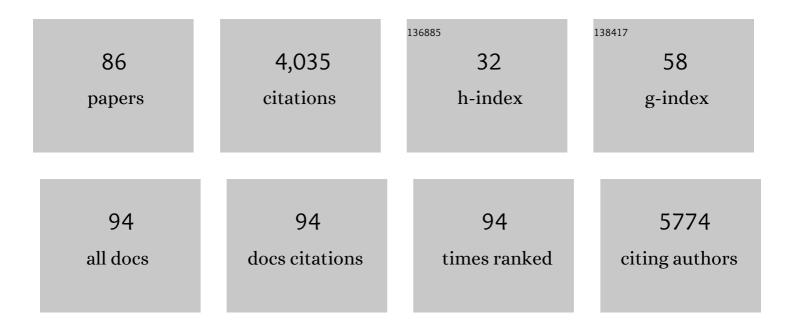
Nan Shen

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Enhanced genome editing to ameliorate a genetic metabolic liver disease through co-delivery of adeno-associated virus receptor. Science China Life Sciences, 2022, 65, 718-730.	2.3	16
2	The CD6/ALCAM pathway promotes lupus nephritis via T cell–mediated responses. Journal of Clinical Investigation, 2022, 132, .	3.9	25
3	P2RY8 variants in lupus patients uncover a role for the receptor in immunological tolerance. Journal of Experimental Medicine, 2022, 219, .	4.2	26
4	Urinary galectin-3 binding protein (G3BP) as a biomarker for disease activity and renal pathology characteristics in lupus nephritis. Arthritis Research and Therapy, 2022, 24, 77.	1.6	4
5	Expanding Roles of Noncoding RNAs in the Pathogenesis of Systemic Lupus Erythematosus. Current Rheumatology Reports, 2022, 24, 64-75.	2.1	2
6	TLR7 gain-of-function genetic variation causes human lupus. Nature, 2022, 605, 349-356.	13.7	208
7	Long non-coding RNA expression profiles in neutrophils revealed potential biomarker for prediction of renal involvement in SLE patients. Rheumatology, 2021, 60, 1734-1746.	0.9	16
8	Health Disparities in Rheumatic Diseases. Rheumatic Disease Clinics of North America, 2021, 47, 119-132.	0.8	7
9	Polymicrobial anaerobic bacterial meningitis secondary to dermal sinus: a case report. Translational Pediatrics, 2021, 10, 3118-3123.	0.5	7
10	Identification of 38 novel loci for systemic lupus erythematosus and genetic heterogeneity between ancestral groups. Nature Communications, 2021, 12, 772.	5.8	128
11	Exploration of the relationship between intestinal flora changes and gut acute graft-versus-host disease after hematopoietic stem cell transplantation. Translational Pediatrics, 2021, 10, 283-295.	0.5	4
12	Benefits of Iron Chelators in the Treatment of Parkinson's Disease. Neurochemical Research, 2021, 46, 1239-1251.	1.6	42
13	The enrichment of neutrophil extracellular traps impair the placentas of systemic lupus erythematosus through accumulating decidual NK cells. Scientific Reports, 2021, 11, 6870.	1.6	8
14	EZH2 Inhibition Interferes With the Activation of Type I Interferon Signaling Pathway and Ameliorates Lupus Nephritis in NZB/NZW F1 Mice. Frontiers in Immunology, 2021, 12, 653989.	2.2	17
15	Fecal microbiota transplantation before hematopoietic stem cell transplantation in a pediatric case of chronic diarrhea with a FOXP3 mutation. Pediatrics and Neonatology, 2021, 62, 172-180.	0.3	8
16	The metabolic hormone leptin promotes the function of TFH cells and supports vaccine responses. Nature Communications, 2021, 12, 3073.	5.8	27
17	Bach2 attenuates IL-2R signaling to control Treg homeostasis and Tfr development. Cell Reports, 2021, 35, 109096.	2.9	14
18	Structural insights into the recognition of histone H3Q5 serotonylation by WDR5. Science Advances, 2021, 7, .	4.7	14

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19	Analysis of gut microbiota alteration and application as an auxiliary prognostic marker for sepsis in children: a pilot study. Translational Pediatrics, 2021, 10, 1647-1657.	0.5	15
20	Glutamine metabolism is essential for the production of IL-17A in γδT cells and skin inflammation. Tissue and Cell, 2021, 71, 101569.	1.0	12
21	Downregulation of Renal Hsa-miR-127-3p Contributes to the Overactivation of Type I Interferon Signaling Pathway in the Kidney of Lupus Nephritis. Frontiers in Immunology, 2021, 12, 747616.	2.2	6
