

Richard J Auchus

List of Publications by Year in descending order

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Version: 2024-02-01

210
papers

17,577
citations

19608

61
h-index

15218

126
g-index

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229
docs citations

229
times ranked

13077
citing authors

#	ARTICLE	IF	CITATIONS
1	Congenital Adrenal Hyperplasiaâ€”Current Insights in Pathophysiology, Diagnostics, and Management. <i>Endocrine Reviews</i> , 2022, 43, 91-159.	8.9	182
2	Salivary microbiome differences in prepubertal children with and without adrenal androgen excess. <i>Pediatric Research</i> , 2022, 91, 1797-1803.	1.1	3
3	Production of 11-Oxygenated Androgens by Testicular Adrenal Rest Tumors. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e272-e280.	1.8	12
4	Crinicerfont Lowers Elevated Hormone Markers in Adults With 21-Hydroxylase Deficiency Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 801-812.	1.8	19
5	Clinical advances in the pharmacotherapy of congenital adrenal hyperplasia. <i>European Journal of Endocrinology</i> , 2022, 186, R1-R14.	1.9	21
6	Cardiac decompensation and promiscuous prenylation of small GTPases in cardiomyocytes in response to local mevalonate pathway disruption^{â€‹}. <i>Journal of Pathology</i> , 2022, 256, 249-252.	2.1	4
7	Randomized Trial of Osilodrostat for the Treatment of Cushing Disease. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e2882-e2895.	1.8	31
8	Recalibrating Interpretations of Aldosterone Assays Across the Physiologic Range: Immunoassay and Liquid Chromatographyâ€”Tandem Mass Spectrometry Measurements Under Multiple Controlled Conditions. <i>Journal of the Endocrine Society</i> , 2022, 6, bvac049.	0.1	12
9	Glucocorticoid Withdrawal Syndrome following treatment of endogenous Cushing Syndrome. <i>Pituitary</i> , 2022, 25, 393-403.	1.6	20
10	The uncommon forms of congenital adrenal hyperplasia. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2022, 29, 263-270.	1.2	9
11	Excess 11-Oxygenated Androgens in Women With Severe Insulin Resistance Are Mediated by Adrenal Insulin Receptor Signaling. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 2626-2635.	1.8	7
12	Salt-Losing 21-Hydroxylase Deficiency Caused by Double Homozygosity for Two â€œMildâ€•Mutations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, e680-e686.	1.8	3
13	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.	1.8	20
14	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.	1.8	11
15	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.	1.8	21
16	Levoketoconazole: a novel treatment for endogenous Cushing's syndrome. <i>Expert Review of Endocrinology and Metabolism</i> , 2021, 16, 159-174.	1.2	8
17	Approach to the Patient with an Incidental Adrenal Mass. <i>Medical Clinics of North America</i> , 2021, 105, 1047-1063.	1.1	4
18	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.	1.2	3

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19	The Dark Side of hormone prescription. <i>Endocrine Connections</i> , 2021, 10, C1-C3.	0.8	2
20	Circadian rhythms of 11-oxygenated C19 steroids and $\hat{1}^5$ -steroid sulfates in healthy men. <i>European Journal of Endocrinology</i> , 2021, 185, K1-K6.	1.9	12
21	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.	1.8	23
22	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.	0.1	23
23	Consensus on diagnosis and management of Cushing's disease: a guideline update. <i>Lancet Diabetes and Endocrinology</i> , 2021, 9, 847-875.	5.5	315
24	Maternal 11-ketoandrostenedione rises through normal pregnancy and is the dominant 11-oxygenated androgen in cord blood. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, , .	1.8	4
25	24-Hour Profiles of 11-Oxygenated C19 Steroids and $\hat{1}^5$ -Steroid Sulfates during Oral and Continuous Subcutaneous Glucocorticoids in 21-Hydroxylase Deficiency. <i>Frontiers in Endocrinology</i> , 2021, 12, 751191.	1.5	10
26	Differences of adrenal-derived androgens in 5 $\hat{1}$ -reductase deficiency versus androgen insensitivity syndrome. <i>Clinical and Translational Science</i> , 2021, , .	1.5	4
27	Sex Hormones and Prostate Cancer. <i>Annual Review of Medicine</i> , 2020, 71, 33-45.	5.0	58
28	The role of adrenal derived androgens in castration resistant prostate cancer. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2020, 197, 105506.	1.2	37
29	Comprehensive Analysis of Steroid Biomarkers for Guiding Primary Aldosteronism Subtyping. <i>Hypertension</i> , 2020, 75, 183-192.	1.3	42
30	Congenital Adrenal Hyperplasia Due to 21-Hydroxylase Deficiency. <i>New England Journal of Medicine</i> , 2020, 383, 1248-1261.	13.9	155
31	Reply to Fl \hat{A} ck 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.	3.3	4
32	A virtual teaching clinic for virtual care during the COVID-19 pandemic. <i>Clinical Diabetes and Endocrinology</i> , 2020, 6, 25.	1.3	14
33	An Innovative Approach to Noninvasive Dynamic Adrenal Testing. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e3808-e3809.	1.8	1
34	Expression in <i>Escherichia Coli</i> , Purification, and Functional Reconstitution of Human Steroid 5 $\hat{1}$ -Reductases. <i>Endocrinology</i> , 2020, 161, .	1.4	4
35	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.	5.5	114
36	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, e3903-e3909.	1.8	15

