Gilles Grateau

List of Publications by Year
in descending order

160	8,733	61857 43	46693 89
papers	citations	h-index	g-index

267	267	267	6984
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A candidate gene for familial Mediterranean fever. Nature Genetics, 1997, 17, 25-31.	9.4	1,402
2	Mutations in the gene encoding mevalonate kinase cause hyper-IgD and periodic fever syndrome. Nature Genetics, 1999, 22, 178-181.	9.4	511
3	Mutations in NALP12 cause hereditary periodic fever syndromes. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 1614-1619.	3.3	331
4	New Mutations of CIAS1 That Are Responsible for Muckle-Wells Syndrome and Familial Cold Urticaria: A Novel Mutation Underlies Both Syndromes. American Journal of Human Genetics, 2002, 70, 1498-1506.	2.6	313
5	Recommendations for the management of autoinflammatory diseases. Annals of the Rheumatic Diseases, 2015, 74, 1636-1644.	0.5	239
6	MEFV-Gene Analysis in Armenian Patients with Familial Mediterranean Fever: Diagnostic Value and Unfavorable Renal Prognosis of the M694V Homozygous Genotypeâ€"Genetic and Therapeutic Implications. American Journal of Human Genetics, 1999, 65, 88-97.	2.6	238
7	Online Registry for Mutations in Hereditary Amyloidosis Including Nomenclature Recommendations. Human Mutation, 2014, 35, E2403-E2412.	1.1	220
8	COVID-19 outcomes in patients with inflammatory rheumatic and musculoskeletal diseases treated with rituximab: a cohort study. Lancet Rheumatology, The, 2021, 3, e419-e426.	2.2	211
9	Identification ofMEFVâ€Independent Modifying Genetic Factors for Familial Mediterranean Fever. American Journal of Human Genetics, 2000, 67, 1136-1143.	2.6	203
10	Mevalonate Kinase Deficiency: A Survey of 50 Patients. Pediatrics, 2011, 128, e152-e159.	1.0	195
11	The enlarging clinical, genetic, and population spectrum of tumor necrosis factor receptor-associated periodic syndrome. Arthritis and Rheumatism, 2002, 46, 2181-2188.	6.7	188
12	Molecular analysis of MVK mutations and enzymatic activity in hyper-IgD and periodic fever syndrome. European Journal of Human Genetics, 2001, 9, 260-266.	1.4	182
13	Clinical significance of P46L and R92Q substitutions in the tumour necrosis factor superfamily 1A gene. Annals of the Rheumatic Diseases, 2006, 65, 1158-1162.	0.5	161
14	Evidence-based recommendations for genetic diagnosis of familial Mediterranean fever. Annals of the Rheumatic Diseases, 2015, 74, 635-641.	0.5	145
15	Impact of human monocyte and macrophage polarization on NLR expression and NLRP3 inflammasome activation. PLoS ONE, 2017, 12, e0175336.	1.1	136
16	Evidence-based recommendations for the practical management of Familial Mediterranean Fever. Seminars in Arthritis and Rheumatism, 2013, 43, 387-391.	1.6	131
17	Glomerular Hyperfiltration in Adult Sickle Cell Anemia. Clinical Journal of the American Society of Nephrology: CJASN, 2010, 5, 756-761.	2.2	130
18	Hemoglobin sickle cell disease complications: a clinical study of 179 cases. Haematologica, 2012, 97, 1136-1141.	1.7	120