22	Evaluation of Molecular Point-of-Care Testing for Respiratory Pathogens in Children With Respiratory Infections: A Retrospective Case-Control Study. Frontiers in Cellular and Infection Microbiology, 2021, 11, 778808.	1.8	3
23	Down-Regulated miR-130a/b Attenuates Rhabdomyosarcoma Proliferation via PPARG. Frontiers in Molecular Biosciences, 2021, 8, 766887.	1.6	2
24	Urine Metabolic Fingerprints Encode Subtypes of Kidney Diseases. Angewandte Chemie, 2020, 132, 1720-1727.	1.6	8
25	Urine Metabolic Fingerprints Encode Subtypes of Kidney Diseases. Angewandte Chemie - International Edition, 2020, 59, 1703-1710.	7.2	99
26	The Genetic Landscape and Epidemiology of Phenylketonuria. American Journal of Human Genetics, 2020, 107, 234-250.	2.6	138
27	Molecular diagnostic practices for infectious gastroenteritis. Chinese Medical Journal, 2020, 133, 1485-1486.	0.9	2
28	Zirconia Hybrid Nanoshells for Nutrient and Toxin Detection. Small, 2020, 16, e2003902.	5.2	37
29	Impact of viral coinfection and macrolide-resistant mycoplasma infection in children with refractory Mycoplasma pneumoniae pneumonia. BMC Infectious Diseases, 2020, 20, 633.	1.3	37
30	Changes in Intestinal Flora and Metabolites in Neonates With Breast Milk Jaundice. Frontiers in Pediatrics, 2020, 8, 177.	0.9	11
31	MicroRNAs in Systemic Lupus Erythematosus: a Perspective on the Path from Biological Discoveries to Clinical Practice. Current Rheumatology Reports, 2020, 22, 17.	2.1	20
32	Inhibition of Glycolysis in Pathogenic TH17 Cells through Targeting a miR-21–Peli1–c-Rel Pathway Prevents Autoimmunity. Journal of Immunology, 2020, 204, 3160-3170.	0.4	17
33	Urinary activated leukocyte cell adhesion molecule as a novel biomarker of lupus nephritis histology. Arthritis Research and Therapy, 2020, 22, 122.	1.6	23
34	PBX1 promotes development of natural killer cells by binding directly to the <i>Nfil3</i> promoter. FASEB Journal, 2020, 34, 6479-6492.	0.2	13
35	MicroRNA-125a-Loaded Polymeric Nanoparticles Alleviate Systemic Lupus Erythematosus by Restoring Effector/Regulatory T Cells Balance. ACS Nano, 2020, 14, 4414-4429.	7.3	53
36	Taurine Metabolism Aggravates the Progression of Lupus by Promoting the Function of Plasmacytoid Dendritic Cells. Arthritis and Rheumatology, 2020, 72, 2106-2117.	2.9	13

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37	Complete clinical remission of invasive Candida infection with CARD9 deficiency after G-CSF treatment. Comparative Immunology, Microbiology and Infectious Diseases, 2020, 70, 101417.	0.7	15
38	De Novo <i>ARID1B</i> mutations cause growth delay associated with aberrant Wnt/β–catenin signaling. Human Mutation, 2020, 41, 1012-1024.	1.1	20
39	PBX1 expression in uterine natural killer cells drives fetal growth. Science Translational Medicine, 2020, 12, .	5.8	54
40	The MicroRNA <i>miR-22</i> Represses Th17 Cell Pathogenicity by Targeting PTEN-Regulated Pathways. ImmunoHorizons, 2020, 4, 308-318.	0.8	6
41	Allelic phenotype values: a model for genotype-based phenotype prediction in phenylketonuria. Genetics in Medicine, 2019, 21, 580-590.	1.1	48
42	Identification of Serum Biomarkers for Systemic Lupus Erythematosus Using a Library of Phage Displayed Random Peptides and Deep Sequencing. Molecular and Cellular Proteomics, 2019, 18, 1851-1863.	2.5	13
43	Interferon-α exacerbates neuropsychiatric phenotypes in lupus-prone mice. Arthritis Research and Therapy, 2019, 21, 205.	1.6	14
44	Excessive CD11c ⁺ Tbet ⁺ B cells promote aberrant T _{FH} differentiation and affinity-based germinal center selection in lupus. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 18550-18560.	3.3	68
