List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	2011 Consensus statement on endomyocardial biopsy from the Association for European Cardiovascular Pathology and the Society for Cardiovascular Pathology. Cardiovascular Pathology, 2012, 21, 245-274.	0.7	423
2	Coronary microvascular dysfunction: mechanisms and functional assessment. Nature Reviews Cardiology, 2015, 12, 48-62.	6.1	377
3	Guidelines for autopsy investigation of sudden cardiac death: 2017 update from the Association for European Cardiovascular Pathology. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2017, 471, 691-705.	1.4	357
4	Leptin induces direct vasodilation through distinct endothelial mechanisms. Diabetes, 2000, 49, 293-297.	0.3	303
5	Consensus statement on surgical pathology of the aorta from the Society for Cardiovascular Pathology and the Association for European Cardiovascular Pathology: I. Inflammatory diseases. Cardiovascular Pathology, 2015, 24, 267-278.	0.7	238
6	Efficient mitochondrial biogenesis drives incomplete penetrance in Leber's hereditary optic neuropathy. Brain, 2014, 137, 335-353.	3.7	229
7	Oestrogens ameliorate mitochondrial dysfunction in Leber's hereditary optic neuropathy. Brain, 2011, 134, 220-234.	3.7	208
8	Consensus statement on surgical pathology of the aorta from the Society for Cardiovascular Pathology and the Association For European Cardiovascular Pathology: II. Noninflammatory degenerative diseases — nomenclature and diagnostic criteria. Cardiovascular Pathology, 2016, 25, 247-257.	0.7	208
9	Cardiac involvement in mitochondrial DNA disease: clinical spectrum, diagnosis, and management. European Heart Journal, 2012, 33, 3023-3033.	1.0	182
10	Are the kinetics of technetium-99m methoxyisobutyl isonitrile affected by cell metabolism and viability?. Circulation, 1990, 82, 1802-1814.	1.6	177
11	Induction of Mitochondrial Biogenesis Is a Maladaptive Mechanism in Mitochondrial Cardiomyopathies. Journal of the American College of Cardiology, 2007, 50, 1362-1369.	1.2	164
12	A homoplasmic mitochondrial transfer Ribonucleic Acid mutation as a cause of maternally inherited hypertrophic cardiomyopathy. Journal of the American College of Cardiology, 2003, 41, 1786-1796.	1.2	161
13	Pathogenic expression of homoplasmic mtDNA mutations needs a complex nuclear–mitochondrial interaction. Trends in Genetics, 2003, 19, 257-262.	2.9	137
14	Human Parvovirus B19 Infection in Infancy Associated with Acute and Chronic Lymphocytic Myocarditis and High Cytokine Levels: Report of 3 Cases and Review. Clinical Infectious Diseases, 2000, 31, 65-69.	2.9	122
15	Heart involvement in AIDS: a prospective study during various stages of the disease. European Heart Journal, 1992, 13, 1452-1459.	1.0	112
16	A Novel mtDNA Point Mutation in Maternally Inherited Cardiomyopathy. Biochemical and Biophysical Research Communications, 1995, 213, 588-593.	1.0	104
17	Mutations of an intronic repeat induce impaired MRE11 expression in primary human cancer with microsatellite instability. Oncogene, 2004, 23, 2640-2647.	2.6	101
18	Gastrointestinal Dysmotility in Mitochondrial Neurogastrointestinal Encephalomyopathy Is Caused by Mitochondrial DNA Depletion. American Journal of Pathology, 2008, 173, 1120-1128.	1.9	100

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19	Diagnostic Value of Endomyocardial Biopsy Guided by Electroanatomic Voltage Mapping in Arrhythmogenic Right Ventricular Cardiomyopathy/Dysplasia. Journal of Cardiovascular Electrophysiology, 2008, 19, 1127-1134.	0.8	96
20	Pathologic evidence of extensive left ventricular involvement in arrhythmogenic right ventricular cardiomyopathy. Human Pathology, 1992, 23, 948-952.	1.1	88
21	Pantothenate kinase-associated neurodegeneration: altered mitochondria membrane potential and defective respiration in Pank2 knock-out mouse model. Human Molecular Genetics, 2012, 21, 5294-5305.	1.4	87
