John W R Schwabe

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#	Paper	IF	Citations
107	Dominant negative mutations in human PPARgamma associated with severe insulin resistance, diabetes mellitus and hypertension. <i>Nature</i> , 1999 , 402, 880-3	50.4	1129
106	The crystal structure of the estrogen receptor DNA-binding domain bound to DNA: how receptors discriminate between their response elements. <i>Cell</i> , 1993 , 75, 567-78	56.2	635
105	Zinc fingers. FASEB Journal, 1995 , 9, 597-604	0.9	504
104	Solution structure of the DNA-binding domain of the oestrogen receptor. <i>Nature</i> , 1990 , 348, 458-61	50.4	416
103	The crystal structure of a two zinc-finger peptide reveals an extension to the rules for zinc-finger/DNA recognition. <i>Nature</i> , 1993 , 366, 483-7	50.4	326
102	Mechanism of corepressor binding and release from nuclear hormone receptors. <i>Genes and Development</i> , 1999 , 13, 3209-16	12.6	323
101	Structural basis for the activation of PPARgamma by oxidized fatty acids. <i>Nature Structural and Molecular Biology</i> , 2008 , 15, 924-31	17.6	321
100	Structure of HDAC3 bound to co-repressor and inositol tetraphosphate. <i>Nature</i> , 2012 , 481, 335-40	50.4	319
99	Mechanism of the nuclear receptor molecular switch. <i>Trends in Biochemical Sciences</i> , 2004 , 29, 317-24	10.3	318
98	A mutation in the thyroid hormone receptor alpha gene. <i>New England Journal of Medicine</i> , 2012 , 366, 243-9	59.2	288
97	STAT6 transcription factor is a facilitator of the nuclear receptor PPARE gulated gene expression in macrophages and dendritic cells. <i>Immunity</i> , 2010 , 33, 699-712	32.3	284
96	Class I HDACs share a common mechanism of regulation by inositol phosphates. <i>Molecular Cell</i> , 2013 , 51, 57-67	17.6	247
95	A death effector domain chain DISC model reveals a crucial role for caspase-8 chain assembly in mediating apoptotic cell death. <i>Molecular Cell</i> , 2012 , 47, 291-305	17.6	237
94	Mutations in the selenocysteine insertion sequence-binding protein 2 gene lead to a multisystem selenoprotein deficiency disorder in humans. <i>Journal of Clinical Investigation</i> , 2010 , 120, 4220-35	15.9	229
93	A dominant-negative peroxisome proliferator-activated receptor gamma (PPARgamma) mutant is a constitutive repressor and inhibits PPARgamma-mediated adipogenesis. <i>Journal of Biological Chemistry</i> , 2000 , 275, 5754-9	5.4	222
92	St John's wort, a herbal antidepressant, activates the steroid X receptor. <i>Journal of Endocrinology</i> , 2000 , 166, R11-6	4.7	200
91	Radical fringe positions the apical ectodermal ridge at the dorsoventral boundary of the vertebrate limb. <i>Nature</i> , 1997 , 386, 360-6	50.4	186

(2017-2006)

90	International Union of Pharmacology. LXVI. Orphan nuclear receptors. <i>Pharmacological Reviews</i> , 2006 , 58, 798-836	22.5	175
89	The role of water in protein-DNA interactions. Current Opinion in Structural Biology, 1997, 7, 126-34	8.1	161
88	Co-operative and Hierarchical Binding of c-FLIP and Caspase-8: A Unified Model Defines How c-FLIP Isoforms Differentially Control Cell Fate. <i>Molecular Cell</i> , 2016 , 61, 834-49	17.6	143
87	Non-DNA binding, dominant-negative, human PPARgamma mutations cause lipodystrophic insulin resistance. <i>Cell Metabolism</i> , 2006 , 4, 303-11	24.6	143
86	Beyond zinc fingers: steroid hormone receptors have a novel structural motif for DNA recognition. <i>Trends in Biochemical Sciences</i> , 1991 , 16, 291-6	10.3	141
85	A dynamic mechanism of nuclear receptor activation and its perturbation in a human disease. Nature Structural Biology, 2003, 10, 136-40		134