45	Lupus-associated atypical memory B cells are mTORC1-hyperactivated and functionally dysregulated. Annals of the Rheumatic Diseases, 2019, 78, 1090-1100.	0.5	61
46	Identification of Renal Long Non-coding RNA RP11-2B6.2 as a Positive Regulator of Type I Interferon Signaling Pathway in Lupus Nephritis. Frontiers in Immunology, 2019, 10, 975.	2.2	52
47	Exosomal hsa-miR199a-3p Promotes Proliferation and Migration in Neuroblastoma. Frontiers in Oncology, 2019, 9, 459.	1.3	39
48	In vitro residual activities in 20 variants of phenylalanine hydroxylase and genotype-phenotype correlation in phenylketonuria patients. Gene, 2019, 707, 239-245.	1.0	7
49	Structure and Degradation of Circular RNAs Regulate PKR Activation in Innate Immunity. Cell, 2019, 177, 865-880.e21.	13.5	543
50	Amino acid signatures of HLA Class-I and II molecules are strongly associated with SLE susceptibility and autoantibody production in Eastern Asians. PLoS Genetics, 2019, 15, e1008092.	1.5	36
51	165â€Identification of serum biomarkers for systemic lupus erythematosus using a library of phage displayed random peptides and deep sequencing. , 2019, , .		0
52	A Diagnostic Panel of DNA Methylation Biomarkers for Lung Adenocarcinoma. Frontiers in Oncology, 2019, 9, 1281.	1.3	30
53	Alterations of the gut microbiome and plasma proteome in Chinese patients with adolescent idiopathic scoliosis. Bone, 2019, 120, 364-370.	1.4	22
54	Deficiency of β-Arrestin 2 in Dendritic Cells Contributes to Autoimmune Diseases. Journal of Immunology, 2019, 202, 407-420.	0.4	8

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55	Mutation spectrum of hyperphenylalaninemia candidate genes and the genotype-phenotype correlation in the Chinese population. Clinica Chimica Acta, 2018, 481, 132-138.	0.5	24
56	Exome-wide association study identifies four novel loci for systemic lupus erythematosus in Han Chinese population. Annals of the Rheumatic Diseases, 2018, 77, 417-417.	0.5	50
57	Heterogeneous clinical spectrum of DNAJC12-deficient hyperphenylalaninemia: from attention deficit to severe dystonia and intellectual disability. Journal of Medical Genetics, 2018, 55, 249-253.	1.5	29
58	Identification of LncRNA Linc00513 Containing Lupus-Associated Genetic Variants as a Novel Regulator of Interferon Signaling Pathway. Frontiers in Immunology, 2018, 9, 2967.	2.2	56
59	Identification of sparganosis based on next-generation sequencing. Infection, Genetics and Evolution, 2018, 66, 256-261.	1.0	13
60	Association of Abnormal Elevations in <scp>IFIT</scp> 3 With Overactive Cyclic <scp>GMP</scp> â€ <scp>AMP</scp> Synthase/Stimulator of Interferon Genes Signaling in Human Systemic Lupus Erythematosus Monocytes. Arthritis and Rheumatology, 2018, 70, 2036-2045.	2.9	57
61	Relationship between genotype, phenylalanine hydroxylase expression and in vitro activity and metabolic phenotype in phenylketonuria. Molecular Genetics and Metabolism, 2018, 125, 86-95.	0.5	31
62	Clinical remission of myopathy with MYH2 deficiency after precision medicine-developed rehabilitation: a case report. American Journal of Translational Research (discontinued), 2018, 10, 3827-3832.	0.0	1
63	A missense variant in NCF1 is associated with susceptibility to multiple autoimmune diseases. Nature Genetics, 2017, 49, 433-437.	9.4	143
64	Biallelic Mutations in DNAJC12 Cause Hyperphenylalaninemia, Dystonia, and Intellectual Disability. American Journal of Human Genetics, 2017, 100, 257-266.	2.6	127