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19	Genetic Linkage of the Muckle-Wells Syndrome to Chromosome 1q44. American Journal of Human Genetics, 1999, 65, 1054-1059.	2.6	118
20	Hereditary Amyloid Cardiomyopathy Caused by a Variant Apolipoprotein A1. American Journal of Pathology, 1999, 154, 221-227.	1.9	114
21	Hereditary renal amyloidosis caused by a new variant lysozyme W64R in a French family. Kidney International, 2002, 61, 907-912.	2.6	108
22	In vivo T cell preactivation in chronic uremic hemodialyzed and non-hemodialyzed patients. Kidney International, 1989, 36, 636-644.	2.6	95
23	Dramatic beneficial effect of interleukin-1 inhibitor treatment in patients with familial Mediterranean fever complicated with amyloidosis and renal failure. Nephrology Dialysis Transplantation, 2012, 27, 1898-1901.	0.4	95
24	Identification and functional consequences of a recurrent NLRP12 missense mutation in periodic fever syndromes. Arthritis and Rheumatism, 2011, 63, 1459-1464.	6.7	91
25	Clinical and histological characteristics of renal AA amyloidosis: a retrospective study of 68 cases with a special interest to amyloid-associated inflammatory response. Human Pathology, 2007, 38, 1798-1809.	1.1	90
26	Role of interleukin-1Î² in NLRP12-associated autoinflammatory disorders and resistance to anti-interleukin-1 therapy. Arthritis and Rheumatism, 2011, 63, 2142-2148.	6.7	88
27	Localized amyloidosis: A survey of 35 French cases. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2005, 12, 239-245.	1.4	73
28	Renal amyloidosis with a frame shift mutation in fibrinogen aalpha-chain gene producing a novel amyloid protein. Blood, 1997, 90, 4799-805.	0.6	70
29	Familial Mediterranean fever in Lebanon: mutation spectrum, evidence for cases in Maronites, Greek orthodoxes, Greek catholics, Syriacs and Chiites and for an association between amyloidosis and M694V and M694I mutations. European Journal of Human Genetics, 2001, 9, 51-55.	1.4	68
30	Schnitzler syndrome: validation and applicability of diagnostic criteria in real-life patients. Allergy: European Journal of Allergy and Clinical Immunology, 2017, 72, 177-182.	2.7	66
31	Diagnostic tools for amyloidosis. Joint Bone Spine, 2002, 69, 538-545.	0.8	63
32	Clinical versus genetic diagnosis of familial Mediterranean fever. QJM - Monthly Journal of the Association of Physicians, 2000, 93, 223-229.	0.2	62
33	Auto inflammatory syndromes: Diagnosis and treatment. Joint Bone Spine, 2007, 74, 544-550.	0.8	62
34	How should we approach classification of autoinflammatory diseases?. Nature Reviews Rheumatology, 2013, 9, 624-629.	3.5	59
35	A new mutation (A546T) of the Î²ig-h3 gene responsible for a French lattice corneal dystrophy type IIIA. American Journal of Ophthalmology, 2000, 129, 248-251.	1.7	58
36	MEFV analysis is of particularly weak diagnostic value for recurrent fevers in Western European Caucasian patients. Arthritis and Rheumatism, 2005, 52, 3603-3605.	6.7	56

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37	The Risk of Familial Mediterranean Fever in MEFV Heterozygotes: A Statistical Approach. PLoS ONE, 2013, 8, e68431.	1.1	56
38	Haemophagocytic syndrome in patients infected with the human immunodeficiency virus: Nine cases and a review. Journal of Infection, 1997, 34, 219-225.	1.7	50
39	NLRP3-associated autoinflammatory diseases: Phenotypic and molecular characteristics of germline versus somatic mutations. Journal of Allergy and Clinical Immunology, 2020, 145, 1254-1261.	1.5	50
40	The relation between familial Mediterranean fever and amyloidosis. Current Opinion in Rheumatology, 2000, 12, 61-64.	2.0	49
41	Longest Form of CCTG Microsatellite Repeat in the Promoter of the CD2BP1/PSTPIP1 Gene Is Associated with Aseptic Abscesses and with Crohn Disease in French Patients. Digestive Diseases and Sciences, 2010, 55, 1681-1688.	1.1	47
42	Osteo-articular manifestations of amyloidosis. Best Practice and Research in Clinical Rheumatology, 2012, 26, 459-475.	1.4	47
43	Validation of the Fautrel classification criteria for adult-onset Stillâ€™s disease. Seminars in Arthritis and Rheumatism, 2018, 47, 578-585.	1.6	47
44	A Clinical Criterion to Exclude the Hyperimmunoglobulin D Syndrome (Mild Mevalonate Kinase) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 46	1.0	45
45	Autoinflammatory diseases: State of the art. Presse Medicale, 2019, 48, e25-e48.	0.8	44
46	Î²-Glucanâ€“induced reprogramming of human macrophages inhibits NLRP3 inflammasome activation in cryopyrinopathies. Journal of Clinical Investigation, 2020, 130, 4561-4573.	3.9	44
47	Prevalence and distribution of MEFV mutations among Arabs from the Maghreb patients suffering from familial Mediterranean fever. Comptes Rendus - Biologies, 2006, 329, 71-74.	0.1	43
48	Specific increase inÂcaspase-1 activity andÂsecretion ofÂIL-1 family cytokines: aÂputative link between mevalonate kinase deficiency andÂinflammation. European Cytokine Network, 2009, 20, 101-107.	1.1	43
49	CIAS1 Mutation in a Patient with Overlap between Muckle-Wells and Chronic Infantile Neurological Cutaneous and Articular Syndromes. Dermatology, 2003, 206, 257-259.	0.9	42
50	Familial Mediterranean Fever in Heterozygotes: Are We Able to Accurately Diagnose the Disease in Very Young Children?. Arthritis and Rheumatism, 2013, 65, 1654-1662.	6.7	40
51	Expression of SAA1, SAA2 and SAA4 genes in human primary monocytes and monocyte-derived macrophages. PLoS ONE, 2019, 14, e0217005.	1.1	40
52	Association of Vasculitis and Familial Mediterranean Fever. Frontiers in Immunology, 2019, 10, 763.	2.2	38
53	AA amyloidosis treated with tocilizumab: case series and updated literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 84-92.	1.4	37
54	AL cardiac amyloidosis and arterial thromboembolic events. Scandinavian Journal of Rheumatology, 2005, 34, 315-319.	0.6	34

