Martin Hrabe de Angelis

List of Publications by Citations

685 papers	46,033 citations	98 h-index	196 g-index
740 ext. papers	56,176 ext. citations	9.3 avg, IF	6.53 L-index

#	Paper	IF	Citations
685	Genetic studies of body mass index yield new insights for obesity biology. *Nature*, **2015**, 518, 197-206	50.4	2687
684	Discovery and refinement of loci associated with lipid levels. *Nature Genetics*, **2013**, 45, 1274-1283	36.3	1904
683	Large-scale association analysis provides insights into the genetic architecture and pathophysiology of type 2 diabetes. *Nature Genetics*, **2012**, 44, 981-90	36.3	1482
682	Defining the role of common variation in the genomic and biological architecture of adult human height. *Nature Genetics*, **2014**, 46, 1173-86	36.3	1339
681	Notch signalling controls pancreatic cell differentiation. *Nature*, **1999**, 400, 877-81	50.4	980
680	New genetic loci link adipose and insulin biology to body fat distribution. *Nature*, **2015**, 518, 187-196	50.4	920
679	Genome-wide trans-ancestry meta-analysis provides insight into the genetic architecture of type 2 diabetes susceptibility. *Nature Genetics*, **2014**, 46, 234-44	36.3	784
678	Human metabolic individuality in biomedical and pharmaceutical research. *Nature*, **2011**, 477, 54-60	50.4	728
677	The genetic architecture of type 2 diabetes. *Nature*, **2016**, 536, 41-47	50.4	704
676	Identification of serum metabolites associated with risk of type 2 diabetes using a targeted metabolomic approach. *Diabetes*, **2013**, 62, 639-48	0.9	634
675	Large-scale association analyses identify new loci influencing glycemic traits and provide insight into the underlying biological pathways. *Nature Genetics*, **2012**, 44, 991-1005	36.3	621
674	High-throughput discovery of novel developmental phenotypes. *Nature*, **2016**, 537, 508-514	50.4	608
673	Genome-wide, large-scale production of mutant mice by ENU mutagenesis. *Nature Genetics*, **2000**, 25, 444-7	36.3	578
672	Mutations in dynein link motor neuron degeneration to defects in retrograde transport. *Science*, **2003**, 300, 808-12	33.3	577
671	Maintenance of somite borders in mice requires the Delta homologue Dll1. *Nature*, **1997**, 386, 717-21	50.4	570
670	Genetics meets metabolomics: a genome-wide association study of metabolite profiles in human serum. *PLoS Genetics*, **2008**, 4, e1000282	6	538
669	Cardioprotection and lifespan extension by the natural polyamine spermidine. *Nature Medicine*, **2016**, 22, 1428-1438	50.5	532

668	A genome-wide perspective of genetic variation in human metabolism. *Nature Genetics*, **2010**, 42, 137-41	36.3	515
667	Novel biomarkers for pre-diabetes identified by metabolomics. *Molecular Systems Biology*, **2012**, 8, 615	12.2	468
666	A humanized version of Foxp2 affects cortico-basal ganglia circuits in mice. *Cell*, **2009**, 137, 961-71	56.2	427
665	Metabolic footprint of diabetes: a multiplatform metabolomics study in an epidemiological setting. *PLoS ONE*, **2010**, 5, e13953	3.7	425
664	An Expanded Genome-Wide Association Study of Type 2 Diabetes in Europeans. *Diabetes*, **2017**, 66, 2888-2902	8.9	414
663	Rare and low-frequency coding variants alter human adult height. *Nature*, **2017**, 542, 186-190	50.4	412
662	Cyclooxygenase-2 controls energy homeostasis in mice by de novo recruitment of brown adipocytes. *Science*, **2010**, 328, 1158-61	33.3	355
661	Aberrant methylation of tRNAs links cellular stress to neuro-developmental disorders. *EMBO Journal*, **2014**, 33, 2020-39	13	331
