Naoki Nishida

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5708032/publications.pdf

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112	1,283	430442	454577
papers	citations	18	30
		h-index	g-index

114	114	114	1473
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A double heterozygous variant in MYH6 and MYH7 associated with hypertrophic cardiomyopathy in a Japanese Family. Journal of Cardiology Cases, 2022, 25, 213-217.	0.2	2
2	Novel histopathological deposition patterns of EGF-containing fibulin-like extracellular matrix protein 1 amyloidosis: an autopsy case exhibiting a possible association between AEFEMP1 amyloidosis and elastic fibres. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 139-140.	1.4	1
3	Histopathology of acute colchicine intoxication: novel findings and their association with clinical manifestations. Journal of Toxicologic Pathology, 2022, 35, 255-262.	0.3	1
4	Patterns of Mixed Pathologies in Down Syndrome. Journal of Alzheimer's Disease, 2022, 87, 595-607.	1.2	8
5	A Novel NKX2-5 Variant in a Child with Left Ventricular Noncompaction, Atrial Septal Defect, Atrioventricular Conduction Disorder, and Syncope. Journal of Clinical Medicine, 2022, 11, 3171.	1.0	2
6	A burden of sarcomere gene variants in fetal-onset patients with left ventricular noncompaction. International Journal of Cardiology, 2021, 328, 122-129.	0.8	5
7	Traumatic rupture of the circle of Willis with closed head injury: Its pathology and possible pathogenesis. Journal of Clinical Forensic and Legal Medicine, 2021, 78, 102114.	0.5	0
8	Clinicopathological features of clinically undiagnosed sporadic transthyretin cardiac amyloidosis: a forensic autopsy-based series. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 125-133.	1.4	6
9	An autopsy case of pure nigropathy with TUBA4A nonsense mutation. Neuropathology and Applied Neurobiology, 2021, 47, 891-893.	1.8	6
10	A Case Report of a Rare Heterozygous Variant in the Desmin Gene Associated With Hypertrophic Cardiomyopathy and Complete Atrioventricular Block. CJC Open, 2021, 3, 1195-1198.	0.7	7
11	Effects of sporadic transthyretin amyloidosis frequently on the gallbladder and the correlation between amyloid deposition in the gallbladder and heart: A forensic autopsyâ€based histopathological evaluation. Pathology International, 2021, 71, 530-537.	0.6	2
12	Two autopsy cases with injuries to the stomach following cardiopulmonary resuscitation. Legal Medicine, 2021, 53, 101916.	0.6	1
13	Sexâ€dependent expression of prostatic markers and hormone receptors in cystic tumor of the atrioventricular node: A histopathological study of three cases. Pathology International, 2021, 71, 141-146.	0.6	2
14	Clinicopathologic Appearance of Advanced Ketoacidosis With Basal Vacuolation in Renal Tubules. Archives of Pathology and Laboratory Medicine, 2021, , .	1.2	0
15	Rescue from Stx2-Producing E.Âcoli-Associated Encephalopathy by Intravenous Injection of Muse Cells in NOD-SCID Mice. Molecular Therapy, 2020, 28, 100-118.	3.7	13
16	Two autopsy cases of sudden unexpected death from Dravet syndrome with novel de novo SCN1A variants. Brain and Development, 2020, 42, 171-178.	0.6	10
17	An autopsy case of sudden unexpected death with loxoprofen sodiumâ€“induced allergic eosinophilic coronary periarteritis. Cardiovascular Pathology, 2020, 44, 107154.	0.7	5
18	An autopsy case of amyloid tubulopathy exhibiting characteristic spheroid-type deposition. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 477, 157-163.	1.4	3

