Guillaume Lefevre

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6906451/publications.pdf

Version: 2024-02-01

50 papers

1,874 citations

236925 25 h-index 42 g-index

57 all docs

57 docs citations

57 times ranked

2846 citing authors

| # | Article | IF | CITATIONS |
|----|---|--------------|--------------|
| 1 | Autoimmune and inflammatory manifestations occur frequently in patients with primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2017, 140, 1388-1393.e8. | 2.9 | 222 |
| 2 | Survival and Prognostic Factors in Systemic Sclerosis–Associated Pulmonary Hypertension: A Systematic Review and Metaâ€Analysis. Arthritis and Rheumatism, 2013, 65, 2412-2423. | 6.7 | 205 |
| 3 | Patient-level analysis of five international cohorts further confirms the efficacy of aspirin for the primary prevention of thrombosis in patients with antiphospholipid antibodies. Autoimmunity Reviews, 2015, 14, 192-200. | 5. 8 | 118 |
| 4 | The Lymphoid Variant of Hypereosinophilic Syndrome. Medicine (United States), 2014, 93, 255-266. | 1.0 | 98 |
| 5 | Prevalence of Anti–RNA Polymerase III Antibodies in Systemic Sclerosis: New Data From a French Cohort and a Systematic Review and Metaâ€Analysis. Arthritis and Rheumatology, 2014, 66, 407-417. | 5 . 6 | 76 |
| 6 | Altered B lymphocyte homeostasis and functions in systemic sclerosis. Autoimmunity Reviews, 2018, 17, 244-255. | 5.8 | 58 |
| 7 | Tocilizumab in Giant Cell Arteritis: A Multicenter Retrospective Study of 34 Patients. Journal of Rheumatology, 2016, 43, 1547-1552. | 2.0 | 56 |
| 8 | (A Critical Appraisal of) Classification of Hypereosinophilic Disorders. Frontiers in Medicine, 2017, 4, 216. | 2.6 | 56 |
| 9 | IL-18 Is Involved in Eosinophil-Mediated Tumoricidal Activity against a Colon Carcinoma Cell Line by Upregulating LFA-1 and ICAM-1. Journal of Immunology, 2015, 195, 2483-2492. | 0.8 | 54 |
| 10 | 18F-FDG PET/CT in patients with amyloid light-chain amyloidosis: case-series and literature review. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 94-98. | 3.0 | 52 |
| 11 | Outcome of Hepatitis E Virus Infection in Patients With Inflammatory Arthritides Treated With Immunosuppressants. Medicine (United States), 2015, 94, e675. | 1.0 | 50 |
| 12 | In Antisynthetase Syndrome, ACPA Are Associated With Severe and Erosive Arthritis. Medicine (United) Tj ETQqC | 0 O rgBT | /Oyerlock 10 |
| 13 | Myocarditis in Patients With Antisynthetase Syndrome. Medicine (United States), 2015, 94, e798. | 1.0 | 47 |
| 14 | "ldiopathic Eosinophilic Vasculitis― Another Side of Hypereosinophilic Syndrome? A Comprehensive Analysis of 117 Cases in Asthma-Free Patients. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 1329-1340.e3. | 3.8 | 40 |
| 15 | Seronegative polyarthritis revealing antisynthetase syndrome: a multicentre study of 40 patients. Rheumatology, 2015, 54, 927-932. | 1.9 | 37 |
| 16 | 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 |
| 17 | Role of B cells in the pathogenesis of systemic sclerosis. Revue De Medecine Interne, 2017, 38, 113-124. | 1.0 | 37 |
| 18 | Epidemiology, clinical picture and longâ€term outcomes of <i>FIP1L1â€PDGFRA</i> â€positive myeloid neoplasm with eosinophilia: Data from 151 patients. American Journal of Hematology, 2020, 95, 1314-1323. | 4.1 | 37 |

