Gilles Grateau

List of Publications by Year in Descending Order

160 papers	6,758 citations	40 h-index	79 g-index
267 ext. papers	8,020 ext. citations	4.6 avg, IF	5.23 L-index

#	Paper	IF	Citations
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	2.7	
159	Tattooing and autoinflammatory diseases: a study among 197 French patients from the JIR cohort.. *Journal of the European Academy of Dermatology and Venereology*, **2021**,	4.6	
158	Could we measure hair colchicine to assess colchicine observance in familial Mediterranean fever?. *Rheumatology*, **2021**, 60, 1563-1564	3.9	1
157	COVID-19 Presenting With Confusion: An Unusual but Suggestive Electroencephalography Pattern of Encephalitis. *Journal of Clinical Neurophysiology*, **2021**, 38, e11-e13	2.2	4
156	Tumour necrosis factor receptor-1 associated periodic syndrome (TRAPS)-related AA amyloidosis: a national case series and systematic review. *Rheumatology*, **2021**, 60, 5775-5784	3.9	4
155	Amyloid Goiter in Familial Mediterranean Fever: Description of 42 Cases from a French Cohort and from Literature Review. *Journal of Clinical Medicine*, **2021**, 10,	5.1	1
154	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	4	0
153	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	2.7	
152	La fièvre méditerranéenne familiale est-elle une situation à risque de développer une forme grave d'infection par la COVID19 ? Résultat d'une étude rétrospective sur 627 patients en période et zone endémique en France. *Revue De Medecine Interne*, **2021**, 42, A60-A61	0.1	78
151	Hemodynamic and biological correlates of glomerular hyperfiltration in sickle cell patients before and under renin-angiotensin system blocker. *Scientific Reports*, **2021**, 11, 11682	4.9	0
150	COVID-19 outcomes in patients with inflammatory rheumatic and musculoskeletal diseases treated with rituximab: a cohort study. *Lancet Rheumatology, The*, **2021**, 3, e419-e426	14.2	81
149	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	3.3	
148	Thyroid disorders in familial Mediterranean fever: think about AA amyloidosis!. *Clinical Rheumatology*, **2021**, 40, 3381-3382	3.9	
147	Fast diagnostic test for familial Mediterranean fever based on a kinase inhibitor. *Annals of the Rheumatic Diseases*, **2021**, 80, 128-132	2.4	6
146	Prescription of interleukin-1 inhibitors in a French adult cohort of familial Mediterranean fever. *European Journal of Internal Medicine*, **2021**, 84, 109-111	3.9	0
145	Infections and AA amyloidosis: An overview. *International Journal of Clinical Practice*, **2021**, 75, e13966	2.9	3
144	"Helicobacter pylori in familial mediterranean fever: A series of 120 patients from literature and from france". *Helicobacter*, **2021**, 26, e12789	4.9	

143	AA amyloidosis complicating monoclonal gammopathies, an unusual feature validating the concept of "monoclonal gammopathy of inflammatory significance"?. *International Journal of Clinical Practice*, **2021**, 75, e14817	2.9	0
142	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	5.3	2
141	Response to Letter to the Editor. *Innate Immunity*, **2020**, 26, 232-233	2.7	
140	Systemic autoinflammatory diseases: Clinical state of the art. *Best Practice and Research in Clinical Rheumatology*, **2020**, 34, 101529	5.3	11
139	AA amyloidosis associated with Fabry disease. *International Journal of Clinical Practice*, **2020**, 74, e13577	2.9	
138	Non-amyloid liver involvement in familial Mediterranean fever: A systematic literature review. *Liver International*, **2020**, 40, 1269-1277	7.9	3
137	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	5.3	2
136	Glucan-induced reprogramming of human macrophages inhibits NLRP3 inflammasome activation in cryopyrinopathies. *Journal of Clinical Investigation*, **2020**, 130, 4561-4573	15.9	20
135	Prognostic Value of Hyponatremia During Acute Painful Episodes in Sickle Cell Disease. *American Journal of Medicine*, **2020**, 133, e465-e482	2.4	0
134	Cause of death and risk factors for mortality in AA amyloidosis: A French retrospective study. *European Journal of Internal Medicine*, **2020**, 82, 130-132	3.9	2
133	Polyarthrite rhumatoïde révélée par une polyadénopathie, une diarrhée et une amylose AA digestive. *Revue Du Rhumatisme (Edition Francaise)*, **2020**, 87, 74-75	0.1	
132	NLRP3-associated autoinflammatory diseases: Phenotypic and molecular characteristics of germline versus somatic mutations. *Journal of Allergy and Clinical Immunology*, **2020**, 145, 1254-1261	11.5	24
131	AA amyloidosis secondary to adult onset Still's disease: About 19 cases. *Seminars in Arthritis and Rheumatism*, **2020**, 50, 156-165	5.3	1
130	Somatic Mosaic NLRP3 Mutations and Inflammasome Activation in Late-Onset Chronic Urticaria. *Journal of Investigative Dermatology*, **2020**, 140, 791-798.e2	4.3	6
129	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*, **2020**,	2.4	13
128	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	5.6	1
127	In familial Mediterranean fever, soluble TREM-1 plasma level is higher in case of amyloidosis. *Innate Immunity*, **2019**, 25, 487-490	2.7	6
126	Autoinflammatory diseases: State of the art. *Presse Medicale*, **2019**, 48, e25-e48	2.2	24