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19	Spheroid-type transthyretin amyloidosis in the gallbladder: A possible histopathological diagnostic clue to prevent overlooking cholecystic latent amyloid deposition. Human Pathology: Case Reports, 2020, 21, 200407.	0.2	1
20	Left Ventricular Noncompaction and Congenital Heart Disease Increases the Risk of Congestive Heart Failure. Journal of Clinical Medicine, 2020, 9, 785.	1.0	23
21	Increased Burden of Ion Channel Gene Variants Is Related to Distinct Phenotypes in Pediatric Patients With Left Ventricular Noncompaction. Circulation Genomic and Precision Medicine, 2020, 13, e002940.	1.6	9
22	Autopsy of a multiple lobar hemorrhage case with amyloidâ€Î²â€related angiitis. Neuropathology, 2020, 40, 280-286.	0.7	9
23	TBX5 R264K acts as a modifier to develop dilated cardiomyopathy in mice independently of T-box pathway. PLoS ONE, 2020, 15, e0227393.	1.1	8
24	Title is missing!. , 2020, 15, e0227393.		0
25	Title is missing!. , 2020, 15, e0227393.		0
26	Title is missing!. , 2020, 15, e0227393.		0
27	Title is missing!. , 2020, 15, e0227393.		0
28	Tau and Amyloid-Î² Pathology in Japanese Forensic Autopsy Series Under 40 Years of Age: Prevalence and Association with APOE Genotype and Suicide Risk. Journal of Alzheimer's Disease, 2019, 72, 641-652.	1.2	8
29	An Autopsy Case of Preclinical/Early Clinical Pick Disease. Journal of Neuropathology and Experimental Neurology, 2019, 78, 971-974.	0.9	1
30	Minimal inflammatory foci of unknown etiology may be a tentative sign of early stage inherited cardiomyopathy. Modern Pathology, 2019, 32, 1281-1290.	2.9	16
31	Clinicopathological and Genetic Profiles of Cases with Myocytes Disarrayâ€"Investigation for Establishing the Autopsy Diagnostic Criteria for Hypertrophic Cardiomyopathy. Journal of Clinical Medicine, 2019, 8, 463.	1.0	11
32	Sudden unexpected death with rare compound heterozygous variants in PRICKLE1. Neurogenetics, 2019, 20, 39-43.	0.7	3
33	A mutant HCN4 channel in a family with bradycardia, left bundle branch block, and left ventricular noncompaction. Heart and Vessels, 2018, 33, 802-819.	0.5	17
34	Autopsy of an Elderly Man With Incidentally Diagnosed TDP-43 Proteinopathy. Alzheimer Disease and Associated Disorders, 2018, 32, 158-161.	0.6	1
35	Endothelin-1 may play an important role in the Fontan circulation. Interactive Cardiovascular and Thoracic Surgery, 2018, 26, 480-486.	0.5	2
36	Lack of modulatory effect of the SCN5A R1193Q polymorphism on cardiac fast Na+ current at body temperature. PLoS ONE, 2018, 13, e0207437.	1.1	4