84	Targeting the CoREST complex with dual histone deacetylase and demethylase inhibitors. <i>Nature Communications</i> , 2018 , 9, 53	17.4	116
83	Aster Proteins Facilitate Nonvesicular Plasma Membrane to ER Cholesterol Transport in Mammalian Cells. <i>Cell</i> , 2018 , 175, 514-529.e20	56.2	116
82	Structural basis for the assembly of the SMRT/NCoR core transcriptional repression machinery. <i>Nature Structural and Molecular Biology</i> , 2011 , 18, 177-84	17.6	114
81	Insights into the activation mechanism of class I HDAC complexes by inositol phosphates. <i>Nature Communications</i> , 2016 , 7, 11262	17.4	113
80	Nuclear hormone receptor co-repressors: structure and function. <i>Molecular and Cellular Endocrinology</i> , 2012 , 348, 440-9	4.4	112
79	The phantom ligand effect: allosteric control of transcription by the retinoid X receptor. <i>Genes and Development</i> , 1997 , 11, 299-308	12.6	112
78	A conserved structural motif reveals the essential transcriptional repression function of Spen proteins and their role in developmental signaling. <i>Genes and Development</i> , 2003 , 17, 1909-20	12.6	111
77	DNA recognition by the oestrogen receptor: from solution to the crystal. <i>Structure</i> , 1993 , 1, 187-204	5.2	96
76	Histone deacetylase (HDAC) 1 and 2 are essential for accurate cell division and the pluripotency of embryonic stem cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 9840-5	11.5	93
75	Coexpression of nuclear receptor partners increases their solubility and biological activities. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997 , 94, 2278-83	11.5	88
74	The oestrogen receptor recognizes an imperfectly palindromic response element through an alternative side-chain conformation. <i>Structure</i> , 1995 , 3, 201-13	5.2	87
73	Targeting Class I Histone Deacetylases in a "Complex" Environment. <i>Trends in Pharmacological Sciences</i> , 2017 , 38, 363-377	13.2	85

72	Solution structures of two zinc-finger domains from SWI5 obtained using two-dimensional 1H nuclear magnetic resonance spectroscopy. A zinc-finger structure with a third strand of beta-sheet. <i>Journal of Molecular Biology</i> , 1992 , 228, 637-51	6.5	78
71	Structural insights into the interaction and activation of histone deacetylase 3 by nuclear receptor corepressors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 6009-14	11.5	76
70	Negative regulation by nuclear receptors: a plethora of mechanisms. <i>Trends in Endocrinology and Metabolism</i> , 2011 , 22, 87-93	8.8	72
69	Residual activity of mutant androgen receptors explains wolffian duct development in the complete androgen insensitivity syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004 , 89, 5815-22	5.6	69
68	The structure of the core NuRD repression complex provides insights into its interaction with chromatin. <i>ELife</i> , 2016 , 5, e13941	8.9	68
67	An evolving understanding of nuclear receptor coregulator proteins. <i>Journal of Molecular Endocrinology</i> , 2013 , 51, T23-36	4.5	63
66	The IDOL-UBE2D complex mediates sterol-dependent degradation of the LDL receptor. <i>Genes and Development</i> , 2011 , 25, 1262-74	12.6	62
65	The structural basis for the specificity of retinoid-X receptor-selective agonists: new insights into the role of helix H12. <i>Journal of Biological Chemistry</i> , 2002 , 277, 11385-91	5.4	59
64	The structure of the ultraspiracle ligand-binding domain reveals a nuclear receptor locked in an inactive conformation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2001 , 98, 1549-54	11.5	59
63	Limbs are moving: where are they going?. <i>Trends in Genetics</i> , 1998 , 14, 229-35	8.5	55
