

Richard J Auchus

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/868633/richard-j-auchus-publications-by-year.pdf>

Version: 2024-04-24

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

200
papers

12,480
citations

52
h-index

108
g-index

228
ext. papers

14,903
ext. citations

6.8
avg, IF

6.9
L-index

#	Paper	IF	Citations
200	Randomized trial of osilodrostat for the treatment of Cushing's disease.. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022 ,	5.6	3
199	Recalibrating Interpretations of Aldosterone Assays Across the Physiologic Range: Immunoassay and Liquid Chromatography-Tandem Mass Spectrometry Measurements Under Multiple Controlled Conditions.. <i>Journal of the Endocrine Society</i> , 2022 , 6, bvac049	0.4	0
198	Glucocorticoid Withdrawal Syndrome following treatment of endogenous Cushing Syndrome.. <i>Pituitary</i> , 2022 , 1	4.3	2
197	The uncommon forms of congenital adrenal hyperplasia. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2022 , 29, 263-270	4	0
196	Approach to the Patient with Primary Aldosteronism: Utility and Limitations of Adrenal Vein Sampling. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, 1195-1208	5.6	6
195	11-Oxygenated Androgens Useful in the Setting of Discrepant Conventional Biomarkers in 21-Hydroxylase Deficiency. <i>Journal of the Endocrine Society</i> , 2021 , 5, bvaa192	0.4	7
194	Clinical advances in the pharmacotherapy of congenital adrenal hyperplasia. <i>European Journal of Endocrinology</i> , 2021 , 186, R1-R14	6.5	0
193	24-Hour Profiles of 11-Oxygenated C Steroids and Steroid Sulfates during Oral and Continuous Subcutaneous Glucocorticoids in 21-Hydroxylase Deficiency. <i>Frontiers in Endocrinology</i> , 2021 , 12, 751191	5.7	1
192	Differences of adrenal-derived androgens in 5 β -reductase deficiency versus androgen insensitivity syndrome. <i>Clinical and Translational Science</i> , 2021 ,	4.9	1
191	Consensus on diagnosis and management of Cushing's disease: a guideline update. <i>Lancet Diabetes and Endocrinology</i> , 2021 , 9, 847-875	18.1	48
190	Crinicerfont Lowers Elevated Hormone Markers in Adults with 21-Hydroxylase Deficiency Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 ,	5.6	5
189	Congenital adrenal hyperplasia - current insights in pathophysiology, diagnostics and management. <i>Endocrine Reviews</i> , 2021 ,	27.2	28
188	Osteoblasts Generate Testosterone From DHEA and Activate Androgen Signaling in Prostate Cancer Cells. <i>Journal of Bone and Mineral Research</i> , 2021 , 36, 1566-1579	6.3	0
187	Clamping Cortisol and Testosterone Mitigates the Development of Insulin Resistance during Sleep Restriction in Men. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, e3436-e3448	5.6	2
186	Tildacerfont in Adults With Classic Congenital Adrenal Hyperplasia: Results from Two Phase 2 Studies. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, e4666-e4679	5.6	6
185	Levoketoconazole: a novel treatment for endogenous Cushing's syndrome. <i>Expert Review of Endocrinology and Metabolism</i> , 2021 , 16, 159-174	4.1	3
184	Salt-Losing 21-Hydroxylase Deficiency Caused by Double Homozygosity for Two "Mild" Mutations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, e680-e686	5.6	2

