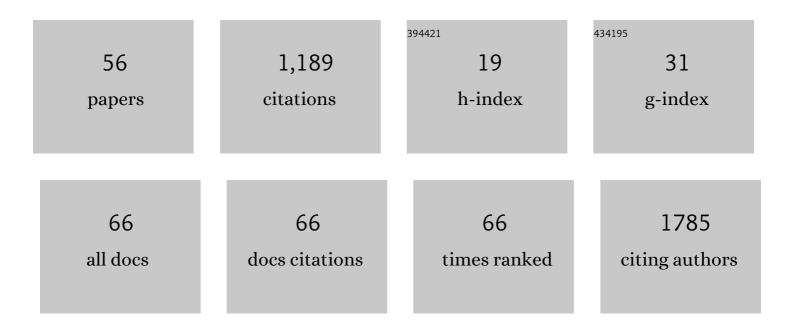


## List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	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
2	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
3	Technique integration of single-cell RNA sequencing with spatially resolved transcriptomics in the tumor microenvironment. Cancer Cell International, 2022, 22, 155.	4.1	2
4	Differential Regulation of the Immune System in Peripheral Blood Following Ischemic Stroke. BioMed Research International, 2022, 2022, 1-15.	1.9	2
5	Genetic variations and clinical spectrum of dystroglycanopathy in a large cohort of Chinese patients. Clinical Genetics, 2021, 99, 384-395.	2.0	19
6	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
7	Thoracic endometriosis presented as catamental hemoptysis: a case series of a rare disease. Current Medical Research and Opinion, 2021, 37, 685-691.	1.9	0
8	Effectiveness of High-risk Human Papillomavirus Testing for Cervical Cancer Screening in China. JAMA Oncology, 2021, 7, 263.	7.1	61
9	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
10	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
11	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
12	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
13	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
14	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
15	Therapeutic Exon Skipping Through a CRISPR-Guided Cytidine Deaminase Rescues Dystrophic Cardiomyopathy in Vivo. Circulation, 2021, 144, 1760-1776.	1.6	26
16	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
17	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
18	Anti-NMDAR encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	106

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19	A Comprehensive Analysis of 2013 Dystrophinopathies in China: A Report From National Rare Disease Center. Frontiers in Neurology, 2020, 11, 572006.	2.4	9
20	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
21	Clinical phenotypes, genotypes and treatment in Chinese dystonia patients with KMT2B variants. Parkinsonism and Related Disorders, 2020, 77, 76-82.	2.2	13
22	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
23	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
24	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
25	Effects of Bisphosphonates on Osteoporosis Induced by Duchenne Muscular Dystrophy: A Prospective Study. Endocrine Practice, 2020, 26, 1477-1485.	2.1	8
26	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
27	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
28	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
29	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
30	GJB1 Mutation-A Disease Spectrum: Report of Case Series. Frontiers in Neurology, 2019, 10, 1406.	2.4	3
31	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
32	<b><i>BAG3</i>-Related Myofibrillar Myopathy Presenting as Hypercapnia: A Case Report and Literature Review</b> . Chinese Medical Sciences Journal, 2019, 36, 1.	0.4	0
33	The Relationship of Circular RNAs With Ovarian Endometriosis. Reproductive Sciences, 2018, 25, 1292-1300.	2.5	31
34	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
35	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
36	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

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37	Identification of Circular RNAs as a Novel Biomarker for Ovarian Endometriosis. Chinese Medical Journal, 2018, 131, 559-566.	2.3	26
38	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
39	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
40	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
41	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
42	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
43	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
44	Menorrhagia and Uterine Volume Associated with Lower Urinary Tract Symptoms in Patients with Adenomyosis. Chinese Medical Journal, 2017, 130, 1552-1556.	2.3	8
45	A Novel AGRN Mutation Leads to Congenital Myasthenic Syndrome Only Affecting Limb-girdle Muscle. Chinese Medical Journal, 2017, 130, 2279-2282.	2.3	15
46	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
47	Adult-onset severe methylenetetrahydrofolate reductase deficiency characterized by reversible spastic paraplegia with a novel mutation. Neurological Sciences, 2016, 37, 1735-1737.	1.9	5
48	Targeted next generation sequencing identifies two novel mutations in SEPN1 in rigid spine muscular dystrophy 1. Oncotarget, 2016, 7, 83843-83849.	1.8	5
49	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
50	Clinical characteristics and management experience of unexpected uterine sarcoma after myomectomy. International Journal of Gynecology and Obstetrics, 2015, 130, 195-199.	2.3	21
51	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
52	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
53	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
54	Anatomical distribution of pelvic deep infiltrating endometriosis and its relationship with pain symptoms. Chinese Medical Journal, 2012, 125, 209-13.	2.3	13

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55	Primary sarcoma of the ovary: clinicopathological characteristics, prognostic factors and evaluation of therapy. Chinese Medical Journal, 2011, 124, 1316-21.	2.3	8
56	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