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37	OR25-03 The Effects of Crinecerfont (NBI-74788), a Novel CRF1 Receptor Antagonist, on Adrenal Androgens and Precursors in Patients with Classic Congenital Adrenal Hyperplasia: Results from A Multiple-Dose Phase 2 Study. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.1	1
38	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, .	0.1	0
39	Sex Differences in 11-Oxygenated Androgen Patterns Across Adulthood. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e2921-e2929.	1.8	48
40	Endocrine causes of hypertension in pregnancy. <i>Gland Surgery</i> , 2020, 9, 69-79.	0.5	21
41	11-Oxygenated androgens in health and disease. <i>Nature Reviews Endocrinology</i> , 2020, 16, 284-296.	4.3	99
42	A Phase 2, Multicenter Study of Nevanimibe for the Treatment of Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 2771-2778.	1.8	19
43	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.	0.8	12
44	Osteoblasts Generate Testosterone From DHEA and Activate Androgen Signaling in Prostate Cancer Cells. <i>Journal of Bone and Mineral Research</i> , 2020, 36, 1566-1579.	3.1	3
45	Abiraterone acetate treatment lowers 11-oxygenated androgens. <i>European Journal of Endocrinology</i> , 2020, 182, 413-421.	1.9	43
46	Androgen excess and diagnostic steroid biomarkers for nonclassic 21-hydroxylase deficiency without cosyntropin stimulation. <i>European Journal of Endocrinology</i> , 2020, 183, 63-71.	1.9	24
47	Endocrine Disturbances Affecting Reproduction. , 2019, , 594-608.e5.		1
48	Analysis of novel heterozygous mutations in the CYP11B2 gene causing congenital aldosterone synthase deficiency and literature review. <i>Steroids</i> , 2019, 150, 108448.	0.8	16
49	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.	3.3	50
50	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.	1.8	51
51	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> , the, 2019, 7, 855-865.	5.5	60
52	Steroid biomarkers in human adrenal disease. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 190, 273-280.	1.2	27
53	11-Oxygenated C19 Steroids Do Not Decline With Age in Women. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 2615-2622.	1.8	74
54	The Clinical Impact of [⁶⁸ Ga]â€•DOTATATE PET/CT for the Diagnosis and Management of Ectopic Adrenocorticotropic Hormone â€• Secreting Tumours. <i>Clinical Endocrinology</i> , 2019, 91, 288-294.	1.2	31