22	Combined treatment with temozolomide and poly(ADP-ribose) polymerase inhibitor enhances survival of mice bearing hematologic malignancy at the central nervous system site. Blood, 2002, 99, 2241-2244.	0.6	83
23	Cardiac mesenchymal stromal cells are a source of adipocytes in arrhythmogenic cardiomyopathy. European Heart Journal, 2016, 37, 1835-1846.	1.0	83
24	Pantethine treatment is effective in recovering the disease phenotype induced by ketogenic diet in a pantothenate kinase-associated neurodegeneration mouse model. Brain, 2014, 137, 57-68.	3.7	78
25	Perindopril and indapamide reverse coronary microvascular remodelling and improve flow in arterial hypertension. Journal of Hypertension, 2011, 29, 364-372.	0.3	77
26	Impaired mitochondrial biogenesis is a common feature to myocardial hypertrophy and end-stage ischemic heart failure. Cardiovascular Pathology, 2016, 25, 103-112.	0.7	77
27	Enhanced ROS production by NADPH oxidase is correlated to changes in antioxidant enzyme activity in human heart failure. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2010, 1802, 331-338.	1.8	76
28	The sexist behaviour of immune checkpoint inhibitors in cancer therapy?. Oncotarget, 2017, 8, 99336-99346.	0.8	76
29	Early histologic findings of pulmonary SARS-CoV-2 infection detected in a surgical specimen. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 743-748.	1.4	69
30	Isoleucyl-tRNA synthetase levels modulate the penetrance of a homoplasmic m.4277T>C mitochondrial tRNAIle mutation causing hypertrophic cardiomyopathy. Human Molecular Genetics, 2012, 21, 85-100.	1.4	67
31	Defining phenotypes and disease progression in sarcomeric cardiomyopathies: contemporary role of clinical investigations. Cardiovascular Research, 2015, 105, 409-423.	1.8	66
32	Heat-shock protein 90: A novel autoantigen in human carotid atherosclerosis. Atherosclerosis, 2009, 207, 74-83.	0.4	64
33	TTC19 Plays a Husbandry Role on UQCRFS1 Turnover in the Biogenesis of Mitochondrial Respiratory Complex III. Molecular Cell, 2017, 67, 96-105.e4.	4.5	64
34	Frequency of development of acute global left ventricular dysfunction in human immunodeficiency virus infection. Journal of the American College of Cardiology, 1994, 24, 1018-1024.	1.2	63
35	Mitochondrial Neurogastrointestinal Encephalomyopathy: Evidence of Mitochondrial DNA Depletion in the Small Intestine. Gastroenterology, 2006, 130, 893-901.	0.6	63
36	Arrhythmogenic right ventricular cardiomyopathy: Clinicopathologic correlation based on a revised definition of pathologic patterns. Human Pathology, 2001, 32, 1078-1086.	1.1	62

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37	Targeting estrogen receptor \hat{l}^2 as preventive therapeutic strategy for Leber's hereditary optic neuropathy. Human Molecular Genetics, 2015, 24, ddv396.	1.4	62
38	Defective <scp>PITRM</scp> 1 mitochondrial peptidase is associated with AÎ ² amyloidotic neurodegeneration. EMBO Molecular Medicine, 2016, 8, 176-190.	3.3	60
39	NADPH oxidase-dependent redox signaling in human heart failure: Relationship between the left and right ventricle. Journal of Molecular and Cellular Cardiology, 2007, 42, 826-834.	0.9	59
40	Juvenile sudden death in a family with polymorphic ventricular arrhythmias caused by a novel RyR2 gene mutation: evidence of specific morphological substrates. Human Pathology, 2005, 36, 761-767.	1.1	58
41	Feasibility of Combined Unipolar and Bipolar Voltage Maps to Improve Sensitivity of Endomyocardial Biopsy. Circulation: Arrhythmia and Electrophysiology, 2015, 8, 625-632.	2.1	58
42	Diagnostic Accuracy of Transthoracic and Multiplane Transesophageal Echocardiography for Valvular Perforation in Acute Infective Endocarditis: Correlation with Anatomic Findings. Clinical Infectious Diseases, 2000, 30, 825-826.	2.9	57
