

Paul T Martin

List of Publications by Year in descending order

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Version: 2024-02-01

90
papers

3,608
citations

109137

35
h-index

143772

57
g-index

91
all docs

91
docs citations

91
times ranked

4046
citing authors

#	ARTICLE	IF	CITATIONS
1	Visualizing Muscle Sialic Acid Expression in the GNE ^{D207VTgGne^{-/-} Cmah^{-/-}} Model of GNE Myopathy: A Comparison of Dietary and Gene Therapy Approaches. <i>Journal of Neuromuscular Diseases</i> , 2022, 9, 53-71.	1.1	6
2	Short-term treatment of golden retriever muscular dystrophy (GRMD) dogs with rAAVrh74.MHCK7.GALGT2 induces muscle glycosylation and utrophin expression but has no significant effect on muscle strength. <i>PLoS ONE</i> , 2021, 16, e0248721.	1.1	7
3	Micro-laminin gene therapy can function as an inhibitor of muscle disease in the dy ^W mouse model of MDC1A. <i>Molecular Therapy - Methods and Clinical Development</i> , 2021, 21, 274-287.	1.8	7
4	Serum Antibodies to N-Glycolylneuraminic Acid Are Elevated in Duchenne Muscular Dystrophy and Correlate with Increased Disease Pathology in Cmah ^{mdx} Mice. <i>American Journal of Pathology</i> , 2021, 191, 1474-1486.	1.9	4
5	Cell migration by swimming: <i>Drosophila</i> adipocytes as a new in vivo model of adhesion-independent motility. <i>Seminars in Cell and Developmental Biology</i> , 2020, 100, 160-166.	2.3	2
6	Expansion of B4GALT7 linkeropathy phenotype to include perinatal lethal skeletal dysplasia. <i>European Journal of Human Genetics</i> , 2019, 27, 1569-1577.	1.4	10
7	Injury Activates a Dynamic Cytoprotective Network to Confer Stress Resilience and Drive Repair. <i>Current Biology</i> , 2019, 29, 3851-3862.e4.	1.8	22
8	rAAVrh74.MCK.GALGT2 Protects against Loss of Hemodynamic Function in the Aging mdx Mouse Heart. <i>Molecular Therapy</i> , 2019, 27, 636-649.	3.7	27
9	Ombitasvir, Paritaprevir, Ritonavir, and Dasabuvir With or Without Ribavirin in Patients With Kidney Disease. <i>Kidney International Reports</i> , 2019, 4, 245-256.	0.4	7
10	Proteolytic and Opportunistic Breaching of the Basement Membrane Zone by Immune Cells during Tumor Initiation. <i>Cell Reports</i> , 2019, 27, 2837-2846.e4.	2.9	36
11	Soluble Heparin Binding Epidermal Growth Factor-Like Growth Factor Is a Regulator of GALGT2 Expression and GALGT2-Dependent Muscle and Neuromuscular Phenotypes. <i>Molecular and Cellular Biology</i> , 2019, 39, .	1.1	12
12	Are High-Dose Steroids Really Necessary in Treatment of Autoimmune Hepatitis?. <i>Clinical Gastroenterology and Hepatology</i> , 2019, 17, 1948-1949.	2.4	0
13	rAAVrh74.MCK.GALGT2 Demonstrates Safety and Widespread Muscle Glycosylation after Intravenous Delivery in C57BL/6J Mice. <i>Molecular Therapy - Methods and Clinical Development</i> , 2019, 15, 305-319.	1.8	15
14	Fat Body Cells Are Motile and Actively Migrate to Wounds to Drive Repair and Prevent Infection. <i>Developmental Cell</i> , 2018, 44, 460-470.e3.	3.1	90
15	An Isolated Limb Infusion Method Allows for Broad Distribution of rAAVrh74.MCK.GALGT2 to Leg Skeletal Muscles in the Rhesus Macaque. <i>Molecular Therapy - Methods and Clinical Development</i> , 2018, 10, 89-104.	1.8	14
16	Live imaging of collagen deposition during skin development and repair in a collagen I GFP fusion transgenic zebrafish line. <i>Developmental Biology</i> , 2018, 441, 4-11.	0.9	43
17	Loss of CMAH during Human Evolution Primed the Monocyte Macrophage Lineage toward a More Inflammatory and Phagocytic State. <i>Journal of Immunology</i> , 2017, 198, 2366-2373.	0.4	37
18	Comparison of Serum rAAV Serotype-Specific Antibodies in Patients with Duchenne Muscular Dystrophy, Becker Muscular Dystrophy, Inclusion Body Myositis, or GNE Myopathy. <i>Human Gene Therapy</i> , 2017, 28, 737-746.	1.4	27

