Michael Lardelli

List of Publications by Year
in descending order

94 papers	4,490 citations	117625 34 h-index	110387 64 g-index
119 all docs	119 docs citations	119 times ranked	5195 citing authors

#	Article	IF	Citations
1	Brain transcriptomes of zebrafish and mouse Alzheimer's disease knock-in models imply early disrupted energy metabolism. DMM Disease Models and Mechanisms, 2022, 15, .	2.4	8
2	Brain Transcriptome Analysis of a Protein-Truncating Mutation in Sortilin-Related Receptor 1 Associated With Early-Onset Familial Alzheimerâ€™s Disease Indicates Early Effects on Mitochondrial and Ribosome Function. Journal of Alzheimer's Disease, 2021, 79, 1105-1119.	2.6	9
3	Transcriptome analyses of 7-day-old zebrafish larvae possessing a familial Alzheimerâ€™s disease-like mutation in psen1 indicate effects on oxidative phosphorylation, ECM and MCM functions, and iron homeostasis. BMC Genomics, 2021, 22, 211.	2.8	8
4	Relevance of a Truncated PRESENILIN 2 Transcript to Alzheimerâ€™s Disease and Neurodegeneration. Journal of Alzheimer's Disease, 2021, 80, 1479-1489.	2.6	4
5	In-Frame and Frameshift Mutations in Zebrafish Presenilin 2 Affect Different Cellular Functions in Young Adult Brains. Journal of Alzheimer's Disease Reports, 2021, 5, 395-404.	2.2	8
6	PRESENILIN 1 Mutations Causing Early-Onset Familial Alzheimerâ€™s Disease or Familial Acne Inversa Differ in Their Effects on Genes Facilitating Energy Metabolism and Signal Transduction. Journal of Alzheimer's Disease, 2021, 82, 327-347.	2.6	9
7	Loss of park7 activity has differential effects on expression of iron responsive element (IRE) gene sets in the brain transcriptome in a zebrafish model of Parkinsonâ€™s disease. Molecular Brain, 2021, 14, 83.	2.6	7
8	Zebrafish Chromosome 14 Gene Differential Expression in the fmr1hu2787 Model of Fragile X Syndrome. Frontiers in Genetics, 2021, 12, 625466.	2.3	4
9	No observed effect on brain vasculature of Alzheimerâ€™s disease-related mutations in the zebrafish presenilin 1 gene. Molecular Brain, 2021, 14, 22.	2.6	1
10	Iron Responsive Element-Mediated Responses to Iron Dyshomeostasis in Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2021, 84, 1597-1630.	2.6	18
11	The evolved divergence of Î³-secretase-susceptibility of homologous proteins Ngfrb and Nradd in zebrafish. BMC Research Notes, 2021, 14, 460.	1.4	0
12	Transcriptome analysis indicates dominant effects on ribosome and mitochondrial function of a premature termination codon mutation in the zebrafish gene psen2. PLoS ONE, 2020, 15, e0232559.	2.5	11
13	Brain transcriptome analysis reveals subtle effects on mitochondrial function and iron homeostasis of mutations in the SORL1 gene implicated in early onset familial Alzheimerâ€™s disease. Molecular Brain, 2020, 13, 142.	2.6	26
14	Accelerated loss of hypoxia response in zebrafish with familial Alzheimerâ€™s disease-like mutation of presenilin 1. Human Molecular Genetics, 2020, 29, 2379-2394.	2.9	12
15	Sorting Out the Role of the Sortilin-Related Receptor 1 in Alzheimerâ€™s Disease. Journal of Alzheimer's Disease Reports, 2020, 4, 123-140.	2.2	22
16	Accelerated brain aging towards transcriptional inversion in a zebrafish model of the K115fs mutation of human PSEN2. PLoS ONE, 2020, 15, e0227258.	2.5	38
17	Brain transcriptome analysis of a familial Alzheimerâ€™s disease-like mutation in the zebrafish presenilin 1 gene implies effects on energy production. Molecular Brain, 2019, 12, 43.	2.6	33
18	Alternative splicing in a presenilin 2 variant associated with Alzheimer disease. Annals of Clinical and Translational Neurology, 2019, 6, 762-777.	3.7	29