43	Valvular perforation in left-sided infective endocarditis: A prospective echocardiographic evaluation and clinical outcome. American Heart Journal, 1997, 134, 656-664.	1.2	53
44	A Western single-center experience with endoscopic submucosal dissection for early gastrointestinal cancers. Gastric Cancer, 2010, 13, 258-263.	2.7	52
45	Nonischemic left ventricular scar and cardiac sudden death in the young. Human Pathology, 2016, 58, 78-89.	1.1	52
46	Sudden cardiac death in younger adults: autopsy diagnosis as a tool for preventive medicine. Human Pathology, 2006, 37, 794-801.	1.1	49
47	PD-L1 Expression in TNBC: A Predictive Biomarker of Response to Neoadjuvant Chemotherapy?. BioMed Research International, 2017, 2017, 1-7.	0.9	49
48	AAV-mediated Liver-specific MPV17 Expression Restores mtDNA Levels and Prevents Diet-induced Liver Failure. Molecular Therapy, 2014, 22, 10-17.	3.7	47
49	Anti–PD-1 and Anti–PD-L1 in Head and Neck Cancer: A Network Meta-Analysis. Frontiers in Immunology, 2021, 12, 705096.	2.2	47
50	Prolyl-isomerase Pin1 controls Notch3 protein expression and regulates T-ALL progression. Oncogene, 2016, 35, 4741-4751.	2.6	45
51	Pathological Findings of HIVâ€Associated Cardiovascular Disease. Annals of the New York Academy of Sciences, 2001, 946, 23-45.	1.8	43
52	The isolated carboxyâ€ŧerminal domain of human mitochondrial leucylâ€ <scp>tRNA</scp> synthetase rescues the pathological phenotype of mitochondrial <scp>tRNA</scp> mutations in human cells. EMBO Molecular Medicine, 2014, 6, 169-182.	3.3	43
53	Mitochondrial myopathy, parkinsonism, and multiple mtDNA deletions in a Sephardic Jewish family. Neurology, 2001, 56, 802-805.	1.5	41
54	Chloroquine supplementation increases the cytotoxic effect of curcumin against Her2/neu overexpressing breast cancer cells <i>in vitro</i> and <i>in vivo</i> in nude mice while counteracts it in immune competent mice. Oncolmmunology, 2017, 6, e1356151.	2.1	41

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55	High-intensity focused ultrasound in breast pathology: non-invasive treatment of benign and malignant lesions. Expert Review of Medical Devices, 2015, 12, 191-199.	1.4	40
56	NOTCH3 inactivation increases triple negative breast cancer sensitivity to gefitinib by promoting EGFR tyrosine dephosphorylation and its intracellular arrest. Oncogenesis, 2018, 7, 42.	2.1	39
57	The Agnostic Role of Site of Metastasis in Predicting Outcomes in Cancer Patients Treated with Immunotherapy. Vaccines, 2020, 8, 203.	2.1	38
58	Heat Shock Proteins and Autoimmunity in Patients with Carotid Atherosclerosis. Annals of the New York Academy of Sciences, 2007, 1107, 1-10.	1.8	37
59	Cardinal vein isomerism. Cardiovascular Pathology, 2002, 11, 149-152.	0.7	36
60	Morphologic evidence of diffuse vascular damage in human and in the experimental model of ethylmalonic encephalopathy. Journal of Inherited Metabolic Disease, 2012, 35, 451-458.	1.7	35
61	A novel LAMP2 mutation associated with severe cardiac hypertrophy and microvascular remodeling in a female with Danon disease: a case report and literature review. Cardiovascular Pathology, 2016, 25, 423-431.	0.7	34
62	FATAL CONGENITAL MYOPATHY AND GASTROINTESTINAL PSEUDO-OBSTRUCTION DUE TO <i>POLG1</i> MUTATIONS. Neurology, 2009, 72, 1103-1105.	1.5	33
63	Cardiomyopathies due to homoplasmic mitochondrial tRNA mutations: morphologic and molecular features. Human Pathology, 2013, 44, 1262-1270.	1.1	32
64	Altered expression of alpha-dystroglycan subunit in human gliomas. Cancer Biology and Therapy, 2006, 5, 441-448.	1.5	31
65	Idiopathic noncirrhotic portal hypertension: current perspectives. Hepatic Medicine: Evidence and Research, 2016, Volume 8, 81-88.	0.9	31
66	Pathogenesis of the deafness-associated A1555G mitochondrial DNA mutation. Biochemical and Biophysical Research Communications, 2002, 293, 521-529.	1.0	30