62	Mutations in TBL1X Are Associated With Central Hypothyroidism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 4564-4573	5.6	54
61	Towards an understanding of protein-DNA recognition. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 1996 , 351, 501-9	5.8	54
60	Tyrosine agonists reverse the molecular defects associated with dominant-negative mutations in human peroxisome proliferator-activated receptor gamma. <i>Endocrinology</i> , 2004 , 145, 1527-38	4.8	51
59	Disruption of the Class IIa HDAC Corepressor Complex Increases Energy Expenditure and Lipid Oxidation. <i>Cell Reports</i> , 2016 , 16, 2802-2810	10.6	48
58	Identification of a novel co-regulator interaction surface on the ligand binding domain of Nurr1 using NMR footprinting. <i>Journal of Biological Chemistry</i> , 2004 , 279, 53338-45	5.4	48
57	Human androgen receptor gene ligand-binding-domain mutations leading to disrupted interaction between the N- and C-terminal domains. <i>Journal of Molecular Endocrinology</i> , 2006 , 36, 361-8	4.5	44
56	NOXA, a sensor of proteasome integrity, is degraded by 26S proteasomes by an ubiquitin-independent pathway that is blocked by MCL-1. <i>Cell Death and Differentiation</i> , 2012 , 19, 1424	- 34 ·7	43
55	Histone deacetylase (HDAC) 1 and 2 complexes regulate both histone acetylation and crotonylation in vivo. <i>Scientific Reports</i> , 2018 , 8, 14690	4.9	41

54	A specific mutation in TBL1XR1 causes Pierpont syndrome. <i>Journal of Medical Genetics</i> , 2016 , 53, 330-7	5.8	40
53	Channels at the catalytic site of glycogen phosphorylase b: binding and kinetic studies with the beta-glycosidase inhibitor D-gluconohydroximo-1,5-lactone N-phenylurethane. <i>Biochemistry</i> , 1988 , 27, 6733-41	3.2	39
52	FERM-dependent E3 ligase recognition is a conserved mechanism for targeted degradation of lipoprotein receptors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 20107-12	11.5	38
51	A dominant negative human peroxisome proliferator-activated receptor (PPAR){alpha} is a constitutive transcriptional corepressor and inhibits signaling through all PPAR isoforms. <i>Endocrinology</i> , 2005 , 146, 1871-82	4.8	38
50	Structural and functional characterization of a cell cycle associated HDAC1/2 complex reveals the structural basis for complex assembly and nucleosome targeting. <i>Nucleic Acids Research</i> , 2015 , 43, 2033	3- 4 4.1	36
49	PROTAC-mediated degradation of class I histone deacetylase enzymes in corepressor complexes. <i>Chemical Communications</i> , 2020 , 56, 4476-4479	5.8	36
48	Mzt1/Tam4, a fission yeast MOZART1 homologue, is an essential component of the Eubulin complex and directly interacts with GCP3(Alp6). <i>Molecular Biology of the Cell</i> , 2013 , 24, 3337-49	3.5	36
47	Recombinant protein expression for structural biology in HEK 293F suspension cells: a novel and accessible approach. <i>Journal of Visualized Experiments</i> , 2014 , e51897	1.6	33
46	Mechanism of Crosstalk between the LSD1 Demethylase and HDAC1 Deacetylase in the CoREST Complex. <i>Cell Reports</i> , 2020 , 30, 2699-2711.e8	10.6	31
45	BIM-mediated membrane insertion of the BAK pore domain is an essential requirement for apoptosis. <i>Cell Reports</i> , 2013 , 5, 409-20	10.6	31
44	Insights into the Recruitment of Class IIa Histone Deacetylases (HDACs) to the SMRT/NCoR Transcriptional Repression Complex. <i>Journal of Biological Chemistry</i> , 2015 , 290, 18237-18244	5.4	30
43	Lysine-14 acetylation of histone H3 in chromatin confers resistance to the deacetylase and demethylase activities of an epigenetic silencing complex. <i>ELife</i> , 2018 , 7,	8.9	29
