.2	0
28	AA amyloidosis complicating monoclonal gammopathies, an unusual feature validating the concept of â€œmonoclonal gammopathy of inflammatory significanceâ€?. International Journal of Clinical Practice, 2021, 75, e14817.	1.7	6
29	Neutrophil-specific gain-of-function mutations in Nlrp3 promote development of cryopyrin-associated periodic syndrome. Journal of Experimental Medicine, 2021, 218, .	8.5	29
30	UBA1 Variations in Neutrophilic Dermatosis Skin Lesions of Patients With VEXAS Syndrome. JAMA Dermatology, 2021, 157, 1349.	4.1	71
31	DADA2 diagnosed in adulthood versus childhood: A comparative study on 306 patients including a systematic literature review and 12 French cases. Seminars in Arthritis and Rheumatism, 2021, 51, 1170-1179.	3.4	14
32	Looking beyond VEXAS: Coexistence of undifferentiated systemic autoinflammatory disease and myelodysplastic syndrome. Seminars in Hematology, 2021, 58, 247-253.	3.4	9
33	Could we measure hair colchicine to assess colchicine observance in familial Mediterranean fever?. Rheumatology, 2021, 60, 1563-1564.	1.9	2
34	AA amyloidosis of unknown origin in New-Caledonia with focus on the association with gout: a consecutive case series of 20 patients. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, , 1-2.	3.0	0
35	Real-Life Indications of Interleukin-1 Blocking Agents in Hereditary Recurrent Fevers: Data From the JIRcohort and a Literature Review. Frontiers in Immunology, 2021, 12, 744780.	4.8	2
36	AA amyloidosis secondary to adult onset Still's disease: About 19 cases. Seminars in Arthritis and Rheumatism, 2020, 50, 156-165.	3.4	9

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37	Somatic Mosaic NLRP3 Mutations and Inflammasome Activation in Late-Onset Chronic Urticaria. Journal of Investigative Dermatology, 2020, 140, 791-798.e2.	0.7	19
38	NLRP3-associated autoinflammatory diseases: Phenotypic and molecular characteristics of germline versus somatic mutations. Journal of Allergy and Clinical Immunology, 2020, 145, 1254-1261.	2.9	50
39	Annals Graphic Medicine - Facing Transitional Care in Still Disease. Annals of Internal Medicine, 2020, 172, W95.	3.9	0
40	Systemic autoinflammatory diseases: Clinical state of the art. Best Practice and Research in Clinical Rheumatology, 2020, 34, 101529.	3.3	34
41	AA amyloidosis associated with Fabry disease. International Journal of Clinical Practice, 2020, 74, e13577.	1.7	1
42	The Use of Interleukine-1 Inhibitors in Familial Mediterranean Fever Patients: A Narrative Review. Frontiers in Immunology, 2020, 11, 971.	4.8	31
43	Nonâ€amyloid liver involvement in familial Mediterranean fever: A systematic literature review. Liver International, 2020, 40, 1269-1277.	3.9	6
44	The association of psychiatric comorbidities with emergency visits and hospitalisations in adult sickleâ€cell patients: a cohort study. British Journal of Haematology, 2020, 189, e21-e23.	2.5	1
45	Clinical and pathological dermatological features of deficiency of adenosine deaminase 2: A multicenter, retrospective, observational study. Journal of the American Academy of Dermatology, 2020, 83, 1794-1798.	1.2	13
46	AA amyloidosis revealing mevalonate kinase deficiency: A report of 20 cases including two new French cases and a comprehensive review of literature. Seminars in Arthritis and Rheumatism, 2020, 50, 1370-1373.	3.4	13
47	Long-Term Follow-Up and Optimization of Interleukin-1 Inhibitors in the Management of Monogenic Autoinflammatory Diseases: Real-Life Data from the JIR Cohort. Frontiers in Pharmacology, 2020, 11, 568865.	3.5	7
48	Cause of death and risk factors for mortality in AA amyloidosis: A French retrospective study. European Journal of Internal Medicine, 2020, 82, 130-132.	2.2	4
49	Î²-Glucanâ€“induced reprogramming of human macrophages inhibits NLRP3 inflammasome activation in cryopyrinopathies. Journal of Clinical Investigation, 2020, 130, 4561-4573.	8.2	44
50	Attacks of TNF-receptor associated periodic syndrome are associated with higher inflammatory markers than familial Mediterranean fever. Clinical and Experimental Rheumatology, 2020, 38 Suppl 127, 131.	0.8	0
51	Rheumatoid arthritis revealed by polyadenopathy, diarrhea and digestive AA amyloidosis. Joint Bone Spine, 2019, 86, 397-398.	1.6	1
52	Epidemiology of Castleman disease associated with AA amyloidosis: description of 2 new cases and literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 197-202.	3.0	7
53	Specific changes in faecal microbiota are associated with familial Mediterranean fever. Annals of the Rheumatic Diseases, 2019, 78, 1398-1404.	0.9	18
54	Neuroinflammatory disorders and mastocytosis: A possible association?. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2878-2881.e1.	3.8	2

