David Launay

List of Publications by Year
in descending order

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298	18,332	14653	17104
papers	citations	66 h-index	122 g-index

392	392	392	15136
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Successful allogeneic hematopoietic stem cell transplantation in patients with VEXAS syndrome: a 2-center experience. Blood Advances, 2022, 6, 998-1003.	5.2	88
2	Antiâ€“programmed death ligand 1 immunotherapies in cancer patients with pre-existing systemic sclerosis: A postmarketed phase IV safety assessment study. European Journal of Cancer, 2022, 160, 134-139.	2.8	10
3	Attenuated androgen discontinuation in patients with hereditary angioedema: a commented case series. Allergy, Asthma and Clinical Immunology, 2022, 18, 4.	2.0	5
4	Effective Antiâ€“SARS-CoV-2 Immune Response in Patients With Clonal Mast Cell Disorders. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 1356-1364.e2.	3.8	2
5	Interstitial pneumonia with autoimmune features: Evaluation of connective tissue disease incidence during follow-up. European Journal of Internal Medicine, 2022, 97, 62-68.	2.2	10
6	Long-term prophylaxis with lanadelumab for HAE: authorization for temporary use in France. Allergy, Asthma and Clinical Immunology, 2022, 18, 30.	2.0	1
7	The letter responds to comment on: Anti-PD(L)1 immunotherapies in patients with cancer and with pre-existing systemic sclerosis: a post-marketed safety assessment study. European Journal of Cancer, 2022, 165, 208-209.	2.8	0
8	Dyslipidemia is insufficiently treated in antiphospholipid syndrome patients. Lupus, 2022, 31, 1379-1384.	1.6	6
9	Simple gene signature to assess murine fibroblast polarization. Scientific Reports, 2022, 12, .	3.3	6
10	Extended myositis-specific and -associated antibodies profile in systemic sclerosis: A cross-sectional study. Joint Bone Spine, 2021, 88, 105048.	1.6	12
11	Serious Infectious Events and Immunoglobulin Replacement Therapy in Patients With Autoimmune Disease Receiving Rituximab: A Retrospective Cohort Study. Clinical Infectious Diseases, 2021, 72, 727-737.	5.8	25
12	Clinical phenotypes of extrapulmonary sarcoidosis: an analysis of a French, multi-ethnic, multicentre cohort. European Respiratory Journal, 2021, 57, 2001160.	6.7	29
13	Hereditary Angioedema with and Without C1-Inhibitor Deficiency in Postmenopausal Women. Journal of Clinical Immunology, 2021, 41, 163-170.	3.8	4
14	Hemodynamic Response to Treatment and Outcomes in Pulmonary Hypertension Associated With Interstitial Lung Disease Versus Pulmonary Arterial Hypertension in Systemic Sclerosis: Data From a Study Identifying Prognostic Factors in Pulmonary Hypertension Associated With Interstitial Lung Disease. Arthritis and Rheumatology, 2021, 73, 295-304.	5.6	26
15	Systemic Pulmonary Events Associated with Myelodysplastic Syndromes: A Retrospective Multicentre Study. Journal of Clinical Medicine, 2021, 10, 1162.	2.4	3
16	18F-FDG positron emission tomography scanning in systemic sclerosis-associated interstitial lung disease: a pilot study. Arthritis Research and Therapy, 2021, 23, 76.	3.5	20
17	Fatal Enterovirus-related Myocarditis in a Patient with Devicâ€™s Syndrome Treated with Rituximab. Cardiac Failure Review, 2021, 7, e09.	3.0	4
18	Time for precision medicine in systemic sclerosis-associated pulmonary arterial hypertension. European Respiratory Journal, 2021, 57, 2100205.	6.7	2

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19	French recommendations for the management of systemic sclerosis. Orphanet Journal of Rare Diseases, 2021, 16, 322.	2.7	37
