

# Nancy Andrews

## List of Publications by Year in descending order

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96  
papers

21,080  
citations

24978

57  
h-index

40881

93  
g-index

96  
all docs

96  
docs citations

96  
times ranked

16544  
citing authors

#	ARTICLE	IF	CITATIONS
1	Metabolic Catastrophe in Mice Lacking Transferrin Receptor in Muscle. <i>EBioMedicine</i> , 2015, 2, 1705-1717.	2.7	62
2	Iron and Copper in Mitochondrial Diseases. <i>Cell Metabolism</i> , 2013, 17, 319-328.	7.2	142
3	Late stage erythroid precursor production is impaired in mice with chronic inflammation. <i>Haematologica</i> , 2012, 97, 1648-1656.	1.7	43
4	Divalent Metal Transporter 1 Regulates Iron-Mediated ROS and Pancreatic $\beta^2$ Cell Fate in Response to Cytokines. <i>Cell Metabolism</i> , 2012, 16, 449-461.	7.2	133
5	Closing the Iron Gate. <i>New England Journal of Medicine</i> , 2012, 366, 376-377.	13.9	31
6	The channel kinase, <i>TRPM7</i> , is required for early embryonic development. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, E225-33.	3.3	153
7	Mutation of Rubie, a Novel Long Non-Coding RNA Located Upstream of <i>Bmp4</i> , Causes Vestibular Malformation in Mice. <i>PLoS ONE</i> , 2012, 7, e29495.	1.1	23
8	Genetic Loss of <i>Tmprss6</i> Increases Effective Erythropoiesis in a Mouse Model of $\beta^2$ -Thalassemia. <i>Blood</i> , 2012, 120, 482-482.	0.6	0
9	Transferrin is a major determinant of hepcidin expression in hypotransferrinemic mice. <i>Blood</i> , 2011, 117, 630-637.	0.6	71
10	An Iron-Clad Role for Proteasomal Degradation. <i>Cell Metabolism</i> , 2011, 14, 281-282.	7.2	1
11	<i>Tmprss6</i> is a genetic modifier of the Hfe-hemochromatosis phenotype in mice. <i>Blood</i> , 2011, 117, 4590-4599.	0.6	80
12	Skeletal muscle hemojuvelin is dispensable for systemic iron homeostasis. <i>Blood</i> , 2011, 117, 6319-6325.	0.6	50
13	Proinflammatory state, hepcidin, and anemia in older persons. <i>Blood</i> , 2010, 115, 3810-3816.	0.6	191
14	Down-regulation of <i>Bmp/Smad</i> signaling by <i>Tmprss6</i> is required for maintenance of systemic iron homeostasis. <i>Blood</i> , 2010, 115, 3817-3826.	0.6	145
15	Hepcidin induction by transgenic overexpression of Hfe does not require the Hfe cytoplasmic tail, but does require hemojuvelin. <i>Blood</i> , 2010, 116, 5679-5687.	0.6	23
16	TRP Channel Regulates EGFR Signaling in Hair Morphogenesis and Skin Barrier Formation. <i>Cell</i> , 2010, 141, 331-343.	13.5	287
17	Ferrit(in)ing Out New Mechanisms in Iron Homeostasis. <i>Cell Metabolism</i> , 2010, 12, 203-204.	7.2	17
18	Hepcidin as a therapeutic tool to limit iron overload and improve anemia in $\beta^2$ -thalassemic mice. <i>Journal of Clinical Investigation</i> , 2010, 120, 4466-4477.	3.9	202