183	Association of Maternal-Neonatal Steroids With Early Pregnancy Endocrine Disrupting Chemicals and Pregnancy Outcomes. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 , 106, 665-687	5.6	4
182	Production of 11-oxygenated androgens by testicular adrenal rest tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021 ,	5.6	1
181	Approach to the Patient with an Incidental Adrenal Mass. <i>Medical Clinics of North America</i> , 2021 , 105, 1047-1063	7	0
180	Intratumoral steroid profiling of adrenal cortisol-producing adenomas by liquid chromatography-mass spectrometry. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2021 , 212, 105924	5.1	0
179	The Dark Side of hormone prescription. <i>Endocrine Connections</i> , 2021 , 10, C1-C3	3.5	0
178	Circadian rhythms of 11-oxygenated C19 steroids and β -steroid sulfates in healthy men. <i>European Journal of Endocrinology</i> , 2021 , 185, K1-K6	6.5	5
177	MON-183 Adrenal Androgen Control and Steroidal Side Effects in Adolescents and Adults with Congenital Adrenal Hyperplasia Treated with Glucocorticoids. <i>Journal of the Endocrine Society</i> , 2020 , 4,	0.4	78
176	Sex Differences in 11-Oxygenated Androgen Patterns Across Adulthood. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	24
175	Endocrine causes of hypertension in pregnancy. <i>Gland Surgery</i> , 2020 , 9, 69-79	2.2	12
174	11-Oxygenated androgens in health and disease. <i>Nature Reviews Endocrinology</i> , 2020 , 16, 284-296	15.2	37
173	A Phase 2, Multicenter Study of Nevanimibe for the Treatment of Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	10
172	Abiraterone acetate treatment lowers 11-oxygenated androgens. <i>European Journal of Endocrinology</i> , 2020 , 182, 413-421	6.5	18
171	Androgen excess and diagnostic steroid biomarkers for nonclassic 21-hydroxylase deficiency without cosyntropin stimulation. <i>European Journal of Endocrinology</i> , 2020 , 183, 63-71	6.5	8
170	The role of adrenal derived androgens in castration resistant prostate cancer. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2020 , 197, 105506	5.1	17
169	Comprehensive Analysis of Steroid Biomarkers for Guiding Primary Aldosteronism Subtyping. <i>Hypertension</i> , 2020 , 75, 183-192	8.5	21
168	Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. <i>New England Journal of Medicine</i> , 2020 , 383, 1248-1261	59.2	59
167	Reply to Fl�k et al.: Alternative androgen pathway biosynthesis drives fetal female virilization in P450 oxidoreductase deficiency. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 14634-14635	11.5	2
166	A virtual teaching clinic for virtual care during the COVID-19 pandemic. <i>Clinical Diabetes and Endocrinology</i> , 2020 , 6, 25	4.7	8

165	Expression in Escherichia Coli, Purification, and Functional Reconstitution of Human Steroid 5 β Reductases. <i>Endocrinology</i> , 2020 , 161,	4.8	3
164	Efficacy and safety of osilodrostat in patients with Cushing's disease (LINC 3): a multicentre phase III study with a double-blind, randomised withdrawal phase. <i>Lancet Diabetes and Endocrinology</i> , 2020 , 8, 748-761	18.1	47
163	11-Oxygenated C19 Steroids Do Not Distinguish the Hyperandrogenic Phenotype of PCOS Daughters from Girls with Obesity. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020 , 105,	5.6	6
162	Sex Hormones and Prostate Cancer. <i>Annual Review of Medicine</i> , 2020 , 71, 33-45	17.4	23
161	The Unique Role of 11-Oxygenated C19 Steroids in Both Premature Adrenarche and Premature Pubarche. <i>Hormone Research in Paediatrics</i> , 2020 , 93, 460-469	3.3	2
160	Efficacy and safety of levoketoconazole in the treatment of endogenous Cushing's syndrome (SONICS): a phase 3, multicentre, open-label, single-arm trial. <i>Lancet Diabetes and Endocrinology</i> , 2019 , 7, 855-865	18.1	34
159	Steroid biomarkers in human adrenal disease. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019 , 190, 273-280	5.1	19
158	11-Oxygenated C19 Steroids Do Not Decline With Age in Women. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 2615-2622	5.6	42
157	The Clinical Impact of [Ga]-DOTATATE PET/CT for the Diagnosis and Management of Ectopic Adrenocorticotrophic Hormone - Secreting Tumours. <i>Clinical Endocrinology</i> , 2019 , 91, 288-294	3.4	19
156	Response to Letter to the Editor: "Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society Clinical Practice Guideline". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 1928	5.6	1
155	Circulating 11-oxygenated androgens across species. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019 , 190, 242-249	5.1	27
154	The "backdoor pathway" of androgen synthesis in human male sexual development. <i>PLoS Biology</i> , 2019 , 17, e3000198	9.7	25
153	Endocrine Disturbances Affecting Reproduction 2019 , 594-608.e5		
152	Analysis of novel heterozygous mutations in the CYP11B2 gene causing congenital aldosterone synthase deficiency and literature review. <i>Steroids</i> , 2019 , 150, 108448	2.8	5
151	Alternative pathway androgen biosynthesis and human fetal female virilization. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 22294-22299	11.5	22
150	Three Discrete Patterns of Primary Aldosteronism Lateralization in Response to Cosyntropin During Adrenal Vein Sampling. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 5867-5876	5.6	27
149	OR16-2 Osilodrostat Treatment in Cushing's Disease (CD): Results from a Phase III, Multicenter, Double-Blind, Randomized Withdrawal Study (LINC 3). <i>Journal of the Endocrine Society</i> , 2019 , 3,	0.4	3
148	SUN-LB064 A Phase 2, Dose-Escalation, Safety and Efficacy Study of Tildacerfont (SPR001) for the Treatment of Patients with Classic Congenital Adrenal Hyperplasia. <i>Journal of the Endocrine Society</i> , 2019 , 3,	0.4	1