660	Animal models of obesity and diabetes mellitus. *Nature Reviews Endocrinology*, **2018**, 14, 140-162	15.2	330
659	Rare variant in scavenger receptor BI raises HDL cholesterol and increases risk of coronary heart disease. *Science*, **2016**, 351, 1166-71	33.3	325
658	Post-stroke inhibition of induced NADPH oxidase type 4 prevents oxidative stress and neurodegeneration. *PLoS Biology*, **2010**, 8, e1000479	9.7	324
657	Exome-wide association study of plasma lipids in >300,000 individuals. *Nature Genetics*, **2017**, 49, 1758-1766	36.3	310
656	Genetic fine mapping and genomic annotation defines causal mechanisms at type 2 diabetes susceptibility loci. *Nature Genetics*, **2015**, 47, 1415-25	36.3	292
655	A comparative phenotypic and genomic analysis of C57BL/6J and C57BL/6N mouse strains. *Genome Biology*, **2013**, 14, R82	18.3	288
654	Ribosomal mutations cause p53-mediated dark skin and pleiotropic effects. *Nature Genetics*, **2008**, 40, 963-70	36.3	285
653	Cytoplasmic thioredoxin reductase is essential for embryogenesis but dispensable for cardiac development. *Molecular and Cellular Biology*, **2005**, 25, 1980-8	4.8	283
652	Multi-ethnic genome-wide association study for atrial fibrillation. *Nature Genetics*, **2018**, 50, 1225-1233	36.3	277
651	Rapamycin extends murine lifespan but has limited effects on aging. *Journal of Clinical Investigation* , **2013**, 123, 3272-91	15.9	267

650	The genetics of blood pressure regulation and its target organs from association studies in 342,415 individuals. *Nature Genetics*, **2016**, 48, 1171-1184	36.3	251
649	An ENU-induced mutation of miR-96 associated with progressive hearing loss in mice. *Nature Genetics*, **2009**, 41, 614-8	36.3	249
648	The mammalian gene function resource: the International Knockout Mouse Consortium. *Mammalian Genome*, **2012**, 23, 580-6	3.2	230
647	The impact of low-frequency and rare variants on lipid levels. *Nature Genetics*, **2015**, 47, 589-97	36.3	229
646	Gene targeting by homologous recombination in mouse zygotes mediated by zinc-finger nucleases. *Proceedings of the National Academy of Sciences of the United States of America*, **2010**, 107, 15022-6	11.5	229
645	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. *Nature Genetics*, **2015**, 47, 1294-1303	36.3	226
644	Trans-ancestry genome-wide association study identifies 12 genetic loci influencing blood pressure and implicates a role for DNA methylation. *Nature Genetics*, **2015**, 47, 1282-1293	36.3	223
643	Impact of common genetic determinants of Hemoglobin A1c on type 2 diabetes risk and diagnosis in ancestrally diverse populations: A transethnic genome-wide meta-analysis. *PLoS Medicine*, **2017**, 14, e1002383	11.6	223
642	Refining the accuracy of validated target identification through coding variant fine-mapping in type 2 diabetes. *Nature Genetics*, **2018**, 50, 559-571	36.3	221
641	The Influence of Age and Sex on Genetic Associations with Adult Body Size and Shape: A Large-Scale Genome-Wide Interaction Study. *PLoS Genetics*, **2015**, 11, e1005378	6	220
640	Interaction between Notch signalling and Lunatic fringe during somite boundary formation in the mouse. *Current Biology*, **1999**, 9, 470-80	6.3	216
639	Mitochondrial glutathione peroxidase 4 disruption causes male infertility. *FASEB Journal*, **2009**, 23, 3233-42	6.4	212
638	Epigenetic germline inheritance of diet-induced obesity and insulin resistance. *Nature Genetics*, **2016**, 48, 497-9	36.3	211
637	Beethoven, a mouse model for dominant, progressive hearing loss DFNA36. *Nature Genetics*, **2002**, 30, 257-8	36.3	207
