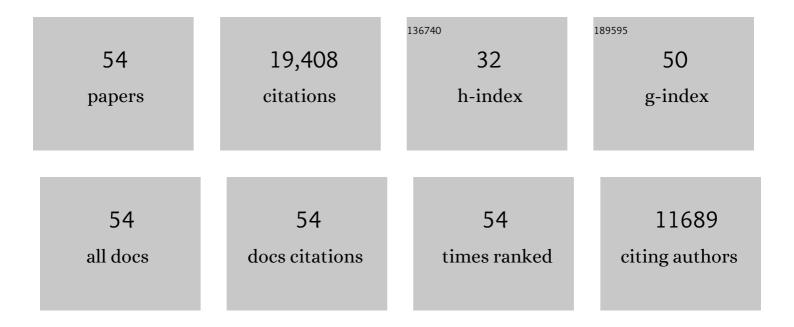
Jeffrey L Myers

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Idiopathic Pulmonary Fibrosis (an Update) and Progressive Pulmonary Fibrosis in Adults: An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2022, 205, e18-e47. | 2.5 | 780 |
| 2 | Integration and Application of Clinical Practice Guidelines for the Diagnosis of Idiopathic Pulmonary Fibrosis and Fibrotic Hypersensitivity Pneumonitis. Chest, 2022, 162, 614-629. | 0.4 | 19 |
| 3 | Utility of a Molecular Classifier as a Complement to High-Resolution Computed Tomography to Identify Usual Interstitial Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 211-220. | 2.5 | 55 |
| 4 | Cryobiopsy for Identification of Usual Interstitial Pneumonia and Other Interstitial Lung Disease Features. Further Lessons from COLDICE, a Prospective Multicenter Clinical Trial. American Journal of Respiratory and Critical Care Medicine, 2021, 203, 1306-1313. | 2.5 | 32 |
| 5 | Severe Acute Respiratory Syndrome Coronavirus 2 Surveillance in Decedents in a Large, Urban Medical Examiner's Office. Clinical Infectious Diseases, 2021, 72, e580-e585. | 2.9 | 4 |
| 6 | Interstitial lung disease pathology in systemic sclerosis. Therapeutic Advances in Musculoskeletal Disease, 2021, 13, 1759720X2110324. | 1.2 | 6 |
| 7 | Update on Diagnosing and Reporting Malignant Pleural Mesothelioma. Acta Medica Academica, 2021, 50, 197. | 0.3 | 5 |
| 8 | A contemporary practical approach to the multidisciplinary management of unclassifiable interstitial lung disease. European Respiratory Journal, 2021, 58, 2100276. | 3.1 | 19 |
| 9 | Thymoliposarcoma with sebaceous differentiation and <i>MDM2</i> amplification. Pathology International, 2021, 71, 633-635. | 0.6 | 0 |
| 10 | The Role of Surgical Lung Biopsy in the Diagnosis of Fibrotic Interstitial Lung Disease: Perspective from the Pulmonary Fibrosis Foundation. Annals of the American Thoracic Society, 2021, 18, 1601-1609. | 1,5 | 8 |
| 11 | In Reply. Archives of Pathology and Laboratory Medicine, 2021, 145, 1326-1327. | 1.2 | 1 |
| 12 | Usual Interstitial Pneumonia is the Most Common Finding in Surgical Lung Biopsies from Patients with Persistent Interstitial Lung Disease Following Infection with SARS-CoV-2. EClinicalMedicine, 2021, 42, 101209. | 3.2 | 33 |
| 13 | Diagnostic accuracy of transbronchial lung cryobiopsy for interstitial lung disease diagnosis (COLDICE): a prospective, comparative study. Lancet Respiratory Medicine,the, 2020, 8, 171-181. | 5.2 | 253 |
| 14 | Frontline Workers in the Backrooms of COVID-19. American Journal of Clinical Pathology, 2020, 154, 286-292. | 0.4 | 3 |
| 15 | Diagnosis of Hypersensitivity Pneumonitis in Adults: An Official ATS/JRS/ALAT Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2020, 202, e36-e69. | 2.5 | 508 |
| 16 | A 53-Year-Old Woman With Dyspnea, Wheezing, and Irreversible Airway Obstruction. Chest, 2020, 157, e189-e192. | 0.4 | 0 |
| 17 | A Molecular Classifier That Identifies Usual Interstitial Pneumonia in Transbronchial Biopsy Specimens of Patients With Interstitial Lung Disease. Chest, 2020, 157, 1391-1392. | 0.4 | 1 |
| 18 | Diffuse alveolar damage (DAD) resulting from coronavirus disease 2019 Infection is Morphologically Indistinguishable from Other Causes of DAD. Histopathology, 2020, 77, 570-578. | 1.6 | 162 |