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19	A Review of the Familial Alzheimerâ€™s Disease Locus PRESENILIN 2 and Its Relationship to PRESENILIN 1. Journal of Alzheimer's Disease, 2018, 66, 1323-1339.	2.6	17
20	The zebrafish orthologue of familial Alzheimerâ€™s disease gene PRESENILIN 2 is required for normal adult melanotic skin pigmentation. PLoS ONE, 2018, 13, e0206155.	2.5	25
21	Degenerate codon mixing for PCR-based manipulation of highly repetitive sequences. BMC Research Notes, 2018, 11, 202.	1.4	1
22	Dysregulation of Neuronal Iron Homeostasis as an Alternative Unifying Effect of Mutations Causing Familial Alzheimerâ€™s Disease. Frontiers in Neuroscience, 2018, 12, 533.	2.8	41
23	Animal Models of Alzheimerâ€™s Disease. , 2017, , 1031-1085.		9
24	Mitochondrion to endoplasmic reticulum apposition length in zebrafish embryo spinal progenitors is unchanged in response to perturbations associated with Alzheimerâ€™s disease. PLoS ONE, 2017, 12, e0179859.	2.5	3
25	The Enemy within: Innate Surveillance-Mediated Cell Death, the Common Mechanism of Neurodegenerative Disease. Frontiers in Neuroscience, 2016, 10, 193.	2.8	30
26	The Zebrafish Equivalent of Alzheimerâ€™s Disease-Associated PRESENILIN Isoform PS2V Regulates Inflammatory and Other Responses to Hypoxic Stress. Journal of Alzheimer's Disease, 2016, 52, 581-608.	2.6	25
27	Evidence For and Against a Pathogenic Role of Reduced Î³-Secretase Activity in Familial Alzheimerâ€™s Disease. Journal of Alzheimer's Disease, 2016, 52, 781-799.	2.6	44
28	Alzheimer's disease-related peptide PS2V plays ancient, conserved roles in suppression of the unfolded protein response under hypoxia and stimulation of Î³-secretase activity. Human Molecular Genetics, 2015, 24, 3662-3678.	2.9	33
29	Analysis of nicastrin gene phylogeny and expression in zebrafish. Development Genes and Evolution, 2015, 225, 171-178.	0.9	4
30	Activity-dependent expression of neuronal PAS domain-containing protein 4 (npas4a) in the developing zebrafish brain. Frontiers in Neuroanatomy, 2014, 8, 148.	1.7	15
31	Hypoxia alters expression of Zebrafish Microtubule-associated protein Tau (mapta, maptb) gene transcripts. BMC Research Notes, 2014, 7, 767.	1.4	14
32	Using the zebrafish model for AlzheimerÃ¢â‚¬â„¢s disease research. Frontiers in Genetics, 2014, 5, 189.	2.3	110
33	The Comparison of Methods for Measuring Oxidative Stress in Zebrafish Brains. Zebrafish, 2014, 11, 248-254.	1.1	11
34	Differential, dominant activation and inhibition of Notch signalling and APP cleavage by truncations of PSEN1 in human disease. Human Molecular Genetics, 2014, 23, 602-617.	2.9	48
35	Identification and expression analysis of the zebrafish orthologues of the mammalian MAP1LC3 gene family. Experimental Cell Research, 2014, 328, 228-237.	2.6	13
36	The Development of an in vivo Î³-Secretase Assay using Zebrafish Embryos. Journal of Alzheimer's Disease, 2013, 36, 521-534.	2.6	11