20	Association between Leflunomide and Pulmonary Hypertension. Annals of the American Thoracic Society, 2021, 18, 1306-1315.	3.2	8
21	IgG4-related disease and hypereosinophilic syndrome: Overlapping phenotypes. Autoimmunity Reviews, 2021, 20, 102889.	5.8	9
22	Thoracic involvement and imaging patterns in IgG4-related disease. European Respiratory Review, 2021, 30, 210078.	7.1	14
23	Passages in culture and stimulation conditions influence protein expression of primary fibroblasts. Proteomics, 2021, , 2100116.	2.2	0
24	Assessment of T-cell polarization onÂthe basis of surface marker expression: Diagnosis and potential therapeutic implications in lymphocytic variant hypereosinophilic syndrome. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1110-1114.e2.	3.8	7
25	Survival Improved in Patients AgedÂâ‰ 70 Years With Systemic Sclerosis-Associated Pulmonary Arterial Hypertension During the Period 2006 to 2017 in France. Chest, 2020, 157, 945-954.	0.8	13
26	Characteristics of patients with systemic sclerosis suffering from a lower limb amputation: Results of a French collaborative study. Journal of Scleroderma and Related Disorders, 2020, 5, 224-230.	1.7	1
27	Adherence to hydroxychloroquine in patients with systemic lupus: Contrasting results and weak correlation between assessment tools. Joint Bone Spine, 2020, 87, 603-610.	1.6	8
28	Early trajectories of skin thickening are associated with severity and mortality in systemic sclerosis. Arthritis Research and Therapy, 2020, 22, 30.	3.5	20
29	Are pregnancies with lupus but without APS of good prognosis?. Autoimmunity Reviews, 2020, 19, 102489.	5.8	3
30	Sarcoidosis diagnosed on granulomas in the explanted heart after transplantation: Results of a French nationwide study. International Journal of Cardiology, 2020, 307, 94-100.	1.7	10
31	Evolution of high-resolution CT-scan in systemic sclerosis-associated interstitial lung disease: Description and prognosis factors. Seminars in Arthritis and Rheumatism, 2020, 50, 1406-1413.	3.4	11
32	Moderate-to-severe eosinophilia induced by treatment with immune checkpoint inhibitors: 37 cases from a national reference center for hypereosinophilic syndromes and the French pharmacovigilance database. OncoImmunology, 2020, 9, 1722022.	4.6	27
33	Protocol for a partially nested randomised controlled trial to evaluate the effectiveness of the scleroderma patient-centered intervention network COVID-19 home-isolation activities together (SPIN-CHAT) program to reduce anxiety among at-risk scleroderma patients. Journal of Psychosomatic Research, 2020, 135, 110132.	2.6	21
34	Consideration of coping strategies for patients suffering from systemic lupus erythematosus: reflection for a personalised practice of patient education. Clinical and Experimental Rheumatology, 2020, 38, 705-712.	0.8	2
35	Hereditary angioedema: Clinical presentation and socioeconomic cost of 200 French patients. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 328-330.	3.8	3
36	Angioedema associated with thrombolysis for ischemic stroke: analysis of a caseâ€control study. Journal of Internal Medicine, 2019, 286, 702-710.	6.0	13

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37	Renal Pathologic Findings in TAFRO Syndrome: Is There a Continuum Between Thrombotic Microangiopathy and Membranoproliferative Glomerulonephritis? A Case Report and Literature Review. Frontiers in Immunology, 2019, 10, 1489.	4.8	24
38	Hereditary angioedema, emergency management of attacks by a call center. European Journal of Internal Medicine, 2019, 67, 42-46.	2.2	6
