

Bernadette Gochuico

List of Publications by Year in descending order

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Version: 2024-02-01

64
papers

3,061
citations

186265
28
h-index

161849
54
g-index

67
all docs

67
docs citations

67
times ranked

3830
citing authors

#	ARTICLE	IF	CITATIONS
1	Progressive Preclinical Interstitial Lung Disease in Rheumatoid Arthritis. Archives of Internal Medicine, 2008, 168, 159.	3.8	319
2	Danazol Treatment for Telomere Diseases. New England Journal of Medicine, 2016, 374, 1922-1931.	27.0	300
3	Effect of pirfenidone on the pulmonary fibrosis of Hermansky-Pudlak syndrome. Molecular Genetics and Metabolism, 2002, 76, 234-242.	1.1	185
4	The clinical spectrum of Erdheim-Chester disease: an observational cohort study. Blood Advances, 2017, 1, 357-366.	5.2	163
5	Early Interstitial Lung Disease in Familial Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 698-705.	5.6	157
6	Genetic and hypoxic alterations of the micro RNA $\alpha 210$ β 1 axis promote iron-sulfur deficiency and pulmonary hypertension. EMBO Molecular Medicine, 2015, 7, 695-713.	6.9	120
7	Identification of Citrullinated Hsp90 Isoforms as Novel Autoantigens in Rheumatoid Arthritis-associated Interstitial Lung Disease. Arthritis and Rheumatism, 2013, 65, 869-879.	6.7	113
8	High attenuation areas on chest computed tomography in community-dwelling adults: the MESA study. European Respiratory Journal, 2016, 48, 1442-1452.	6.7	110
9	Interstitial Lung Disease and Pulmonary Fibrosis in Hermansky-Pudlak Syndrome Type 2, an Adaptor Protein-3 Complex Disease. Molecular Medicine, 2012, 18, 56-64.	4.4	86
10	Erdheim-Chester Disease: A Rare Multisystem Histiocytic Disorder Associated with Interstitial Lung Disease. American Journal of the Medical Sciences, 2001, 321, 66-75.	1.1	84
11	Epithelial Stress and Apoptosis Underlie Hermansky-Pudlak Syndrome-associated Interstitial Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2010, 182, 207-219.	5.6	83
12	Alveolar Macrophage Dysregulation in Hermansky-Pudlak Syndrome Type 1. American Journal of Respiratory and Critical Care Medicine, 2009, 180, 1114-1121.	5.6	71
13	Hermansky-Pudlak syndrome: Mutation update. Human Mutation, 2020, 41, 543-580.	2.5	65
14	Pirfenidone for the treatment of Hermansky-Pudlak syndrome pulmonary fibrosis. Molecular Genetics and Metabolism, 2011, 103, 128-134.	1.1	64
15	Cannabinoid CB1 receptor overactivity contributes to the pathogenesis of idiopathic pulmonary fibrosis. JCI Insight, 2017, 2, .	5.0	59
16	Early Alveolar Epithelial Dysfunction Promotes Lung Inflammation in a Mouse Model of Hermansky-Pudlak Syndrome. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 449-458.	5.6	56
17	Chitinase 3-like 1 and its receptors in Hermansky-Pudlak syndrome-associated lung disease. Journal of Clinical Investigation, 2015, 125, 3178-3192.	8.2	54
18	ADP-ribosyltransferase-specific Modification of Human Neutrophil Peptide-1. Journal of Biological Chemistry, 2006, 281, 17054-17060.	3.4	52