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37	Clinical and Echocardiographic Impact of Tafazzin Variants on Dilated Cardiomyopathy Phenotype in Left Ventricular Non-Compaction Patients in Early Infancy. Circulation Journal, 2018, 82, 2609-2618.	0.7	7
38	Fetal closed head injuries following maternal motor vehicle accident. Medicine (United States), 2018, 97, e13133.	0.4	6
39	Sudden unexpected death with primary adrenal lymphoma. Legal Medicine, 2018, 35, 25-28.	0.6	3
40	Sarcomere gene variants act as a genetic trigger underlying the development of left ventricular noncompaction. Pediatric Research, 2018, 84, 733-742.	1.1	26
41	The TNNI3 Arg192His mutation in a 13-year-old girl with left ventricular noncompaction. Journal of Cardiology Cases, 2018, 18, 33-36.	0.2	7
42	Latent pathogenicity of the G38S polymorphism of KCNE1ÂK+ channel modulator. Heart and Vessels, 2017, 32, 186-192.	0.5	6
43	Epilepsyâ€related sudden unexpected death: targeted molecular analysis of inherited heart disease genes using nextâ€generation DNA sequencing. Brain Pathology, 2017, 27, 292-304.	2.1	42
44	Incipient progressive supranuclear palsy is more common than expected and may comprise clinicopathological subtypes: a forensic autopsy series. Acta Neuropathologica, 2017, 133, 809-823.	3.9	58
45	Intramyocardial bronchogenic cyst: histological appearance and a review of the literature. Cardiovascular Pathology, 2017, 28, 64-67.	0.7	6
46	Argyrophilic grain disease in a 46-year-old male suicide victim. Journal of the Neurological Sciences, 2017, 380, 223-225.	0.3	6
47	A Wide and Specific Spectrum of Genetic Variants and Genotypeâ€“Phenotype Correlations Revealed by Nextâ€Generation Sequencing in Patients with Left Ventricular Noncompaction. Journal of the American Heart Association, 2017, 6, .	1.6	46
48	An Autopsy Case of Sudden Unexpected Death of a Young Adult in a Hot Bath: Molecular Analysis Using Next-Generation DNA Sequencing. Clinical Medicine Insights: Case Reports, 2017, 10, 117954761770288.	0.3	6
49	Sudden unexpected death in early Parkinson's disease: neurogenic or cardiac death?. Cardiovascular Pathology, 2017, 30, 19-22.	0.7	16
50	A Case of Sudden Unexpected Death with the Presence of Multiple Myocardial Bridges. International Journal of Clinical Cardiology, 2017, 4, .	0.1	1
51	Pseudoaneurysmal defect of mitralâ€aortic intervalvular fibrosa is likely to be a cerebral embolic source. An autopsy study in remote period from the stroke. Pathology International, 2016, 66, 472-474.	0.6	0
52	Left ventricular nonâ€compaction revealed by aortic regurgitation due to Kawasaki disease in a boy with LDB3 mutation. Pediatrics International, 2016, 58, 797-800.	0.2	4
53	SCN5A(K817E), a novel Brugada syndromeâ€“associated mutation that alters the activation gating of NaV1.5 channel. Heart Rhythm, 2016, 13, 1113-1120.	0.3	10
54	Postmortem genetic analysis of sudden unexplained death syndrome under 50 years of age: A next-generation sequencing study. Heart Rhythm, 2016, 13, 1544-1551.	0.3	40

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55	A novel ACTC1 mutation in a young boy with left ventricular noncompaction and arrhythmias. HeartRhythm Case Reports, 2016, 2, 92-97.	0.2	3
56	Anomalous origin of the right coronary artery evaluated with multidetector computed tomography and its clinical relevance. Journal of Cardiology, 2016, 68, 196-201.	0.8	11
57	Neuropathologic Features of Suicide Victims Who Presented With Acute Poststroke Depression. Journal of Neuropathology and Experimental Neurology, 2015, 74, 401-410.	0.9	17
58	The Susceptibilities of Human Ether-Ã-Go-Go-Related Gene Channel with the G487R Mutation to Arrhythmogenic Factors. Biological and Pharmaceutical Bulletin, 2015, 38, 781-784.	0.6	1
59	A Novel MYH7 Gene Mutation in a Fetus With Left Ventricular Noncompaction. Canadian Journal of Cardiology, 2015, 31, 103.e1-103.e3.	0.8	11
60	Anomalous origin of the right coronary artery from the left coronary sinus with an intramural course: comparison between sudden-death and non-sudden-death cases. Cardiovascular Pathology, 2015, 24, 154-159.	0.7	14
61	A case of sudden death after Japanese encephalitis vaccination. Legal Medicine, 2015, 17, 279-282.	0.6	5
62	Pathological features of preclinical or early clinical stages of corticobasal degeneration: a comparison with advanced cases. Neuropathology and Applied Neurobiology, 2015, 41, 893-905.	1.8	24
63	An autopsy case of pneumococcal Waterhouse-Friderichsen syndrome with possible functional asplenia/hyposplenia. International Journal of Clinical and Experimental Pathology, 2015, 8, 7518-25.	0.5	10
64	High takeoff of the left main coronary artery at autopsy after sudden unexpected death in a male. Pathology, 2014, 46, 361-364.	0.3	4
65	An Autopsy Case of Acute and Nonalcoholic Thiamine-Deficient Encephalopathy. European Neurology, 2014, 71, 230-232.	0.6	2
66	A590T mutation in KCNQ1 C-terminal helix D decreases IKs channel trafficking and function but not Yotiao interaction. Journal of Molecular and Cellular Cardiology, 2014, 72, 273-280.	0.9	8
67	Identification and characterization of a novel genetic mutation with prolonged QT syndrome in an unexplained postoperative death. International Journal of Legal Medicine, 2014, 128, 105-115.	1.2	10
68	Sudden unexpected death owing to unilateral medial medullary infarction with early involvement of the respiratory center. Legal Medicine, 2014, 16, 146-149.	0.6	0
69	Glycine/Serine Polymorphism at Position 38 Influences KCNE1 Subunitâ€™s Modulatory Actions on Rapid and Slow Delayed Rectifier K⁺ Currents. Circulation Journal, 2014, 78, 610-618.	0.7	12
70	An autopsy case of infantile-onset vanishing white matter disease related to an EIF2B2 mutation (V85E) in a hemizygous region. International Journal of Clinical and Experimental Pathology, 2014, 7, 3355-62.	0.5	4
71	Medical neglect death due to acute lymphoblastic leukaemia: an autopsy case report. Fukuoka Acta Medica, 2014, 105, 234-40.	0.1	0
72	14-3-3Îµ Gene variants in a Japanese patient with left ventricular noncompaction and hypoplasia of the corpus callosum. Gene, 2013, 515, 173-180.	1.0	15