636	Delta-Notch signaling controls the generation of neurons/glia from neural stem cells in a stepwise process. *Development (Cambridge)*, **2003**, 130, 1391-402	6.6	206
635	Targeted estrogen delivery reverses the metabolic syndrome. *Nature Medicine*, **2012**, 18, 1847-56	50.5	201
634	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. *Nature Genetics*, **2014**, 46, 826-36	36.3	199
633	FMT-XCT: in vivo animal studies with hybrid fluorescence molecular tomography-X-ray computed tomography. *Nature Methods*, **2012**, 9, 615-20	21.6	192

632	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. *Nature Genetics*, **2018**, 50, 26-41	36.3	186
631	Modulation of the notch signaling by Mash1 and Dlx1/2regulates sequential specification and differentiation of progenitor cell types in the subcortical telencephalon. *Development (Cambridge)*, **2002**, 129, 5029-5040	6.6	184
630	Maternal and fetal genetic effects on birth weight and their relevance to cardio-metabolic risk factors. *Nature Genetics*, **2019**, 51, 804-814	36.3	181
629	Trans-ancestry meta-analyses identify rare and common variants associated with blood pressure and hypertension. *Nature Genetics*, **2016**, 48, 1151-1161	36.3	181
628	The Notch ligand Jagged1 is required for inner ear sensory development. *Proceedings of the National Academy of Sciences of the United States of America*, **2001**, 98, 3873-8	11.5	180
627	Large-scale analyses of common and rare variants identify 12 new loci associated with atrial fibrillation. *Nature Genetics*, **2017**, 49, 946-952	36.3	176
626	The European dimension for the mouse genome mutagenesis program. *Nature Genetics*, **2004**, 36, 925-7	36.3	176
625	Expression of Delta1 and Serrate1 (Jagged1) in the mouse inner ear. *Mechanisms of Development*, **1999**, 84, 169-72	1.7	176
624	Effects of G-protein mutations on skin color. *Nature Genetics*, **2004**, 36, 961-8	36.3	170
623	Towards frailty biomarkers: Candidates from genes and pathways regulated in aging and age-related diseases. *Ageing Research Reviews*, **2018**, 47, 214-277	12	160
622	Introducing the German Mouse Clinic: open access platform for standardized phenotyping. *Nature Methods*, **2005**, 2, 403-4	21.6	148
621	Inflammation and mitochondrial fatty acid beta-oxidation link obesity to early tumor promotion. *Proceedings of the National Academy of Sciences of the United States of America*, **2009**, 106, 3354-9	11.5	147
620	Systematic Evaluation of Pleiotropy Identifies 6 Further Loci Associated With Coronary Artery Disease. *Journal of the American College of Cardiology*, **2017**, 69, 823-836	15.1	146
619	Fibroblast Growth Factor 21-Metabolic Role in Mice and Men. *Endocrine Reviews*, **2017**, 38, 468-488	27.2	146
618	6 Somitogenesis. *Current Topics in Developmental Biology*, **1997**, 225-287	5.3	146
617	Disease model discovery from 3,328 gene knockouts by The International Mouse Phenotyping Consortium. *Nature Genetics*, **2017**, 49, 1231-1238	36.3	145
616	Neuronal 3,5,3-triiodothyronine (T3) uptake and behavioral phenotype of mice deficient in Mct8, the neuronal T3 transporter mutated in Allan-Herndon-Dudley syndrome. *Journal of Neuroscience*, **2009**, 29, 9439-49	6.6	143
615	Expression pattern of G protein-coupled receptor 30 in LacZ reporter mice. *Endocrinology*, **2009**, 150, 1722-30	4.8	143

614	Targeted disruption of the Walker-Warburg syndrome gene Pomt1 in mouse results in embryonic lethality. *Proceedings of the National Academy of Sciences of the United States of America*, **2004**, 101, 14126-31	11.5	133
