

David A Lomas

List of Publications by Year in descending order

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95
papers

9,387
citations

71102

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99
docs citations

99
times ranked

8269
citing authors

#	ARTICLE	IF	CITATIONS
1	Hepatobiliary phenotypes of adults with alpha-1 antitrypsin deficiency. <i>Gut</i> , 2022, 71, 415-423.	12.1	28
2	Alpha-1 Antitrypsin MZ Heterozygosity Is an Endotype of Chronic Obstructive Pulmonary Disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2022, 205, 313-323.	5.6	21
3	The molecular species responsible for α_1 antitrypsin deficiency are suppressed by a small molecule chaperone. <i>FEBS Journal</i> , 2021, 288, 2222-2237.	4.7	8
4	Conversion of the death inhibitor ARC to a killer activates pancreatic β cell death in diabetes. <i>Developmental Cell</i> , 2021, 56, 747-760.e6.	7.0	8
5	InforMing the PATHway of COPD Treatment (IMPACT) trial: fibrinogen levels predict risk of moderate or severe exacerbations. <i>Respiratory Research</i> , 2021, 22, 130.	3.6	9
6	The Importance of N186 in the Alpha-1-Antitrypsin Shutter Region Is Revealed by the Novel Bologna Deficiency Variant. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5668.	4.1	5
7	Quantitative 18 F-fluorodeoxyglucose positron emission tomography/computed tomography to assess pulmonary inflammation in COPD. <i>ERJ Open Research</i> , 2021, 7, 00699-2020.	2.6	2
8	Scaling Concepts in Serpin Polymer Physics. <i>Materials</i> , 2021, 14, 2577.	2.9	4
9	The development of highly potent and selective small molecule correctors of Z α_1 -antitrypsin misfolding. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 41, 127973.	2.2	9
10	Development of a small molecule that corrects misfolding and increases secretion of Z α_1 antitrypsin. <i>EMBO Molecular Medicine</i> , 2021, 13, e13167.	6.9	33
11	The structural basis for Z α_1 -antitrypsin polymerization in the liver. <i>Science Advances</i> , 2020, 6, .	10.3	26
12	High-resolution ex vivo NMR spectroscopy of human Z α_1 -antitrypsin. <i>Nature Communications</i> , 2020, 11, 6371.	12.8	15
13	Heme metabolism genes Downregulated in COPD Cachexia. <i>Respiratory Research</i> , 2020, 21, 100.	3.6	4
14	Alpha α_1 -Antitrypsin Deficiency. <i>New England Journal of Medicine</i> , 2020, 382, 1443-1455.	27.0	269
15	Intrahepatic heteropolymerization of M and Z alpha-1-antitrypsin. <i>JCI Insight</i> , 2020, 5, .	5.0	16
16	<i>C. elegans</i> expressing D76N β 2-microglobulin: a model for in vivo screening of drug candidates targeting amyloidosis. <i>Scientific Reports</i> , 2019, 9, 19960.	3.3	14
17	Calcium signalling in mammalian cell lines expressing wild type and mutant human α_1 -Antitrypsin. <i>Scientific Reports</i> , 2019, 9, 17293.	3.3	3
18	Characterisation of a type II functionally-deficient variant of alpha-1-antitrypsin discovered in the general population. <i>PLoS ONE</i> , 2019, 14, e0206955.	2.5	13

