Jonathan J Lyons

List of Publications by Year in descending order

65 papers	3,710 citations	186265 28 h-index	133252 59 g-index
68 all docs	68 docs citations	68 times ranked	4784 citing authors

#	Article	IF	Citations
1	Defining baseline variability of serum tryptase levels improves accuracy in identifying anaphylaxis. Journal of Allergy and Clinical Immunology, 2022, 149, 1010-1017.e10.	2.9	38
2	Elevated Basal Serum Tryptase: Disease Distribution and Variability in a Regional Health System. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 2424-2435.e5.	3.8	19
3	Resolving the genetics of human tryptases: implications for health, disease, and clinical use as a biomarker. Current Opinion in Allergy and Clinical Immunology, 2022, 22, 143-152.	2.3	6
4	Hereditary alpha-tryptasemia despite normal tryptase-encoding gene copy number owing to copy number loss in trans. Annals of Allergy, Asthma and Immunology, 2022, 128, 460-461.	1.0	4
5	Hereditary alpha-tryptasemia modifies clinical phenotypes among individuals with congenital hypermobility disorders. Human Genetics and Genomics Advances, 2022, 3, 100094.	1.7	1
6	Incorporating Tryptase Genotyping Into the Workup and Diagnosis of Mast Cell Diseases and Reactions. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 1964-1973.	3.8	17
7	Heritable risk for severe anaphylaxis associated with increased Î±-tryptaseâ€“encoding germline copy number at TPSAB1. Journal of Allergy and Clinical Immunology, 2021, 147, 622-632.	2.9	137
8	Patients with mast cell activation symptoms and elevated baseline serum tryptase level have unique bone marrow morphology. Journal of Allergy and Clinical Immunology, 2021, 147, 1497-1501.e1.	2.9	34
9	NovelPGM3compound heterozygous variants with IgEâ€related dermatitis, lymphopenia, without syndromic features. Pediatric Allergy and Immunology, 2021, 32, 566-575.	2.6	6
10	Distinct Small Intestine Mast Cell Histologic Changes in Patients With Hereditary Alpha-tryptasemia and Mast Cell Activation Syndrome. American Journal of Surgical Pathology, 2021, 45, 997-1004.	3.7	24
11	Clinical Impact of Inherited and Acquired Genetic Variants in Mastocytosis. International Journal of Molecular Sciences, 2021, 22, 411.	4.1	21
12	SASH3variants cause a novel form of X-linked combined immunodeficiency with immune dysregulation. Blood, 2021, 138, 1019-1033.	1.4	28
13	Hereditary Alpha-Tryptasemia: a Commonly Inherited Modifier of Anaphylaxis. Current Allergy and Asthma Reports, 2021, 21, 33.	5.3	19
14	The Genetic Basis and Clinical Impact of Hereditary Alpha-Tryptasemia. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2235-2242.	3.8	38
15	Selecting the Right Criteria and Proper Classification to Diagnose Mast Cell Activation Syndromes: A Critical Review. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3918-3928.	3.8	33
16	Assessment of Osteoporosis and Fracture Risk in Mastocytosis within a North American Cohort. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 4459-4467.e10.	3.8	6
17	Routine KIT p.D816V screening identifies clonal mast cell disease in patients with Hymenoptera allergy regularly missed using baseline tryptase levels alone. Journal of Allergy and Clinical Immunology, 2021, 148, 621-626.e7.	2.9	27
18	Clinical relevance of inherited genetic differences in human tryptases. Annals of Allergy, Asthma and Immunology, 2021, 127, 638-647.	1.0	30