42	Transient expression in HEK 293 cells: an alternative to E. coli for the production of secreted and intracellular mammalian proteins. <i>Methods in Molecular Biology</i> , 2015 , 1258, 209-22	1.4	28
41	Towards an understanding of the structure and function of MTA1. <i>Cancer and Metastasis Reviews</i> , 2014 , 33, 857-67	9.6	28
40	A novel albumin gene mutation (R222I) in familial dysalbuminemic hyperthyroxinemia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E1381-6	5.6	26
39	Contrasting Phenotypes in Resistance to Thyroid Hormone Alpha Correlate with Divergent Properties of Thyroid Hormone Receptor Il Mutant Proteins. <i>Thyroid</i> , 2017 , 27, 973-982	6.2	25
38	The steroid/nuclear receptors: from three-dimensional structure to complex function. <i>Vitamins and Hormones</i> , 1994 , 49, 1-47	2.5	25
37	Molecular determinants of the balance between co-repressor and co-activator recruitment to the retinoic acid receptor. <i>Journal of Biological Chemistry</i> , 2003 , 278, 43797-806	5.4	24

36	Maternal isodisomy for chromosome 9 causing homozygosity for a novel FOXE1 mutation in syndromic congenital hypothyroidism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 4031-	· 6 ^{5.6}	22
35	A variant NuRD complex containing PWWP2A/B excludes MBD2/3 to regulate transcription at active genes. <i>Nature Communications</i> , 2018 , 9, 3798	17.4	22
34	A Pharmacogenetic Approach to the Treatment of Patients With Mutations. <i>Diabetes</i> , 2018 , 67, 1086-10	092 9	21
33	Nuclear receptors: the evolution of diversity. <i>Science Signaling</i> , 2004 , 2004, pe4	8.8	20
32	Transcriptional control: how nuclear receptors get turned on. Current Biology, 1996, 6, 372-4	6.3	20
31	Two de novo mutations in the AR gene cause the complete androgen insensitivity syndrome in a pair of monozygotic twins. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002 , 87, 1057-61	5.6	19
30	Heme binding to human CLOCK affects interactions with the E-box. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 19911-19916	11.5	16
29	Comparison of the molecular consequences of different mutations at residue 754 and 690 of the androgen receptor (AR) and androgen insensitivity syndrome (AIS) phenotype. <i>Clinical Endocrinology</i> , 2009 , 71, 253-60	3.4	15
28	What is evolution playing at?. Current Biology, 1993, 3, 628-30	6.3	15
27	Diverse nucleosome Site-Selectivity among histone deacetylase complexes. <i>ELife</i> , 2020 , 9,	8.9	15
27 26	Diverse nucleosome Site-Selectivity among histone deacetylase complexes. <i>ELife</i> , 2020 , 9, Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77		15
26	Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77	3.8	14
26	Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77 Genetic disorders of nuclear receptors. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1181-1192 The MiDAC histone deacetylase complex is essential for embryonic development and has a unique	3.8	14
26 25 24	Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77 Genetic disorders of nuclear receptors. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1181-1192 The MiDAC histone deacetylase complex is essential for embryonic development and has a unique multivalent structure. <i>Nature Communications</i> , 2020 , 11, 3252 The ansamycin antibiotic, rifamycin SV, inhibits BCL6 transcriptional repression and forms a	3.8 15.9 17.4	14 14 13