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|----|--|-----|-----------|
| 19 | Risk factors for severe bacterial infections in patients with systemic autoimmune diseases receiving rituximab. Clinical Rheumatology, 2014, 33, 799-805. | 2.2 | 35 |
| 20 | Serum free light chains of immunoglobulins as biomarkers for systemic sclerosis characteristics, activity and severity. Autoimmunity Reviews, 2014, 13, 974-980. | 5.8 | 34 |
| 21 | Severe chronic primary neutropenia in adults: report on a series of 108 patients. Blood, 2015, 126, 1643-1650. | 1.4 | 32 |
| 22 | Thrombotic events during long-term follow-up of obstetric antiphospholipid syndrome patients. Lupus, 2011, 20, 861-865. | 1.6 | 30 |
| 23 | Digestive-tract sarcoidosis. Medicine (United States), 2016, 95, e4279. | 1.0 | 30 |
| 24 | Proinflammatory B-cell profile in the early phases of MS predicts an active disease. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e431. | 6.0 | 29 |
| 25 | 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. Oncolmmunology, 2020, 9, 1722022. | 4.6 | 27 |
| 26 | Alopecia induced by tumour necrosis factor-alpha antagonists: description of 52 cases and disproportionality analysis in a nationwide pharmacovigilance database. Rheumatology, 2014, 53, 1465-1469. | 1.9 | 26 |
| 27 | Value of the Overall Pneumococcal Polysaccharide Response in the Diagnosis of Primary Humoral Immunodeficiencies. Frontiers in Immunology, 2017, 8, 1862. | 4.8 | 26 |
| 28 | 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 |
| 29 | 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 |
| 30 | Hypereosinophilia with asthma and systemic (non-vasculitic) manifestations: Eosinophilic granulomatosis with polyangiitis or hypereosinophilic syndrome?. Autoimmunity Reviews, 2017, 16, 208-209. | 5.8 | 17 |
| 31 | 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 |
| 32 | Relevance of Partitioning DLCO to Detect Pulmonary Hypertension in Systemic Sclerosis. PLoS ONE, 2013, 8, e78001. | 2.5 | 15 |
| 33 | Effective immune restoration after immunosuppressant discontinuation in a lupus patient presenting progressive multifocal leukoencephalopathy. Journal of the Neurological Sciences, 2009, 287, 246-249. | 0.6 | 14 |
| 34 | 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 |
| 35 | Episodic angioedema with eosinophilia (Gleich syndrome) in children: A clinical review. Pediatric Allergy and Immunology, 2020, 31, 297-302. | 2.6 | 14 |
| 36 | Pernicious anemia presenting as catatonia: correlating vitamin B12 levels and catatonic symptoms. General Hospital Psychiatry, 2015, 37, 273.e5-273.e7. | 2.4 | 13 |

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| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 37 | Distal ischemia as the initial presentation of hypereosinophilic syndrome-related arterial involvement: A case study and literature review. Autoimmunity Reviews, 2019, 18, 828-830. | 5.8 | 13 |
| 38 | FIP1L1-PDGFRA-Associated Hypereosinophilic Syndrome as a Treatable Cause of Watershed Infarction. Stroke, 2021, 52, e605-e609. | 2.0 | 10 |
| 39 | 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 |
| 40 | Immunoglobulin G (IgG) and IgG subclass reference intervals in children, using Optilite® reagents. Clinical Chemistry and Laboratory Medicine, 2018, 56, 1319-1327. | 2.3 | 8 |
| 41 | Successful Heart Transplantation for Unreversible Endomyocardial Fibrosis Related to FIP1L1-PDGFRA Chronic Eosinophilic Leukemia. Transplantation, 2015, 99, e176-e177. | 1.0 | 7 |
| 42 | Subcutaneous Immunoglobulin Therapy Prevents Systemic Capillary Leak Syndrome Attack. American Journal of Medicine, 2016, 129, e77-e78. | 1.5 | 7 |
| 43 | 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 |
| 44 | An appraisal of the frequency and severity of noninfectious manifestations in primary immunodeficiencies: AAstudy of a national retrospective cohort of 1375 patients over 10 years. Journal of Allergy and Clinical Immunology, 2022, 149, 2116-2125. | 2.9 | 7 |
| 45 | Recurrent cardiac arrest due to eosinophilia-related coronary vasospasm successfully treated by benralizumab. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3497-3499.e1. | 3.8 | 6 |
| 46 | Long term use of metformin in idiopathic cyclic edema, report of thirteen cases and review of the literature. Pharmacological Research, 2017, 119, 237-239. | 7.1 | 4 |
| 47 | Association Between Baseline Therapy and Flare Reduction in Mepolizumab-Treated Patients With Hypereosinophilic Syndrome. Frontiers in Immunology, 2022, 13, 840974. | 4.8 | 3 |
| 48 | Reply to Gilchrist et al. and to Musher. Clinical Infectious Diseases, 2018, 66, 637-638. | 5.8 | 1 |
| 49 | Classical pathway activity C3c, C4 and C1-inhibitor protein reference intervals determination in EDTA plasma. Biochemia Medica, 2019, 29, 559-569. | 2.7 | 1 |
| 50 | CXCL13 is expressed in various haematological disorders other than angioimmunoblastic T-cell lymphoma. Pathology Research and Practice, 2020, 216, 153004. | 2.3 | 0 |