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19	Deletion of <i>Pofut1</i> in Mouse Skeletal Myofibers Induces Muscle Aging-Related Phenotypes in <i>cis</i> and in <i>trans</i> . <i>Molecular and Cellular Biology</i> , 2017, 37, .	1.1	11
20	Macrophage Functions in Tissue Patterning and Disease: New Insights from the Fly. <i>Developmental Cell</i> , 2017, 40, 221-233.	3.1	79
21	Induction of T-Cell Infiltration and Programmed Death Ligand 2 Expression by Adeno-Associated Virus in Rhesus Macaque Skeletal Muscle and Modulation by Prednisone. <i>Human Gene Therapy</i> , 2017, 28, 493-509.	1.4	17
22	Hepatitis C virus increases the risk of kidney disease among HIV-positive patients: Systematic review and meta-analysis. <i>Journal of Medical Virology</i> , 2016, 88, 487-497.	2.5	34
23	N-terminal β -Dystroglycan (β DG-N): A Potential Serum Biomarker for Duchenne Muscular Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 2016, 3, 247-260.	1.1	10
24	Corpse Engulfment Generates a Molecular Memory that Primes the Macrophage Inflammatory Response. <i>Cell</i> , 2016, 165, 1658-1671.	13.5	160
25	Wound repair: a showcase for cell plasticity and migration. <i>Current Opinion in Cell Biology</i> , 2016, 42, 29-37.	2.6	165
26	Accurate Reconstruction of Cell and Particle Tracks from 3D Live Imaging Data. <i>Cell Systems</i> , 2016, 3, 102-107.	2.9	8
27	Systems Analysis of the Dynamic Inflammatory Response to Tissue Damage Reveals Spatiotemporal Properties of the Wound Attractant Gradient. <i>Current Biology</i> , 2016, 26, 1975-1989.	1.8	48
28	B4GALNT2 (GALGT2) Gene Therapy Reduces Skeletal Muscle Pathology in the FKRP P448L Mouse Model of Limb Girdle Muscular Dystrophy 2I. <i>American Journal of Pathology</i> , 2016, 186, 2429-2448.	1.9	36
29	New treatment for hepatitis C in chronic kidney disease, dialysis, and transplant. <i>Kidney International</i> , 2016, 89, 988-994.	2.6	40
30	Novel Perspectives on the Hepatitis B Virus Vaccine in the Chronic Kidney Disease Population. <i>International Journal of Artificial Organs</i> , 2015, 38, 625-631.	0.7	11
31	Ephrin-Bs Drive Junctional Downregulation and Actin Stress Fiber Disassembly to Enable Wound Re-epithelialization. <i>Cell Reports</i> , 2015, 13, 1380-1395.	2.9	60
32	A role for Galgt1 in skeletal muscle regeneration. <i>Skeletal Muscle</i> , 2015, 5, 3.	1.9	13
33	Deletion of Galgt2 (B4Galnt2) Reduces Muscle Growth in Response to Acute Injury and Increases Muscle Inflammation and Pathology in Dystrophin-Deficient Mice. <i>American Journal of Pathology</i> , 2015, 185, 2668-2684.	1.9	20
34	A Comparative Study of N-glycolylneuraminic Acid (Neu5Gc) and Cytotoxic T Cell (CT) Carbohydrate Expression in Normal and Dystrophin-Deficient Dog and Human Skeletal Muscle. <i>PLoS ONE</i> , 2014, 9, e88226.	1.1	19
35	The Unravelling Link between Chronic Kidney Disease and Hepatitis C Infection. <i>New Journal of Science</i> , 2014, 2014, 1-9.	1.0	10
36	Vascular Delivery of rAAVrh74.MCK.GALGT2 to the Gastrocnemius Muscle of the Rhesus Macaque Stimulates the Expression of Dystrophin and Laminin β 2 Surrogates. <i>Molecular Therapy</i> , 2014, 22, 713-724.	3.7	61