39	Oral Plasma Kallikrein Inhibitor BCX7353 is Safe and Effective as an On-Demand Treatment of Angioedema Attacks in Hereditary Angioedema (HAE) Patients: Results of the ZENITH-1 Trial. Journal of Allergy and Clinical Immunology, 2019, 143, AB36.	2.9	4
40	Phenotypes Determined by Cluster Analysis and Their Survival in the Prospective European Scleroderma Trials and Research Cohort of Patients With Systemic Sclerosis. Arthritis and Rheumatology, 2019, 71, 1553-1570.	5.6	75
41	Idiopathic inflammatory myopathies: state of the art on clinical practice guidelines. RMD Open, 2019, 4, e000784.	3.8	19
42	Survival and prognosis factors in systemic sclerosis: data of a French multicenter cohort, systematic review, and meta-analysis of the literature. Arthritis Research and Therapy, 2019, 21, 86.	3.5	114
43	Systemic sclerosis: state of the art on clinical practice guidelines. RMD Open, 2019, 4, e000782.	3.8	91
44	Combined pulmonary fibrosis and emphysema in systemic sclerosis: A syndrome associated with heavy morbidity and mortality. Seminars in Arthritis and Rheumatism, 2019, 49, 98-104.	3.4	33
45	Ulnar Artery Occlusion and Severity Markers of Vasculopathy in Systemic Sclerosis: A Multicenter Crossâ€Sectional Study. Arthritis and Rheumatology, 2019, 71, 983-990.	5.6	25
46	Remission of Refractory Systemic-Onset Juvenile Idiopathic Arthritis After Treatment With Siltuximab. Journal of Clinical Rheumatology, 2019, 25, e40-e42.	0.9	3
47	Reply to Gilchrist et al. and to Musher. Clinical Infectious Diseases, 2018, 66, 637-638.	5.8	1
48	Association between Rheumatoid Arthritis and Pulmonary Hypertension: Data from the French Pulmonary Hypertension Registry. Respiration, 2018, 95, 244-250.	2.6	17
49	Efficacy and safety of rituximab in systemic sclerosis: French retrospective study and literature review. Autoimmunity Reviews, 2018, 17, 582-587.	5.8	74
50	Reliability and Validity of Three Versions of the Brief Fear of Negative Evaluation Scale in Patients With Systemic Sclerosis: A Scleroderma Patientâ€Centered Intervention Network Cohort Study. Arthritis Care and Research, 2018, 70, 1646-1652.	3.4	8
51	Altered B lymphocyte homeostasis and functions in systemic sclerosis. Autoimmunity Reviews, 2018, 17, 244-255.	5.8	58
52	Validation of the Social Appearance Anxiety Scale in Patients With Systemic Sclerosis: A Scleroderma Patientâ€Centered Intervention Network Cohort Study. Arthritis Care and Research, 2018, 70, 1557-1562.	3.4	17
53	Proinflammatory B-cell profile in the early phases of MS predicts an active disease. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e431.	6.0	29
54	Long-term efficacy of remission-maintenance regimens for ANCA-associated vasculitides. Annals of the Rheumatic Diseases, 2018, 77, 1150-1156.	0.9	139

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55	Factor structure and convergent validity of the Derriford Appearance Scale-24 using standard scoring versus treating â€˜not applicableâ€™ responses as missing data: a Scleroderma Patient-centered Intervention Network (SPIN) cohort study. BMJ Open, 2018, 8, e018641.	1.9	4
56	Evaluation of avoralstat, an oral kallikrein inhibitor, in a Phase 3 hereditary angioedema prophylaxis trial: The ‹scp›OPuS‹/scp›â€¢Âstudy. Allergy: European Journal of Allergy and Clinical Immunology, 2018, 73, 1871-1880.	5.7	31
57	Specialist Advice Support for Management of Severe Hereditary Angioedema Attacks: A Multicenter Cluster-Randomized Controlled Trial. Annals of Emergency Medicine, 2018, 72, 194-203.e1.	0.6	6
58	Anticoagulation withdrawal in antiphospholipid syndrome: a retrospective matched-control study. Lupus, 2018, 27, 357-364.	1.6	14