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73	Distribution of myofibroblast and tenascinâ€‹<scp>C</scp> in cystic adventitial disease: Comparison with ganglion. Pathology International, 2013, 63, 591-598.	0.6	7
74	Impaired nerve fiber regeneration in axotomized peripheral nerves in streptozotocinâ€‹diabetic rats. Journal of Diabetes Investigation, 2013, 4, 533-539.	1.1	9
75	Surgical Repair of Left Ventricular Noncompaction in a Patient with a Novel Mutation of the Myosin Heavy Chain 7 Gene. Tohoku Journal of Experimental Medicine, 2012, 228, 301-304.	0.5	5
76	A Novel Missense Mutation Causing a G487R Substitution in the S2â€“S3 Loop of Human etherâ€Ââ€goâ€goâ€Related Gene Channel. Journal of Cardiovascular Electrophysiology, 2012, 23, 1246-1253.	0.8	5
77	Determination of fluoride in human whole blood and urine by gas chromatography-mass spectrometry. Forensic Toxicology, 2008, 26, 23-26.	1.4	26
78	Immunohistochemical study of thyroid transcription factor-1 and surfactant-associated protein A for investigation of peripheral airway structure in perinatal fatality. Legal Medicine, 2008, 10, 96-100.	0.6	2
79	Indication and limitations of using palatal rugae for personal identification in edentulous cases. Forensic Science International, 2008, 176, 178-182.	1.3	46
80	Patent Ductus Arteriosus with Infective Endocarditis at Age 92. Internal Medicine, 2008, 47, 263-268.	0.3	18
81	The sudden and unexpected death of a female-to-male transsexual patient. Journal of Clinical Forensic and Legal Medicine, 2007, 14, 382-386.	0.5	7
82	Pathological demonstration of rapid involvement into the subcutaneous tissue in a case of fatal hydrofluoric acid burns. Forensic Science International, 2007, 167, 49-52.	1.3	25
83	Fatal hemopericardium caused by retrograde acute abdominal aortic dissection: An autopsy report and morphological consideration. International Journal of Cardiology, 2006, 112, 253-255.	0.8	6
84	Relationship between cardiopulmonary resuscitation and injuries of the cardiac conduction system: Pathological features and pathogenesis of such injuries*. Critical Care Medicine, 2006, 34, 363-367.	0.4	19
85	Simple and sensitive determination of free and total morphine in human liver and kidney using gas chromatographyâ€“mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2006, 830, 359-363.	1.2	19
86	Sudden unexpected death of a 17-year-old male infected with the influenza virus. Legal Medicine, 2005, 7, 51-57.	0.6	8
87	Risk factors of sudden death in the Japanese hot bath in the senior population. Forensic Science International, 2005, 149, 151-158.	1.3	69
88	Two adult cases of congenital atresia of the left coronary ostiumâ€”comparison of a sudden death case with a long-term survival case. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2005, 447, 742-746.	1.4	25
89	1. An Autopsy Case of Fatal Arrhythmia Induced by Injuries of the Atrioventricular Conduction System: A case report. Medicine, Science and the Law, 2004, 44, 353-358.	0.6	6
90	3. A Fatal Case of Poisoning by Lidocaine Overdosage â€“ Analysis of Lidocaine in Formalin-Fixed Tissues. Medicine, Science and the Law, 2004, 44, 266-271.	0.6	24