125	Expression of SAA1, SAA2 and SAA4 genes in human primary monocytes and monocyte-derived macrophages. *PLoS ONE*, **2019**, 14, e0217005	3.7	25
124	Monoclonal Gammopathy, Arthralgias, and Recurrent Fever Syndrome: A New Autoinflammatory Syndrome?. *Journal of Rheumatology*, **2019**, 46, 1535-1539	4.1	4
123	Association of Vasculitis and Familial Mediterranean Fever. *Frontiers in Immunology*, **2019**, 10, 763	8.4	20
122	Plasma histamine elevation in a large cohort of sickle cell disease patients. *British Journal of Haematology*, **2019**, 186, 125-129	4.5	1
121	The missing link between familial Mediterranean fever and recurrent aseptic meningitis. *Pediatrics and Neonatology*, **2019**, 60, 350	1.8	
120	Rheumatoid arthritis revealed by polyadenopathy, diarrhea and digestive AA amyloidosis. *Joint Bone Spine*, **2019**, 86, 397-398	2.9	1
119	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	2.7	4
118	Specific changes in faecal microbiota are associated with familial Mediterranean fever. *Annals of the Rheumatic Diseases*, **2019**, 78, 1398-1404	2.4	5
117	The NLRP3 p.A441V Mutation in -AID Pathogenesis: Functional Consequences, Phenotype-Genotype Correlations and Evidence for a Recurrent Mutational Event. *ACR Open Rheumatology*, **2019**, 1, 267-276	3.5	4
116	Commentary to "A 44-year-old female with familial Mediterranean fever, cardiomyopathy and end stage renal disease" by Magaki et al. *Brain Pathology*, **2019**, 29, 311	6	1
115	Association entre fi□re m□diterran□enne familiale et scl□ose en plaque. S□ie de 19 cas issus de la cohorte fran□ise et d□ne revue syst□matique de la litt□ature. *Revue De Medecine Interne*, **2019**, 40, A88	0.1	
114	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	4.1	19
113	Validation of the Fautrel classification criteria for adult-onset Still's disease. *Seminars in Arthritis and Rheumatism*, **2018**, 47, 578-585	5.3	24
112	Clinical and multi-omics cross-phenotyping of patients with autoimmune and autoinflammatory diseases: the observational TRANSIMMUNOM protocol. *BMJ Open*, **2018**, 8, e021037	3	12
111	Association of hidradenitis suppurativa and familial Mediterranean fever: A case series of 6 patients. *Joint Bone Spine*, **2017**, 84, 159-162	2.9	20
110	AA amyloidosis is an emerging cause of nephropathy in obese patients. *European Journal of Internal Medicine*, **2017**, 39, e18-e20	3.9	7
109	Association d□ne maladie de Verneuil □une fi□re m□diterran□enne familiale´: 6´cas. *Revue Du Rhumatisme (Edition Francaise)*, **2017**, 84, 543-546	0.1	
108	Renin-angiotensin system blockade promotes a cardio-renal protection in albuminuric homozygous sickle cell patients. *British Journal of Haematology*, **2017**, 179, 820-828	4.5	12