613	Oscillating expression of c-Hey2 in the presomitic mesoderm suggests that the segmentation clock may use combinatorial signaling through multiple interacting bHLH factors. *Developmental Biology*, **2000**, 227, 91-103	3.1	133
612	Prevalence of sexual dimorphism in mammalian phenotypic traits. *Nature Communications*, **2017**, 8, 15475	17.4	130
611	EMPReSS: standardized phenotype screens for functional annotation of the mouse genome. *Nature Genetics*, **2005**, 37, 1155	36.3	130
610	Exome sequencing of 20,791 cases of type 2 diabetes and 24,440 controls. *Nature*, **2019**, 570, 71-76	50.4	129
609	Effects of ENU dosage on mouse strains. *Mammalian Genome*, **2000**, 11, 484-8	3.2	128
608	Calcitonin controls bone formation by inhibiting the release of sphingosine 1-phosphate from osteoclasts. *Nature Communications*, **2014**, 5, 5215	17.4	127
607	Direct production of mouse disease models by embryo microinjection of TALENs and oligodeoxynucleotides. *Proceedings of the National Academy of Sciences of the United States of America*, **2013**, 110, 3782-7	11.5	122
606	Rare variants in PPARG with decreased activity in adipocyte differentiation are associated with increased risk of type 2 diabetes. *Proceedings of the National Academy of Sciences of the United States of America*, **2014**, 111, 13127-32	11.5	121
605	Autoimmunity and inflammation due to a gain-of-function mutation in phospholipase C gamma 2 that specifically increases external Ca2+ entry. *Immunity*, **2005**, 22, 451-65	32.3	121
604	Mouse large-scale phenotyping initiatives: overview of the European Mouse Disease Clinic (EUMODIC) and of the Wellcome Trust Sanger Institute Mouse Genetics Project. *Mammalian Genome*, **2012**, 23, 600-10	3.2	116
603	Dominant-negative effects of a novel mutated Ins2 allele causes early-onset diabetes and severe beta-cell loss in Munich Ins2C95S mutant mice. *Diabetes*, **2007**, 56, 1268-76	0.9	116
602	Loss of mitochondrial peptidase Clpp leads to infertility, hearing loss plus growth retardation via accumulation of CLPX, mtDNA and inflammatory factors. *Human Molecular Genetics*, **2013**, 22, 4871-87	5.6	114
601	Chemical Hybridization of Glucagon and Thyroid Hormone Optimizes Therapeutic Impact for Metabolic Disease. *Cell*, **2016**, 167, 843-857.e14	56.2	114
600	ER stress-mediated apoptosis in a new mouse model of osteogenesis imperfecta. *PLoS Genetics*, **2008**, 4, e7	6	111
599	Genetics of dark skin in mice. *Genes and Development*, **2003**, 17, 214-28	12.6	110
598	Post-publication sharing of data and tools. *Nature*, **2009**, 461, 171-3	50.4	109
597	Secretome profiling of primary human skeletal muscle cells. *Biochimica Et Biophysica Acta - Proteins and Proteomics*, **2014**, 1844, 1011-7	4	107

596	Analysis of mammalian gene function through broad-based phenotypic screens across a consortium of mouse clinics. *Nature Genetics*, **2015**, 47, 969-978	36.3	106
595	Genome-wide meta-analysis of 241,258 adults accounting for smoking behaviour identifies novel loci for obesity traits. *Nature Communications*, **2017**, 8, 14977	17.4	105
594	Neuroprotection after stroke by targeting NOX4 as a source of oxidative stress. *Antioxidants and Redox Signaling*, **2013**, 18, 1418-27	8.4	105
593	Steroids in teleost fishes: A functional point of view. *Steroids*, **2015**, 103, 123-44	2.8	104
592	Mouse phenotyping. *Methods*, **2011**, 53, 120-35	4.6	103
591	Genome-wide physical activity interactions in adiposity - A meta-analysis of 200,452 adults. *PLoS Genetics*, **2017**, 13, e1006528	6	103