59	Pulmonary Arterial Hypertension Associated With Systemic Lupus Erythematosus. Chest, 2018, 153, 143-151.	0.8	68
60	Limited Exercise Capacity in Patients with Systemic Sclerosis: Identifying Contributing Factors with Cardiopulmonary Exercise Testing. Journal of Rheumatology, 2018, 45, 95-102.	2.0	14
61	Prevalence and Clinical Associations of Antiphospholipid Antibodies in Systemic Sclerosis: New Data From a French Cross-Sectional Study, Systematic Review, and Meta-Analysis. Frontiers in Immunology, 2018, 9, 2457.	4.8	17
62	Antiphospholipid Syndrome With Isolated Isotype M Anticardiolipin and/or Anti-B2GPI Antibody Is Associated With Stroke. Stroke, 2018, 49, 2770-2772.	2.0	13
63	Haemodynamics and serial risk assessment in systemic sclerosis associated pulmonary arterial hypertension. European Respiratory Journal, 2018, 52, 1800678.	6.7	60
64	Whole-Body Distribution and Clinical Association of Telangiectases in Systemic Sclerosis. JAMA Dermatology, 2018, 154, 796.	4.1	14
65	Clinical phenotypes and survival of pre-capillary pulmonary hypertension in systemic sclerosis. PLoS ONE, 2018, 13, e0197112.	2.5	47
66	Plasminogen gene mutation with normal C1 inhibitor hereditary angioedema: Three additional French families. Allergy: European Journal of Allergy and Clinical Immunology, 2018, 73, 2237-2239.	5.7	40
67	Bleeding complications and antithrombotic treatment in 264 pregnancies in antiphospholipid syndrome. Lupus, 2018, 27, 1679-1686.	1.6	25
68	Effectiveness of icatibant for treatment of hereditary angioedema attacks is not affected by body weight: findings from the Icatibant Outcome Survey, a cohort observational study. Clinical and Translational Allergy, 2018, 8, 11.	3.2	3
69	Hypertension pulmonaire et connectivites. Revue Du Rhumatisme Monographies, 2018, 85, 210-220.	0.0	0
70	Schnitzler syndrome co-occurring with idiopathic multicentric Castleman disease that responds to anti-IL-1 therapy: A case report and clue to pathophysiology. Current Research in Translational Medicine, 2018, 66, 83-86.	1.8	9
71	Efficacy of anti-TNF alpha in severe and refractory major vessel involvement of Behcet's disease: A multicenter observational study of 18 patients. Clinical Immunology, 2018, 197, 54-59.	3.2	51
72	SAT0472â€…Ulnar occlusion is a marker of global vascular damage in systemic sclerosis: results from a monocentric prospective study of 99 patients. , 2018, , .		0

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73	Role of B cells in the pathogenesis of systemic sclerosis. Revue De Medecine Interne, 2017, 38, 113-124.	1.0	37
74	Diagnosis of primary antibody and complement deficiencies in young adults after a first invasive bacterial infection. Clinical Microbiology and Infection, 2017, 23, 576.e1-576.e5.	6.0	25
75	Intravenous immunoglobulins in systemic sclerosis: Data from a French nationwide cohort of 46 patients and review of the literature. Autoimmunity Reviews, 2017, 16, 377-384.	5.8	49
76	Extensive levamisole-induced vasculitis. Clinical and Experimental Dermatology, 2017, 42, 413-415.	1.3	10
77	Hereditary angioedema with normal C1 inhibitor: clinical characteristics and treatment response with plasma-derived human C1 inhibitor concentrate (BerinertÂ®) in a French cohort. European Journal of Dermatology, 2017, 27, 155-159.	0.6	24
78	Haemodynamically proven pulmonary hypertension in a patient with GATA2 deficiency-associated pulmonary alveolar proteinosis and fibrosis. European Respiratory Journal, 2017, 49, 1700178.	6.7	9