107	VLITL is a major cross-β-sheet signal for fibrinogen Aα-chain frameshift variants. *Blood*, **2017**, 130, 2799-2807	12.2	15
106	A survey of resistance to colchicine treatment for French patients with familial Mediterranean fever. *Orphanet Journal of Rare Diseases*, **2017**, 12, 54	4.2	16
105	Concordance between CRP and SAA in familial Mediterranean fever during attack-free period: A study of 218 patients. *Clinical Biochemistry*, **2017**, 50, 206-209	3.5	17
104	Schnitzler syndrome: validation and applicability of diagnostic criteria in real-life patients. *Allergy: European Journal of Allergy and Clinical Immunology*, **2017**, 72, 177-182	9.3	38
103	Real-World Experience and Impact of Canakinumab in Cryopyrin-Associated Periodic Syndrome: Results From a French Observational Study. *Arthritis Care and Research*, **2017**, 69, 903-911	4.7	10
102	Efficacité des anti-Il1´dans le TRAPS´: expérience du centre de référence et revue de la littérature. *Revue De Medecine Interne*, **2017**, 38, A105-A106	0.1	
101	Impact of human monocyte and macrophage polarization on NLR expression and NLRP3 inflammasome activation. *PLoS ONE*, **2017**, 12, e0175336	3.7	99
100	THU0567 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	2.4	
99	Jejunoileitis in Adult-onset Familial Mediterranean Fever in Japan. *Internal Medicine*, **2016**, 55, 1237	1.1	
98	Evidence-based recommendations for genetic diagnosis of familial Mediterranean fever. *Annals of the Rheumatic Diseases*, **2015**, 74, 635-41	2.4	112
97	Recommendations for the management of autoinflammatory diseases. *Annals of the Rheumatic Diseases*, **2015**, 74, 1636-44	2.4	179
96	Musculoskeletal symptoms in patients with cryopyrin-associated periodic syndromes: a large database study. *Arthritis and Rheumatology*, **2015**, 67, 3027-36	9.5	19
95	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	2.7	28
94	Prevalence and correlates of metabolic acidosis among patients with homozygous sickle cell disease. *Clinical Journal of the American Society of Nephrology: CJASN*, **2014**, 9, 648-53	6.9	21
93	Fièvres récurrentes héréditaires. Réponses au prétest. *Annales De Dermatologie Et De Venereologie* , **2014**, 141, 546	0.3	
92	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, 231-7	2.7	6
91	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, 66-8	2.7	2
90	Online registry for mutations in hereditary amyloidosis including nomenclature recommendations. *Human Mutation*, **2014**, 35, E2403-12	4.7	162