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37	Novel evidence on hepatitis C virus-associated glomerular disease. <i>Kidney International</i> , 2014, 86, 466-469.	2.6	12
38	Active and passive immunization strategies based on the SDPM1 peptide demonstrate pre-clinical efficacy in the APP ^{swE} PSEN1 ^{dE9} mouse model for Alzheimer's disease. <i>Neurobiology of Disease</i> , 2014, 62, 31-43.	2.1	5
39	N-Glycolylneuraminic acid deficiency worsens cardiac and skeletal muscle pathophysiology in β -sarcoglycan-deficient mice. <i>Glycobiology</i> , 2013, 23, 833-843.	1.3	16
40	Identification of New Dystroglycan Complexes in Skeletal Muscle. <i>PLoS ONE</i> , 2013, 8, e73224.	1.1	34
41	Absence of Hepatitis B Resistance Mutants before Introduction of Oral Antiviral Therapy. <i>ISRN Hepatology</i> , 2013, 2013, 1-5.	0.9	2
42	A Method to Produce and Purify Recombinant Full-Length Recombinant Alpha Dystroglycan: Analysis of N- and O-Linked Monosaccharide Composition in CHO Cells with or without LARGE Overexpression. <i>PLOS Currents</i> , 2013, 5, .	1.4	4
43	Sarcospan-dependent Akt activation is required for utrophin expression and muscle regeneration. <i>Journal of Cell Biology</i> , 2012, 197, 1009-1027.	2.3	54
44	Distinct contributions of Galgt1 and Galgt2 to carbohydrate expression and function at the mouse neuromuscular junction. <i>Molecular and Cellular Neurosciences</i> , 2012, 51, 112-126.	1.0	11
45	The Evidence-Based Epidemiology of HCV-Associated Kidney Disease. <i>International Journal of Artificial Organs</i> , 2012, 35, 621-628.	0.7	5
46	Induction of a regenerative microenvironment in skeletal muscle is sufficient to induce embryonal rhabdomyosarcoma in p53-deficient mice. <i>Journal of Pathology</i> , 2012, 226, 40-49.	2.1	22
47	Comparative Proteomic Profiling of Dystroglycan-Associated Proteins in Wild Type, <i>mdx</i> , and <i>Galgt2</i> Transgenic Mouse Skeletal Muscle. <i>Journal of Proteome Research</i> , 2012, 11, 4413-4424.	1.8	41
48	Management of Chronic Hepatitis B in Special Populations: Immunosuppressed Patients and Chronic Kidney Disease. <i>Current Hepatitis Reports</i> , 2011, 10, 269-276.	0.3	0
49	Role of extracellular matrix proteins and their receptors in the development of the vertebrate neuromuscular junction. <i>Developmental Neurobiology</i> , 2011, 71, 982-1005.	1.5	127
50	Immunization with the SDPM1 peptide lowers amyloid plaque burden and improves cognitive function in the APP ^{swE} PSEN1(A246E) transgenic mouse model of Alzheimer's disease. <i>Neurobiology of Disease</i> , 2010, 39, 409-422.	2.1	10
51	Chronic Kidney Disease after Liver Transplantation: Recent Evidence. <i>International Journal of Artificial Organs</i> , 2010, 33, 803-811.	0.7	31
52	Therapy with Nucleos(t)ide Analogues: Current Role in Dialysis Patients. <i>International Journal of Artificial Organs</i> , 2010, 33, 329-338.	0.7	12
53	A Human-Specific Deletion in Mouse <i>Cmah</i> Increases Disease Severity in the <i>mdx</i> Model of Duchenne Muscular Dystrophy. <i>Science Translational Medicine</i> , 2010, 2, 42ra54.	5.8	91
54	Mice Lacking Dystrophin or β -Sarcoglycan Spontaneously Develop Embryonal Rhabdomyosarcoma with Cancer-Associated p53 Mutations and Alternatively Spliced or Mutant Mdm2 Transcripts. <i>American Journal of Pathology</i> , 2010, 176, 416-434.	1.9	45