147	Adrenal Vein Sampling Lateralization Despite Mineralocorticoid Receptor Antagonists Exposure in Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 487-492	5.6	18
146	Introduction to the 2018 Keith L. Parker Award Lecture, William E. Rainey, PhD. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019 , 188, 131-133	5.1	0
145	Predicted Benign and Synonymous Variants in Cause Primary Adrenal Insufficiency Through Missplicing. <i>Journal of the Endocrine Society</i> , 2019 , 3, 201-221	0.4	13
144	Structural and Functional Biology of Aldo-Keto Reductase Steroid-Transforming Enzymes. <i>Endocrine Reviews</i> , 2019 , 40, 447-475	27.2	32
143	Germ cell neoplasia in situ complicating 17 β -hydroxysteroid dehydrogenase type 3 deficiency. <i>Molecular and Cellular Endocrinology</i> , 2019 , 489, 3-8	4.4	2
142	Androgen Biosynthesis and Gene Defects 2019 , 713-720		
141	Adrenocorticotropin Acutely Regulates Pregnenolone Sulfate Production by the Human Adrenal In Vivo and In Vitro. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018 , 103, 320-327	5.6	17
140	Catalytic modulation of human cytochromes P450 17A1 and P450 11B2 by phospholipid. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2018 , 181, 63-72	5.1	5
139	Strategies that athletes use to avoid detection of androgenic-anabolic steroid doping and sanctions. <i>Molecular and Cellular Endocrinology</i> , 2018 , 464, 28-33	4.4	15
138	Mifepristone in the treatment of the ectopic adrenocorticotrophic hormone syndrome. <i>Clinical Endocrinology</i> , 2018 , 89, 570-576	3.4	6
137	The Rise, Fall, and Resurrection of 11-Oxygenated Androgens in Human Physiology and Disease. <i>Hormone Research in Paediatrics</i> , 2018 , 89, 284-291	3.3	27
136	Primary Aldosteronism: Practical Approach to Diagnosis and Management. <i>Circulation</i> , 2018 , 138, 823-833	6.7	66
135	HSD3B1(1245A>C) variant regulates dueling abiraterone metabolite effects in prostate cancer. <i>Journal of Clinical Investigation</i> , 2018 , 128, 3333-3340	15.9	32
134	Exhaled nitric oxide and vascular endothelial growth factor as predictors of cold symptoms after stress. <i>Biological Psychology</i> , 2018 , 132, 116-124	3.2	7
133	11-Ketotestosterone Is the Dominant Circulating Bioactive Androgen During Normal and Premature Adrenarche. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018 , 103, 4589-4598	5.6	52
132	Human Urinary mRNA as a Biomarker of Cardiovascular Disease. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002213	5.2	13
131	Phase 2 Randomized, Placebo-Controlled Clinical Trial of Recombinant Human Growth Hormone (rhGH) During Rehabilitation From Traumatic Brain Injury. <i>Frontiers in Endocrinology</i> , 2018 , 9, 520	5.7	8
130	Congenital Adrenal Hyperplasia Due to Steroid 21-Hydroxylase Deficiency: An Endocrine Society Clinical Practice Guideline. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018 , 103, 4043-4088	5.6	371