89	How should we approach classification of autoinflammatory diseases?. *Nature Reviews Rheumatology*, **2013**, 9, 624-9	8.1	39
88	Familial Mediterranean fever in heterozygotes: are we able to accurately diagnose the disease in very young children?. *Arthritis and Rheumatism*, **2013**, 65, 1654-62		22
87	Evidence-based recommendations for the practical management of Familial Mediterranean Fever. *Seminars in Arthritis and Rheumatism*, **2013**, 43, 387-91	5.3	103
86	Light drinking has positive public health consequences. *Annals of Oncology*, **2013**, 24, 1420-1	10.3	1
85	The risk of familial Mediterranean fever in MEFV heterozygotes: a statistical approach. *PLoS ONE*, **2013**, 8, e68431	3.7	46
84	High lactate dehydrogenase levels at admission for painful vaso-occlusive crisis is associated with severe outcome in adult SCD patients. *Clinical Biochemistry*, **2012**, 45, 1578-82	3.5	23
83	Osteo-articular manifestations of amyloidosis. *Best Practice and Research in Clinical Rheumatology*, **2012**, 26, 459-75	5.3	37
82	Hemoglobin sickle cell disease complications: a clinical study of 179 cases. *Haematologica*, **2012**, 97, 1136-41	6.4	97
81	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, 1898-901	4.3	85
80	Synchrotron x-ray microdiffraction reveals intrinsic structural features of amyloid deposits in situ. *Biophysical Journal*, **2011**, 101, 486-93	2.9	7
79	Manifestations ostéoarticulaires des amyloses. *Revue Du Rhumatisme Monographies*, **2011**, 78, 279-285	0	
78	Is procalcitonin a marker of invasive bacterial infection in acute sickle-cell vaso-occlusive crisis?. *Infection*, **2011**, 39, 41-5	5.8	9
77	Identification and functional consequences of a recurrent NLRP12 missense mutation in periodic fever syndromes. *Arthritis and Rheumatism*, **2011**, 63, 1459-64		74
76	Role of interleukin-1β in NLRP12-associated autoinflammatory disorders and resistance to anti-interleukin-1 therapy. *Arthritis and Rheumatism*, **2011**, 63, 2142-8		72
75	Existe-t-il des traitements de l'amylose inflammatoire ?. *Revue Du Rhumatisme (Edition Francaise)*, **2011**, 78, 1-3	0.1	
74	Mevalonate kinase deficiency: a survey of 50 patients. *Pediatrics*, **2011**, 128, e152-9	7.4	160
73	Glomerular hyperfiltration in adult sickle cell anemia: a frequent hemolysis associated feature. *Clinical Journal of the American Society of Nephrology: CJASN*, **2010**, 5, 756-61	6.9	96
72	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, 1681-8	4	40

71	Autoinflammatory conditions: when to suspect? How to treat?. *Best Practice and Research in Clinical Rheumatology*, **2010**, 24, 401-11	5.3	19
70	Functional consequences of a germline mutation in the leucine-rich repeat domain of NLRP3 identified in an atypical autoinflammatory disorder. *Arthritis and Rheumatism*, **2010**, 62, 1176-85		24
69	Specific increase in caspase-1 activity and secretion of IL-1 family cytokines: a putative link between mevalonate kinase deficiency and inflammation. *European Cytokine Network*, **2009**, 20, 101-7	3.3	36
68	Involvement of the modifier gene of a human Mendelian disorder in a negative selection process. *PLoS ONE*, **2009**, 4, e7676	3.7	7
67	A clinical criterion to exclude the hyperimmunoglobulin D syndrome (mild mevalonate kinase deficiency) in patients with recurrent fever. *Journal of Rheumatology*, **2009**, 36, 1677-81	4.1	41
66	Pharmacological inhibitors of the mevalonate pathway activate pro-IL-1 processing and IL-1 release by human monocytes. *European Cytokine Network*, **2009**, 20, 112-20	3.3	23
65	Mutations in NALP12 cause hereditary periodic fever syndromes. *Proceedings of the National Academy of Sciences of the United States of America*, **2008**, 105, 1614-9	11.5	266
64	Fifth International Congress on Familial Mediterranean Fever and Systemic Autoinflammatory Diseases. *Expert Review of Clinical Immunology*, **2008**, 4, 425-8	5.1	6
63	Respiratory rate: the neglected vital sign. *Medical Journal of Australia*, **2008**, 189, 531-2	4	7
62	Cellulitis due to Myroides odoratimimus in a patient with alcoholic cirrhosis. *Clinical and Experimental Dermatology*, **2008**, 33, 97-8	1.8	25
61	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, 490-9	4	9
60	Compression de la moelle épinière thoracique révélant une granulomatose de Wegener chez une patiente ayant un diagnostic initial de micropolyangéite. *Revue Du Rhumatisme (Edition Francaise)*, **2007**, 74, 680-682	0.1	
59	Rhumatismes auto-inflammatoires: stratégie diagnostique et traitement. *Revue Du Rhumatisme (Edition Francaise)*, **2007**, 74, 945-952	0.1	
58	Thoracic spinal cord compression indicating Wegener's granulomatosis in a patient with a previous presumptive diagnosis of microscopic polyangiitis. *Joint Bone Spine*, **2007**, 74, 382-4	2.9	17
57	Auto inflammatory syndromes: Diagnosis and treatment. *Joint Bone Spine*, **2007**, 74, 544-50	2.9	52
56	Unusual sites of Salmonella osteoarthritis in patients with sickle cell disease: two cases. *Clinical Rheumatology*, **2007**, 26, 1356-8	3.9	5
55	Tuberculosis in adult patients with sickle cell disease. *Journal of Infection*, **2007**, 55, 439-44	18.9	9
54	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, 1798-809	3.7	65

