Sophie Georgin-Lavialle

List of Publications by Year
in descending order

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	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
2	UBA1 Variations in Neutrophilic Dermatosis Skin Lesions of Patients With VEXAS Syndrome. JAMA Dermatology, 2021, 157, 1349.	4.1	71
3	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
4	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
5	Validation of the Fautrel classification criteria for adult-onset Stillâ€™s disease. Seminars in Arthritis and Rheumatism, 2018, 47, 578-585.	3.4	47
6	Autoinflammatory diseases: State of the art. Presse Medicale, 2019, 48, e25-e48.	1.9	44
7	Î²-Glucanâ€“induced reprogramming of human macrophages inhibits NLRP3 inflammasome activation in cryopyrinopathies. Journal of Clinical Investigation, 2020, 130, 4561-4573.	8.2	44
8	Systemic autoinflammatory diseases: Clinical state of the art. Best Practice and Research in Clinical Rheumatology, 2020, 34, 101529.	3.3	34
9	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
10	The Use of Interleukine-1 Inhibitors in Familial Mediterranean Fever Patients: A Narrative Review. Frontiers in Immunology, 2020, 11, 971.	4.8	31
11	Neutrophil-specific gain-of-function mutations in Nlrp3 promote development of cryopyrin-associated periodic syndrome. Journal of Experimental Medicine, 2021, 218, .	8.5	29
12	Association of hidradenitis suppurativa and familial Mediterranean fever: A case series of 6 patients. Joint Bone Spine, 2017, 84, 159-162.	1.6	28
13	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
14	Building a transitional care checklist in rheumatology: A Delphi-like survey. Joint Bone Spine, 2018, 85, 435-440.	1.6	24
15	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
16	Assessment and effective targeting of Interleukin-1 in multicentric reticulohistyocytosis. Joint Bone Spine, 2015, 82, 280-283.	1.6	19
17	Somatic Mosaic NLRP3 Mutations and Inflammasome Activation in Late-Onset Chronic Urticaria. Journal of Investigative Dermatology, 2020, 140, 791-798.e2.	0.7	19
18	Specific changes in faecal microbiota are associated with familial Mediterranean fever. Annals of the Rheumatic Diseases, 2019, 78, 1398-1404.	0.9	18

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19	LACC1 deficiency links juvenile arthritis with autophagy and metabolism in macrophages. Journal of Experimental Medicine, 2021, 218, .	8.5	17
20	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
21	Fast diagnostic test for familial Mediterranean fever based on a kinase inhibitor. Annals of the Rheumatic Diseases, 2021, 80, 128-132.	0.9	16
22	Remdesivir for COVID-19 in Europe: will it provide value for money?. Lancet Respiratory Medicine,the, 2021, 9, 127-128.	10.7	14
23	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
24	Complete Remission of Schnitzler Syndrome and WaldenstrÃ¶m Macroglobulinemia under Rituximab-Cyclophosphamide-Dexamethasone. Dermatology, 2015, 230, 18-22.	2.1	13
25	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
26	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
27	Spondyloarthritis associated with familial Mediterranean fever: successful treatment with anakinra. Rheumatology, 2017, 56, 167-169.	1.9	12
28	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
29	Infections and AA amyloidosis: An overview. International Journal of Clinical Practice, 2021, 75, e13966.	1.7	12
30	AA amyloidosis is an emerging cause of nephropathy in obese patients. European Journal of Internal Medicine, 2017, 39, e18-e20.	2.2	11
31	Boundaries between familial Mediterranean fever and juvenile spondyloarthritis: Analysis of three French retrospective cohorts. Joint Bone Spine, 2018, 85, 733-739.	1.6	11
32	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
33	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
34	AA amyloidosis secondary to adult onset Still's disease: About 19 cases. Seminars in Arthritis and Rheumatism, 2020, 50, 156-165.	3.4	9
35	Chronic hepatic involvement in the clinical spectrum of A20 haploinsufficiency. Liver International, 2021, 41, 1894-1900.	3.9	9
36	Looking beyond VEXAS: Coexistence of undifferentiated systemic autoinflammatory disease and myelodysplastic syndrome. Seminars in Hematology, 2021, 58, 247-253.	3.4	9

#	Article	IF	Citations
37	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
38	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
39	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
40	Plasma histamine elevation in a large cohort of sickle cell disease patients. British Journal of Haematology, 2019, 186, 125-129.	2.5	7
41	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
42	Neurological manifestations in mevalonate kinase deficiency: A systematic review. Molecular Genetics and Metabolism, 2022, 136, 85-93.	1.1	7
43	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
44	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. Journal of Rheumatology, 2019, 46, 1535-1539.	2.0	6
45	Nonâ€amyloid liver involvement in familial Mediterranean fever: A systematic literature review. Liver International, 2020, 40, 1269-1277.	3.9	6
46	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
47	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
48	UBA1 gene mutation in giant cell arteritis. Clinical Rheumatology, 2022, 41, 1257-1259.	2.2	6
49	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
50	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
51	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
52	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
53	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
54	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

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55	Neuroinflammatory disorders and mastocytosis: A possible association?. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2878-2881.e1.	3.8	2
56	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
57	Could we measure hair colchicine to assess colchicine observance in familial Mediterranean fever?. Rheumatology, 2021, 60, 1563-1564.	1.9	2
58	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
59	Isolated Pericardial Infiltration Without Myocardial Involvement inÂLight-Chainâ€“Related Amyloidosis. Annals of Thoracic Surgery, 2017, 103, e255-e257.	1.3	1
60	Rheumatoid arthritis revealed by polyadenopathy, diarrhea and digestive AA amyloidosis. Joint Bone Spine, 2019, 86, 397-398.	1.6	1
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	AA amyloidosis associated with Fabry disease. International Journal of Clinical Practice, 2020, 74, e13577.	1.7	1
63	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
64	â€œHelicobacter pylori in familial mediterranean fever: A series of 120 patients from literature and from franceâ€• Helicobacter, 2021, 26, e12789.	3.5	1
65	Is neutrophilic dermatosis a manifestation of familial Mediterranean fever?. Scandinavian Journal of Rheumatology, 2022, 51, 42-49.	1.1	1
66	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
67	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
68	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
69	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
70	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
71	Annals Graphic Medicine - Facing Transitional Care in Still Disease. Annals of Internal Medicine, 2020, 172, W95.	3.9	0
72	Daily multidisciplinary COVID-19 meeting: Experiences from a French university hospital. Respiratory Medicine and Research, 2021, 79, 100828.	0.6	0

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73	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
74	Thyroid disorders in familial Mediterranean fever: think about AA amyloidosis!. Clinical Rheumatology, 2021, 40, 3381-3382.	2.2	0
75	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
76	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
77	On the Determinants of IDO Activity in Patients With Familial Mediterranean Fever. Modern Rheumatology, 2022, , .	1.8	0
78	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