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55	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, 1928-1928.	1.8	1
56	Circulating 11-oxygenated androgens across species. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 190, 242-249.	1.2	46
57	The "backdoor pathway" of androgen synthesis in human male sexual development. <i>PLoS Biology</i> , 2019, 17, e3000198.	2.6	54
58	Adrenal Vein Sampling Lateralization Despite Mineralocorticoid Receptor Antagonists Exposure in Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 487-492.	1.8	40
59	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.	1.2	1
60	Predicted Benign and Synonymous Variants in CYP11A1 Cause Primary Adrenal Insufficiency Through Missplicing. <i>Journal of the Endocrine Society</i> , 2019, 3, 201-221.	0.1	27
61	Structural and Functional Biology of Aldo-Keto Reductase Steroid-Transforming Enzymes. <i>Endocrine Reviews</i> , 2019, 40, 447-475.	8.9	73
62	Germ cell neoplasia in situ complicating 17 β -hydroxysteroid dehydrogenase type 3 deficiency. <i>Molecular and Cellular Endocrinology</i> , 2019, 489, 3-8.	1.6	5
63	Androgen Biosynthesis and Gene Defects. , 2019, , 713-720.		0
64	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, .	0.1	3
65	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, .	0.1	1
66	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.	1.8	24
67	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.	1.2	8
68	Strategies that athletes use to avoid detection of androgenic-anabolic steroid doping and sanctions. <i>Molecular and Cellular Endocrinology</i> , 2018, 464, 28-33.	1.6	21
69	Exhaled nitric oxide and vascular endothelial growth factor as predictors of cold symptoms after stress. <i>Biological Psychology</i> , 2018, 132, 116-124.	1.1	11
70	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.	1.8	73
71	Human Urinary mRNA as a Biomarker of Cardiovascular Disease. <i>Circulation Genomic and Precision Medicine</i> , 2018, 11, e002213.	1.6	25
72	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.	1.5	11

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73	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.	1.8	667
74	Mifepristone in the treatment of the ectopic adrenocorticotrophic hormone syndrome. <i>Clinical Endocrinology</i> , 2018, 89, 570-576.	1.2	12
75	The Rise, Fall, and Resurrection of 11-Oxygenated Androgens in Human Physiology and Disease. <i>Hormone Research in Paediatrics</i> , 2018, 89, 284-291.	0.8	40
76	Primary Aldosteronism. <i>Circulation</i> , 2018, 138, 823-835.	1.6	113
77	HSD3B1(1245A>C) variant regulates dueling abiraterone metabolite effects in prostate cancer. <i>Journal of Clinical Investigation</i> , 2018, 128, 3333-3340.	3.9	43
78	Adrenocortical carcinoma in a 17th-century girl. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 165, 109-113.	1.2	4
79	Steroid 17-hydroxylase and 17,20-lyase deficiencies, genetic and pharmacologic. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 165, 71-78.	1.2	159
80	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.	1.2	26
81	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.	4.3	30
82	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.	1.8	84
83	Steroidogenic Metabolism of Galeterone Reveals a Diversity of Biochemical Activities. <i>Cell Chemical Biology</i> , 2017, 24, 825-832.e6.	2.5	41
84	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.	5.5	595
85	Molecular Recognition in Mitochondrial Cytochromes P450 That Catalyze the Terminal Steps of Corticosteroid Biosynthesis. <i>Biochemistry</i> , 2017, 56, 2282-2293.	1.2	20
86	Clinical significance of 11-oxygenated androgens. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2017, 24, 252-259.	1.2	60
87	A high rate of novel CYP11B1 mutations in Saudi Arabia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 174, 217-224.	1.2	7
88	The Current Status and Evolution of Hormone Testing in the Digital Age. <i>Endocrinology and Metabolism Clinics of North America</i> , 2017, 46, xvii-xix.	1.2	0
89	Metabolic, Reproductive, and Neurologic Abnormalities in Agpat1-Null Mice. <i>Endocrinology</i> , 2017, 158, 3954-3973.	1.4	20
90	Discordance between imaging and immunohistochemistry in unilateral primary aldosteronism. <i>Clinical Endocrinology</i> , 2017, 87, 665-672.	1.2	68