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55	Pharmacological inhibitors ofÂtheÂmevalonate pathway activate pro-IL-1 processing andÂIL-1 release byÂhuman monocytes. European Cytokine Network, 2009, 20, 112-120.	1.1	34
56	Systemic autoinflammatory diseases: Clinical state of the art. Best Practice and Research in Clinical Rheumatology, 2020, 34, 101529.	1.4	34
57	Cellulitis due to Myroides odoratimimus in a patient with alcoholic cirrhosis. Clinical and Experimental Dermatology, 2007, 33, 071202194819001-???.	0.6	33
58	Intrafamilial variable phenotypic expression of a CIAS1 mutation: from Muckle-Wells to chronic infantile neurological cutaneous and articular syndrome. Journal of Rheumatology, 2005, 32, 747-51.	1.0	33
59	A survey of resistance to colchicine treatment for French patients with familial Mediterranean fever. Orphanet Journal of Rare Diseases, 2017, 12, 54.	1.2	32
60	High lactate dehydrogenase levels at admission for painful vaso-occlusive crisis is associated with severe outcome in adult SCD patients. Clinical Biochemistry, 2012, 45, 1578-1582.	0.8	31
61	Radionuclide Exploration of Dialysis Amyloidosis: Preliminary Experience. American Journal of Kidney Diseases, 1988, 11, 231-237.	2.1	30
62	Prevalence and Correlates of Metabolic Acidosis among Patients with Homozygous Sickle Cell Disease. Clinical Journal of the American Society of Nephrology: CJASN, 2014, 9, 648-653.	2.2	30
63	Concordance between CRP and SAA in familial Mediterranean fever during attack-free period: A study of 218 patients. Clinical Biochemistry, 2017, 50, 206-209.	0.8	30
64	Mutational spectrum in the MEFV and TNFRSF1A genes in patients suffering from AA amyloidosis and recurrent inflammatory attacks. Nephrology Dialysis Transplantation, 2002, 17, 1212-1217.	0.4	29
65	Association of hidradenitis suppurativa and familial Mediterranean fever: A case series of 6 patients. Joint Bone Spine, 2017, 84, 159-162.	0.8	28
66	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.	0.5	28
67	Short-term effects of parathyroidectomy on plasma biochemistry in chronic uremia. Kidney International, 1989, 36, 120-126.	2.6	27
68	Autoinflammatory conditions: when to suspect? How to treat?. Best Practice and Research in Clinical Rheumatology, 2010, 24, 401-411.	1.4	27
69	Functional consequences of a germline mutation in the leucineâ€rich repeat domain of NLRP3 identified in an atypical autoinflammatory disorder. Arthritis and Rheumatism, 2010, 62, 1176-1185.	6.7	27
70	Recurring episodes of meningitis (Mollaret's meningitis) with one showing an association with herpes simplex virus type 2. Journal of Infection, 1996, 32, 247-248.	1.7	25
71	Intrafamilial segregation analysis of the p.E148Q MEFV allele in familial Mediterranean fever. Annals of the Rheumatic Diseases, 2006, 65, 1154-1157.	0.5	24
72	Musculoskeletal Symptoms in Patients With Cryopyrinâ€Associated Periodic Syndromes: A Large Database Study. Arthritis and Rheumatology, 2015, 67, 3027-3036.	2.9	24