53	Autoinflammatory diseases. *Acta Clinica Belgica*, **2006**, 61, 264-9	1.8	10
52	Clinical significance of P46L and R92Q substitutions in the tumour necrosis factor superfamily 1A gene. *Annals of the Rheumatic Diseases*, **2006**, 65, 1158-62	2.4	141
51	Intrafamilial segregation analysis of the p.E148Q MEFV allele in familial Mediterranean fever. *Annals of the Rheumatic Diseases*, **2006**, 65, 1154-7	2.4	20
50	Prevalence and distribution of MEFV mutations among Arabs from the Maghreb patients suffering from familial Mediterranean fever. *Comptes Rendus - Biologies*, **2006**, 329, 71-4	1.4	35
49	Fi□vres r□urrentes g□□iques. *Revue De Medecine Interne*, **2006**, 27, S259-S260	0.1	0
48	Diffuse large and small bowel necrosis in catastrophic antiphospholipid syndrome. *European Journal of Gastroenterology and Hepatology*, **2006**, 18, 1011-4	2.2	22
47	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, 239-45	2.7	60
46	AL cardiac amyloidosis and arterial thromboembolic events. *Scandinavian Journal of Rheumatology*, **2005**, 34, 315-9	1.9	28
45	MEFV analysis is of particularly weak diagnostic value for recurrent fevers in Western European Caucasian patients. *Arthritis and Rheumatism*, **2005**, 52, 3603-5		48
44	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	4.1	30
43	Allogenic bone marrow transplantation: not a treatment yet for familial Mediterranean fever. *Blood* , **2003**, 102, 409	2.2	12
42	CIAS1 mutation in a patient with overlap between Muckle-Wells and chronic infantile neurological cutaneous and articular syndromes. *Dermatology*, **2003**, 206, 257-9	4.4	35
41	Musculoskeletal disorders in secondary amyloidosis and hereditary fevers. *Best Practice and Research in Clinical Rheumatology*, **2003**, 17, 929-44	5.3	10
40	TNFRSF1A-associated periodic syndrome (TRAPS), Muckle-Wells syndrome (MWS) and renal amyloidosis. *Journal of Nephrology*, **2003**, 16, 435-7	4.8	15
39	Le curli du coli : une vari□□physiologique d□amylose. *Medecine/Sciences*, **2002**, 18, 664-664		
38	Diagnostic tools for amyloidosis. *Joint Bone Spine*, **2002**, 69, 538-45	2.9	53
37	The enlarging clinical, genetic, and population spectrum of tumor necrosis factor receptor-associated periodic syndrome. *Arthritis and Rheumatism*, **2002**, 46, 2181-8		147
36	Moyens paracliniques du diagnostic des amyloses. *Revue Du Rhumatisme (Edition Francaise)*, **2002**, 69, 1172-1180	0.1	3