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91	The Public Health Consequences of Performance-Enhancing Substances. JAMA - Journal of the American Medical Association, 2017, 318, 1983.	3.8	4
92	An International Consortium Update: Pathophysiology, Diagnosis, and Treatment of Polycystic Ovarian Syndrome in Adolescence. Hormone Research in Paediatrics, 2017, 88, 371-395.	0.8	282
93	Phase II trial of pazopanib in advanced/progressive malignant pheochromocytoma and paraganglioma. Endocrine, 2017, 57, 220-225.	1.1	40
94	Obesity-Induced Infertility in Male Mice Is Associated With Disruption of Crisp4 Expression and Sperm Fertilization Capacity. Endocrinology, 2017, 158, 2930-2943.	1.4	26
95	Electrochemistry of cytochrome P450 17 α -hydroxylase/17,20-lyase (P450c17). Molecular and Cellular Endocrinology, 2017, 441, 62-67.	1.6	4
96	Mild Adrenal Cortisol Excess. , 2017, , 181-197.		2
97	Mifepristone Improves Octreotide Efficacy in Resistant Ectopic Cushing's Syndrome. Case Reports in Endocrinology, 2016, 2016, 1-5.	0.2	4
98	Serum Cortisol-to-Cortisone Ratio and Blood Pressure in Severe Obesity before and after Weight Loss. CardioRenal Medicine, 2016, 6, 1-7.	0.7	7
99	Classics in Cardiovascular Endocrinology: Aldosterone Action Beyond Electrolytes. Endocrinology, 2016, 157, 429-431.	1.4	4
100	Redirecting abiraterone metabolism to fine-tune prostate cancer anti-androgen therapy. Nature, 2016, 533, 547-551.	13.7	138
101	Mutated KCNJ5 activates the acute and chronic regulatory steps in aldosterone production. Journal of Molecular Endocrinology, 2016, 57, 1-11.	1.1	35
102	Instability of the Human Cytochrome P450 Reductase A287P Variant Is the Major Contributor to Its Antley-Bixler Syndrome-like Phenotype. Journal of Biological Chemistry, 2016, 291, 20487-20502.	1.6	26
103	Age-dependent Increases in Adrenal Cytochrome b5 and Serum 5-Androstenediol-3-sulfate. Journal of Clinical Endocrinology and Metabolism, 2016, 101, 4585-4593.	1.8	34
104	Cytochrome b5 Activates the 17,20-Lyase Activity of Human Cytochrome P450 17A1 by Increasing the Coupling of NADPH Consumption to Androgen Production. Biochemistry, 2016, 55, 4356-4365.	1.2	37
105	Opposing Effects of Cyclooxygenase-2 (COX-2) on Estrogen Receptor β (ER β) Response to 5 α -Reductase Inhibition in Prostate Epithelial Cells. Journal of Biological Chemistry, 2016, 291, 14747-14760.	1.6	8
106	Influence of race/ethnicity on cardiovascular risk factors in polycystic ovary syndrome, the Dallas Heart Study. Clinical Endocrinology, 2016, 85, 92-99.	1.2	31
107	Mechanism of 17 α ,20-Lyase and New Hydroxylation Reactions of Human Cytochrome P450 17A1. Journal of Biological Chemistry, 2016, 291, 17143-17164.	1.6	54
108	Impaired 17,20-Lyase Activity in Male Mice Lacking Cytochrome b5 in Leydig Cells. Molecular Endocrinology, 2016, 30, 469-478.	3.7	13