67	Idiopathic Non Cirrhotic Portal Hypertension and Spleno-Portal Axis Abnormalities in Patients with Severe Primary Antibody Deficiencies. Journal of Immunology Research, 2014, 2014, 1-8.	0.9	30
68	Maternally inherited cardiomyopathy: clinical and molecular characterization of a large kindred harboring the A4300G point mutation in mitochondrial deoxyribonucleic acid. Journal of the American College of Cardiology, 1999, 33, 1584-1589.	1.2	29
69	Lonidamine Causes Inhibition of Angiogenesis-Related Endothelial Cell Functions. Neoplasia, 2004, 6, 513-522.	2.3	29
70	Cardiac involvement in consecutive unselected hospitalized COVID-19 population: In-hospital evaluation and one-year follow-up. International Journal of Cardiology, 2021, 339, 235-242.	0.8	28
71	Clinical course of cardiomyopathy in HIV-infected patients with or without encephalopathy related to the myocardial expression of tumour necrosis factor-α and nitric oxide synthase. Aids, 2000, 14, 827-838.	1.0	27
72	Myositis in primary Sjögren's syndrome: data from a multicentre cohort. Clinical and Experimental Rheumatology, 2015, 33, 457-64.	0.4	27

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73	Mapping genetic determinants of coronary microvascular remodeling in the spontaneously hypertensive rat. Basic Research in Cardiology, 2013, 108, 316.	2.5	26
74	Left Ventricular Outflow Tract Obstruction in Atrioventricular Septal Defects: A Pathologic and Morphometric Evaluation. Clinical Cardiology, 1991, 14, 513-521.	0.7	25
75	A multiple retinoic acid antagonist induces conotruncal anomalies, including transposition of the great arteries, in mice. Cardiovascular Pathology, 2006, 15, 194-202.	0.7	25
76	Histomorphometric features predict 1-year outcome of patients with idiopathic dilated cardiomyopathy considered to be at low priority for cardiac transplantation. American Heart Journal, 1994, 128, 316-325.	1.2	24
77	Reciprocal congenic lines for a major stroke QTL on rat chromosome 1. Physiological Genomics, 2006, 27, 108-113.	1.0	23
78	Folic acid and methionine in the prevention of teratogen-induced congenital defects in mice. Cardiovascular Pathology, 2009, 18, 100-109.	0.7	23
79	Cystic adventitial degeneration of the popliteal artery: Lectin histochemical study. European Journal of Vascular Surgery, 1994, 8, 16-19.	0.9	22
80	Nonischemic Left Ventricular Scar. Circulation, 2014, 130, e180-2.	1.6	22
81	Can MRI Biomarkers Predict Triple-Negative Breast Cancer?. Diagnostics, 2020, 10, 1090.	1.3	22
82	The pattern of desmin filaments in myocardial disarray. Human Pathology, 1995, 26, 262-266.	1.1	21
83	New derivatives of the antimalarial drug Pyrimethamine in the control of melanoma tumor growth: an in vitro and in vivo study. Journal of Experimental and Clinical Cancer Research, 2016, 35, 137.	3.5	21
84	Increased Expression of Thyroid Hormone Receptor Isoforms in End-Stage Human Congestive Heart Failure. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 2080-2084.	1.8	21
85	Detection of deleted mitochondrial DNA in Kearns-Sayre syndrome using laser capture microdissection. Human Pathology, 2003, 34, 1058-1061.	1.1	20
86	Platelet-derived growth factor C and calpain-3 are modulators of human melanoma cell invasiveness. Oncology Reports, 2013, 30, 2887-2896.	1.2	20
87	Sudden cardiac death in an Italian competitive athlete: Pre-participation screening and cardiovascular emergency care are both essential. International Journal of Cardiology, 2016, 206, 84-86.	0.8	20
88	Short peptides from leucyl-tRNA synthetase rescue disease-causing mitochondrial tRNA point mutations. Human Molecular Genetics, 2016, 25, 903-915.	1.4	19
89	CD73 expression and pathologic response to neoadjuvant chemotherapy in triple negative breast cancer. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 476, 569-576.	1.4	19
