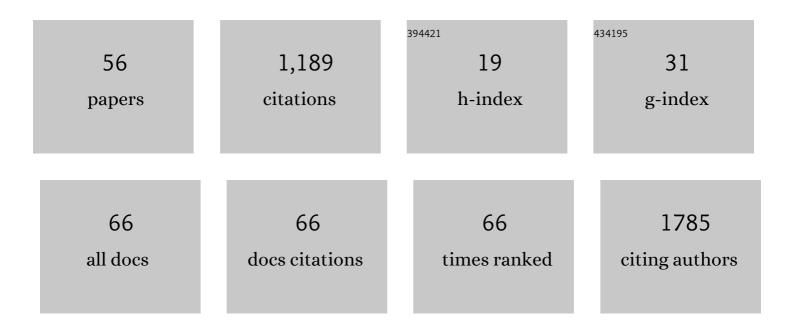


List of Publications by Year in descending order

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Υι Πλι

#	Article	IF	CITATIONS
1	Clinical Outcomes in Duchenne Muscular Dystrophy: A Study of 5345 Patients from the TREAT-NMD DMD Global Database. Journal of Neuromuscular Diseases, 2017, 4, 293-306.	2.6	125
2	Anti-NMDAR encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	106
3	Comparison of myelin oligodendrocyte glycoprotein (MOG)-antibody disease and AQP4-IgG-positive neuromyelitis optica spectrum disorder (NMOSD) when they co-exist with anti-NMDA (N-methyl-D-aspartate) receptor encephalitis. Multiple Sclerosis and Related Disorders, 2018, 20, 144-152.	2.0	89
4	Targeted next-generation sequencing as a comprehensive test for patients with and female carriers of DMD/BMD: a multi-population diagnostic study. European Journal of Human Genetics, 2014, 22, 110-118.	2.8	66
5	Effectiveness of High-risk Human Papillomavirus Testing for Cervical Cancer Screening in China. JAMA Oncology, 2021, 7, 263.	7.1	61
6	Whole exome sequencing identified a novel <i><scp>DAG</scp>1</i> mutation in a patient with rare, mild and late age of onset muscular dystrophyâ€dystroglycanopathy. Journal of Cellular and Molecular Medicine, 2019, 23, 811-818.	3.6	56
7	Muscle and Heart Function Restoration in a Limb Girdle Muscular Dystrophy 2I (LGMD2I) Mouse Model by Systemic FKRP Gene Delivery. Molecular Therapy, 2014, 22, 1890-1899.	8.2	46
8	Single-molecule optical mapping enables quantitative measurement of D4Z4 repeats in facioscapulohumeral muscular dystrophy (FSHD). Journal of Medical Genetics, 2020, 57, 109-120.	3.2	43
9	Risk factors for postoperative recurrence of ovarian endometriosis: long-term follow-up of 358 women. Journal of Ovarian Research, 2019, 12, 79.	3.0	42
10	A review of the risk factors, genetics and treatment of endometriosis in Chinese women: a comparative update. Reproductive Health, 2018, 15, 82.	3.1	37
11	Surgical outcomes in patients with anti-N-methyl D-aspartate receptor encephalitis with ovarian teratoma. American Journal of Obstetrics and Gynecology, 2019, 221, 485.e1-485.e10.	1.3	35
12	The Relationship of Circular RNAs With Ovarian Endometriosis. Reproductive Sciences, 2018, 25, 1292-1300.	2.5	31
13	A comprehensive genetic diagnosis of Chinese muscular dystrophy and congenital myopathy patients by targeted next-generation sequencing. Neuromuscular Disorders, 2015, 25, 617-624.	0.6	29
14	Exonic rearrangements in <i>DMD</i> in Chinese Han individuals affected with Duchenne and Becker muscular dystrophies. Human Mutation, 2020, 41, 668-677.	2.5	29
15	Identification of Circular RNAs as a Novel Biomarker for Ovarian Endometriosis. Chinese Medical Journal, 2018, 131, 559-566.	2.3	26
16	Therapeutic Exon Skipping Through a CRISPR-Guided Cytidine Deaminase Rescues Dystrophic Cardiomyopathy in Vivo. Circulation, 2021, 144, 1760-1776.	1.6	26
17	Amelioration of Muscle and Nerve Pathology in LAMA2 Muscular Dystrophy by AAV9-Mini-Agrin. Molecular Therapy - Methods and Clinical Development, 2018, 9, 47-56.	4.1	24
18	Clinical characteristics and management experience of unexpected uterine sarcoma after myomectomy. International Journal of Gynecology and Obstetrics, 2015, 130, 195-199.	2.3	21