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109	Cortisol response to acute stress in asthma: Moderation by depressive mood. <i>Physiology and Behavior</i> , 2016, 159, 20-26.	1.0	9
110	Global Disorders of Sex Development Update since 2006: Perceptions, Approach and Care. <i>Hormone Research in Paediatrics</i> , 2016, 85, 158-180.	0.8	852
111	The Metabolism, Analysis, and Targeting of Steroid Hormones in Breast and Prostate Cancer. <i>Hormones and Cancer</i> , 2016, 7, 149-164.	4.9	62
112	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, 1174-1180.	1.8	43
113	Adrenal-derived 11-oxygenated 19-carbon steroids are the dominant androgens in classic 21-hydroxylase deficiency. <i>European Journal of Endocrinology</i> , 2016, 174, 601-609.	1.9	168
114	Rapid kinetic methods to dissect steroidogenic cytochrome P450 reaction mechanisms. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2016, 161, 13-23.	1.2	8
115	In Reply. <i>Oncologist</i> , 2015, 20, e14-e14.	1.9	0
116	Conversion of abiraterone to D4A drives anti-tumour activity in prostate cancer. <i>Nature</i> , 2015, 523, 347-351.	13.7	221
117	The next 150 years of congenital adrenal hyperplasia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2015, 153, 63-71.	1.2	57
118	Adrenal Steroidogenesis and Congenital Adrenal Hyperplasia. <i>Endocrinology and Metabolism Clinics of North America</i> , 2015, 44, 275-296.	1.2	121
119	Human Cytochrome P450 21A2, the Major Steroid 21-Hydroxylase. <i>Journal of Biological Chemistry</i> , 2015, 290, 13128-13143.	1.6	74
120	Bone Morphogenetic Protein-4 (BMP4): A Paracrine Regulator of Human Adrenal C19 Steroid Synthesis. <i>Endocrinology</i> , 2015, 156, 2530-2540.	1.4	20
121	Aldosterone and Salt Loading Independently Exacerbate the Exercise Pressor Reflex in Rats. <i>Hypertension</i> , 2015, 66, 627-633.	1.3	11
122	Management considerations for the adult with congenital adrenal hyperplasia. <i>Molecular and Cellular Endocrinology</i> , 2015, 408, 190-197.	1.6	54
123	Genetic Forms of Adrenal Insufficiency. <i>Endocrine Practice</i> , 2015, 21, 395-399.	1.1	13
124	Making water-soluble integral membrane proteins in vivo using an amphipathic protein fusion strategy. <i>Nature Communications</i> , 2015, 6, 6826.	5.8	30
125	The Classic and Nonclassic Congenital Adrenal Hyperplasias. <i>Endocrine Practice</i> , 2015, 21, 383-389.	1.1	29
126	Profiles of 21-Carbon Steroids in 21-hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 2283-2290.	1.8	65

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127	The diverse chemistry of cytochrome P450 17A1 (P450c17, CYP17A1). <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2015, 151, 52-65.	1.2	94
128	P450 Enzymes in Steroid Processing. , 2015, , 851-879.		23
129	Mechanistic Scrutiny Identifies a Kinetic Role for Cytochrome b5 Regulation of Human Cytochrome P450c17 (CYP17A1, P450 17A1). <i>PLoS ONE</i> , 2015, 10, e0141252.	1.1	28
130	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, 646-649.	1.1	3
131	An Expert Consensus Statement on Use of Adrenal Vein Sampling for the Subtyping of Primary Aldosteronism. <i>Hypertension</i> , 2014, 63, 151-160.	1.3	475
132	Use of Prednisone With Abiraterone Acetate in Metastatic Castration-Resistant Prostate Cancer. <i>Oncologist</i> , 2014, 19, 1231-1240.	1.9	78
133	Adrenal Androgens and Androgen Precursorsâ€™ Definition, Synthesis, Regulation and Physiologic Actions. , 2014, 4, 1369-1381.		80
134	Androstenedione Is the Preferred Androgen Source in Hormone Refractory Prostate Cancerâ€™ Letter. <i>Clinical Cancer Research</i> , 2014, 20, 4971-4971.	3.2	9
135	Hypotension following Patent Ductus Arteriosus Ligation: The Role of Adrenal Hormones. <i>Journal of Pediatrics</i> , 2014, 164, 1449-1455.e1.	0.9	46
136	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.	1.2	29
137	Steroid Assays and Endocrinology: Best Practices for Basic Scientists. <i>Endocrinology</i> , 2014, 155, 2049-2051.	1.4	46
138	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.	1.4	13
139	Epoxidation Activities of Human Cytochromes P450c17 and P450c21. <i>Biochemistry</i> , 2014, 53, 7531-7540.	1.2	9
140	Catalytically Relevant Electrostatic Interactions of Cytochrome P450c17 (CYP17A1) and Cytochrome b5. <i>Journal of Biological Chemistry</i> , 2014, 289, 33838-33849.	1.6	32
141	Abiraterone Acetate to Lower Androgens in Women With Classic 21-Hydroxylase Deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, 2763-2770.	1.8	64
142	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, 317-322.	0.5	87
143	A Gain-of-Function Mutation in DHT Synthesis in Castration-Resistant Prostate Cancer. <i>Cell</i> , 2013, 154, 1074-1084.	13.5	257
144	Approach to the Patient: The Adult With Congenital Adrenal Hyperplasia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2013, 98, 2645-2655.	1.8	107