79	Strains Responsible for Invasive Meningococcal Disease in Patients With Terminal Complement Pathway Deficiencies. Journal of Infectious Diseases, 2017, 215, 1331-1338.	4.0	35
80	Clinical characteristics and response to attack treatment with BerinertÂ® of drug induced angioedema in the French Cohort COBRA.. Journal of Allergy and Clinical Immunology, 2017, 139, AB236.	2.9	0
81	Hereditary angioedema with normal C1 inhibitor in a French cohort: Clinical characteristics and response to treatment with icatibant. Immunity, Inflammation and Disease, 2017, 5, 29-36.	2.7	27
82	Specific Polysaccharide Antibody Deficiency Revealed by Severe Bacterial Infections in Adulthood: A Report on 11 Cases. Clinical Infectious Diseases, 2017, 65, 328-331.	5.8	15
83	The Icatibant Outcome Survey: experience of hereditary angioedema management from six European countries. Journal of the European Academy of Dermatology and Venereology, 2017, 31, 1214-1222.	2.4	21
84	Normal PAI-2 level in French FXII-HAE patients. Journal of Allergy and Clinical Immunology, 2017, 139, 1719-1720.	2.9	5
85	Longâ€term safety of icatibant treatment of patients with angioedema in realâ€world clinical practice. Allergy: European Journal of Allergy and Clinical Immunology, 2017, 72, 994-998.	5.7	16
86	Update of EULAR recommendations for the treatment of systemic sclerosis. Annals of the Rheumatic Diseases, 2017, 76, 1327-1339.	0.9	794
87	Pulmonary hypertension in systemic sclerosis: different phenotypes. European Respiratory Review, 2017, 26, 170056.	7.1	97
88	Combined measurement of carbon monoxide and nitric oxide lung transfer does not improve the identification of pulmonary hypertension in systemic sclerosis. European Respiratory Journal, 2017, 50, 1701008.	6.7	10
89	OP0236â€…Benefit and safety of antithrombotic treatment in 264 pregnancies in patients with antiphospholipid syndrome. , 2017, , .		0
90	An international SUrvey on non-iNvaSive tecHniques to assess the mIcrocirculation in patients with RayNaudâ€™s phEnomenon (SUNSHINE survey). Rheumatology International, 2017, 37, 1879-1890.	3.0	33

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91	Diagnosis and treatment of upper airway oedema caused by acute angio-oedema in the emergency department: a French consensus statement. European Journal of Emergency Medicine, 2017, 24, 318-325.	1.1	6
92	Role of Stromelysin 2 (Matrix Metalloproteinase 10) as a Novel Mediator of Vascular Remodeling Underlying Pulmonary Hypertension Associated With Systemic Sclerosis. Arthritis and Rheumatology, 2017, 69, 2209-2221.	5.6	17
93	Biologics in myelodysplastic syndrome-related systemic inflammatory and autoimmune diseases: French multicenter retrospective study of 29 patients. Autoimmunity Reviews, 2017, 16, 903-910.	5.8	32
94	Central nervous system involvement in eosinophilic granulomatosis with polyangiitis (Churg-Strauss): Report of 26 patients and review of the literature. Autoimmunity Reviews, 2017, 16, 963-969.	5.8	70
95	Persistent triple antiphospholipid antibody positivity as a strong risk factor of first thrombosis, in a long-term follow-up study of patients without history of thrombosis or obstetrical morbidity. Lupus, 2017, 26, 163-169.	1.6	53
96	B Cell Homeostasis and Functional Properties Are Altered in an Hypochlorous Acid-Induced Murine Model of Systemic Sclerosis. Frontiers in Immunology, 2017, 8, 53.	4.8	14
97	Value of the Overall Pneumococcal Polysaccharide Response in the Diagnosis of Primary Humoral Immunodeficiencies. Frontiers in Immunology, 2017, 8, 1862.	4.8	26