129	Adrenocortical carcinoma in a 17th-century girl. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017 , 165, 109-113	5.1	3
128	Steroid 17-hydroxylase and 17,20-lyase deficiencies, genetic and pharmacologic. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017 , 165, 71-78	5.1	99
127	Development and validation of a novel LC-MS/MS method for simultaneous determination of abiraterone and its seven steroidal metabolites in human serum: Innovation in separation of diastereoisomers without use of a chiral column. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017 , 172, 231-239	5.1	20
126	Identification of Unique Antigenic Determinants in the Amino Terminus of IA-2 (ICA512) in Childhood and Adult Autoimmune Diabetes: New Biomarker Development. <i>Diabetes Care</i> , 2017 , 40, 561-568	14.6	22
125	11-Oxygenated Androgens Are Biomarkers of Adrenal Volume and Testicular Adrenal Rest Tumors in 21-Hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 2701-2710	5.6	51
124	Steroidogenic Metabolism of Galeterone Reveals a Diversity of Biochemical Activities. <i>Cell Chemical Biology</i> , 2017 , 24, 825-832.e6	8.2	31
123	Outcomes after adrenalectomy for unilateral primary aldosteronism: an international consensus on outcome measures and analysis of remission rates in an international cohort. <i>Lancet Diabetes and Endocrinology</i> , 2017 , 5, 689-699	18.1	355
122	Molecular Recognition in Mitochondrial Cytochromes P450 That Catalyze the Terminal Steps of Corticosteroid Biosynthesis. <i>Biochemistry</i> , 2017 , 56, 2282-2293	3.2	16
121	Clinical significance of 11-oxygenated androgens. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2017 , 24, 252-259	4	38
120	A high rate of novel CYP11B1 mutations in Saudi Arabia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017 , 174, 217-224	5.1	3
119	Metabolic, Reproductive, and Neurologic Abnormalities in Agpat1-Null Mice. <i>Endocrinology</i> , 2017 , 158, 3954-3973	4.8	9
118	Discordance between imaging and immunohistochemistry in unilateral primary aldosteronism. <i>Clinical Endocrinology</i> , 2017 , 87, 665-672	3.4	41
117	An International Consortium Update: Pathophysiology, Diagnosis, and Treatment of Polycystic Ovarian Syndrome in Adolescence. <i>Hormone Research in Paediatrics</i> , 2017 , 88, 371-395	3.3	166
116	Phase II trial of pazopanib in advanced/progressive malignant pheochromocytoma and paraganglioma. <i>Endocrine</i> , 2017 , 57, 220-225	4	33
115	Obesity-Induced Infertility in Male Mice Is Associated With Disruption of Crisp4 Expression and Sperm Fertilization Capacity. <i>Endocrinology</i> , 2017 , 158, 2930-2943	4.8	19
114	Electrochemistry of cytochrome P450 17 β hydroxylase/17,20-lyase (P450c17). <i>Molecular and Cellular Endocrinology</i> , 2017 , 441, 62-67	4.4	3
113	Mild Adrenal Cortisol Excess 2017 , 181-197		1
112	Age-dependent Increases in Adrenal Cytochrome b5 and Serum 5-Androstenediol-3-sulfate. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 4585-4593	5.6	26

111	Cytochrome b5 Activates the 17,20-Lyase Activity of Human Cytochrome P450 17A1 by Increasing the Coupling of NADPH Consumption to Androgen Production. <i>Biochemistry</i> , 2016 , 55, 4356-65	3.2	32
110	Opposing Effects of Cyclooxygenase-2 (COX-2) on Estrogen Receptor α Response to 5 α -Reductase Inhibition in Prostate Epithelial Cells. <i>Journal of Biological Chemistry</i> , 2016 , 291, 14747-60	5.4	7
109	Influence of race/ethnicity on cardiovascular risk factors in polycystic ovary syndrome, the Dallas Heart Study. <i>Clinical Endocrinology</i> , 2016 , 85, 92-9	3.4	24
108	Mechanism of 17 β -Lyase and New Hydroxylation Reactions of Human Cytochrome P450 17A1: 18O LABELING AND OXYGEN SURROGATE EVIDENCE FOR A ROLE OF A PERFERRYL OXYGEN. <i>Journal of Biological Chemistry</i> , 2016 , 291, 17143-64	5.4	41
107	Impaired 17,20-Lyase Activity in Male Mice Lacking Cytochrome b5 in Leydig Cells. <i>Molecular Endocrinology</i> , 2016 , 30, 469-78		11
106	Cortisol response to acute stress in asthma: Moderation by depressive mood. <i>Physiology and Behavior</i> , 2016 , 159, 20-6	3.5	8
105	Global Disorders of Sex Development Update since 2006: Perceptions, Approach and Care. <i>Hormone Research in Paediatrics</i> , 2016 , 85, 158-80	3.3	379
104	The Metabolism, Analysis, and Targeting of Steroid Hormones in Breast and Prostate Cancer. <i>Hormones and Cancer</i> , 2016 , 7, 149-64	5	48
103	Single-Dose Study of a Corticotropin-Releasing Factor Receptor-1 Antagonist in Women With 21-Hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 1174-80	5.6	32
102	Adrenal-derived 11-oxygenated 19-carbon steroids are the dominant androgens in classic 21-hydroxylase deficiency. <i>European Journal of Endocrinology</i> , 2016 , 174, 601-9	6.5	120
101	Rapid kinetic methods to dissect steroidogenic cytochrome P450 reaction mechanisms. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2016 , 161, 13-23	5.1	5
100	Mifepristone Improves Octreotide Efficacy in Resistant Ectopic Cushing's Syndrome. <i>Case Reports in Endocrinology</i> , 2016 , 2016, 8453801	1.2	4
99	Classics in Cardiovascular Endocrinology: Aldosterone Action Beyond Electrolytes. <i>Endocrinology</i> , 2016 , 157, 429-31	4.8	3
98	Redirecting abiraterone metabolism to fine-tune prostate cancer anti-androgen therapy. <i>Nature</i> , 2016 , 533, 547-51	50.4	116
97	Mutated KCNJ5 activates the acute and chronic regulatory steps in aldosterone production. <i>Journal of Molecular Endocrinology</i> , 2016 , 57, 1-11	4.5	26
96	Instability of the Human Cytochrome P450 Reductase A287P Variant Is the Major Contributor to Its Antley-Bixler Syndrome-like Phenotype. <i>Journal of Biological Chemistry</i> , 2016 , 291, 20487-502	5.4	16
95	Management considerations for the adult with congenital adrenal hyperplasia. <i>Molecular and Cellular Endocrinology</i> , 2015 , 408, 190-7	4.4	37
94	Genetic forms of adrenal insufficiency. <i>Endocrine Practice</i> , 2015 , 21, 395-9	3.2	10