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19	Can we keep the "academic" in academic medicine?. Journal of Clinical Investigation, 2010, 120, 390-393.	3.9	4
20	ABCs of erythroid mitochondrial iron uptake. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 16012-16013.	3.3	7
21	Iron Is Essential for Neuron Development and Memory Function in Mouse Hippocampus. Journal of Nutrition, 2009, 139, 672-679.	1.3	159
22	Genes determining blood cell traits. Nature Genetics, 2009, 41, 1161-1162.	9.4	29
23	Scara5 Is a Ferritin Receptor Mediating Non-Transferrin Iron Delivery. Developmental Cell, 2009, 16, 35-46.	3.1	264
24	Mutations in TMPRSS6 cause iron-refractory iron deficiency anemia (IRIDA). Nature Genetics, 2008, 40, 569-571.	9.4	586
25	Chapter 6 Iron Homeostasis and Erythropoiesis. Current Topics in Developmental Biology, 2008, 82, 141-167.	1.0	50
26	The Transferrin Receptor Modulates Hfe-Dependent Regulation of Hepcidin Expression. Cell Metabolism, 2008, 7, 205-214.	7.2	315
27	Deficiency of heme-regulated eIF2 $\alpha$ kinase decreases hepcidin expression and splenic iron in HFE $^{-/-}$ mice. Haematologica, 2008, 93, 753-756.	1.7	20
28	Deletion of <i>Trpm7</i> Disrupts Embryonic Development and Thymopoiesis Without Altering Mg <sup>2+</sup> Homeostasis. Science, 2008, 322, 756-760.	6.0	379
29	Climbing through Medicine's Glass Ceiling. New England Journal of Medicine, 2007, 357, 1887-1889.	13.9	42
30	Hepcidin antimicrobial peptide transgenic mice exhibit features of the anemia of inflammation. Blood, 2007, 109, 4038-4044.	0.6	162
31	Of mice and iron: ferroportin disease. Blood, 2007, 109, 4115-4115.	0.6	1
32	When Is a Heme Transporter Not a Heme Transporter? When It's a Folate Transporter. Cell Metabolism, 2007, 5, 5-6.	7.2	29
33	Ineffective erythropoiesis in $\beta^2$ -thalassemia is characterized by increased iron absorption mediated by down-regulation of hepcidin and up-regulation of ferroportin. Blood, 2007, 109, 5027-5035.	0.6	277
34	Genetic variation in Mon1a affects protein trafficking and modifies macrophage iron loading in mice. Nature Genetics, 2007, 39, 1025-1032.	9.4	61
35	Transferrin receptor 1 is a cellular receptor for New World haemorrhagic fever arenaviruses. Nature, 2007, 446, 92-96.	13.7	374
36	Iron Homeostasis. Annual Review of Physiology, 2007, 69, 69-85.	5.6	557

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37	Modulation of bone morphogenetic protein signaling in vivo regulates systemic iron balance. <i>Journal of Clinical Investigation</i> , 2007, 117, 1933-1939.	3.9	401
38	The function of heme-regulated eIF2 $\alpha$ kinase in murine iron homeostasis and macrophage maturation. <i>Journal of Clinical Investigation</i> , 2007, 117, 3296-3305.	3.9	81
39	Chronic hepcidin induction causes hyposideremia and alters the pattern of cellular iron accumulation in hemochromatotic mice. <i>Blood</i> , 2006, 107, 2952-2958.	0.6	75
40	The Ins and Outs of Iron Homeostasis. <i>Physiology</i> , 2006, 21, 115-123.	1.6	69
41	Bone morphogenetic protein signaling by hemojuvelin regulates hepcidin expression. <i>Nature Genetics</i> , 2006, 38, 531-539.	9.4	921
42	Hereditary Hemochromatosis Protein, HFE, Interaction with Transferrin Receptor 2 Suggests a Molecular Mechanism for Mammalian Iron Sensing. <i>Journal of Biological Chemistry</i> , 2006, 281, 28494-28498.	1.6	297
43	Anemia of inflammation: the hepcidin link. <i>Current Opinion in Hematology</i> , 2005, 12, 107-111.	1.2	169
44	Haptoglobin modifies the hemochromatosis phenotype in mice. <i>Blood</i> , 2005, 105, 3353-3355.	0.6	36
45	A mutation in <i>Sec15l1</i> causes anemia in hemoglobin deficit ( <i>hbd</i> ) mice. <i>Nature Genetics</i> , 2005, 37, 1270-1273.	9.4	86
46	Iron in Skin of Mice with Three Etiologies of Systemic Iron Overload. <i>Journal of Investigative Dermatology</i> , 2005, 125, 1200-1205.	0.3	19
47	Case 21-2005. <i>New England Journal of Medicine</i> , 2005, 353, 189-198.	13.9	11
48	Understanding Heme Transport. <i>New England Journal of Medicine</i> , 2005, 353, 2508-2509.	13.9	47
49	The iron exporter ferroportin/ <i>Slc40a1</i> is essential for iron homeostasis. <i>Cell Metabolism</i> , 2005, 1, 191-200.	7.2	1,006
50	Molecular control of iron metabolism. <i>Best Practice and Research in Clinical Haematology</i> , 2005, 18, 159-169.	0.7	78
51	A mouse model of juvenile hemochromatosis. <i>Journal of Clinical Investigation</i> , 2005, 115, 2187-2191.	3.9	319
52	Complexity of CNC Transcription Factors As Revealed by Gene Targeting of the <i>Nrf3</i> Locus. <i>Molecular and Cellular Biology</i> , 2004, 24, 3286-3294.	1.1	87
53	The molecular regulation of iron metabolism. <i>The Hematology Journal</i> , 2004, 5, 373-380.	2.0	34
54	Transferrin is required for early T-cell differentiation. <i>Immunology</i> , 2004, 112, 543-549.	2.0	56