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19	Small intestinal immunopathology and GI-associated antibody formation in hereditary alpha-tryptasemia. Journal of Allergy and Clinical Immunology, 2021, 148, 813-821.e7.	2.9	17
20	On the complexities of tryptase geneticsÂand impact on clinical phenotypes. Journal of Allergy and Clinical Immunology, 2021, 148, 1342-1343.	2.9	9
21	Mast cell tryptases in allergic inflammation and immediate hypersensitivity. Current Opinion in Immunology, 2021, 72, 94-106.	5.5	26
22	Inherited and acquired determinants of serum tryptase levels in humans. Annals of Allergy, Asthma and Immunology, 2021, 127, 420-426.	1.0	26
23	Updated Diagnostic Criteria and Classification of Mast Cell Disorders: A Consensus Proposal. HemaSphere, 2021, 5, e646.	2.7	128
24	Skewed Lymphocyte Subpopulations and Associated Phenotypes in Patients with Mastocytosis. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 292-301.e2.	3.8	5
25	Compound Heterozygous PGM3 Mutations in a Thai Patient with a Specific Antibody Deficiency Requiring Monthly IVIG Infusions. Journal of Clinical Immunology, 2020, 40, 227-231.	3.8	4
26	Clinical response to omalizumab in patients with hereditary Î±-tryptasemia. Annals of Allergy, Asthma and Immunology, 2020, 124, 99-100.e1.	1.0	18
27	Targeting Mast Cells with Biologics. Immunology and Allergy Clinics of North America, 2020, 40, 667-685.	1.9	14
28	Hymenoptera venom-induced anaphylaxis and hereditary alpha-tryptasemia. Current Opinion in Allergy and Clinical Immunology, 2020, 20, 431-437.	2.3	30
29	Usefulness of testing for hereditary alpha tryptasemia in symptomatic patients with elevated tryptase. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 2066-2067.	3.8	19
30	Clinical Approach to a Patient with Elevated Serum Tryptase: Implications of Acute Versus Basally Elevated Levels. , 2020, , 35-54.		3
31	Impact of naturally forming human Î±/Î²-tryptase heterotetramers in the pathogenesis of hereditary Î±-tryptasemia. Journal of Experimental Medicine, 2019, 216, 2348-2361.	8.5	85
32	Why the 20% + 2 Tryptase Formula Is a Diagnostic Gold Standard for Severe Systemic Mast Cell Activation and Mast Cell Activation Syndrome. International Archives of Allergy and Immunology, 2019, 180, 44-51.	2.1	87
33	GATA-2â€“deficient mast cells limit IgE-mediated immediate hypersensitivity reactions in human subjects. Journal of Allergy and Clinical Immunology, 2019, 144, 613-617.e14.	2.9	21
34	Mast cell activation in the context of elevated basal serum tryptase: genetics and presentations. Current Allergy and Asthma Reports, 2019, 19, 55.	5.3	18
35	The clinical and mechanistic intersection of primary atopic disorders and inborn errors of growth and metabolism. Immunological Reviews, 2019, 287, 135-144.	6.0	12
36	Angioedema and High Tryptase: To Marrow or Not to Marrow?. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 752.	3.8	0