93	Making water-soluble integral membrane proteins in vivo using an amphipathic protein fusion strategy. <i>Nature Communications</i> , 2015 , 6, 6826	17.4	17
92	The classic and nonclassic congenital adrenal hyperplasias. <i>Endocrine Practice</i> , 2015 , 21, 383-9	3.2	20
91	Profiles of 21-Carbon Steroids in 21-hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015 , 100, 2283-90	5.6	53
90	The diverse chemistry of cytochrome P450 17A1 (P450c17, CYP17A1). <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2015 , 151, 52-65	5.1	55
89	In reply. <i>Oncologist</i> , 2015 , 20, e14	5.7	
88	Conversion of abiraterone to D4A drives anti-tumour activity in prostate cancer. <i>Nature</i> , 2015 , 523, 347-51	5.4	190
87	The next 150 years of congenital adrenal hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2015 , 153, 63-71	5.1	43
86	Adrenal steroidogenesis and congenital adrenal hyperplasia. <i>Endocrinology and Metabolism Clinics of North America</i> , 2015 , 44, 275-96	5.5	69
85	Human Cytochrome P450 21A2, the Major Steroid 21-Hydroxylase: STRUCTURE OF THE ENZYME-PROGESTERONE SUBSTRATE COMPLEX AND RATE-LIMITING C-H BOND CLEAVAGE. <i>Journal of Biological Chemistry</i> , 2015 , 290, 13128-43	5.4	55
84	Bone Morphogenetic Protein-4 (BMP4): A Paracrine Regulator of Human Adrenal C19 Steroid Synthesis. <i>Endocrinology</i> , 2015 , 156, 2530-40	4.8	16
83	Aldosterone and Salt Loading Independently Exacerbate the Exercise Pressor Reflex in Rats. <i>Hypertension</i> , 2015 , 66, 627-33	8.5	10
82	Mechanistic Scrutiny Identifies a Kinetic Role for Cytochrome b5 Regulation of Human Cytochrome P450c17 (CYP17A1, P450 17A1). <i>PLoS ONE</i> , 2015 , 10, e0141252	3.7	22
81	P450 Enzymes in Steroid Processing 2015 , 851-879		10
80	Serum Cortisol-to-Cortisone Ratio and Blood Pressure in Severe Obesity before and after Weight Loss. <i>CardioRenal Medicine</i> , 2015 , 6, 1-7	2.8	5
79	A-ring modified steroidal azoles retaining similar potent and slowly reversible CYP17A1 inhibition as abiraterone. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2014 , 143, 1-10	5.1	25
78	Two surfaces of cytochrome b5 with major and minor contributions to CYP3A4-catalyzed steroid and nifedipine oxygenation chemistries. <i>Archives of Biochemistry and Biophysics</i> , 2014 , 541, 53-60	4.1	13
77	Epoxidation activities of human cytochromes P450c17 and P450c21. <i>Biochemistry</i> , 2014 , 53, 7531-40	3.2	7
76	Catalytically relevant electrostatic interactions of cytochrome P450c17 (CYP17A1) and cytochrome b5. <i>Journal of Biological Chemistry</i> , 2014 , 289, 33838-49	5.4	25