98	TAFRO Syndrome in Caucasians: A Case Report and Review of the Literature. Frontiers in Medicine, 2017, 4, 149.	2.6	30
99	Predictors of lung function test severity and outcome in systemic sclerosis-associated interstitial lung disease. PLoS ONE, 2017, 12, e0181692.	2.5	68
100	Factors associated with the 6-minute walk distance in patients with systemic sclerosis. Arthritis Research and Therapy, 2017, 19, 279.	3.5	22
101	Breakthrough attacks in patients with hereditary angioedema receiving long-term prophylaxis are responsive to icatibant: findings from the Icatibant Outcome Survey. Allergy, Asthma and Clinical Immunology, 2017, 13, 31.	2.0	14
102	Joint and tendon involvement predict disease progression in systemic sclerosis: a EUSTAR prospective study. Annals of the Rheumatic Diseases, 2016, 75, 103-109.	0.9	93
103	A gender gap in primary and secondary heart dysfunctions in systemic sclerosis: a EUSTAR prospective study. Annals of the Rheumatic Diseases, 2016, 75, 163-169.	0.9	82
104	Mycophenolate mofetil following cyclophosphamide in worsening systemic sclerosis-associated interstitial lung disease. Journal of Scleroderma and Related Disorders, 2016, 1, 234-240.	1.7	14
105	Hereditary angioedema with normal C1 inhibitor and factor XII mutation: a series of 57 patients from the French National Center of Reference for Angioedema. Clinical and Experimental Immunology, 2016, 185, 332-337.	2.6	42
106	Neutrophilic Dermatoses in Antineutrophil Cytoplasmic Antibody-Associated Vasculitis. Medicine (United States), 2016, 95, e2957.	1.0	29
107	Pulmonary arterial hypertension in idiopathic inflammatory myopathies. Medicine (United States), 2016, 95, e4911.	1.0	40
108	Prediction of improvement in skin fibrosis in diffuse cutaneous systemic sclerosis: a EUSTAR analysis. Annals of the Rheumatic Diseases, 2016, 75, 1743-1748.	0.9	68

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109	Efficacy of sildenafil on ischaemic digital ulcer healing in systemic sclerosis: the placebo-controlled SEDUCE study. Annals of the Rheumatic Diseases, 2016, 75, 1009-1015.	0.9	112
110	FRI0299â€...Pulmonary Arterial Hypertension in Idiopathic Inflammatory Myopathies: Data from The French Pulmonary Hypertension Registry and Review of The Literature. Annals of the Rheumatic Diseases, 2016, 75, 543.2-543.	0.9	0
111	Reemergence of Splenectomy for ITP Second-line Treatment?. Annals of Surgery, 2016, 264, 772-777.	4.2	23
112	A New Era in the Treatment of Scleroderma-associated Interstitial Lung Disease?. Journal of Rheumatology, 2016, 43, 1619-1621.	2.0	0
113	N-terminal pro-brain natriuretic peptide is a strong predictor of mortality in systemic sclerosis. International Journal of Cardiology, 2016, 223, 385-389.	1.7	16
114	Misdiagnosis trends in patients with hereditary angioedema from the real-world clinical setting. Annals of Allergy, Asthma and Immunology, 2016, 117, 394-398.	1.0	78
115	Mastocytosis among elderly patients. Medicine (United States), 2016, 95, e3901.	1.0	6
116	Digestive-tract sarcoidosis. Medicine (United States), 2016, 95, e4279.	1.0	30
117	A nationwide study of acquired C1-inhibitor deficiency in France. Medicine (United States), 2016, 95, e4363.	1.0	64
118	Longâ€Term Outcomes Among Participants in the WEGENT Trial of Remissionâ€Maintenance Therapy for Granulomatosis With Polyangiitis (Wegener's) or Microscopic Polyangiitis. Arthritis and Rheumatology, 2016, 68, 690-701.	5.6	101
119	Special considerations in pregnant systemic sclerosis patients. Expert Review of Clinical Immunology, 2016, 12, 1161-1173.	3.0	19