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73	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.	1.0	23
74	Diffuse large and small bowel necrosis in catastrophic antiphospholipid syndrome. European Journal of Gastroenterology and Hepatology, 2006, 18, 1011-1014.	0.8	22
75	Somatic Mosaic NLRP3 Mutations and Inflammasome Activation in Late-Onset Chronic Urticaria. Journal of Investigative Dermatology, 2020, 140, 791-798.e2.	0.3	19
76	Thoracic spinal cord compression indicating Wegener's granulomatosis in a patient with a previous presumptive diagnosis of microscopic polyangiitis. Joint Bone Spine, 2007, 74, 382-384.	0.8	18
77	Specific changes in faecal microbiota are associated with familial Mediterranean fever. Annals of the Rheumatic Diseases, 2019, 78, 1398-1404.	0.5	18
78	VLITL is a major cross-Î²-sheet signal for fibrinogen AÎ±-chain frameshift variants. Blood, 2017, 130, 2799-2807.	0.6	17
79	Clinical and multi-omics cross-phenotyping of patients with autoimmune and autoinflammatory diseases: the observational TRANSIMMUNOM protocol. BMJ Open, 2018, 8, e021037.	0.8	17
80	TNFRSF1A-associated periodic syndrome (TRAPS), Muckle-Wells syndrome (MWS) and renal amyloidosis. Journal of Nephrology, 2003, 16, 435-7.	0.9	17
81	Variants of Alport's syndrome. Pediatric Nephrology, 1987, 1, 419-421.	0.9	16
82	Genetically determined recurrent fevers. Current Opinion in Immunology, 2001, 13, 539-542.	2.4	16
83	Fast diagnostic test for familial Mediterranean fever based on a kinase inhibitor. Annals of the Rheumatic Diseases, 2021, 80, 128-132.	0.5	16
84	Fibrinogen Act chain mutation (Arg554 Leu) associated with hereditary renal amyloidosis in a French family. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 1998, 5, 279-284.	1.4	15
85	Musculoskeletal disorders in secondary amyloidosis and hereditary fevers. Best Practice and Research in Clinical Rheumatology, 2003, 17, 929-944.	1.4	15
86	Hereditary fevers. Current Opinion in Rheumatology, 1999, 11, 75-78.	2.0	14
87	Analysis of the NOD2/CARD15 Gene in Patients Affected with the Aseptic Abscesses Syndrome with or without Inflammatory Bowel Disease. Digestive Diseases and Sciences, 2008, 53, 490-499.	1.1	14
88	Realâ€World Experience and Impact of Canakinumab in Cryopyrinâ€Associated Periodic Syndrome: Results From a French Observational Study. Arthritis Care and Research, 2017, 69, 903-911.	1.5	14
89	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.	1.6	14
90	Post-surgical deterioration of renal function in primary hyperoxaluria. Nephrology Dialysis Transplantation, 1987, 1, 261-4.	0.4	14