75	Abiraterone acetate to lower androgens in women with classic 21-hydroxylase deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, 2763-70	5.6	45
74	Fertility in patients with genetic deficiencies of cytochrome P450c17 (CYP17A1): combined 17-hydroxylase/17,20-lyase deficiency and isolated 17,20-lyase deficiency. <i>Fertility and Sterility</i> , 2014 , 101, 317-22	4.8	55
73	Paradoxical Results after Inadvertent Use of Cosyntropin [Adrenocorticotropin Hormone (1-24)] Rather than Acthrel (Ovine Corticotropin Releasing Hormone) during Inferior Petrosal Sinus Sampling. <i>Endocrine Practice</i> , 2014 , 20, 646-9	3.2	3
72	An expert consensus statement on use of adrenal vein sampling for the subtyping of primary aldosteronism. <i>Hypertension</i> , 2014 , 63, 151-60	8.5	338
71	Use of prednisone with abiraterone acetate in metastatic castration-resistant prostate cancer. <i>Oncologist</i> , 2014 , 19, 1231-40	5.7	57
70	Adrenal androgens and androgen precursors-definition, synthesis, regulation and physiologic actions. <i>Comprehensive Physiology</i> , 2014 , 4, 1369-81	7.7	50
69	Androstenedione is the preferred androgen source in hormone refractory prostate cancer--letter. <i>Clinical Cancer Research</i> , 2014 , 20, 4971	12.9	9
68	Hypotension following patent ductus arteriosus ligation: the role of adrenal hormones. <i>Journal of Pediatrics</i> , 2014 , 164, 1449-55.e1	3.6	39
67	A gain-of-function mutation in DHT synthesis in castration-resistant prostate cancer. <i>Cell</i> , 2013 , 154, 1074-1084	56.2	210
66	Approach to the patient: the adult with congenital adrenal hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2013 , 98, 2645-55	5.6	84
65	Mass spectrometry theory and application to adrenal diseases. <i>Molecular and Cellular Endocrinology</i> , 2013 , 371, 201-7	4.4	25
64	The action of cytochrome b(5) on CYP2E1 and CYP2C19 activities requires anionic residues D58 and D65. <i>Biochemistry</i> , 2013 , 52, 210-20	3.2	25
63	Introduction to the 2012 Keith L. Parker memorial lecturer: Walter L. Miller, MD. <i>Molecular and Cellular Endocrinology</i> , 2013 , 371, 2-4	4.4	0
62	Secondary Adrenal Insufficiency 2013 , 32-46		
61	Gene mutations that promote adrenal aldosterone production, sodium retention, and hypertension. <i>The Application of Clinical Genetics</i> , 2013 , 7, 1-13	3.1	2
60	Mifepristone, a glucocorticoid receptor antagonist, produces clinical and metabolic benefits in patients with Cushing's syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, 2039-49	5.6	323
59	Minor activities and transition state properties of the human steroid hydroxylases cytochromes P450c17 and P450c21, from reactions observed with deuterium-labeled substrates. <i>Biochemistry</i> , 2012 , 51, 7064-77	3.2	29
58	Synthesis of halogenated pregnanes, mechanistic probes of steroid hydroxylases CYP17A1 and CYP21A2. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2012 , 128, 38-50	5.1	7