590	Novel genetic markers associate with atrial fibrillation risk in Europeans and Japanese. *Journal of the American College of Cardiology*, **2014**, 63, 1200-1210	15.1	102
589	Sequence interpretation. Functional annotation of mouse genome sequences. *Science*, **2001**, 291, 1251-5	33.3	101
588	The power of gene-based rare variant methods to detect disease-associated variation and test hypotheses about complex disease. *PLoS Genetics*, **2015**, 11, e1005165	6	98
587	Leveraging cross-species transcription factor binding site patterns: from diabetes risk loci to disease mechanisms. *Cell*, **2014**, 156, 343-58	56.2	96
586	Common variants at 12q15 and 12q24 are associated with infant head circumference. *Nature Genetics*, **2012**, 44, 532-538	36.3	94
585	Acute dietary fat intake initiates alterations in energy metabolism and insulin resistance. *Journal of Clinical Investigation*, **2017**, 127, 695-708	15.9	93
584	Cytokine response of primary human myotubes in an in vitro exercise model. *American Journal of Physiology - Cell Physiology*, **2013**, 305, C877-86	5.4	88
583	Distinct signatures of host-microbial meta-metabolome and gut microbiome in two C57BL/6 strains under high-fat diet. *ISME Journal*, **2014**, 8, 2380-96	11.9	87
582	Zebrafish and steroids: what do we know and what do we need to know?. *Journal of Steroid Biochemistry and Molecular Biology*, **2013**, 137, 165-73	5.1	85
581	ADAMTS-7 inhibits re-endothelialization of injured arteries and promotes vascular remodeling through cleavage of thrombospondin-1. *Circulation*, **2015**, 131, 1191-201	16.7	84
580	Genome-wide association study of kidney function decline in individuals of European descent. *Kidney International*, **2015**, 87, 1017-29	9.9	83
579	Towards better mouse models: enhanced genotypes, systemic phenotyping and envirotype modelling. *Nature Reviews Genetics*, **2009**, 10, 371-80	30.1	83

578	Node and midline defects are associated with left-right development in Delta1 mutant embryos. *Development (Cambridge)*, **2003**, 130, 3-13	6.6	83
577	A large scale hearing loss screen reveals an extensive unexplored genetic landscape for auditory dysfunction. *Nature Communications*, **2017**, 8, 886	17.4	81
576	Large scale ENU screens in the mouse: genetics meets genomics. *Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis*, **1998**, 400, 25-32	3.3	80
575	Restless legs syndrome-associated intronic common variant in Meis1 alters enhancer function in the developing telencephalon. *Genome Research*, **2014**, 24, 592-603	9.7	79
574	Roquin Suppresses the PI3K-mTOR Signaling Pathway to Inhibit T Helper Cell Differentiation and Conversion of Treg to Tfr Cells. *Immunity*, **2017**, 47, 1067-1082.e12	32.3	78
573	A genomic approach to therapeutic target validation identifies a glucose-lowering GLP1R variant protective for coronary heart disease. *Science Translational Medicine*, **2016**, 8, 341ra76	17.5	77
572	Circulating lysophosphatidylcholines are markers of a metabolically benign nonalcoholic fatty liver. *Diabetes Care*, **2013**, 36, 2331-8	14.6	77
571	Generation and characterization of dickkopf3 mutant mice. *Molecular and Cellular Biology*, **2006**, 26, 2317-26	4.8	77
570	Identification and functional characterization of G6PC2 coding variants influencing glycemic traits define an effector transcript at the G6PC2-ABCB11 locus. *PLoS Genetics*, **2015**, 11, e1004876	6	76
569	Effects of metformin on metabolite profiles and LDL cholesterol in patients with type 2 diabetes. *Diabetes Care*, **2015**, 38, 1858-67	14.6	76
568	52 Genetic Loci Influencing Myocardial Mass. *Journal of the American College of Cardiology*, **2016**, 68, 1435-1448	15.1	76