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19	Eight years after an international workshop on myotonic dystrophy patient registries: case study of a global collaboration for a rare disease. Orphanet Journal of Rare Diseases, 2018, 13, 155.	2.7	19
20	Genetic variations and clinical spectrum of dystroglycanopathy in a large cohort of Chinese patients. Clinical Genetics, 2021, 99, 384-395.	2.0	19
21	Microbiome of the lower genital tract in Chinese women with endometriosis by 16s-rRNA sequencing technique: a pilot study. Annals of Translational Medicine, 2020, 8, 1440-1440.	1.7	19
22	Pregnancy outcomes in women with infertility and coexisting endometriosis and adenomyosis after laparoscopic surgery: a long-term retrospective follow-up study. BMC Pregnancy and Childbirth, 2021, 21, 383.	2.4	15
23	A Novel AGRN Mutation Leads to Congenital Myasthenic Syndrome Only Affecting Limb-girdle Muscle. Chinese Medical Journal, 2017, 130, 2279-2282.	2.3	15
24	Serum exosomes can restore cellular function <i>in vitro</i> and be used for diagnosis in dysferlinopathy. Theranostics, 2018, 8, 1243-1255.	10.0	14
25	Clinical phenotypes, genotypes and treatment in Chinese dystonia patients with KMT2B variants. Parkinsonism and Related Disorders, 2020, 77, 76-82.	2.2	13
26	Anatomical distribution of pelvic deep infiltrating endometriosis and its relationship with pain symptoms. Chinese Medical Journal, 2012, 125, 209-13.	2.3	13
27	Factors associated with deep infiltrating endometriosis versus ovarian endometrioma in China: a subgroup analysis from the FEELING study. BMC Women's Health, 2018, 18, 205.	2.0	12
28	An LMNB1 Duplication Caused Adult-Onset Autosomal Dominant Leukodystrophy in Chinese Family: Clinical Manifestations, Neuroradiology and Genetic Diagnosis. Frontiers in Molecular Neuroscience, 2017, 10, 215.	2.9	11
29	Fertility Outcomes After Laparoscopic Cystectomy in Infertile Patients with StageÂIII–IV Endometriosis: a Cohort with 6–10Âyears of Follow-up. Advances in Therapy, 2020, 37, 2159-2168.	2.9	10
30	Ultrasound-guided nusinersen administration for spinal muscular atrophy patients with severe scoliosis: an observational study. Orphanet Journal of Rare Diseases, 2021, 16, 274.	2.7	10
31	A Comprehensive Analysis of 2013 Dystrophinopathies in China: A Report From National Rare Disease Center. Frontiers in Neurology, 2020, 11, 572006.	2.4	9
32	Menorrhagia and Uterine Volume Associated with Lower Urinary Tract Symptoms in Patients with Adenomyosis. Chinese Medical Journal, 2017, 130, 1552-1556.	2.3	8
33	Economic evaluation of cervical cancer screening strategies in urban China. Chinese Journal of Cancer Research: Official Journal of China Anti-Cancer Association, Beijing Institute for Cancer Research, 2019, 31, 974-983.	2.2	8
34	Effects of Bisphosphonates on Osteoporosis Induced by Duchenne Muscular Dystrophy: A Prospective Study. Endocrine Practice, 2020, 26, 1477-1485.	2.1	8
35	Primary sarcoma of the ovary: clinicopathological characteristics, prognostic factors and evaluation of therapy. Chinese Medical Journal, 2011, 124, 1316-21.	2.3	8
36	A novel mutation in NF1 is associated with diverse intra-familial phenotypic variation and astrocytoma in a Chinese family. Journal of Clinical Neuroscience, 2016, 31, 182-184.	1.5	7