57	Adrenal disorders in pregnancy. <i>Nature Reviews Endocrinology</i> , 2012 , 8, 668-78	15.2	24
56	The Adrenal Vein Sampling International Study (AVIS) for identifying the major subtypes of primary aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, 1606-14	5.6	243
55	Clinical and biochemical consequences of CYP17A1 inhibition with abiraterone given with and without exogenous glucocorticoids in castrate men with advanced prostate cancer. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, 507-16	5.6	199
54	Effect of KCNJ5 mutations on gene expression in aldosterone-producing adenomas and adrenocortical cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, E1567-72	5.6	112
53	Defects in androgen biosynthesis causing 46,XY disorders of sexual development. <i>Seminars in Reproductive Medicine</i> , 2012 , 30, 417-26	1.4	41
52	Abiraterone inhibits 3 β hydroxysteroid dehydrogenase: a rationale for increasing drug exposure in castration-resistant prostate cancer. <i>Clinical Cancer Research</i> , 2012 , 18, 3571-9	12.9	79
51	Human steroid biosynthesis for the oncologist. <i>Journal of Investigative Medicine</i> , 2012 , 60, 495-503	2.9	45
50	The physiology and biochemistry of adrenarche. <i>Endocrine Development</i> , 2011 , 20, 20-27		30
49	The molecular biology, biochemistry, and physiology of human steroidogenesis and its disorders. <i>Endocrine Reviews</i> , 2011 , 32, 81-151	27.2	1314
48	Why human cytochrome P450c21 is a progesterone 21-hydroxylase. <i>Biochemistry</i> , 2011 , 50, 3968-74	3.2	19
47	The life and scientific contributions of Keith L. Parker, 1953-2008. <i>Molecular and Cellular Endocrinology</i> , 2011 , 336, 191-2	4.4	1
46	CYP17A1 intron mutation causing cryptic splicing in 17 β hydroxylase deficiency. <i>PLoS ONE</i> , 2011 , 6, e25492	3.7	15
45	Primary aldosteronism and a Texas two-step. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2011 , 12, 37-42	10.5	6
44	Steroid profiling by gas chromatography-mass spectrometry and high performance liquid chromatography-mass spectrometry for adrenal diseases. <i>Hormones and Cancer</i> , 2011 , 2, 324-32	5	49
43	Dihydrotestosterone synthesis bypasses testosterone to drive castration-resistant prostate cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 13728-33	11.5	255
42	Gene expression profiles in aldosterone-producing adenomas and adjacent adrenal glands. <i>European Journal of Endocrinology</i> , 2011 , 164, 613-9	6.5	47
41	Combined 17 β hydroxylase/17,20-lyase deficiency due to p.R96W mutation in the CYP17 gene in a Brazilian patient. <i>Arquivos Brasileiros De Endocrinologia E Metabologia</i> , 2010 , 54, 744-8		5
40	Guidelines for the Development of Comprehensive Care Centers for Congenital Adrenal Hyperplasia: Guidance from the CARES Foundation Initiative. <i>International Journal of Pediatric Endocrinology (Springer)</i> , 2010 , 2010, 275213	1.5	20

39	Management of the adult with congenital adrenal hyperplasia. <i>International Journal of Pediatric Endocrinology (Springer)</i> , 2010 , 2010, 614107	1.5	3
38	3beta-hydroxysteroid dehydrogenase is a possible pharmacological target in the treatment of castration-resistant prostate cancer. <i>Endocrinology</i> , 2010 , 151, 3514-20	4.8	60
37	Reversible sympathetic overactivity in hypertensive patients with primary aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 4756-61	5.6	55
36	46,XX DSD: the masculinised female. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2010 , 24, 219-42	6.5	22
35	Congenital adrenal hyperplasia in adults. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2010 , 17, 210-6	4	18
34	Non-traditional metabolic pathways of adrenal steroids. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2009 , 10, 27-32	10.5	13
33	Rapid cortisol assays improve the success rate of adrenal vein sampling for primary aldosteronism. <i>Annals of Surgery</i> , 2009 , 249, 318-21	7.8	75
32	Associations among androgens, estrogens, and natriuretic peptides in young women: observations from the Dallas Heart Study. <i>Journal of the American College of Cardiology</i> , 2007 , 49, 109-16	15.1	122
31	Phenotypic variability in 17beta-hydroxysteroid dehydrogenase-3 deficiency and diagnostic pitfalls. <i>Clinical Endocrinology</i> , 2007 , 67, 20-8	3.4	99
30	"Subclinical Cushing's syndrome" is not subclinical: improvement after adrenalectomy in 9 patients. <i>Surgery</i> , 2007 , 142, 900-5; discussion 905.e1	3.6	68
29	Primary aldosteronism. <i>Current Cardiology Reports</i> , 2007 , 9, 447-52	4.2	2
28	Measurement of 18-hydroxycorticosterone during adrenal vein sampling for primary aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007 , 92, 2648-51	5.6	28
27	Position statement: Utility, limitations, and pitfalls in measuring testosterone: an Endocrine Society position statement. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007 , 92, 405-13	5.6	879
26	Arginine 276 controls the directional preference of AKR1C9 (rat liver 3alpha-hydroxysteroid dehydrogenase) in human embryonic kidney 293 cells. <i>Endocrinology</i> , 2006 , 147, 1591-7	4.8	21
25	Human cytochrome b5 requires residues E48 and E49 to stimulate the 17,20-lyase activity of cytochrome P450c17. <i>Biochemistry</i> , 2006 , 45, 755-62	3.2	60
24	Miscellaneous endocrine causes of hypertension. <i>Current Cardiology Reports</i> , 2005 , 7, 418-24	4.2	1
23	Minireview: cellular redox state regulates hydroxysteroid dehydrogenase activity and intracellular hormone potency. <i>Endocrinology</i> , 2005 , 146, 2531-8	4.8	86
22	The rise in adrenal androgen biosynthesis: adrenarche. <i>Seminars in Reproductive Medicine</i> , 2004 , 22, 337-44	4.7	136