#	ARTICLE	IF	CITATIONS
19	Expanding the phenotype of COPA syndrome: a kindred with typical and atypical features. <i>Journal of Medical Genetics</i> , 2019, 56, 778-782.	3.2	49
20	Natural History of Pulmonary Fibrosis in Two Subjects With the Same Telomerase Mutation. <i>Chest</i> , 2011, 139, 1203-1209.	0.8	47
21	Glucose Transporter-1 Distribution in Fibrotic Lung Disease. <i>Chest</i> , 2013, 143, 1685-1691.	0.8	47
22	Automated Quantification of High-Resolution CT Scan Findings in Individuals at Risk for Pulmonary Fibrosis. <i>Chest</i> , 2011, 140, 1590-1597.	0.8	46
23	Mononuclear Phagocytes and Airway Epithelial Cells: Novel Sources of Matrix Metalloproteinase-8 (MMP-8) in Patients with Idiopathic Pulmonary Fibrosis. <i>PLoS ONE</i> , 2014, 9, e97485.	2.5	42
24	Dysregulation of Galectin-3. Implications for Hermansky-Pudlak Syndrome Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2014, 50, 605-613.	2.9	42
25	Galectin-3 Interacts with the CHI3L1 Axis and Contributes to Hermansky-Pudlak Syndrome Lung Disease. <i>Journal of Immunology</i> , 2018, 200, 2140-2153.	0.8	38
26	Circulating Fibrocytes as Biomarker of Prognosis in Hermansky-Pudlak Syndrome. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2014, 190, 1395-1401.	5.6	36
27	Transforming Growth Factor- β 1 Downregulates Vascular Endothelial Growth Factor-D Expression in Human Lung Fibroblasts via the Jun NH2-Terminal Kinase Signaling Pathway. <i>Molecular Medicine</i> , 2014, 20, 120-134.	4.4	33
28	Identification of <i>Helicobacter pylori</i> VacA in human lung and its effects on lung cells. <i>Biochemical and Biophysical Research Communications</i> , 2015, 460, 721-726.	2.1	32
29	Extensive Citrullination Promotes Immunogenicity of HSP90 through Protein Unfolding and Exposure of Cryptic Epitopes. <i>Journal of Immunology</i> , 2016, 197, 1926-1936.	0.8	32
30	Clinical, Molecular, and Cellular Features of Non-Puerto Rican Hermansky-Pudlak Syndrome Patients of Hispanic Descent. <i>Journal of Investigative Dermatology</i> , 2011, 131, 2394-2400.	0.7	31
31	Anti-citrullinated heat shock protein 90 antibodies identified in bronchoalveolar lavage fluid are a marker of lung-specific immune responses. <i>Clinical Immunology</i> , 2014, 155, 60-70.	3.2	30
32	Clinical management and outcomes of patients with Hermansky-Pudlak syndrome pulmonary fibrosis evaluated for lung transplantation. <i>PLoS ONE</i> , 2018, 13, e0194193.	2.5	29
33	MAP1LC3B overexpression protects against Hermansky-Pudlak syndrome type-1-induced defective autophagy in vitro. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2016, 310, L519-L531.	2.9	25
34	Clinical and molecular phenotyping of a child with Hermansky-Pudlak syndrome-7, an uncommon genetic type of HPS. <i>Molecular Genetics and Metabolism</i> , 2017, 120, 378-383.	1.1	25
35	CA-125 in Disease Progression and Treatment of Lymphangiioleiomyomatosis. <i>Chest</i> , 2018, 153, 339-348.	0.8	25
36	Prolonged treatment with open-label pirfenidone in Hermansky-Pudlak syndrome pulmonary fibrosis. <i>Molecular Genetics and Metabolism</i> , 2018, 125, 168-173.	1.1	24