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55	Limites nosologiques entre fiÃ¨vre mÃ©diterranÃ©enne familiale et spondylarthrite juvÃ©nileÂ: analyse de trois cohortes rÃ©trospectives franÃ§aises. Revue Du Rhumatisme (Edition Francaise), 2019, 86, 504-510.	0.0	0
56	Positive Impact of Expert Reference Center Validation on Performance of Next-Generation Sequencing for Genetic Diagnosis of Autoinflammatory Diseases. Journal of Clinical Medicine, 2019, 8, 1729.	2.4	9
57	Autoinflammatory diseases: State of the art. Presse Medicale, 2019, 48, e25-e48.	1.9	44
58	Autoinflammation secondaire Ã des dÃ©fauts dâ€™ubiquitination dans la voie NFKBÂ: haploinsuffisance de A20 (HA20) et dÃ©ficit en Otuline (OtulinopÃ©nie). Revue Du Rhumatisme (Edition Francaise), 2019, 86, 358-366.	0.0	0
59	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. Journal of Rheumatology, 2019, 46, 1535-1539.	2.0	6
60	Plasma histamine elevation in a large cohort of sickle cell disease patients. British Journal of Haematology, 2019, 186, 125-129.	2.5	7
61	Commentary to â€œA 44â€yearâ€old female with familial Mediterranean fever, cardiomyopathy and end stage renal diseaseâ€•by Magaki et al.. Brain Pathology, 2019, 29, 311-311.	4.1	1
62	Do we need the PFAPA syndrome in adults with non-monogenic periodic fevers?. Annals of the Rheumatic Diseases, 2019, , annrheumdis-2019-216827.	0.9	3
63	Boundaries between familial Mediterranean fever and juvenile spondyloarthritis: Analysis of three French retrospective cohorts. Joint Bone Spine, 2018, 85, 733-739.	1.6	11
64	A decision tree for the genetic diagnosis of deficiency of adenosine deaminase 2 (DADA2): a French reference centres experience. European Journal of Human Genetics, 2018, 26, 960-971.	2.8	65
65	Efficacy of Continuous Interleukin 1 Blockade in Mevalonate Kinase Deficiency: A Multicenter Retrospective Study in 13 Adult Patients and Literature Review. Journal of Rheumatology, 2018, 45, 425-429.	2.0	23
66	Validation of the Fautrel classification criteria for adult-onset Stillâ€™s disease. Seminars in Arthritis and Rheumatism, 2018, 47, 578-585.	3.4	47
67	Building a transitional care checklist in rheumatology: A Delphi-like survey. Joint Bone Spine, 2018, 85, 435-440.	1.6	24
68	Minor salivary gland biopsy is more effective than normal appearing skin biopsy for amyloid detection in systemic amyloidosis: A prospective monocentric study. European Journal of Internal Medicine, 2018, 57, e20-e21.	2.2	12
69	Association of hidradenitis suppurativa and familial Mediterranean fever: A case series of 6 patients. Joint Bone Spine, 2017, 84, 159-162.	1.6	28
70	Isolated Pericardial Infiltration Without Myocardial Involvement inÂLight-Chainâ€“Related Amyloidosis. Annals of Thoracic Surgery, 2017, 103, e255-e257.	1.3	1
71	AA amyloidosis is an emerging cause of nephropathy in obese patients. European Journal of Internal Medicine, 2017, 39, e18-e20.	2.2	11
72	A survey of resistance to colchicine treatment for French patients with familial Mediterranean fever. Orphanet Journal of Rare Diseases, 2017, 12, 54.	2.7	32

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73	Spondyloarthritis associated with familial Mediterranean fever: successful treatment with anakinra. Rheumatology, 2017, 56, 167-169.	1.9	12
74	THU0567â€…Comparative Performance of Fautrel and Yamaguchi Criteria for The Classification of Patients with Adult Onset Still Disease: Preliminary Results:. Annals of the Rheumatic Diseases, 2016, 75, 397.1-397.	0.9	0
75	Decreased tryptophan and increased kynurenine levels in mastocytosis associated with digestive symptoms. Allergy: European Journal of Allergy and Clinical Immunology, 2016, 71, 416-420.	5.7	4
76	Evaluating the Impact of Computerized Provider Order Entry on Medical Students Training at Bedside: A Randomized Controlled Trial. PLoS ONE, 2015, 10, e0138094.	2.5	7
77	Complete Remission of Schnitzler Syndrome and WaldenstrÃ¶m Macroglobulinemia under Rituximab-Cyclophosphamide-Dexamethasone. Dermatology, 2015, 230, 18-22.	2.1	13
78	Assessment and effective targeting of Interleukin-1 in multicentric reticulohistyocytosis. Joint Bone Spine, 2015, 82, 280-283.	1.6	19