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91	Allogenic bone marrow transplantation: not a treatment yet for familial Mediterranean fever. Blood, 2003, 102, 409-409.	0.6	13
92	Reninâ€angiotensin system blockade promotes a cardioâ€renal protection in albuminuric homozygous sickle cell patients. British Journal of Haematology, 2017, 179, 820-828.	1.2	13
93	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.	1.6	13
94	Infections and AA amyloidosis: An overview. International Journal of Clinical Practice, 2021, 75, e13966.	0.8	12
95	Is procalcitonin a marker of invasive bacterial infection in acute sickle-cell vaso-occlusive crisis?. Infection, 2011, 39, 41-45.	2.3	11
96	AA amyloidosis is an emerging cause of nephropathy in obese patients. European Journal of Internal Medicine, 2017, 39, e18-e20.	1.0	11
97	Tumour necrosis factor receptor-1 associated periodic syndrome (TRAPS)-related AA amyloidosis: a national case series and systematic review. Rheumatology, 2021, 60, 5775-5784.	0.9	11
98	AUTOINFLAMMATORY DISEASES. Acta Clinica Belgica, 2006, 61, 264-269.	0.5	10
99	Tuberculosis in adult patients with sickle cell disease. Journal of Infection, 2007, 55, 439-444.	1.7	10
100	Respiratory rate: the neglected vital sign. Medical Journal of Australia, 2008, 189, 531-532.	0.8	10
101	Pulmonary hypertension in a case of dermatomyositis. Journal of Rheumatology, 1993, 20, 1452-3.	1.0	10
102	Autoinflammatory syndromes. Presse Medicale, 2019, 48, e21-e23.	0.8	9
103	AA amyloidosis secondary to adult onset Still's disease: About 19 cases. Seminars in Arthritis and Rheumatism, 2020, 50, 156-165.	1.6	9
104	Synchrotron X-Ray Microdiffraction Reveals Intrinsic Structural Features of Amyloid Deposits In Situ. Biophysical Journal, 2011, 101, 486-493.	0.2	8
105	Cystic fibrosis and AA amyloidosis: a survey in the French cystic fibrosis network. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 231-237.	1.4	8
106	The NLRP3 p.A441V Mutation in NLRP3 â€AID Pathogenesis: Functional Consequences, Phenotypeâ€Genotype Correlations and Evidence for a Recurrent Mutational Event. ACR Open Rheumatology, 2019, 1, 267-276.	0.9	8
107	AA amyloidosis complicating cryopyrin-associated periodic syndrome: a study of 86 cases including 23 French patients and systematic review. Rheumatology, 2022, 61, 4827-4834.	0.9	8
108	A novel variant of transthyretin (Glu42Asp) associated with sporadic late-onset cardiac amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 1998, 5, 285-287.	1.4	7

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109	Unusual sites of Salmonella osteoarthritis in patients with sickle cell disease: two cases. Clinical Rheumatology, 2007, 26, 1356-1358.	1.0	7
110	Involvement of the Modifier Gene of a Human Mendelian Disorder in a Negative Selection Process. PLoS ONE, 2009, 4, e7676.	1.1	7
111	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.	1.4	7
112	In familial Mediterranean fever, soluble TREM-1 plasma level is higher in case of amyloidosis. Innate Immunity, 2019, 25, 487-490.	1.1	7
113	Plasma histamine elevation in a large cohort of sickle cell disease patients. British Journal of Haematology, 2019, 186, 125-129.	1.2	7
114	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.	1.6	7
115	Neurological manifestations in mevalonate kinase deficiency: A systematic review. Molecular Genetics and Metabolism, 2022, 136, 85-93.	0.5	7
116	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.	1.8	7
117	Fifth International Congress on Familial Mediterranean Fever and Systemic Autoinflammatory Diseases. Expert Review of Clinical Immunology, 2008, 4, 425-428.	1.3	6
118	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. Journal of Rheumatology, 2019, 46, 1535-1539.	1.0	6
119	Nonâ€amyloid liver involvement in familial Mediterranean fever: A systematic literature review. Liver International, 2020, 40, 1269-1277.	1.9	6
120	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.	0.9	6
121	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.	0.8	6
122	Transthyretin mutation (TTRGly47Ala) associated with familial amyloid polyneuropathy in a French family. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2002, 9, 272-275.	1.4	5
123	Moyens paracliniques du diagnostic des amyloses. Revue Du Rhumatisme (Edition Francaise), 2002, 69, 1172-1180.	0.0	5
124	Hemodynamic and biological correlates of glomerular hyperfiltration in sickle cell patients before and under reninâ€“angiotensin system blocker. Scientific Reports, 2021, 11, 11682.	1.6	5
125	COVID-19 Presenting With Confusion: An Unusual but Suggestive Electroencephalography Pattern of Encephalitis. Journal of Clinical Neurophysiology, 2021, 38, e11-e13.	0.9	5
126	Lateâ€onset familial amyloid polyneuropathy with the TTR Met 30 mutation in France. Clinical Genetics, 1993, 43, 143-145.	1.0	4