120	Dedicated call center (SOS-HAE) for hereditary angioedema attacks: study protocol for a randomised controlled trial. Trials, 2016, 17, 225.	1.6	5
121	Efficacy of Anti-TNFÎ± in Severe and Refractory Neuro-Behcet Disease. Medicine (United States), 2016, 95, e3550.	1.0	43
122	Midostaurin in Advanced Systemic Mastocytosis. New England Journal of Medicine, 2016, 374, 2605-2606.	27.0	54
123	Digital ulcers predict a worse disease course in patients with systemic sclerosis. Annals of the Rheumatic Diseases, 2016, 75, 681-686.	0.9	111
124	Mast cells' involvement in inflammation pathways linked to depression: evidence in mastocytosis. Molecular Psychiatry, 2016, 21, 1511-1516.	7.9	64
125	Systemic inflammatory and autoimmune manifestations associated with myelodysplastic syndromes and chronic myelomonocytic leukaemia: a French multicentre retrospective study. Rheumatology, 2016, 55, 291-300.	1.9	170
126	Dermatomyositis With or Without Anti-Melanoma Differentiation-Associated Gene 5 Antibodies. American Journal of Pathology, 2016, 186, 691-700.	3.8	78

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127	Repositioning Antitubercular 6-Nitro-2,3-dihydroimidazo[2,1-b][1,3]oxazoles for Neglected Tropical Diseases: Structureâ€“Activity Studies on a Preclinical Candidate for Visceral Leishmaniasis. Journal of Medicinal Chemistry, 2016, 59, 2530-2550.	6.4	46
128	Current Approaches to the Treatment of Systemic-Sclerosis-Associated Pulmonary Arterial Hypertension (SSc-PAH). Current Rheumatology Reports, 2016, 18, 10.	4.7	38
129	A prospective study of the 6â€…min walk test as a surrogate marker for haemodynamics in two independent cohorts of treatment-naÃ¯ve systemic sclerosis-associated pulmonary arterial hypertension. Annals of the Rheumatic Diseases, 2016, 75, 1457-1465.	0.9	16
130	Spectrum and Prognosis of Noninfectious Renal Mixed Cryoglobulinemic GN. Journal of the American Society of Nephrology: JASN, 2016, 27, 1213-1224.	6.1	44
131	Gentle blood aspiration and tube cushioning reduce pneumatic tube system interference in lactate dehydrogenase assays. Annals of Clinical Biochemistry, 2016, 53, 295-297.	1.6	9
132	Triggers and Prodromal Symptoms of Angioedema Attacks in Patients With Hereditary Angioedema. Journal of Investigational Allergology and Clinical Immunology, 2016, 26, 383-386.	1.3	57
133	Hereditary Angioedema: Clinical Characteristics and Treatment Response with Berinert in a French Cohort in Patient Under 16 Years Old. Journal of Allergy and Clinical Immunology, 2015, 135, AB193.	2.9	1
134	Determinants of Hydroxychloroquine Blood Concentration Variations in Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2015, 67, 2176-2184.	5.6	118
135	SAT0543â€…Igg4-Related Disease: Clinical and Biological Characteristics in 90 Patients from a Large Multicentric National Registry. Annals of the Rheumatic Diseases, 2015, 74, 857.1-857.	0.9	0
136	FRI0509â€…Chromosome 22Q11.2 Deletion Syndrome (Digeorge Syndrome) and Autoimmunity: A French Retrospective Pediatric Study of 15 Cases. Annals of the Rheumatic Diseases, 2015, 74, 613.2-613.	0.9	0
137	Distinctive Patterns of Transthyretin Amyloid in Salivary Tissue. American Journal of Surgical Pathology, 2015, 39, 1035-1044.	3.7	18
138	Analysis of characteristics associated with reinjection of icatibant: Results from the Icatibant Outcome Survey. Allergy and Asthma Proceedings, 2015, 36, 399-406.	2.2	19
139	A French National Survey on Clotting Disorders in Mastocytosis. Medicine (United States), 2015, 94, e1414.	1.0	17
140	OP0059â€…Phenotypes Determined by Cluster Analysis and their Survival in the Prospective Eustar Cohort of Patients with Systemic Sclerosis. Annals of the Rheumatic Diseases, 2015, 74, 90.1-90.	0.9	1