90	Relation of complex ventricular arrhythmias to presenting features and prognosis in dilated cardiomyopathy. International Journal of Cardiology, 1990, 29, 47-54.	0.8	18

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91	AECVP and SCVP 2009 Recommendations for Training in Cardiovascular Pathology. Cardiovascular Pathology, 2010, 19, 129-135.	0.7	18
92	Oncocytic glioblastoma: a glioblastoma showing oncocytic changes and increased mitochondrial DNA copy number. Human Pathology, 2013, 44, 1867-1876.	1.1	15
93	Endomyocardial Biopsy Guided by Electroanatomic Voltage Mapping in Arrhythmogenic Right Ventricular Cardiomyopathy: A Case Report. Journal of Cardiovascular Electrophysiology, 2007, 18, 991-993.	0.8	14
94	Endomyocardial biopsy findings in patients with ventricular arrhythmias of unknown origin. Cardiovascular Pathology, 1996, 5, 139-144.	0.7	13
95	Transcriptional Network Analysis for the Regulation of Left Ventricular Hypertrophy and Microvascular Remodeling. Journal of Cardiovascular Translational Research, 2013, 6, 931-944.	1.1	13
96	Ontogenetic Pattern of Thyroid Hormone Receptor Expression in the Human Testis. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 3453-3457.	1.8	13
97	Novel compound mutations in the mitochondrial translation elongation factor (TSFM) gene cause severe cardiomyopathy with myocardial fibro-adipose replacement. Scientific Reports, 2019, 9, 5108.	1.6	12
98	Breast cancer subtypes affect the nodal response after neoadjuvant chemotherapy in locally advanced breast cancer: Are we ready to endorse axillary conservation?. Breast Journal, 2019, 25, 273-277.	0.4	12
99	Evaluation of Gastrointestinal mtDNA Depletion in Mitochondrial Neurogastrointestinal Encephalomyopathy (MNGIE). Methods in Molecular Biology, 2011, 755, 223-232.	0.4	11
100	Pathologic Evidence of Arrhythmogenic Cardiomyopathy and Myocarditis in Two Siblings. Cardiovascular Pathology, 1998, 7, 39-46.	0.7	10
101	Investigating Patterns of Immune Interaction in Ovarian Cancer: Probing the O-glycoproteome by the Macrophage Galactose-Like C-Type Lectin (MGL). Cancers, 2020, 12, 2841.	1.7	10
102	Tissue Immune Profile: A Tool to Predict Response to Neoadjuvant Therapy in Triple Negative Breast Cancer. Cancers, 2020, 12, 2648.	1.7	10
103	Standard of Care and Promising New Agents for the Treatment of Mesenchymal Triple-Negative Breast Cancer. Cancers, 2021, 13, 1080.	1.7	10
104	Circulating CD137+ T Cells Correlate with Improved Response to Anti-PD1 Immunotherapy in Patients with Cancer. Clinical Cancer Research, 2022, 28, 1027-1037.	3.2	10
105	External Quality Assessment (EQA) program for the preanalytical and analytical immunohistochemical determination of HER2 in breast cancer: an experience on a regional scale. Journal of Experimental and Clinical Cancer Research, 2013, 32, 58.	3.5	9
106	Anti-aminoacyl-tRNA synthetase-related myositis and dermatomyositis: clues for differential diagnosis on muscle biopsy. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 477-487.	1.4	9
107	Myocardial fibrosis: morphologic patterns and role of imaging in diagnosis and prognostication. Cardiovascular Pathology, 2022, 56, 107391.	0.7	9
108	Comparison between electroanatomic and pathologic findings in a patient with arrhythmogenic right ventricular cardiomyopathy/dysplasia treated with orthotopic cardiac transplant. Heart Rhythm, 2010, 7, 828-831.	0.3	8

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109	Effect of different drug classes on reverse remodeling of intramural coronary arterioles in the spontaneously hypertensive rat. Microcirculation, 2017, 24, e12298.	1.0	8
110	Role of radiomics in predicting lung cancer spread through air spaces in a heterogeneous dataset. Translational Lung Cancer Research, 2022, 11, 560-571.	1.3	8