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37	Impairment of Alveolar Macrophage Transcription in Idiopathic Pulmonary Fibrosis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2007, 175, 1151-1157.	5.6	23
38	Neurological manifestations of Erdheim-Chester Disease. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 497-506.	3.7	23
39	The Immunome in Two Inherited Forms of Pulmonary Fibrosis. <i>Frontiers in Immunology</i> , 2018, 9, 76.	4.8	19
40	Hermansky-Pudlak syndrome pulmonary fibrosis: a rare inherited interstitial lung disease. <i>European Respiratory Review</i> , 2021, 30, 200193.	7.1	18
41	Natural killer cell activity and dysfunction in Hermansky-Pudlak syndrome. <i>British Journal of Haematology</i> , 2017, 176, 118-123.	2.5	17
42	Matrix metalloproteinase activity in the lung is increased in Hermansky-Pudlak syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 162.	2.7	17
43	Bleomycin Induces Drug Efflux in Lungs. A Pitfall for Pharmacological Studies of Pulmonary Fibrosis. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2020, 62, 178-190.	2.9	16
44	CB ₁ R and iNOS are distinct players promoting pulmonary fibrosis in Hermansky-Pudlak syndrome. <i>Clinical and Translational Medicine</i> , 2021, 11, e471.	4.0	16
45	Immunophenotypic and Ultrastructural Analysis of Mast Cells in Hermansky-Pudlak Syndrome Type-1: A Possible Connection to Pulmonary Fibrosis. <i>PLoS ONE</i> , 2016, 11, e0159177.	2.5	15
46	Hermansky-Pudlak syndrome and oculocutaneous albinism in Chinese children with pigmentation defects and easy bruising. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 52.	2.7	13
47	Automated Digital Quantification of Pulmonary Fibrosis in Human Histopathology Specimens. <i>Frontiers in Medicine</i> , 2021, 8, 607720.	2.6	13
48	Genetic variants associated with Hermansky-Pudlak syndrome. <i>Platelets</i> , 2020, 31, 544-547.	2.3	12
49	Cellular and molecular defects in a patient with Hermansky-Pudlak syndrome type 5. <i>PLoS ONE</i> , 2017, 12, e0173682.	2.5	11
50	Clinical and Histopathologic Features of Interstitial Lung Disease in Erdheim-Chester Disease. <i>Journal of Clinical Medicine</i> , 2018, 7, 243.	2.4	11
51	In vitro functional correction of Hermansky-Pudlak Syndrome type-1 by lentiviral-mediated gene transfer. <i>Molecular Genetics and Metabolism</i> , 2015, 114, 62-65.	1.1	10
52	Inflammatory bowel disease in Hermansky-Pudlak syndrome: a retrospective single-centre cohort study. <i>Journal of Internal Medicine</i> , 2021, 290, 129-140.	6.0	10
53	Identification of a novel mutation in HPS6 in a patient with hemophilia B and oculocutaneous albinism. <i>Molecular Genetics and Metabolism</i> , 2016, 119, 284-287.	1.1	9
54	Severe bleeding with subclinical oculocutaneous albinism in a patient with a novel HPS6 missense variant. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 2819-2823.	1.2	7

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55	Hermansky-Pudlak syndrome-2 alters mitochondrial homeostasis in the alveolar epithelium of the lung. <i>Respiratory Research</i> , 2021, 22, 49.	3.6	5
56	Prevalence of Hypothyroidism in Patients With Erdheim-Chester Disease. <i>JAMA Network Open</i> , 2020, 3, e2019169.	5.9	4
57	Pituitary Imaging Abnormalities and Related Endocrine Disorders in Erdheim-Chester Disease. <i>Cancers</i> , 2021, 13, 4126.	3.7	4
58	Diagnosis of Chediak Higashi disease in a 67-year old woman. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 3007-3013.	1.2	4
59	A Mixed Blood-Lymphatic Endothelial Cell Phenotype in Lymphangioliomyomatosis and Idiopathic Pulmonary Fibrosis but Not in Kaposi's Sarcoma or Tuberous Sclerosis Complex. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2022, 66, 337-340.	2.9	3
60	Dysregulated myosin in Hermansky-Pudlak syndrome lung fibroblasts is associated with increased cell motility. <i>Respiratory Research</i> , 2022, 23, .	3.6	2
61	The curse of idiopathic. <i>Journal of Inherited Metabolic Disease</i> , 2018, 41, 3-4.	3.6	1
62	Progressive pulmonary fibrosis in a murine model of Hermansky-Pudlak syndrome. <i>Respiratory Research</i> , 2022, 23, 112.	3.6	1
63	A comprehensive, multidisciplinary, precision medicine approach to discover effective therapy for an undiagnosed, progressive, fibroinflammatory disease. <i>Translational Research</i> , 2020, 215, 31-40.	5.0	0
64	Novel Hermansky-Pudlak Syndrome Type 6 Missense Variant Associated with Subclinical Oculocutaneous Albinism and Mild Bleeding. <i>Blood</i> , 2018, 132, 1153-1153.	1.4	0