141	An Optimized Fluorescence-Based Bidimensional Immunoproteomic Approach for Accurate Screening of Autoantibodies. PLoS ONE, 2015, 10, e0132142.	2.5	9
142	Value of systolic pulmonary arterial pressure as a prognostic factor of death in the systemic sclerosis EUSTAR population. Rheumatology, 2015, 54, 1262-1269.	1.9	25
143	Seronegative polyarthritis revealing antisynthetase syndrome: a multicentre study of 40 patients. Rheumatology, 2015, 54, 927-932.	1.9	37
144	CD3-CD4+ lymphoid variant of hypereosinophilic syndrome: nodal and extranodal histopathological and immunophenotypic features of a peripheral indolent clonal T-cell lymphoproliferative disorder. Haematologica, 2015, 100, 1086-95.	3.5	37

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145	ActualitÃ©s en allergologieÂ: mÃ©decine interne. Revue Francaise D'allergologie, 2015, 55, 207-209.	0.2	0
146	Treatment of HAE Attacks in the Icatibant Outcome Survey: An Analysis of Icatibant Self-Administration versus Administration by Health Care Professionals. International Archives of Allergy and Immunology, 2015, 167, 21-28.	2.1	37
147	Factors associated with hospital admission in hereditary angioedema attacks: a multicenter prospective study. Annals of Allergy, Asthma and Immunology, 2015, 114, 499-503.	1.0	14
148	Prediction of worsening of skin fibrosis in patients with diffuse cutaneous systemic sclerosis using the EUSTAR database. Annals of the Rheumatic Diseases, 2015, 74, 1124-1131.	0.9	96
149	Aminothiazoles: Hit to lead development to identify antileishmanial agents. European Journal of Medicinal Chemistry, 2015, 102, 582-593.	5.5	34
150	A comparison between nailfold capillaroscopy patterns in adulthood in juvenile and adult-onset systemic sclerosis: A EUSTAR exploratory study. Microvascular Research, 2015, 102, 19-24.	2.5	13
151	Nitroimidazo-oxazole compound DNDI-VL-2098: an orally effective preclinical drug candidate for the treatment of visceral leishmaniasis. Journal of Antimicrobial Chemotherapy, 2015, 70, 518-527.	3.0	56
152	ASXL1 but Not TET2 Mutations Adversely Impact Overall Survival of Patients Suffering Systemic Mastocytosis with Associated Clonal Hematologic Non-Mast-Cell Diseases. PLoS ONE, 2014, 9, e85362.	2.5	65
153	FRI0479â€…Serum Free Light Chains of Immunoglobulins Are Associated with Disease Activity in Systemic Sclerosis: A Prospective and Controlled Study. Annals of the Rheumatic Diseases, 2014, 73, 560.3-560.	0.9	0
154	AB0614â€…Prognostic Factors of Functional Outcome in Systemic Sclerosis-Associated Interstitial Lung Disease. Annals of the Rheumatic Diseases, 2014, 73, 1008.4-1009.	0.9	1
155	In vitro metabolism, disposition, preclinical pharmacokinetics and prediction of human pharmacokinetics of DNDI-VL-2098, a potential oral treatment for Visceral Leishmaniasis. European Journal of Pharmaceutical Sciences, 2014, 65, 147-155.	4.0	29
156	OP0096â€…Relevance of the 6-Minute Walking Test in Assessing the Severity of Pulmonary Arterial Hypertension Associated with Systemic Sclerosis, without Interstitial Lung Disease. Annals of the Rheumatic Diseases, 2014, 73, 97.3-98.	0.9	0
157	Tranexamic acid as maintenance treatment for non-histaminergic angioedema: analysis of efficacy and safety in 37 patients. Clinical and Experimental Immunology, 2014, 178, 112-117.	2.6	58
158	To Spare or Not to Spareâ€¦? The Aortic Valve in Scleroderma and Aortic Root Aneurysm. Journal of Cardiac Surgery, 2014, 29, 55-58.	0.7	0
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