65	Quantitative Susceptibility Mapping of the Substantia Nigra in Parkinson's Disease. Applied Magnetic Resonance, 2017, 48, 533-544.	0.6	13
66	Epigenetic regulator CXXC5 recruits DNA demethylase Tet2 to regulate TLR7/9-elicited IFN response in pDCs. Journal of Experimental Medicine, 2017, 214, 1471-1491.	4.2	81
67	The role of long non-coding RNAs in rheumatic diseases. Nature Reviews Rheumatology, 2017, 13, 657-669.	3.5	65
68	T-bet+CD11c+ B cells are critical for antichromatin immunoglobulin G production in the development of lupus. Arthritis Research and Therapy, 2017, 19, 225.	1.6	58
69	MiR-125a Is a critical modulator for neutrophil development. PLoS Genetics, 2017, 13, e1007027.	1.5	19
70	SENP7 Potentiates cGAS Activation by Relieving SUMO-Mediated Inhibition of Cytosolic DNA Sensing. PLoS Pathogens, 2017, 13, e1006156.	2.1	81
71	Identification of Cyclinâ€Ðependent Kinase 1 as a Novel Regulator of Type I Interferon Signaling in Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2016, 68, 1222-1232.	2.9	35
72	Dendritic Cells in Systemic Lupus Erythematosus: From Pathogenic Players to Therapeutic Tools. Mediators of Inflammation, 2016, 2016, 1-12.	1.4	43

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73	Phenotypic and functional alterations of pDCs in lupus-prone mice. Scientific Reports, 2016, 6, 20373.	1.6	23
74	In vitro residual activity of phenylalanine hydroxylase variants and correlation with metabolic phenotypes in PKU. Gene, 2016, 594, 138-143.	1.0	15
75	Identification of the long noncoding RNA NEAT1 as a novel inflammatory regulator acting through MAPK pathway in human lupus. Journal of Autoimmunity, 2016, 75, 96-104.	3.0	233
76	High-density genotyping of immune-related loci identifies new SLE risk variants in individuals with Asian ancestry. Nature Genetics, 2016, 48, 323-330.	9.4	219
77	Co-expression of phenylalanine hydroxylase variants and effects of interallelic complementation on in vitro enzyme activity and genotype-phenotype correlation. Molecular Genetics and Metabolism, 2016, 117, 328-335.	0.5	33
78	miR-744 enhances type I interferon signaling pathway by targeting PTP1B in primary human renal mesangial cells. Scientific Reports, 2015, 5, 12987.	1.6	23
79	Association of large intergenic noncoding RNA expression with disease activity and organ damage in systemic lupus erythematosus. Arthritis Research and Therapy, 2015, 17, 131.	1.6	92
80	Type I Interferon Inhibition of MicroRNAâ€146a Maturation Through Upâ€Regulation of Monocyte Chemotactic Protein–Induced Protein 1 in Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2015, 67, 3209-3218.	2.9	51
81	Berberine may rescue <i>Fusobacterium nucleatum</i> -induced colorectal tumorigenesis by modulating the tumor microenvironment. Oncotarget, 2015, 6, 32013-32026.	0.8	108
82	Lupus risk variants in the PXK locus alter B-cell receptor internalization. Frontiers in Genetics, 2015, 5, 450.	1.1	25
83	miRNAs in the Pathogenesis of Systemic Lupus Erythematosus. International Journal of Molecular Sciences, 2015, 16, 9557-9572.	1.8	55
84	Probiotics <i>Clostridium butyricum</i> and <i>Bacillus subtilis</i> ameliorate intestinal tumorigenesis. Future Microbiology, 2015, 10, 1433-1445.	1.0	82
85	Expression, localization, and clinical application of the RNA binding domain of U1-70kD in HEp-2 cells. Annals of Clinical and Laboratory Science, 2015, 45, 134-9.	0.2	0
86	Dysregulated expression of interleukin-23 and interleukin-12 subunits in systemic lupus erythematosus patients. Modern Rheumatology, 2007, 17, 220-223.	0.9	32