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|----|--|-----|-----------|
| 19 | Postmortem Lung Findings in a Patient With Asthma and Coronavirus Disease 2019. Chest, 2020, 158, e99-e101. | 0.4 | 79 |
| 20 | Humoral immune responses mediate the development of a restrictive phenotype of chronic lung allograft dysfunction. JCI Insight, 2020, 5, . | 2.3 | 17 |
| 21 | The 2018 Diagnosis of Idiopathic Pulmonary Fibrosis Guidelines: Surgical Lung Biopsy for Radiological Pattern of Probable Usual Interstitial Pneumonia Is Not Mandatory. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1089-1092. | 2.5 | 45 |
| 22 | Use of a molecular classifier to identify usual interstitial pneumonia in conventional transbronchial lung biopsy samples: a prospective validation study. Lancet Respiratory Medicine,the, 2019, 7, 487-496. | 5.2 | 119 |
| 23 | The characterisation of interstitial lungÂdisease multidisciplinary team meetings:ÂaÂglobal study. ERJ Open Research, 2019, 5, 00209-2018. | 1.1 | 49 |
| 24 | Reply to Moodley and to Ravaglia et al American Journal of Respiratory and Critical Care Medicine, 2019, 199, 667-669. | 2.5 | 0 |
| 25 | Hypersensitivity Pneumonitis. Chest, 2019, 155, 699-711. | 0.4 | 148 |
| 26 | Cryobiopsy versus open lung biopsy in the diagnosis of interstitial lung disease (COLDICE): protocol of a multicentre study. BMJ Open Respiratory Research, 2019, 6, e000443. | 1.2 | 17 |
| 27 | The utility of SDHB and FH immunohistochemistry in patients evaluated for hereditary paraganglioma-pheochromocytoma syndromes. Human Pathology, 2018, 71, 47-54. | 1.1 | 39 |
| 28 | Detection of 6 TFEB-amplified renal cell carcinomas and 25 renal cell carcinomas with MITF translocations: systematic morphologic analysis of 85 cases evaluated by clinical TFE3 and TFEB FISH assays. Modern Pathology, 2018, 31, 179-197. | 2.9 | 73 |
| 29 | Primary mammary analogue secretory carcinoma of the lung: a case report. Human Pathology, 2018, 74, 109-113. | 1.1 | 35 |
| 30 | A Review of Smoking-Related Interstitial Fibrosis, Respiratory Bronchiolitis, and Desquamative Interstitial Pneumonia: Overlapping Histology and Confusing Terminology. Archives of Pathology and Laboratory Medicine, 2018, 142, 1177-1181. | 1.2 | 20 |
| 31 | Diagnosis of Idiopathic Pulmonary Fibrosis. An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2018, 198, e44-e68. | 2.5 | 2,678 |
| 32 | Development and validation of a radiological diagnosis model for hypersensitivity pneumonitis. European Respiratory Journal, 2018, 52, 1800443. | 3.1 | 55 |
| 33 | A Standardized Diagnostic Ontology for Fibrotic Interstitial Lung Disease. An International Working Group Perspective. American Journal of Respiratory and Critical Care Medicine, 2017, 196, 1249-1254. | 2.5 | 166 |
| 34 | The diagnosis of idiopathic pulmonary fibrosis: current and future approaches. Lancet Respiratory Medicine,the, 2017, 5, 61-71. | 5.2 | 79 |
| 35 | Diagnosis and Treatment of Fibrotic Hypersensitivity Pneumonia. Where We Stand and Where We Need to Go. American Journal of Respiratory and Critical Care Medicine, 2017, 196, 690-699. | 2.5 | 176 |