26 25 24 23	Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77 Genetic disorders of nuclear receptors. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1181-1192 The MiDAC histone deacetylase complex is essential for embryonic development and has a unique multivalent structure. <i>Nature Communications</i> , 2020 , 11, 3252 The ansamycin antibiotic, rifamycin SV, inhibits BCL6 transcriptional repression and forms a complex with the BCL6-BTB/POZ domain. <i>PLoS ONE</i> , 2014 , 9, e90889 A novel CYP11B2 gene mutation in an Asian family with aldosterone synthase deficiency. <i>Journal of</i>	3.8 15.9 17.4 3.7	14 14 13
26 25 24 23 22	Histone deacetylase 3 indirectly modulates tubulin acetylation. <i>Biochemical Journal</i> , 2015 , 472, 367-77 Genetic disorders of nuclear receptors. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1181-1192 The MiDAC histone deacetylase complex is essential for embryonic development and has a unique multivalent structure. <i>Nature Communications</i> , 2020 , 11, 3252 The ansamycin antibiotic, rifamycin SV, inhibits BCL6 transcriptional repression and forms a complex with the BCL6-BTB/POZ domain. <i>PLoS ONE</i> , 2014 , 9, e90889 A novel CYP11B2 gene mutation in an Asian family with aldosterone synthase deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009 , 94, 914-9	3.8 15.9 17.4 3.7 5.6	14 14 13 13

(2015-1993)

18	The cocrystal structures of two zinc-stabilized DNA-binding domains illustrate different ways of achieving sequence-specific DNA recognition. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 1993 , 58, 141-7	3.9	8
17	HDAC3 deacetylates the DNA mismatch repair factor MutSIto stimulate triplet repeat expansions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 23597-23605	5 11.5	8
16	Cryo-EM structural analysis of FADD:Caspase-8 complexes defines the catalytic dimer architecture for co-ordinated control of cell fate. <i>Nature Communications</i> , 2021 , 12, 819	17.4	8
15	Transcriptional repression by nuclear receptors: mechanisms and role in disease. <i>Biochemical Society Transactions</i> , 2000 , 28, 390	5.1	7
14	Selective Aster inhibitors distinguish vesicular and nonvesicular sterol transport mechanisms. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021 , 118,	11.5	6
13	Expression and Purification of Protein Complexes Suitable for Structural Studies Using Mammalian HEK 293F Cells. <i>Current Protocols in Protein Science</i> , 2017 , 90, 5.28.1-5.28.16	3.1	5
12	The topology of chromatin-binding domains in the NuRD deacetylase complex. <i>Nucleic Acids Research</i> , 2020 , 48, 12972-12982	20.1	4
11	Backbone resonance assignment of the BCL6-BTB/POZ domain. <i>Biomolecular NMR Assignments</i> , 2018 , 12, 47-50	0.7	3
10	Radical fringe positions the apical ectodermal ridge at the dorsoventral boundary of the vertebrate limb. <i>Nature</i> , 1997 , 388, 906-906	50.4	3
9	Signal transduction: fast lane to transcriptional activation. <i>Current Biology</i> , 1998 , 8, R765-7	6.3	3
8	Histone H2B Deacylation Selectivity: Exploring Chromatin's Dark Matter with an Engineered Sortase <i>Journal of the American Chemical Society</i> , 2022 ,	16.4	3
7	Distinctly differentor really much the same?. Current Biology, 1992, 2, 237-9	6.3	2
6	Hyperthyroxinemia and Hypercortisolemia due to Familial Dysalbuminemia. <i>Thyroid</i> , 2020 , 30, 1681-168	346.2	1
5	Structure-guided approach to relieving transcriptional repression inResistance to Thyroid Hormone [Implectual of the control o	4.8	O
4	Nuclear receptor: co-repressor interactions. <i>Biochemical Society Transactions</i> , 2000 , 28, A63-A63	5.1	
3	DNA between the sheets. <i>Current Biology</i> , 1992 , 2, 661-3	6.3	
2	Analysis of CoREST Complex-Chromatin Interactions with Chemical Tools. FASEB Journal, 2018, 32, 524.	7 0.9	
1	Assembly and Regulation of Nuclear Receptor Corepressor Complexes 2015 , 155-175		