567	Novel multiple sclerosis susceptibility loci implicated in epigenetic regulation. *Science Advances*, **2016**, 2, e1501678	14.3	75
566	Mitochondrial dysfunction and decrease in body weight of a transgenic knock-in mouse model for TDP-43. *Journal of Biological Chemistry*, **2014**, 289, 10769-10784	5.4	72
565	Creatine improves health and survival of mice. *Neurobiology of Aging*, **2008**, 29, 1404-11	5.6	72
564	Identification of coexpressed gene clusters in a comparative analysis of transcriptome and proteome in mouse tissues. *Proceedings of the National Academy of Sciences of the United States of America*, **2005**, 102, 8621-6	11.5	71
563	Common genetic variation in the human FNDC5 locus, encoding the novel muscle-derived browning factor irisin, determines insulin sensitivity. *PLoS ONE*, **2013**, 8, e61903	3.7	71
562	Spinal poly-GA inclusions in a C9orf72 mouse model trigger motor deficits and inflammation without neuron loss. *Acta Neuropathologica*, **2017**, 134, 241-254	14.3	70
561	Urocortin 3 modulates social discrimination abilities via corticotropin-releasing hormone receptor type 2. *Journal of Neuroscience*, **2010**, 30, 9103-16	6.6	69

560	Androgen metabolism via 17beta-hydroxysteroid dehydrogenase type 3 in mammalian and non-mammalian vertebrates: comparison of the human and the zebrafish enzyme. *Journal of Molecular Endocrinology*, **2005**, 35, 305-16	4.5	69
559	Peri-conceptional obesogenic exposure induces sex-specific programming of disease susceptibilities in adult mouse offspring. *Biochimica Et Biophysica Acta - Molecular Basis of Disease*, **2014**, 1842, 304-17	6.9	68
558	Large-scale genome-wide analysis identifies genetic variants associated with cardiac structure and function. *Journal of Clinical Investigation*, **2017**, 127, 1798-1812	15.9	68
557	Pathophysiology-based subphenotyping of individuals at elevated risk for type 2 diabetes. *Nature Medicine*, **2021**, 27, 49-57	50.5	68
556	Toxicity modelling of Plk1-targeted therapies in genetically engineered mice and cultured primary mammalian cells. *Nature Communications*, **2011**, 2, 395	17.4	67
555	Expression of the mouse Delta1 gene during organogenesis and fetal development. *Mechanisms of Development*, **1999**, 84, 165-8	1.7	67
554	Systemic first-line phenotyping. *Methods in Molecular Biology*, **2009**, 530, 463-509	1.4	67
553	EuroPhenome: a repository for high-throughput mouse phenotyping data. *Nucleic Acids Research*, **2010**, 38, D577-85	20.1	66
552	A Myo6 mutation destroys coordination between the myosin heads, revealing new functions of myosin VI in the stereocilia of mammalian inner ear hair cells. *PLoS Genetics*, **2008**, 4, e1000207	6	66
551	Exome-wide association study reveals novel susceptibility genes to sporadic dilated cardiomyopathy. *PLoS ONE*, **2017**, 12, e0172995	3.7	66
550	Epigenetic alterations in longevity regulators, reduced life span, and exacerbated aging-related pathology in old father offspring mice. *Proceedings of the National Academy of Sciences of the United States of America*, **2018**, 115, E2348-E2357	11.5	65
549	Osteopontin deficiency protects against obesity-induced hepatic steatosis and attenuates glucose production in mice. *Diabetologia*, **2011**, 54, 2132-42	10.3	65
548	Requirement of the RNA-editing enzyme ADAR2 for normal physiology in mice. *Journal of Biological Chemistry*, **2011**, 286, 18614-22	5.4	64
547	The dominant alopecia phenotypes Bareskin, Rex-denuded, and Reduced Coat 2 are caused by mutations in gasdermin 3. *Genomics*, **2004**, 84, 824-35	4.3	62