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127	Genotypic diagnosis of familial Mediterranean fever (FMF) using new microsatellite markers: example of two extensive non-Ashkenazi Jewish pedigrees.. Journal of Medical Genetics, 1997, 34, 375-381.	1.5	4
128	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.	1.0	4
129	Cause of death and risk factors for mortality in AA amyloidosis: A French retrospective study. European Journal of Internal Medicine, 2020, 82, 130-132.	1.0	4
130	Amyloidosis physiopathology. Joint Bone Spine, 2000, 67, 164-70.	0.8	4
131	Recurrent hepatic hematoma due to familial lysozyme amyloidosis resolves with conservative management. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2014, 21, 66-68.	1.4	3
132	Manifestations ostÃ©oarticulaires des amyloses. Revue Du Rhumatisme Monographies, 2011, 78, 279-285.	0.0	2
133	Light drinking has positive public health consequences. Annals of Oncology, 2013, 24, 1420-1421.	0.6	2
134	Prescription of interleukin-1 inhibitors in a French adult cohort of familial Mediterranean fever. European Journal of Internal Medicine, 2021, 84, 109-111.	1.0	2
135	Could we measure hair colchicine to assess colchicine observance in familial Mediterranean fever?. Rheumatology, 2021, 60, 1563-1564.	0.9	2
136	Mosaic variants in TNFRSF1A: an emerging cause of tumour necrosis factor receptor-associated periodic syndrome. Rheumatology, 2022, 62, 473-479.	0.9	2
137	Breakthroughs in the genetics of hereditary fevers. European Journal of Internal Medicine, 2000, 11, 242-244.	1.0	1
138	FiÃ¨vres rÃ©currentes gÃ©nÃ©tiques. Revue De Medecine Interne, 2006, 27, S259-S260.	0.6	1
139	Rheumatoid arthritis revealed by polyadenopathy, diarrhea and digestive AA amyloidosis. Joint Bone Spine, 2019, 86, 397-398.	0.8	1
140	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.	2.1	1
141	AA amyloidosis associated with Fabry disease. International Journal of Clinical Practice, 2020, 74, e13577.	0.8	1
142	â€œHelicobacter pylori in familial mediterranean fever: A series of 120 patients from literature and from franceâ€• Helicobacter, 2021, 26, e12789.	1.6	1
143	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.	0.9	1
144	Prognostic Value of Hyponatremia During Acute Painful Episodes in Sickle Cell Disease. American Journal of Medicine, 2020, 133, e465-e482.	0.6	1

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145	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, .	1.3	1
146	Study of a restriction fragment length polymorphism for serum amyloid P gene in rheumatoid arthritis with amyloidosis. Journal of Rheumatology, 1991, 18, 994-6.	1.0	1
147	La toxoplasmose cÃ©rÃ©brale au cours du SIDA. Revue De Medecine Interne, 1990, 11, S116.	0.6	0
148	Hypertension artÃ©rielle pulmonaire associÃ©e Ã une dermatomyosite. Revue De Medecine Interne, 1992, 13, S180.	0.6	0
149	Kidney biopsy complicated by perirenal haematoma induces flare of systemic lupus erythematosus: two cases. Nephrology Dialysis Transplantation, 1996, 11, 2299-2301.	0.4	0
150	35. Fibrinogen biosynthesis by renal mesangial cells. A possible role in renal amyloid deposits. Blood Coagulation and Fibrinolysis, 1998, 9, 686.	0.5	0
151	Une nouvelle famille d'amylose Ã lysozyme associÃ©e Ã un variant gÃ©nÃ©tique original. Revue De Medecine Interne, 2001, 22, 22-23.	0.6	0
152	Le curli du coli : une variÃ©tÃ© physiologique dâ€™amylose. Medecine/Sciences, 2002, 18, 664-664.	0.0	0
153	Jejunoileitis in Adult-onset Familial Mediterranean Fever in Japan. Internal Medicine, 2016, 55, 1237-1237.	0.3	0
154	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.	0.5	0
155	The missing link between familial Mediterranean fever and recurrent aseptic meningitis. Pediatrics and Neonatology, 2019, 60, 350.	0.3	0
156	Response to Letter to the Editor. Innate Immunity, 2020, 26, 232-233.	1.1	0
157	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.	1.4	0
158	Thyroid disorders in familial Mediterranean fever: think about AA amyloidosis!. Clinical Rheumatology, 2021, 40, 3381-3382.	1.0	0
159	Percutaneous drainage of retroperitoneal abscesses.. Radiology, 1989, 170, 280-281.	3.6	0
160	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.	1.4	0