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19	Once-Daily Single-Inhaler Triple versus Dual Therapy in Patients with COPD. <i>New England Journal of Medicine</i> , 2018, 378, 1671-1680.	27.0	823
20	Fibrinogen does not relate to cardiovascular or muscle manifestations in COPD: cross-sectional data from the ERICA study. <i>Thorax</i> , 2018, 73, 1182-1185.	5.6	9
21	Heteropolymerization of α_1 -antitrypsin mutants in cell models mimicking heterozygosity. <i>Human Molecular Genetics</i> , 2018, 27, 1785-1793.	2.9	24
22	An automated high-throughput system for phenotypic screening of chemical libraries on <i>C. elegans</i> and parasitic nematodes. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2018, 8, 8-21.	3.4	71
23	In Vitro Approaches for the Assessment of Serpin Polymerization. <i>Methods in Molecular Biology</i> , 2018, 1826, 87-107.	0.9	0
24	α_1 -Antitrypsin Polymerizes in Alveolar Macrophages of Smokers With and Without α_1 -Antitrypsin Deficiency. <i>Chest</i> , 2018, 154, 607-616.	0.8	22
25	The p38 mitogen activated protein kinase inhibitor losmapimod in chronic obstructive pulmonary disease patients with systemic inflammation, stratified by fibrinogen: A randomised double-blind placebo-controlled trial. <i>PLoS ONE</i> , 2018, 13, e0194197.	2.5	23
26	New Therapeutic Targets for Alpha-1 Antitrypsin Deficiency. <i>Chronic Obstructive Pulmonary Diseases (Miami, Fla)</i> , 2018, 5, 233-243.	0.7	18
27	Taking out the JuNK to treat α_1 -antitrypsin deficiency. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2017, 14, 201-202.	17.8	1
28	The pathological Trento variant of alpha α_1 -antitrypsin (E75V) shows nonclassical behaviour during polymerization. <i>FEBS Journal</i> , 2017, 284, 2110-2126.	4.7	23
29	Body mass index change in gastrointestinal cancer and chronic obstructive pulmonary disease is associated with Dedicator of Cytokines 1. <i>Journal of Cachexia, Sarcopenia and Muscle</i> , 2017, 8, 428-436.	7.3	13
30	Sex-Based Genetic Association Study Identifies <i>CELSR1</i> as a Possible Chronic Obstructive Pulmonary Disease Risk Locus among Women. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2017, 56, 332-341.	2.9	28
31	An antibody that prevents serpin polymerisation acts by inducing a novel allosteric behaviour. <i>Biochemical Journal</i> , 2016, 473, 3269-3290.	3.7	15
32	Update on alpha-1 antitrypsin deficiency: New therapies. <i>Journal of Hepatology</i> , 2016, 65, 413-424.	3.7	66
33	The endoplasmic reticulum remains functionally connected by vesicular transport after its fragmentation in cells expressing α_1 -antitrypsin. <i>FASEB Journal</i> , 2016, 30, 4083-4097.	0.5	22
34	An ECLIPSE View of Alpha-1 Antitrypsin Deficiency. <i>Annals of the American Thoracic Society</i> , 2016, 13, S326-S331.	3.2	11
35	α_1 -Antitrypsin deficiency. <i>Nature Reviews Disease Primers</i> , 2016, 2, 16051.	30.5	215
36	Circulating desmosine levels do not predict emphysema progression but are associated with cardiovascular risk and mortality in COPD. <i>European Respiratory Journal</i> , 2016, 47, 1365-1373.	6.7	64

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37	Polymers of Z α 1-antitrypsin are secreted in cell models of disease. <i>European Respiratory Journal</i> , 2016, 47, 1005-1009.	6.7	41
38	Deficiency Mutations of Alpha-1 Antitrypsin. Effects on Folding, Function, and Polymerization. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2016, 54, 71-80.	2.9	31
39	Does Protease-Antiprotease Imbalance Explain Chronic Obstructive Pulmonary Disease?. <i>Annals of the American Thoracic Society</i> , 2016, 13 Suppl 2, S130-7.	3.2	25
40	An integrative approach combining ion mobility mass spectrometry, X-ray crystallography, and nuclear magnetic resonance spectroscopy to study the conformational dynamics of α 1-antitrypsin upon ligand binding. <i>Protein Science</i> , 2015, 24, 1301-1312.	7.6	37
41	Characterising the association of latency with α 1-antitrypsin polymerisation using a novel monoclonal antibody. <i>International Journal of Biochemistry and Cell Biology</i> , 2015, 58, 81-91.	2.8	26
42	Circulating Soluble Receptor for Advanced Glycation End Products (sRAGE) as a Biomarker of Emphysema and the RAGE Axis in the Lung. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2015, 192, 785-792.	5.6	82
43	An antibody raised against a pathogenic serpin variant induces mutant-like behaviour in the wild-type protein. <i>Biochemical Journal</i> , 2015, 468, 99-108.	3.7	22
44	A single-chain variable fragment intrabody prevents intracellular polymerization of Z α 1-antitrypsin while allowing its antiproteinase activity. <i>FASEB Journal</i> , 2015, 29, 2667-2678.	0.5	44
45	The TRiC/CCT Chaperone Is Implicated in Alzheimer's Disease Based on Patient GWAS and an RNAi Screen in Δ 2-Expressing <i>Caenorhabditis elegans</i> . <i>PLoS ONE</i> , 2014, 9, e102985.	2.5	34
46	Altered native stability is the dominant basis for susceptibility of α 1-antitrypsin mutants to polymerization. <i>Biochemical Journal</i> , 2014, 460, 103-119.	3.7	25
47	Circulating polymers in α 1-antitrypsin deficiency. <i>European Respiratory Journal</i> , 2014, 43, 1501-1504.	6.7	69
48	S51 Circulating Desmosine Relates To Cardiovascular Comorbidity, Coronary Artery Calcification Score (cacs), Systemic Inflammation And Mortality In Patients With Copd. <i>Thorax</i> , 2014, 69, A28-A29.	5.6	1
49	Coronary artery calcification is increased in patients with COPD and associated with increased morbidity and mortality. <i>Thorax</i> , 2014, 69, 718-723.	5.6	151
50	The molecular and cellular pathology of α 1-antitrypsin deficiency. <i>Trends in Molecular Medicine</i> , 2014, 20, 116-127.	6.7	98
51	Lessons from ECLIPSE: a review of COPD biomarkers. <i>Thorax</i> , 2014, 69, 666-672.	5.6	125
52	Susceptibility to Chronic Mucus Hypersecretion, a Genome Wide Association Study. <i>PLoS ONE</i> , 2014, 9, e91621.	2.5	25
53	p53 and Translation Attenuation Regulate Distinct Cell Cycle Checkpoints during Endoplasmic Reticulum (ER) Stress. <i>Journal of Biological Chemistry</i> , 2013, 288, 7606-7617.	3.4	35
54	Twenty Years of Polymers: A Personal Perspective on Alpha-1 Antitrypsin Deficiency. <i>COPD: Journal of Chronic Obstructive Pulmonary Disease</i> , 2013, 10, 17-25.	1.6	27