546	Calcium oxalate stone formation in the inner ear as a result of an Slc26a4 mutation. *Journal of Biological Chemistry*, **2010**, 285, 21724-35	5.4	61
545	The redox environment triggers conformational changes and aggregation of hIAPP in Type II Diabetes. *Scientific Reports*, **2017**, 7, 44041	4.9	60
544	eIF6 coordinates insulin sensitivity and lipid metabolism by coupling translation to transcription. *Nature Communications*, **2015**, 6, 8261	17.4	60
543	Every-other-day feeding extends lifespan but fails to delay many symptoms of aging in mice. *Nature Communications*, **2017**, 8, 155	17.4	60

542	Large-scale cis- and trans-eQTL analyses identify thousands of genetic loci and polygenic scores that regulate blood gene expression. *Nature Genetics*, **2021**, 53, 1300-1310	36.3	60
541	Type 2 diabetes alters metabolic and transcriptional signatures of glucose and amino acid metabolism during exercise and recovery. *Diabetologia*, **2015**, 58, 1845-54	10.3	59
540	Discovery and Fine-Mapping of Glycaemic and Obesity-Related Trait Loci Using High-Density Imputation. *PLoS Genetics*, **2015**, 11, e1005230	6	59
539	DJ-1-deficient mice show less TH-positive neurons in the ventral tegmental area and exhibit non-motoric behavioural impairments. *Genes, Brain and Behavior*, **2010**, 9, 305-17	3.6	59
538	Missing-in-metastasis MIM/MTSS1 promotes actin assembly at intercellular junctions and is required for integrity of kidney epithelia. *Journal of Cell Science*, **2011**, 124, 1245-55	5.3	59
537	Efficient isolation of pure and functional mitochondria from mouse tissues using automated tissue disruption and enrichment with anti-TOM22 magnetic beads. *PLoS ONE*, **2013**, 8, e82392	3.7	59
536	Mutation in the betaA3/A1-crystallin encoding gene Cryba1 causes a dominant cataract in the mouse. *Genomics*, **1999**, 62, 67-73	4.3	58
535	Applying the ARRIVE Guidelines to an In Vivo Database. *PLoS Biology*, **2015**, 13, e1002151	9.7	56
534	MFAP4 Promotes Vascular Smooth Muscle Migration, Proliferation and Accelerates Neointima Formation. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **2016**, 36, 122-33	9.4	56
533	Identification of biomarkers for apoptosis in cancer cell lines using metabolomics: tools for individualized medicine. *Journal of Internal Medicine*, **2013**, 274, 425-39	10.8	56
532	Distinct regulatory elements direct delta1 expression in the nervous system and paraxial mesoderm of transgenic mice. *Mechanisms of Development*, **2000**, 95, 23-34	1.7	55
531	Modeling disease mutations by gene targeting in one-cell mouse embryos. *Proceedings of the National Academy of Sciences of the United States of America*, **2012**, 109, 9354-9	11.5	54
530	Fibroblast growth factor 21 is elevated in metabolically unhealthy obesity and affects lipid deposition, adipogenesis, and adipokine secretion of human abdominal subcutaneous adipocytes. *Molecular Metabolism*, **2015**, 4, 519-27	8.8	53
529	Noncanonical thyroid hormone signaling mediates cardiometabolic effects in vivo. *Proceedings of the National Academy of Sciences of the United States of America*, **2017**, 114, E11323-E11332	11.5	53
528	Cytochrome c oxidase subunit 4 isoform 2-knockout mice show reduced enzyme activity, airway hyporeactivity, and lung pathology. *FASEB Journal*, **2012**, 26, 3916-30	0.9	53
527	Functional inactivation of the genome-wide association study obesity gene neuronal growth regulator 1 in mice causes a body mass phenotype. *PLoS ONE*, **2012**, 7, e41537	3.7	53