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145	Mass spectrometry theory and application to adrenal diseases. <i>Molecular and Cellular Endocrinology</i> , 2013, 371, 201-207.	1.6	28
146	The Action of Cytochrome <i>c</i> ₅ on CYP2E1 and CYP2C19 Activities Requires Anionic Residues D58 and D65. <i>Biochemistry</i> , 2013, 52, 210-220.	1.2	27
147	Introduction to the 2012 Keith L. Parker Memorial Lecturer: Walter L. Miller, MD. <i>Molecular and Cellular Endocrinology</i> , 2013, 371, 2-4.	1.6	1
148	Gene mutations that promote adrenal aldosterone production, sodium retention, and hypertension. <i>The Application of Clinical Genetics</i> , 2013, 7, 1.	1.4	4
149	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, 507-516.	1.8	234
150	Effect of <i>KCNJ5</i> Mutations on Gene Expression in Aldosterone-Producing Adenomas and Adrenocortical Cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012, 97, E1567-E1572.	1.8	130
151	Congenital Adrenal Hyperplasia—More Dogma Bites the Dust. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012, 97, 772-775.	1.8	23
152	Defects in Androgen Biosynthesis Causing 46,XY Disorders of Sexual Development. <i>Seminars in Reproductive Medicine</i> , 2012, 30, 417-426.	0.5	46
153	Abiraterone Inhibits 3 β -Hydroxysteroid Dehydrogenase: A Rationale for Increasing Drug Exposure in Castration-Resistant Prostate Cancer. <i>Clinical Cancer Research</i> , 2012, 18, 3571-3579.	3.2	87
154	Human Steroid Biosynthesis for the Oncologist. <i>Journal of Investigative Medicine</i> , 2012, 60, 495-503.	0.7	54
155	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, 2039-2049.	1.8	409
156	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, 7064-7077.	1.2	31
157	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.	1.2	8
158	Adrenal disorders in pregnancy. <i>Nature Reviews Endocrinology</i> , 2012, 8, 668-678.	4.3	37
159	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, 1606-1614.	1.8	310
160	Inhibition of 3 β -hydroxysteroid dehydrogenase by abiraterone as a rationale for dose escalation in castration-resistant prostate cancer. <i>Journal of Clinical Oncology</i> , 2012, 30, 209-209.	0.8	0
161	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, 13728-13733.	3.3	303
162	The Physiology and Biochemistry of Adrenarche. <i>Endocrine Development</i> , 2011, 20, 20-27.	1.3	38

#	ARTICLE	IF	CITATIONS
163	The Molecular Biology, Biochemistry, and Physiology of Human Steroidogenesis and Its Disorders. <i>Endocrine Reviews</i> , 2011, 32, 81-151.	8.9	1,743
164	Why Human Cytochrome P450c21 Is a Progesterone 21-Hydroxylase. <i>Biochemistry</i> , 2011, 50, 3968-3974.	1.2	24
165	The life and scientific contributions of Keith L. Parker, 1953–2008. <i>Molecular and Cellular Endocrinology</i> , 2011, 336, 191-192.	1.6	5
166	CYP17A1 Intron Mutation Causing Cryptic Splicing in 17 β -Hydroxylase Deficiency. <i>PLoS ONE</i> , 2011, 6, e25492.	1.1	19
167	Primary aldosteronism and a Texas two-step. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2011, 12, 37-42.	2.6	7
168	Steroid Profiling by Gas Chromatography–Mass Spectrometry and High Performance Liquid Chromatography–Mass Spectrometry for Adrenal Diseases. <i>Hormones and Cancer</i> , 2011, 2, 324-332.	4.9	63
169	Gene expression profiles in aldosterone-producing adenomas and adjacent adrenal glands. <i>European Journal of Endocrinology</i> , 2011, 164, 613-619.	1.9	54
170	Congenital adrenal hyperplasia in adults. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2010, 17, 210-216.	1.2	24
171	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, 744-748.	1.3	8
172	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, 1-17.	1.6	35
173	Management of the Adult with Congenital Adrenal Hyperplasia. <i>International Journal of Pediatric Endocrinology (Springer)</i> , 2010, 2010, 1-9.	1.6	5
174	3 β -Hydroxysteroid Dehydrogenase Is a Possible Pharmacological Target in the Treatment of Castration-Resistant Prostate Cancer. <i>Endocrinology</i> , 2010, 151, 3514-3520.	1.4	71
175	Reversible Sympathetic Overactivity in Hypertensive Patients with Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 4756-4761.	1.8	65
176	46,XX DSD: the masculinised female. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2010, 24, 219-242.	2.2	33
177	Non-traditional metabolic pathways of adrenal steroids. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2009, 10, 27-32.	2.6	15
178	Rapid Cortisol Assays Improve the Success Rate of Adrenal Vein Sampling for Primary Aldosteronism. <i>Annals of Surgery</i> , 2009, 249, 318-321.	2.1	95
179	Measurement of 18-Hydroxycorticosterone during Adrenal Vein Sampling for Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 2648-2651.	1.8	35
180	Utility, Limitations, and Pitfalls in Measuring Testosterone: An Endocrine Society Position Statement. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 405-413.	1.8	1,048