35	Hereditary renal amyloidosis caused by a new variant lysozyme W64R in a French family. *Kidney International*, **2002**, 61, 907-12	9.9	90
34	Mutational spectrum in the MEFV and TNFRSF1A genes in patients suffering from AA amyloidosis and recurrent inflammatory attacks. *Nephrology Dialysis Transplantation*, **2002**, 17, 1212-7	4.3	27
33	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, 272-5	2.7	3
32	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, 1498-506	11	262
31	Genetically determined recurrent fevers. *Current Opinion in Immunology*, **2001**, 13, 539-42	7.8	12
30	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, 51-5	5.3	57
29	Molecular analysis of MVK mutations and enzymatic activity in hyper-IgD and periodic fever syndrome. *European Journal of Human Genetics*, **2001**, 9, 260-6	5.3	138
28	The relation between familial Mediterranean fever and amyloidosis. *Current Opinion in Rheumatology*, **2000**, 12, 61-4	5.3	43
27	Clinical versus genetic diagnosis of familial Mediterranean fever. *QJM - Monthly Journal of the Association of Physicians*, **2000**, 93, 223-9	2.7	51
26	A new mutation (A546T) of the betaig-h3 gene responsible for a French lattice corneal dystrophy type IIIA. *American Journal of Ophthalmology*, **2000**, 129, 248-51	4.9	53
25	Breakthroughs in the genetics of hereditary fevers. *European Journal of Internal Medicine*, **2000**, 11, 242-244	3.9	1
24	Identification ofMEFV-Independent Modifying Genetic Factors for Familial Mediterranean Fever. *American Journal of Human Genetics*, **2000**, 67, 1136-1143	11	188
23	Amyloidosis physiopathology. *Joint Bone Spine*, **2000**, 67, 164-70	2.9	4
22	Mutations in the gene encoding mevalonate kinase cause hyper-IgD and periodic fever syndrome. International Hyper-IgD Study Group. *Nature Genetics*, **1999**, 22, 178-81	36.3	427
21	MEFV-Gene analysis in armenian patients with Familial Mediterranean fever: diagnostic value and unfavorable renal prognosis of the M694V homozygous genotype-genetic and therapeutic implications. *American Journal of Human Genetics*, **1999**, 65, 88-97	11	209
20	Genetic linkage of the Muckle-Wells syndrome to chromosome 1q44. *American Journal of Human Genetics*, **1999**, 65, 1054-9	11	93
19	Hereditary amyloid cardiomyopathy caused by a variant apolipoprotein A1. *American Journal of Pathology*, **1999**, 154, 221-7	5.8	98
18	Hereditary fevers. *Current Opinion in Rheumatology*, **1999**, 11, 75-8	5.3	11

17	Fibrinogen A alpha 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, 279-84	2.7	11
16	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, 285-7	2.7	5
15	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, 375-81	5.8	3
14	A candidate gene for familial Mediterranean fever. *Nature Genetics*, **1997**, 17, 25-31	36.3	1169
13	Haemophagocytic syndrome in patients infected with the human immunodeficiency virus: nine cases and a review. *Journal of Infection*, **1997**, 34, 219-25	18.9	41
12	Renal amyloidosis with a frame shift mutation in fibrinogen aalpha-chain gene producing a novel amyloid protein. *Blood*, **1997**, 90, 4799-805	2.2	65
11	Kidney biopsy complicated by perirenal haematoma induces flare of systemic lupus erythematosus: two cases. *Nephrology Dialysis Transplantation*, **1996**, 11, 2299-301	4.3	
10	Recurring episodes of meningitis (Mollaret's meningitis) with one showing an association with herpes simplex virus type 2. *Journal of Infection*, **1996**, 32, 247-8	18.9	21
9	Late-onset familial amyloid polyneuropathy with the TTR Met 30 mutation in France. *Clinical Genetics*, **1993**, 43, 143-5	4	3
8	Pulmonary hypertension in a case of dermatomyositis. *Journal of Rheumatology*, **1993**, 20, 1452-3	4.1	9
7	Thrombose de la veine jugulaire interne au cours d'un syndrome d'hyperstimulation ovarienne. *Revue De Medecine Interne*, **1992**, 13, S494	0.1	
6	Study of a restriction fragment length polymorphism for serum amyloid P gene in rheumatoid arthritis with amyloidosis. *Journal of Rheumatology*, **1991**, 18, 994-6	4.1	1
5	Short-term effects of parathyroidectomy on plasma biochemistry in chronic uremia. *Kidney International*, **1989**, 36, 120-6	9.9	20
4	In vivo T cell preactivation in chronic uremic hemodialyzed and non-hemodialyzed patients. *Kidney International*, **1989**, 36, 636-44	9.9	74
3	Radionuclide exploration of dialysis amyloidosis: preliminary experience. *American Journal of Kidney Diseases*, **1988**, 11, 231-7	7.4	26
2	Variants of Alport's syndrome. *Pediatric Nephrology*, **1987**, 1, 419-21	3.2	11
1	Post-surgical deterioration of renal function in primary hyperoxaluria. *Nephrology Dialysis Transplantation*, **1987**, 1, 261-4	4.3	12