526	The novel mouse mutation Oblivion inactivates the PMCA2 pump and causes progressive hearing loss. *PLoS Genetics*, **2008**, 4, e1000238	6	53
525	Profiling at mRNA, protein, and metabolite levels reveals alterations in renal amino acid handling and glutathione metabolism in kidney tissue of Pept2-/- mice. *Physiological Genomics*, **2007**, 28, 301-10	3.6	53

524	N-acyl Taurines and Acylcarnitines Cause an Imbalance in Insulin Synthesis and Secretion Provoking □Cell Dysfunction in Type 2 Diabetes. *Cell Metabolism*, **2017**, 25, 1334-1347.e4	24.6	52
523	SLIRP Regulates the Rate of Mitochondrial Protein Synthesis and Protects LRPPRC from Degradation. *PLoS Genetics*, **2015**, 11, e1005423	6	52
522	MIM-Induced Membrane Bending Promotes Dendritic Spine Initiation. *Developmental Cell*, **2015**, 33, 644-59	10.2	52
521	Conductivity of single-stranded and double-stranded deoxyribose nucleic acid under ambient conditions: The dominance of water. *Applied Physics Letters*, **2006**, 88, 102102	3.4	52
520	Variations of eye size parameters among different strains of mice. *Mammalian Genome*, **2006**, 17, 851-7	3.2	52
519	Oxalate-induced chronic kidney disease with its uremic and cardiovascular complications in C57BL/6 mice. *American Journal of Physiology - Renal Physiology*, **2016**, 310, F785-F795	4.3	51
518	Hyperoxaluria Requires TNF Receptors to Initiate Crystal Adhesion and Kidney Stone Disease. *Journal of the American Society of Nephrology: JASN*, **2017**, 28, 761-768	12.7	51
517	Bacterial encapsulins as orthogonal compartments for mammalian cell engineering. *Nature Communications*, **2018**, 9, 1990	17.4	51
516	Genetic loci associated with heart rate variability and their effects on cardiac disease risk. *Nature Communications*, **2017**, 8, 15805	17.4	50
515	Molecular Integration of Incretin and Glucocorticoid Action Reverses Immunometabolic Dysfunction and Obesity. *Cell Metabolism*, **2017**, 26, 620-632.e6	24.6	50
514	Calcineurin Links Mitochondrial Elongation with Energy Metabolism. *Cell Metabolism*, **2015**, 22, 838-50	24.6	50
513	Neuronal expression of glucosylceramide synthase in central nervous system regulates body weight and energy homeostasis. *PLoS Biology*, **2013**, 11, e1001506	9.7	50
512	Characterization of phospholipase C gamma enzymes with gain-of-function mutations. *Journal of Biological Chemistry*, **2009**, 284, 23083-93	5.4	50
511	Signs of the principle body axes prior to primitive streak formation in the rabbit embryo. *Anatomy and Embryology*, **1995**, 192, 159-69		50
510	Expression of Tas1 taste receptors in mammalian spermatozoa: functional role of Tas1r1 in regulating basal Ca□+ and cAMP concentrations in spermatozoa. *PLoS ONE*, **2012**, 7, e32354	3.7	50
509	A robust and reliable non-invasive test for stress responsivity in mice. *Frontiers in Behavioral Neuroscience*, **2014**, 8, 125	3.5	49
508	Transcriptome and proteome analysis of early embryonic mouse brain development. *Proteomics*, **2008**, 8, 1257-65	4.8	49
507	Sex-dependent susceptibility to Listeria monocytogenes infection is mediated by differential interleukin-10 production. *Infection and Immunity*, **2005**, 73, 5952-60	3.7	49

506	Large-scale phenotyping of an accurate genetic mouse model of JNCL identifies novel early pathology outside the central nervous system. *PLoS ONE*, **2012**, 7, e38310	3.7	49
505	Association of Atopic Dermatitis with Cardiovascular Risk Factors and Diseases. *Journal of Investigative Dermatology*, **2017**, 137, 1074-1081	4.3	48
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