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55	An Hfe-dependent pathway mediates hyposideremia in response to lipopolysaccharide-induced inflammation in mice. <i>Nature Genetics</i> , 2004, 36, 481-485.	9.4	108
56	Balancing Acts. <i>Cell</i> , 2004, 117, 285-297.	13.5	1,544
57	Contributions of $\beta_2$ -microglobulin-dependent molecules and lymphocytes to iron regulation: insights from HfeRag1 <sup>-/-</sup> and $\beta_2mRag1$ <sup>-/-</sup> double knock-out mice. <i>Blood</i> , 2004, 103, 2847-2849.	0.6	31
58	Hepcidin, a candidate modifier of the hemochromatosis phenotype in mice. <i>Blood</i> , 2004, 103, 2841-2843.	0.6	46
59	Pathophysiologic mechanisms of anemia of chronic disease. <i>Postgraduate Medicine</i> , 2004, 116, 017-022.	0.9	0
60	Iron metabolism in mice with partial frataxin deficiency. <i>Cerebellum</i> , 2003, 2, 146-153.	1.4	5
61	Regulatory defects in liver and intestine implicate abnormal hepcidin and Cybrd1 expression in mouse hemochromatosis. <i>Nature Genetics</i> , 2003, 34, 102-107.	9.4	274
62	Constitutive hepcidin expression prevents iron overload in a mouse model of hemochromatosis. <i>Nature Genetics</i> , 2003, 34, 97-101.	9.4	284
63	Probucol prevents early coronary heart disease and death in the high-density lipoprotein receptor SR-BI/apolipoprotein E double knockout mouse. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 7283-7288.	3.3	132
64	Hfe deficiency increases susceptibility to cardiotoxicity and exacerbates changes in iron metabolism induced by doxorubicin. <i>Blood</i> , 2003, 102, 2574-2580.	0.6	139
65	Transferrin receptor 1 is differentially required in lymphocyte development. <i>Blood</i> , 2003, 102, 3711-3718.	0.6	103
66	Regulation of iron absorption in Hfe mutant mice. <i>Blood</i> , 2002, 100, 1465-1469.	0.6	78
67	Inappropriate expression of hepcidin is associated with iron refractory anemia: implications for the anemia of chronic disease. <i>Blood</i> , 2002, 100, 3776-3781.	0.6	572
68	A genetic view of iron homeostasis. <i>Seminars in Hematology</i> , 2002, 39, 227-234.	1.8	42
69	Metal transporters and disease. <i>Current Opinion in Chemical Biology</i> , 2002, 6, 181-186.	2.8	94
70	The other physician-scientist problem: Where have all the young girls gone?. <i>Nature Medicine</i> , 2002, 8, 439-441.	15.2	78
71	Iron-dependent regulation of the divalent metal ion transporter. <i>FEBS Letters</i> , 2001, 509, 309-316.	1.3	269
72	Mining copper transport genes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2001, 98, 6543-6545.	3.3	32