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37	Mechanism hypotheses for the electrophysiological manifestations of two cases of endplate acetylcholinesterase deficiency related congenital myasthenic syndrome. Journal of Clinical Neuroscience, 2018, 48, 229-232.	1.5	6
38	Clinical characteristics difference between early and late recurrence of ovarian endometriosis after laparoscopic cystectomy. Archives of Gynecology and Obstetrics, 2020, 302, 905-913.	1.7	6
39	A Novel Dystrophin Deletion Mutation in a Becker Muscular Dystrophy Patient With Early-Onset Dilated Cardiomyopathy. Canadian Journal of Cardiology, 2014, 30, 956.e1-956.e3.	1.7	5
40	Adult-onset severe methylenetetrahydrofolate reductase deficiency characterized by reversible spastic paraplegia with a novel mutation. Neurological Sciences, 2016, 37, 1735-1737.	1.9	5
41	Preventive therapeutic options for postoperative recurrence of ovarian endometrioma: gonadotropin-releasing hormone agonist with or without levonorgestrel intrauterine system insertion. Archives of Gynecology and Obstetrics, 2021, 303, 533-539.	1.7	5
42	Sequence and Structure Characteristics of 22 Deletion Breakpoints in Intron 44 of the DMD Gene Based on Long-Read Sequencing. Frontiers in Genetics, 2021, 12, 638220.	2.3	5
43	Hysterectomy and bilateral adnexectomy using transvaginal natural orifice transluminal endoscopic surgery: The role of <scp>multichannel</scp> abdominal <scp>PORT</scp> and vaginal support ring. Journal of Obstetrics and Gynaecology Research, 2021, 47, 2521-2528.	1.3	5
44	Cost-effectiveness Analysis of Dienogest Compared With Gonadotropin-Releasing Hormone Agonist After Conservative Surgery for Endometriosis in China. Clinical Therapeutics, 2021, 43, 1276-1284.e1.	2.5	5
45	Targeted next generation sequencing identifies two novel mutations in SEPN1 in rigid spine muscular dystrophy 1. Oncotarget, 2016, 7, 83843-83849.	1.8	5
46	Clinical outcomes and immune reconstitution in 103 advanced AIDS patients undergoing 12-month highly active antiretroviral therapy. Chinese Medical Journal, 2006, 119, 1677-82.	2.3	5
47	Not Having Been Breastfed May Protect Chinese Women From Developing Deep Infiltrating Endometriosis: Results From Subgroup Analyses of the FEELING Study. Reproductive Sciences, 2019, 26, 1158-1167.	2.5	4
48	Dropped head syndrome: a rare adverse drug reaction identified in the FDA adverse event reporting system and review of case reports in the literature. Expert Opinion on Drug Safety, 2022, 21, 1329-1336.	2.4	4
49	GJB1 Mutation-A Disease Spectrum: Report of Case Series. Frontiers in Neurology, 2019, 10, 1406.	2.4	3
50	Rare <i>NOTCH3</i> Variants in a Chinese Population-Based Cohort and Its Relationship With Cerebral Small Vessel Disease. Stroke, 2021, 52, 3918-3925.	2.0	3
51	Technique integration of single-cell RNA sequencing with spatially resolved transcriptomics in the tumor microenvironment. Cancer Cell International, 2022, 22, 155.	4.1	2
52	The clinical features and long-term surgical outcomes of different types of abdominal wall endometriosis. Archives of Gynecology and Obstetrics, 2023, 307, 163-168.	1.7	2
53	Differential Regulation of the Immune System in Peripheral Blood Following Ischemic Stroke. BioMed Research International, 2022, 2022, 1-15.	1.9	2
54	Steno-occlusive cerebral arteriopathy in patients with glycogen storage disease type I. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 434-435.	1.9	0

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55	Thoracic endometriosis presented as catamental hemoptysis: a case series of a rare disease. Current Medical Research and Opinion, 2021, 37, 685-691.	1.9	Ο
56	<i>BAG3</i>-Related Myofibrillar Myopathy Presenting as Hypercapnia: A Case Report and Literature Review . Chinese Medical Sciences Journal, 2019, 36, 1.	0.4	0