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37	Models of Alzheimerâ€™s Disease. , 2013, , 595-632.		0
38	The Guinea Pig as a Model for Sporadic Alzheimerâ€™s Disease (AD): The Impact of Cholesterol Intake on Expression of AD-Related Genes. PLoS ONE, 2013, 8, e66235.	2.5	42
39	Regular Care and Maintenance of a Zebrafish (Danio rerio) Laboratory: An Introduction. Journal of Visualized Experiments, 2012, , e4196.	0.3	189
40	The BACE1-PSEN-AÎ²PP Regulatory Axis has an Ancient Role in Response to Low Oxygen/Oxidative Stress. Journal of Alzheimer's Disease, 2012, 28, 515-530.	2.6	50
41	Robust homeostasis of Presenilin1 protein levels by transcript regulation. Neuroscience Letters, 2012, 519, 14-19.	2.1	6
42	Evaluation of Color Preference in Zebrafish for Learning and Memory. Journal of Alzheimer's Disease, 2012, 28, 459-469.	2.6	104
43	Zebrafish as a tool in Alzheimer's disease research. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2011, 1812, 346-352.	3.8	60
44	Transgenic Zebrafish Recapitulating tbx16 Gene Early Developmental Expression. PLoS ONE, 2011, 6, e21559.	2.5	17
45	The response of HMGA1 to changes in oxygen availability is evolutionarily conserved. Experimental Cell Research, 2011, 317, 1503-1512.	2.6	23
46	Identification and expression analysis of the zebrafish orthologue of Klotho. Development Genes and Evolution, 2011, 221, 179-186.	0.9	12
47	The Peak of the Oil Age â€“ Analyzing the world oil production Reference Scenario in World Energy Outlook 2008. Energy Policy, 2010, 38, 1398-1414.	8.8	254
48	A hyperactive sleeping beauty transposase enhances transgenesis in zebrafish embryos. BMC Research Notes, 2010, 3, 282.	1.4	6
49	Cryptic organisation within an apparently irregular rostrocaudal distribution of interneurons in the embryonic zebrafish spinal cord. Experimental Cell Research, 2010, 316, 3292-3303.	2.6	6
50	A Zebrafish Melanophore Model of AmyloidÎ² Toxicity. Zebrafish, 2010, 7, 155-159.	1.1	21
51	Altering Presenilin Gene Activity in Zebrafish Embryos Causes Changes in Expression of Genes with Potential Involvement in Alzheimer's Disease Pathogenesis. Journal of Alzheimer's Disease, 2009, 16, 133-147.	2.6	25
52	Mining the Data on Coal. Science, 2009, 324, 880-881.	12.6	5
53	Zebrafish aplnra functions in epiboly. BMC Research Notes, 2009, 2, 231.	1.4	8
54	Independent and cooperative action of Psen2 with Psen1 in zebrafish embryos. Experimental Cell Research, 2009, 315, 2791-2801.	2.6	47

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55	Selective neuronal requirement for huntingtin in the developing zebrafish. Human Molecular Genetics, 2009, 18, 4830-4842.	2.9	47
56	Complex Splicing and Neural Expression of Duplicated Tau Genes in Zebrafish Embryos. Journal of Alzheimer's Disease, 2009, 18, 305-317.	2.6	46
57	Interference with splicing of Presenilin transcripts has potent dominant negative effects on Presenilin activity. Human Molecular Genetics, 2008, 17, 402-412.	2.9	48
58	Evolutionary and Expression Analysis of the Zebrafish Deubiquitylating Enzyme, Usp9. Zebrafish, 2007, 4, 95-101.	1.1	13
59	Huntingtin-deficient zebrafish exhibit defects in iron utilization and development. Human Molecular Genetics, 2007, 16, 1905-1920.	2.9	136
60	Alzheimer disease: Amyloidogenesis, the presenilins and animal models. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2007, 1772, 285-297.	3.8	119
61	The protein tyrosine phosphatase Pez regulates TGFÎ², epithelialâ€“mesenchymal transition, and organ development. Journal of Cell Biology, 2007, 178, 1223-1235.	5.2	76
62	A Rapid Apoptosis Assay Measuring Relative Acridine Orange Fluorescence in Zebrafish Embryos. Zebrafish, 2007, 4, 113-116.	1.1	115
63	Scientists need to confront economists about peak oil. Nature, 2007, 446, 257-257.	27.8	4
64	Zebrafish Angiotensin II Receptor-like 1a (agtrl1a) is expressed in migrating hypoblast, vasculature, and in multiple embryonic epithelia. Gene Expression Patterns, 2007, 7, 258-265.	0.8	30
65	Contribution of mGluR and Fmr1 functional pathways to neurite morphogenesis, craniofacial development and fragile X syndrome. Human Molecular Genetics, 2006, 15, 3446-3458.	2.9	117
66	Zebrafish fgfr1 is a member of the fgf8 synexpression group and is required for fgf8 signalling at the midbrain-hindbrain boundary. Development Genes and Evolution, 2004, 214, 285-95.	0.9	55
67	Expression of three zebrafish orthologs of human FMR1-related genes and their phylogenetic relationships. Development Genes and Evolution, 2004, 214, 567-574.	0.9	36
68	Expression of three spalt (sal) gene homologues in zebrafish embryos. Development Genes and Evolution, 2003, 213, 35-43.	0.9	22
69	The evolutionary relationships of zebrafish genes tbx6 , tbx16 / spadetail and mga. Development Genes and Evolution, 2003, 213, 519-522.	0.9	17
70	Expression Analysis of a Tyrosinase Promoter Sequence in Zebrafish. Pigment Cell & Melanoma Research, 2003, 16, 117-126.	3.6	15
71	Developmental control of Presenilin1 expression, endoproteolysis, and interaction in zebrafish embryos. Experimental Cell Research, 2003, 289, 124-132.	2.6	47
72	Generation and PCR Screening of Bacteriophage Î» Sublibraries Enriched for Rare Clones the		3