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55	Genetic Defects in Muscular Dystrophy. <i>Methods in Enzymology</i> , 2010, 479, 291-322.	0.4	21
56	Terlipressin for Hepatorenal Syndrome: A Meta-Analysis of Randomized Trials. <i>International Journal of Artificial Organs</i> , 2009, 32, 133-140.	0.7	66
57	Overexpression of <i>Galgt2</i> in skeletal muscle prevents injury resulting from eccentric contractions in both mdx and wild-type mice. <i>American Journal of Physiology - Cell Physiology</i> , 2009, 296, C476-C488.	2.1	78
58	Embryonic overexpression of <i>Galgt2</i> inhibits skeletal muscle growth via activation of myostatin signaling. <i>Muscle and Nerve</i> , 2009, 39, 25-41.	1.0	7
59	The synaptic CT carbohydrate modulates binding and expression of extracellular matrix proteins in skeletal muscle: Partial dependence on utrophin. <i>Molecular and Cellular Neurosciences</i> , 2009, 41, 448-463.	1.0	35
60	Overexpression of <i>Galgt2</i> Reduces Dystrophic Pathology in the Skeletal Muscles of Alpha Sarcoglycan-Deficient Mice. <i>American Journal of Pathology</i> , 2009, 175, 235-247.	1.9	47
61	O-fucosylation of muscle agrin determines its ability to cluster acetylcholine receptors. <i>Molecular and Cellular Neurosciences</i> , 2008, 39, 452-464.	1.0	34
62	Muscular dystrophy associated with β -dystroglycan deficiency in Sphynx and Devon Rex cats. <i>Neuromuscular Disorders</i> , 2008, 18, 942-952.	0.3	36
63	Long-term enhancement of skeletal muscle mass and strength by single gene administration of myostatin inhibitors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 4318-4322.	3.3	235
64	Congenital Muscular Dystrophies Involving the O-Mannose Pathway. <i>Current Molecular Medicine</i> , 2007, 7, 417-425.	0.6	61
65	Postnatal overexpression of the CT GalNAc transferase inhibits muscular dystrophy in mdx mice without altering muscle growth or neuromuscular development: Evidence for a utrophin-independent mechanism. <i>Neuromuscular Disorders</i> , 2007, 17, 209-220.	0.3	45
66	Overexpression of the Cytotoxic T Cell (CT) Carbohydrate Inhibits Muscular Dystrophy in the dyW Mouse Model of Congenital Muscular Dystrophy 1A. <i>American Journal of Pathology</i> , 2007, 171, 181-199.	1.9	54
67	Mechanisms of Disease: congenital muscular dystrophies—glycosylation takes center stage. <i>Nature Clinical Practice Neurology</i> , 2006, 2, 222-230.	2.7	42
68	The Congenital Muscular Dystrophies: Recent Advances and Molecular Insights. <i>Pediatric and Developmental Pathology</i> , 2006, 9, 427-443.	0.5	60
69	The Dystroglycanopathies: The New Disorders of O-Linked Glycosylation. <i>Seminars in Pediatric Neurology</i> , 2005, 12, 152-158.	1.0	65
70	Transgenic Overexpression of Dystroglycan Does Not Inhibit Muscular Dystrophy in mdx Mice. <i>American Journal of Pathology</i> , 2004, 164, 711-718.	1.9	24
71	Glycobiology of the neuromuscular junction. <i>Journal of Neurocytology</i> , 2003, 32, 915-929.	1.6	61
72	Treatment of hepatitis C. <i>Current Hepatitis Reports</i> , 2003, 2, 3-8.	0.3	1