21	Overview of dehydroepiandrosterone biosynthesis. <i>Seminars in Reproductive Medicine</i> , 2004 , 22, 281-8	1.4	61
20	Two prevalent CYP17 mutations and genotype-phenotype correlations in 24 Brazilian patients with 17-hydroxylase deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004 , 89, 49-60	5.6	160
19	Adrenarche - physiology, biochemistry and human disease. <i>Clinical Endocrinology</i> , 2004 , 60, 288-96	3.4	241
18	Two intronic mutations cause 17-hydroxylase deficiency by disrupting splice acceptor sites: direct demonstration of aberrant splicing and absent enzyme activity by expression of the entire CYP17 gene in HEK-293 cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004 , 89, 43-8	5.6	30
17	The backdoor pathway to dihydrotestosterone. <i>Trends in Endocrinology and Metabolism</i> , 2004 , 15, 432-8	8.8	229
16	Human 17beta-hydroxysteroid dehydrogenases types 1, 2, and 3 catalyze bi-directional equilibrium reactions, rather than unidirectional metabolism, in HEK-293 cells. <i>Archives of Biochemistry and Biophysics</i> , 2004 , 429, 50-9	4.1	50
15	CYP17 mutation E305G causes isolated 17,20-lyase deficiency by selectively altering substrate binding. <i>Journal of Biological Chemistry</i> , 2003 , 278, 48563-9	5.4	96
14	5alpha-androstane-3alpha,17beta-diol is formed in tammar wallaby pouch young testes by a pathway involving 5alpha-pregnane-3alpha,17alpha-diol-20-one as a key intermediate. <i>Endocrinology</i> , 2003 , 144, 575-80	4.8	146
13	The 17, 20-lyase activity of cytochrome p450c17 from human fetal testis favors the delta5 steroidogenic pathway. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003 , 88, 3762-6	5.6	130
12	Aldo is back: recent advances and unresolved controversies in hyperaldosteronism. <i>Current Opinion in Nephrology and Hypertension</i> , 2003 , 12, 153-8	3.5	12
11	5alpha-reduced C21 steroids are substrates for human cytochrome P450c17. <i>Archives of Biochemistry and Biophysics</i> , 2003 , 418, 151-60	4.1	74
10	The enantiomer of progesterone (ent-progesterone) is a competitive inhibitor of human cytochromes P450c17 and P450c21. <i>Archives of Biochemistry and Biophysics</i> , 2003 , 409, 134-44	4.1	36
9	Towards a unifying mechanism for CYP17 mutations that cause isolated 17,20-lyase deficiency. <i>Endocrine Research</i> , 2002 , 28, 443-7	1.9	11
8	The genetics, pathophysiology, and management of human deficiencies of P450c17. <i>Endocrinology and Metabolism Clinics of North America</i> , 2001 , 30, 101-19, vii	5.5	182
7	Molecular modeling of the hamster adrenal P450C17. <i>Endocrine Research</i> , 2000 , 26, 723-8	1.9	1
6	Estrogen: consequences and implications of human mutations in synthesis and action. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1999 , 84, 4677-94	5.6	308
5	Molecular modeling of human P450c17 (17alpha-hydroxylase/17,20-lyase): insights into reaction mechanisms and effects of mutations. <i>Molecular Endocrinology</i> , 1999 , 13, 1169-82		167
4	P450c17 mutations R347H and R358Q selectively disrupt 17,20-lyase activity by disrupting interactions with P450 oxidoreductase and cytochrome b5. <i>Molecular Endocrinology</i> , 1999 , 13, 167-75		157

3	Cytochrome b5 augments the 17,20-lyase activity of human P450c17 without direct electron transfer. <i>Journal of Biological Chemistry</i> , 1998 , 273, 3158-65	5-4	407
2	The regulation of 17,20 lyase activity. <i>Steroids</i> , 1997 , 62, 133-42	2.8	192
1	The genetic and functional basis of isolated 17,20-lyase deficiency. <i>Nature Genetics</i> , 1997 , 17, 201-5	36.3	277