#	Article	IF	Citations
73	Nonspecific, Nested Suppression PCR Method for Isolation of Unknown Flanking DNA ("Cold-Start) Tj ETQq1 1 0.784314 rgBT /Overlock		1
74	Identification of a second presenilin gene in zebrafish with similarity to the human Alzheimer's disease gene presenilin2. Development Genes and Evolution, 2002, 212, 486-490.	0.9	58
75	The identity and distribution of neural cells expressing the mesodermal determinant spadetail. BMC Developmental Biology, 2002, 2, 9.	2.1	19
76	The structure and function of vertebrate fibroblast growth factor receptor 1. International Journal of Developmental Biology, 2002, 46, 393-400.	0.6	66
77	Characterization and Developmental Expression of the Amphioxus Homolog of Notch (AmphiNotch): Evolutionary Conservation of Multiple Expression Domains in Amphioxus and Vertebrates. Developmental Biology, 2001, 232, 493-507.	2.0	52
78	Simple, Directional cDNA Cloning for In Situ Transcript Hybridization Screens. BioTechniques, 2001, 31, 938-946.	1.8	6
79	Tyrosinase gene expression in zebrafish embryos. Development Genes and Evolution, 2001, 211, 150-153.	0.9	71
80	Evolutionary analysis of vertebrate Notch genes. Development Genes and Evolution, 2001, 211, 350-354.	0.9	35
81	Distinct and regulated expression of Notch receptors in hematopoietic lineages and during myeloid differentiation. European Journal of Immunology, 2001, 31, 3240-3247.	2.9	49
82	Nonspecific, Nested Suppression PCR Method for Isolation of Unknown Flanking DNA. BioTechniques, 2000, 28, 895-902.	1.8	12
83	Three novel Notch genes in zebrafish: implications for vertebrate Notch gene evolution and function. Development Genes and Evolution, 1997, 207, 51-63.	0.9	126
84	Sequence of zebrafish fibulin-1 and its expression in developing heart and other embryonic organs. Development Genes and Evolution, 1997, 207, 340-351.	0.9	16
85	Generation and PCR Screening of Bacteriophage h Sublibraries Enriched for Rare Clones (the) Tj ETQq1 1 0.784314 rgBT /Overlock 10		1
86	Expression of the Notch 3 intracellular domain in mouse central nervous system progenitor cells is lethal and leads to disturbed neural tube development. Mechanisms of Development, 1996, 59, 177-190.	1.7	104
87	Discriminatory Frontier policy. Nature, 1995, 376, 12-12.	27.8	1
88	Residents' rights. Nature, 1995, 378, 330-330.	27.8	0
89	Expression of Notch 1, 2 and 3 is regulated by epithelial-mesenchymal interactions and retinoic acid in the developing mouse tooth and associated with determination of ameloblast cell fate.. Journal of Cell Biology, 1995, 130, 407-418.	5.2	170
90	Nestin mRNA expression correlates with the central nervous system progenitor cell state in many, but not all, regions of developing central nervous system. Developmental Brain Research, 1995, 84, 109-129.	1.7	453

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91	Complementary and combinatorial patterns of Notch gene family expression during early mouse development. Mechanisms of Development, 1995, 53, 357-368.	1.7	167
92	The novel Notch homologue mouse Notch 3 lacks specific epidermal growth factor-repeats and is expressed in proliferating neuroepithelium. Mechanisms of Development, 1994, 46, 123-136.	1.7	302
93	The Human NOTCH1, 2, and 3 Genes Are Located at Chromosome Positions 9q34, 1p13-p11, and 19p13.2-p13.1 in Regions of Neoplasia-Associated Translocation. Genomics, 1994, 24, 253-258.	2.9	93
94	Motch A and Motch Bâ€"Two Mouse Notch Homologues Coexpressed in a Wide Variety of Tissues. Experimental Cell Research, 1993, 204, 364-372.	2.6	133