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73	Overexpression of the CT GalNAc transferase inhibits muscular dystrophy in a cleavage-resistant dystroglycan mutant mouse. <i>Biochemical and Biophysical Research Communications</i> , 2003, 302, 831-836.	1.0	10
74	Role of transcription factors in skeletal muscle and the potential for pharmacological manipulation. <i>Current Opinion in Pharmacology</i> , 2003, 3, 300-308.	1.7	13
75	Inhibition of dystroglycan cleavage causes muscular dystrophy in transgenic mice. <i>Neuromuscular Disorders</i> , 2003, 13, 365-375.	0.3	54
76	Identification of peptides that specifically bind A β 1-40 amyloid in vitro and amyloid plaques in Alzheimer's disease brain using phage display. <i>Neurobiology of Disease</i> , 2003, 14, 146-156.	2.1	23
77	Glycobiology of neuromuscular disorders. <i>Glycobiology</i> , 2003, 13, 67R-75.	1.3	70
78	Dystroglycan glycosylation and its role in matrix binding in skeletal muscle. <i>Glycobiology</i> , 2003, 13, 55R-66.	1.3	83
79	Glycobiology of the synapse. <i>Glycobiology</i> , 2002, 12, 1R-7.	1.3	62
80	Overexpression of the cytotoxic T cell GalNAc transferase in skeletal muscle inhibits muscular dystrophy in mdx mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002, 99, 5616-5621.	3.3	137
81	Overexpression of the CT GalNAc Transferase in Skeletal Muscle Alters Myofiber Growth, Neuromuscular Structure, and Laminin Expression. <i>Developmental Biology</i> , 2002, 242, 58-73.	0.9	74
82	Definition of pre- and postsynaptic forms of the CT carbohydrate antigen at the neuromuscular junction: ubiquitous expression of the CT antigens and the CT GalNAc transferase in mouse tissues. <i>Molecular Brain Research</i> , 2002, 109, 146-160.	2.5	40
83	Modulation of Agrin Binding and Activity by the CT and Related Carbohydrate Antigens. <i>Molecular and Cellular Neurosciences</i> , 2002, 19, 539-551.	1.0	29
84	Tests for acute and chronic viral hepatitis. <i>Postgraduate Medicine</i> , 2000, 107, 123-130.	0.9	14
85	Liver disease. <i>Postgraduate Medicine</i> , 2000, 107, 95-96.	0.9	0
86	N-Acetyllactosamine and the CT Carbohydrate Antigen Mediate Agrin-Dependent Activation of MuSK and Acetylcholine Receptor Clustering in Skeletal Muscle. <i>Molecular and Cellular Neurosciences</i> , 2000, 15, 380-397.	1.0	40
87	Distribution of ten laminin chains in dystrophic and regenerating muscles. <i>Neuromuscular Disorders</i> , 1999, 9, 423-433.	0.3	77
88	Distinct Structures and Functions of Related Pre- and Postsynaptic Carbohydrates at the Mammalian Neuromuscular Junction. <i>Molecular and Cellular Neurosciences</i> , 1999, 13, 105-118.	1.0	58
89	Synaptic Integrins in Developing, Adult, and Mutant Muscle: Selective Association of β 1, β 7A, and β 7B Integrins with the Neuromuscular Junction. <i>Developmental Biology</i> , 1996, 174, 125-139.	0.9	162
90	Management of HCV in Dialysis Patients. , 0, , 50-54.		0