111	Felodipine protects human atrial muscle from hypoxia–reoxygenation dysfunction: a force–frequency relationship study in an in vitro model of stunning. International Journal of Cardiology, 1997, 62, 107-132.	0.8	7
112	Exogenous peptides are able to penetrate human cell and mitochondrial membranes, stabilize mitochondrial tRNA structures, and rescue severe mitochondrial defects. FASEB Journal, 2020, 34, 7675-7686.	0.2	6
113	Exploring the Ability of LARS2 Carboxy-Terminal Domain in Rescuing the MELAS Phenotype. Life, 2021, 11, 674.	1.1	6
114	Functional anatomy of the human vagina. Journal of Endocrinological Investigation, 2003, 26, 92-6.	1.8	6
115	Pathological examination of breast cancer samples before and after neoadjuvant therapy: recommendations from the Italian Group for the Study of Breast Pathology - Italian Society of Pathology (GIPaM-SIAPeC). Pathologica, 2022, 114, 104-110.	1.3	6
116	Cardiac pathologic findings in 3 unusual cases of sudden cardiac death related to anorexiant drugs. Human Pathology, 2017, 69, 101-109.	1.1	5
117	Prognostic impact of spread through air spaces in lung adenocarcinoma. Interactive Cardiovascular and Thoracic Surgery, 2022, 34, 1011-1015.	0.5	5
118	Diagnosis of Myocarditis Mimicking Arrhythmogenic Right Ventricular Cardiomyopathy. Journal of the American College of Cardiology, 2009, 54, 664-665.	1.2	4
119	The phenotypic expression of mitochondrial tRNA-mutations can be modulated by either mitochondrial leucyl-tRNA synthetase or the C-terminal domain thereof. Frontiers in Genetics, 2015, 6, 113.	1.1	4
120	Mitochondrial Energetics and Ca2+-Activated ATPase in Obstructive Hypertrophic Cardiomyopathy. Journal of Clinical Medicine, 2020, 9, 1799.	1.0	4
121	Eosinophilic Infiltration Immediately Following Transplantation. Cardiovascular Pathology, 1999, 8, 297-299.	0.7	3
122	Neuromuscular relaxants in non-cardiac surgery after cardiomyoplasty. Canadian Journal of Anaesthesia, 1998, 45, 324-327.	0.7	2
123	Authors' Response: Expression of TR Isoforms in Failing Human Heart. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5089a-5090.	1.8	2
124	Myopathy Complicating Lupus Pregnancy. Journal of Clinical Rheumatology, 2013, 19, 132-133.	0.5	2
125	Coronary atherosclerosis and sudden cardiac death in the young: another face of the culprit, another way of striking?. International Journal of Cardiology, 2018, 264, 28-29.	0.8	2
126	Diagnosis of arrhythmogenic right ventricular cardiomyopathy: the role of endomyocardial biopsy guided by electroanatomic voltage map. Europace, 2009, 11, 970-970.	0.7	1

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127	An unusual manifestation of IgG4-related disease. Rheumatology, 2018, 57, 1305-1307.	0.9	1
128	An Unusual Ulcer. International Journal of Lower Extremity Wounds, 2018, 17, 290-294.	0.6	1
129	NADPH oxidase is related with lipid peroxidation and redox-sensitive kinase activation in human failing hearts. Journal of Molecular and Cellular Cardiology, 2007, 42, S153.	0.9	Ο
130	Mitochondrial tRNA mutations manifest not only as hypertrophic cardiomyopathy but also as noncompaction—reply. Human Pathology, 2014, 45, 1791-1792.	1.1	0
131	FRI0500â€Evaluation of the Role of Fractalkine Chemokine CX3CL1 and Its Receptor CX3CR1 in Inflammatory Myopathies: Table 1. Annals of the Rheumatic Diseases, 2014, 73, 568.1-568.	0.5	0
132	AB0624â€High Levels of Proinflammatory Biomarkers in Patients with Idiopathic Inflammatory Myopathies. Annals of the Rheumatic Diseases, 2014, 73, 1012.1-1012.	0.5	0
133	SAT0305â€Histology of minor salivary glands in patients with sjÖgren's syndrome, association with clinical and laboratory aspects. , 2017, , .		0
134	Results of Adrenalectomy for Isolated, Metachronous Metastasis of Breast Cancer: A Retrospective Cohort Study. Frontiers in Surgery, 2021, 8, 671424.	0.6	0
135	"Protenuria in SLE: Is it always lupus?― Lupus, 2021, 30, 664-668.	0.8	0