| 36 | Usual Interstitial Pneumonia Can Be Detected in Transbronchial Biopsies Using Machine Learning. Annals of the American Thoracic Society, 2017, 14, 1646-1654. | 1.5 | 77 |

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|----|---|-----|-----------|
| 37 | Multicentre evaluation of multidisciplinary team meeting agreement on diagnosis in diffuse parenchymal lung disease: a case-cohort study. Lancet Respiratory Medicine,the, 2016, 4, 557-565. | 5.2 | 337 |
| 38 | Acute Exacerbation of Idiopathic Pulmonary Fibrosis. An International Working Group Report. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 265-275. | 2.5 | 1,006 |
| 39 | Histopathology of Explanted Lungs From Patients With a Diagnosis of Pulmonary Sarcoidosis. Chest, 2016, 149, 499-507. | 0.4 | 38 |
| 40 | Classification of usual interstitial pneumonia in patients with interstitial lung disease: assessment of a machine learning approach using high-dimensional transcriptional data. Lancet Respiratory Medicine,the, 2015, 3, 473-482. | 5.2 | 112 |
| 41 | An Official ATS/ERS/JRS/ALAT Clinical Practice Guideline: Treatment of Idiopathic Pulmonary Fibrosis. An Update of the 2011 Clinical Practice Guideline. American Journal of Respiratory and Critical Care Medicine, 2015, 192, e3-e19. | 2.5 | 1,521 |
| 42 | Evaluation of Napsin A, TTF-1, p63, p40, and CK5/6 Immunohistochemical Stains in Pulmonary Neuroendocrine Tumors. American Journal of Clinical Pathology, 2014, 142, 320-324. | 0.4 | 48 |
| 43 | Response. Chest, 2014, 145, 434. | 0.4 | Ο |
| 44 | An Official American Thoracic Society/European Respiratory Society Statement: Update of the International Multidisciplinary Classification of the Idiopathic Interstitial Pneumonias. American Journal of Respiratory and Critical Care Medicine, 2013, 188, 733-748. | 2.5 | 3,134 |
| 45 | Hypersensitivity Pneumonia: Role of Surgical Lung Biopsy. Archives of Pathology and Laboratory Medicine, 2012, 136, 889-895. | 1.2 | 11 |
| 46 | Life-Long Learning and Self-Assessment. Archives of Pathology and Laboratory Medicine, 2012, 136, 851-853. | 1.2 | 6 |
| 47 | Hypersensitivity pneumonia: the role of lung biopsy in diagnosis and management. Modern Pathology, 2012, 25, S58-S67. | 2.9 | 65 |
| 48 | An Official ATS/ERS/JRS/ALAT Statement: Idiopathic Pulmonary Fibrosis: Evidence-based Guidelines for Diagnosis and Management. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 788-824. | 2.5 | 6,033 |
| 49 | Role of Surgical Lung Biopsy in Separating Chronic Hypersensitivity Pneumonia From Usual Interstitial Pneumonia/Idiopathic Pulmonary Fibrosis*: Analysis of 31 Biopsies From 15 Patients. Chest, 2008, 134, 126-132. | 0.4 | 112 |
| 50 | Acute Exacerbations of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 636-643. | 2.5 | 996 |
| 51 | Idiopathic Pulmonary Fibrosis in Patients Younger Than 50 Years. Mayo Clinic Proceedings, 2005, 80, 37-40. | 1.4 | 39 |
| 52 | Idiopathic Pulmonary Fibrosis in Patients Younger Than 50 Years. Mayo Clinic Proceedings, 2005, 80, 37-40. | 1.4 | 30 |
| 53 | Airway-centered Interstitial Fibrosis. American Journal of Surgical Pathology, 2004, 28, 62-68. | 2.1 | 120 |
| 54 | Organizing Diffuse Alveolar Damage Associated With Progressive Systemic Sclerosis. Mayo Clinic Proceedings, 1997, 72, 639-642. | 1.4 | 39 |