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37	Primary atopic disorders. Journal of Experimental Medicine, 2018, 215, 1009-1022.	8.5	74
38	Persistent tryptase elevation in a patient with Gaucher disease. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 697-699.	3.8	13
39	A common haplotype containing functional CACNA1H variants is frequently coinherited with increased TPSAB1 copy number. Genetics in Medicine, 2018, 20, 503-512.	2.4	25
40	First Identification of an Inherited TPSAB1 Quintuplication in a Patient with Clonal Mast Cell Disease. Journal of Clinical Immunology, 2018, 38, 457-459.	3.8	58
41	Hereditary Alpha Tryptasemia. Immunology and Allergy Clinics of North America, 2018, 38, 483-495.	1.9	116
42	Prospective phenotyping of NGLY1-CDDG, the first congenital disorder of deglycosylation. Genetics in Medicine, 2017, 19, 160-168.	2.4	124
43	Detection of phosphoglucomutase-3 deficiency by lectin-based flow cytometry. Journal of Allergy and Clinical Immunology, 2017, 140, 291-294.e4.	2.9	10
44	ERBIN deficiency links STAT3 and TGF-Î² pathway defects with atopy in humans. Journal of Experimental Medicine, 2017, 214, 669-680.	8.5	70
45	Germline hypomorphic CARD11 mutations in severe atopic disease. Nature Genetics, 2017, 49, 1192-1201.	21.4	174
46	TGF-Î² pathway activation primes naÃ¯ve lymphocytes to support atopic phenotypes in humans. Journal of Allergy and Clinical Immunology, 2017, 139, AB93.	2.9	0
47	TPSAB1 Genotype and Effects of Pro-Tryptases on Human Mast Cell-Associated Disorders. Journal of Allergy and Clinical Immunology, 2017, 139, AB376.	2.9	1
48	PD-L1 up-regulation restrains Th17 cell differentiation in STAT3 loss- and STAT1 gain-of-function patients. Journal of Experimental Medicine, 2017, 214, 2523-2533.	8.5	55
49	Editorial: Stressing out mast cells via CRF 1. Journal of Leukocyte Biology, 2017, 102, 1284-1285.	3.3	1
50	Food allergies can persist after myeloablative hematopoietic stem cell transplantation in dedicator of cytokinesis 8â€“deficient patients. Journal of Allergy and Clinical Immunology, 2016, 137, 1895-1898.e5.	2.9	30
51	Elevated basal serum tryptase identifies a multisystem disorder associated with increased TPSAB1 copy number. Nature Genetics, 2016, 48, 1564-1569.	21.4	279
52	Loss-of-function mutations in TNFAIP3 leading to A20 haploinsufficiency cause an early-onset autoinflammatory disease. Nature Genetics, 2016, 48, 67-73.	21.4	513
53	GATA3 haploinsufficiency does not block allergic sensitization or atopic disease. Journal of Allergy and Clinical Immunology, 2016, 137, 627-629.e2.	2.9	2
54	Diminution of signal transducer and activator of transcription 3 signaling inhibits vascular permeability and anaphylaxis. Journal of Allergy and Clinical Immunology, 2016, 138, 187-199.	2.9	56

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55	FOXP3+ Tregs require WASP to restrain Th2-mediated food allergy. Journal of Clinical Investigation, 2016, 126, 4030-4044.	8.2	53
56	Convergence of Clinical and Cellular Phenotypes Among Patients with STAT3 and ERBB2IP Mutations. Journal of Allergy and Clinical Immunology, 2015, 135, AB152.	2.9	0
57	Early-onset lymphoproliferation and autoimmunity caused by germline STAT3 gain-of-function mutations. Blood, 2015, 125, 591-599.	1.4	436
58	Glycans Instructing Immunity: The Emerging Role of Altered Glycosylation in Clinical Immunology. Frontiers in Pediatrics, 2015, 3, 54.	1.9	56
59	Distinct Cutaneous Manifestations and Cold-Induced Leukocyte Activation Associated WithPLCG2Mutations. JAMA Dermatology, 2015, 151, 627.	4.1	55
60	Tu1385 Functional Gastrointestinal Disorders in Families With Basal Elevated Serum Tryptase. Gastroenterology, 2015, 148, S-876.	1.3	1
61	Atopic Dermatitis in Children. Immunology and Allergy Clinics of North America, 2015, 35, 161-183.	1.9	173
62	Autosomal recessive phosphoglucomutase 3 (PGM3) mutations link glycosylation defects to atopy, immune deficiency, autoimmunity, and neurocognitive impairment. Journal of Allergy and Clinical Immunology, 2014, 133, 1400-1409.e5.	2.9	193
63	Mendelian inheritance of elevated serum tryptase associated with atopy and connective tissue abnormalities. Journal of Allergy and Clinical Immunology, 2014, 133, 1471-1474.	2.9	110
64	Mendelian Inheritance Of Elevated Tryptase Associated With Atopy and Connective Tissue Abnormalities. Journal of Allergy and Clinical Immunology, 2014, 133, AB165.	2.9	0
65	Systemic hypersensitivity reaction mimicking anaphylaxis after first filgrastim administration in a healthy donor. Transfusion, 2013, 53, 1146-1147.	1.6	10