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55	Reactive centre loop mutants of α_1 -antitrypsin reveal position-specific effects on intermediate formation along the polymerization pathway. <i>Bioscience Reports</i> , 2013, 33, .	2.4	24
56	Endoplasmic reticulum polymers impair luminal protein mobility and sensitize to cellular stress in α_1 -antitrypsin deficiency. <i>Hepatology</i> , 2013, 57, 2049-2060.	7.3	108
57	An Oral Inhibitor of p38 MAP Kinase Reduces Plasma Fibrinogen in Patients With Chronic Obstructive Pulmonary Disease. <i>Journal of Clinical Pharmacology</i> , 2012, 52, 416-424.	2.0	99
58	Structural Dynamics Associated with Intermediate Formation in an Archetypal Conformational Disease. <i>Structure</i> , 2012, 20, 504-512.	3.3	33
59	Targeted gene correction of α_1 -antitrypsin deficiency in induced pluripotent stem cells. <i>Nature</i> , 2011, 478, 391-394.	27.8	635
60	Characterisation of serpin polymers in vitro and in vivo. <i>Methods</i> , 2011, 53, 255-266.	3.8	31
61	Therapeutic target-site variability in α_1 -antitrypsin characterized at high resolution. <i>Acta Crystallographica Section F: Structural Biology Communications</i> , 2011, 67, 1492-1497.	0.7	39
62	The Serpinopathies. <i>Methods in Enzymology</i> , 2011, 501, 421-466.	1.0	35
63	A novel monoclonal antibody to characterize pathogenic polymers in liver disease associated with α_1 -antitrypsin deficiency. <i>Hepatology</i> , 2010, 52, 1078-1088.	7.3	138
64	Defining the mechanism of polymerization in the serpinopathies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 17146-17151.	7.1	135
65	25th Anniversary Transatlantic Airway Conference: Protein Misfolding and Obstructive Lung Disease. <i>Proceedings of the American Thoracic Society</i> , 2010, 7, 343-345.	3.5	1
66	Conformational Pathology of the Serpins: Themes, Variations, and Therapeutic Strategies. <i>Annual Review of Biochemistry</i> , 2009, 78, 147-176.	11.1	239
67	Crystallographic and Cellular Characterisation of Two Mechanisms Stabilising the Native Fold of α_1 -Antitrypsin: Implications for Disease and Drug Design. <i>Journal of Molecular Biology</i> , 2009, 387, 857-868.	4.2	34
68	Sex Differences in Emphysema and Airway Disease in Smokers. <i>Chest</i> , 2009, 136, 1480-1488.	0.8	88
69	The intracellular accumulation of polymeric neuroserpin explains the severity of the dementia FENIB. <i>Human Molecular Genetics</i> , 2008, 17, 1527-1539.	2.9	95
70	Genetic predisposition to chronic obstructive pulmonary disease: advances in α_1 -antitrypsin deficiency and the serpinopathies. <i>Clinical Medicine</i> , 2007, 7, 446-447.	1.9	7
71	Parker B. Francis Lectureship. Antitrypsin Deficiency, the Serpinopathies, and Chronic Obstructive Pulmonary Disease. <i>Proceedings of the American Thoracic Society</i> , 2006, 3, 499-501.	3.5	11
72	The Selective Advantage of α_1 -Antitrypsin Deficiency. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2006, 173, 1072-1077.	5.6	79