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73	Expression of the DMT1 (NRAMP2/DCT1) iron transporter in mice with genetic iron overload disorders. <i>Blood</i> , 2001, 97, 1138-1140.	0.6	90
74	Uroporphyrin in Hfe mutant mice given 5-aminolevulinic acid: A new model of Fe-mediated porphyria cutanea tarda. <i>Hepatology</i> , 2001, 33, 406-412.	3.6	20
75	A mutation in a mitochondrial transmembrane protein is responsible for the pleiotropic hematological and skeletal phenotype of flexed-tail ( <i>f/f</i> ) mice. <i>Genes and Development</i> , 2001, 15, 652-657.	2.7	84
76	A mouse model of familial porphyria cutanea tarda. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2001, 98, 259-64.	3.3	30
77	Autosomal-dominant hemochromatosis is associated with a mutation in the ferroportin (SLC11A3) gene. <i>Journal of Clinical Investigation</i> , 2001, 108, 619-623.	3.9	429
78	Inherited iron overload disorders. <i>Current Opinion in Pediatrics</i> , 2000, 12, 596-602.	1.0	10
79	Positional cloning of zebrafish ferroportin1 identifies a conserved vertebrate iron exporter. <i>Nature</i> , 2000, 403, 776-781.	13.7	1,491
80	Comparison of the Interactions of Transferrin Receptor and Transferrin Receptor 2 with Transferrin and the Hereditary Hemochromatosis Protein HFE. <i>Journal of Biological Chemistry</i> , 2000, 275, 38135-38138.	1.6	214
81	Ectopic Expression of Transcription Factor NF-E2 Alters the Phenotype of Erythroid and Monoblastoid Cells. <i>Journal of Biological Chemistry</i> , 2000, 275, 25292-25298.	1.6	13
82	The iron transporter DMT1. <i>International Journal of Biochemistry and Cell Biology</i> , 1999, 31, 991-994.	1.2	216
83	Transferrin receptor is necessary for development of erythrocytes and the nervous system. <i>Nature Genetics</i> , 1999, 21, 396-399.	9.4	510
84	Disorders of Iron Metabolism. <i>New England Journal of Medicine</i> , 1999, 341, 1986-1995.	13.9	1,693
85	Commentary on. <i>Journal of Pediatric Hematology/Oncology</i> , 1999, 21, 353-355.	0.3	4
86	Molecular insights into mechanisms of iron transport. <i>Current Opinion in Hematology</i> , 1999, 6, 61.	1.2	29
87	Molecules in focus The NF-E2 transcription factor. <i>International Journal of Biochemistry and Cell Biology</i> , 1998, 30, 429-432.	1.2	78
88	Nramp2 is mutated in the anemic Belgrade (b) rat: Evidence of a role for Nramp2 in endosomal iron transport. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1998, 95, 1148-1153.	3.3	898
89	Molecular Characterization and Localization of the Human MAFG Gene. <i>Genomics</i> , 1997, 44, 147-149.	1.3	10
90	The Maf transcription factors: regulators of differentiation. <i>Trends in Biochemical Sciences</i> , 1997, 22, 437-441.	3.7	254

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91	p45 NF-E2 regulates expression of thromboxane synthase in megakaryocytes. EMBO Journal, 1997, 16, 5654-5661.	3.5	73
92	Multiple Proteins Interact with the Nuclear Inhibitory Protein Repressor Element in the Human Interleukin-3 Promoter. Journal of Biological Chemistry, 1995, 270, 24572-24579.	1.6	16
93	Structure and regulation of the chicken erythroid $\hat{\Gamma}$ -aminolevulinate synthase gene. Nucleic Acids Research, 1994, 22, 1226-1233.	6.5	14
94	Novel Bacterial P-Type ATPases with Histidine-Rich Heavy-Metal-Associated Sequences. Biochemical and Biophysical Research Communications, 1994, 205, 1644-1650.	1.0	23
95	A rapid micropreparation technique for extraction of DNA-binding proteins from limiting numbers of mammalian cells. Nucleic Acids Research, 1991, 19, 2499-2499.	6.5	2,268
96	Phage T3 DNA contains an exact copy of the 23 base-pair phage T7 RNA polymerase promoter sequence. Journal of Molecular Biology, 1981, 147, 41-53.	2.0	20