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91	Conduction system abnormalities in alcoholics with asymptomatic valvular disease who suffer sudden death. Legal Medicine, 2003, 5, 212-219.	0.6	12
92	Subendocardial small infarct in the superior ventricular septum as a cause of sudden death. Forensic Science International, 2003, 138, 62-67.	1.3	5
93	Unexpected death due to right-sided infective endocarditis in a methamphetamine abuser. Legal Medicine, 2003, 5, 65-68.	0.6	8
94	Intracerebral hemorrhage as the cause of death in a severely burned body. Legal Medicine, 2003, 5, 108-109.	0.6	0
95	Sudden Unexpected Death of a Methamphetamine Abuser with Cardiopulmonary Abnormalities. Medicine, Science and the Law, 2003, 43, 267-271.	0.6	30
96	Two cases of sudden death in obese psychiatric patients with microscopic cardiopulmonary abnormalities. Fukuoka Acta Medica, 2003, 94, 66-74.	0.1	2
97	Acute cerebral infarction caused by congenital hypoplasty of cerebral artery in a severe burn case. Legal Medicine, 2002, 4, 119-122.	0.6	1
98	HPLC simultaneous determination of glycerol and mannitol in human tissues for forensic analysis. Forensic Science International, 2002, 125, 127-133.	1.3	16
99	Distribution of tetracaine and its metabolite in rabbits after high versus normal spinal anesthesia. Forensic Science International, 2001, 124, 130-136.	1.3	6
100	Identification of reddish alcoholic beverages by GC/MS using aroma components as indicators. Legal Medicine, 2001, 3, 237-240.	0.6	3
101	Cerebral Artery Thrombosis as a Cause of Striatocapsular Infarction. Cerebrovascular Diseases, 2000, 10, 151-154.	0.8	13
102	Prevalence of Renal Artery Stenosis in Autopsy Patients With Stroke. Stroke, 2000, 31, 61-65.	1.0	108
103	Sudden death of an infant with bronchopulmonary dysplasia and bilateral cervical lymphadenopathyâ€"a vagal death?. Legal Medicine, 2000, 2, 106-109.	0.6	0
104	The anatomical location of conduction system. Legal Medicine, 2000, 2, 123.	0.6	0
105	Sudden unexpected death with dysplastic change in the atrioventricular node artery. Legal Medicine, 2000, 2, 216-220.	0.6	6
106	Histopathological characterization of aortic intimal sarcoma with multiple tumor emboli. Pathology International, 2000, 50, 923-927.	0.6	26
107	Recurrent incompetence of repaired floppy mitral valves and the severity of myxomatous degeneration. Surgery Today, 2000, 30, 497-502.	0.7	3
108	Coronary atherosclerosis and interventions: Pathological sequences and restenosis. Pathology International, 1999, 49, 273-290.	0.6	55

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109	Sutured Wound in Myxomatous Mitral Valve One Month After Repair Surgery: Report of Two Cases. Surgery Today, 1999, 29, 811-812.	0.7	1
110	Two cases of mesothelial/monocytic incidental cardiac excrescences of the heart. Pathology International, 1998, 48, 641-644.	0.6	12
111	A Huge Coronary Aneurysm Resulting from a Coronary Artery-to-Left Ventricle Fistula.. Internal Medicine, 1998, 37, 366-369.	0.3	9
112	An unusual heterotopia of pyloric glands of the stomach with inverted downgrowth. Pathology International, 1993, 43, 192-197.	0.6	11