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73	Molecular mousetraps, α_1 -antitrypsin deficiency and the serpinopathies. <i>Clinical Medicine</i> , 2005, 5, 249-257.	1.9	45
74	Latent S49P Neuroserpin Forms Polymers in the Dementia Familial Encephalopathy with Neuroserpin Inclusion Bodies. <i>Journal of Biological Chemistry</i> , 2005, 280, 13735-13741.	3.4	51
75	Polymers of Z α_1 -Antitrypsin Co-Localize with Neutrophils in Emphysematous Alveoli and Are Chemotactic in Vivo. <i>American Journal of Pathology</i> , 2005, 166, 377-386.	3.8	180
76	Mutants of Neuroserpin That Cause Dementia Accumulate as Polymers within the Endoplasmic Reticulum. <i>Journal of Biological Chemistry</i> , 2004, 279, 28283-28291.	3.4	102
77	Targeting a Surface Cavity of α_1 -Antitrypsin to Prevent Conformational Disease. <i>Clinical Science</i> , 2003, 104, 57P-57P.	0.0	0
78	Serpinopathies and the conformational dementias. <i>Nature Reviews Genetics</i> , 2002, 3, 759-768.	16.3	211
79	α_1 -Antitrypsin polymerization and the serpinopathies: pathobiology and prospects for therapy. <i>Journal of Clinical Investigation</i> , 2002, 110, 1585-1590.	8.2	199
80	α_1 -Antitrypsin polymerization and the serpinopathies: pathobiology and prospects for therapy. <i>Journal of Clinical Investigation</i> , 2002, 110, 1585-1590.	8.2	134
81	Polymerization of Plasminogen Activator Inhibitor-1. <i>Journal of Biological Chemistry</i> , 2001, 276, 9115-9122.	3.4	52
82	A novel polymorphism (471C>T) in alpha-1-antitrypsin in a patient with asthma. <i>Human Mutation</i> , 2001, 17, 155-156.	2.5	5
83	Characterization of a new variant of α_1 -antitrypsin EJohannesburg (H15N) in association with asthma. <i>Human Mutation</i> , 2001, 17, 156-156.	2.5	7
84	The Serpins Are an Expanding Superfamily of Structurally Similar but Functionally Diverse Proteins. <i>Journal of Biological Chemistry</i> , 2001, 276, 33293-33296.	3.4	1,069
85	Topography of a 2.0 Å... structure of α_1 -antitrypsin reveals targets for rational drug design to prevent conformational disease. <i>Protein Science</i> , 2000, 9, 1274-1281.	7.6	177
86	Familial dementia caused by polymerization of mutant neuroserpin. <i>Nature</i> , 1999, 401, 376-379.	27.8	342
87	A Kinetic Mechanism for the Polymerization of α_1 -Antitrypsin. <i>Journal of Biological Chemistry</i> , 1999, 274, 9548-9555.	3.4	205
88	A 2.6 Å structure of a serpin polymer and implications for conformational disease 1 Edited by R. Huber. <i>Journal of Molecular Biology</i> , 1999, 293, 449-455.	4.2	116
89	Heteropolymerization of S, I, and Z α_1 -antitrypsin and liver cirrhosis. <i>Journal of Clinical Investigation</i> , 1999, 103, 999-1006.	8.2	172
90	Lung Polymers in Z α_1 -Antitrypsin Deficiency-related Emphysema. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 1998, 18, 670-674.	2.9	132

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91	$\hat{\pm}1$ -Antitrypsin Deficiency. Chest, 1996, 110, 243S-247S.	0.8	52
92	Structural explanation for the deficiency of S $\hat{\pm}1$ -antitrypsin. Nature Structural Biology, 1996, 3, 910-911.	9.7	87
93	$\hat{\pm}1$ -Antitrypsin Mmalton (Phe52-deleted) Forms Loop-Sheet Polymers in Vivo.. Journal of Biological Chemistry, 1995, 270, 16864-16870.	3.4	133
94	Effect of the Z mutation on the physical and inhibitory properties of .alpha.1-antitrypsin. Biochemistry, 1993, 32, 500-508.	2.5	229
95	The mechanism of Z $\hat{\pm}1$ -antitrypsin accumulation in the liver. Nature, 1992, 357, 605-607.	27.8	1,020