#	ARTICLE	IF	CITATIONS
181	Associations Among Androgens, Estrogens, and Natriuretic Peptides in Young Women. <i>Journal of the American College of Cardiology</i> , 2007, 49, 109-116.	1.2	156
182	Phenotypic variability in 17 β -hydroxysteroid dehydrogenase-3 deficiency and diagnostic pitfalls. <i>Clinical Endocrinology</i> , 2007, 67, 20-28.	1.2	115
183	“Subclinical Cushing’s syndrome” is not subclinical: improvement after adrenalectomy in 9 patients. <i>Surgery</i> , 2007, 142, 900-905.e1.	1.0	78
184	Primary aldosteronism. <i>Current Cardiology Reports</i> , 2007, 9, 447-452.	1.3	2
185	Human Cytochrome b5 Requires Residues E48 and E49 to Stimulate the 17,20-Lyase Activity of Cytochrome P450c17. <i>Biochemistry</i> , 2006, 45, 755-762.	1.2	66
186	Arginine 276 Controls the Directional Preference of AKR1C9 (Rat Liver 3 β -Hydroxysteroid) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 542 Td	1.4	23
187	Miscellaneous endocrine causes of hypertension. <i>Current Cardiology Reports</i> , 2005, 7, 418-424.	1.3	1
188	Minireview: Cellular Redox State Regulates Hydroxysteroid Dehydrogenase Activity and Intracellular Hormone Potency. <i>Endocrinology</i> , 2005, 146, 2531-2538.	1.4	95
189	The Rise in Adrenal Androgen Biosynthesis: Adrenarche. <i>Seminars in Reproductive Medicine</i> , 2004, 22, 337-347.	0.5	159
190	Overview of Dehydroepiandrosterone Biosynthesis. <i>Seminars in Reproductive Medicine</i> , 2004, 22, 281-288.	0.5	70
191	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.	1.8	194
192	Adrenarche - physiology, biochemistry and human disease. <i>Clinical Endocrinology</i> , 2004, 60, 288-296.	1.2	279
193	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, 43-48.	1.8	33
194	The backdoor pathway to dihydrotestosterone. <i>Trends in Endocrinology and Metabolism</i> , 2004, 15, 432-438.	3.1	275
195	Human 17 β -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, 50-59.	1.4	57
196	5 α -reduced C21 steroids are substrates for human cytochrome P450c17. <i>Archives of Biochemistry and Biophysics</i> , 2003, 418, 151-160.	1.4	86
197	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, 134-144.	1.4	39
198	CYP17 Mutation E305G Causes Isolated 17,20-Lyase Deficiency by Selectively Altering Substrate Binding. <i>Journal of Biological Chemistry</i> , 2003, 278, 48563-48569.	1.6	116

#	ARTICLE	IF	CITATIONS
199	5 α -Androstane-3 β ,17 β -Diol Is Formed in Tammar Wallaby Pouch Young Testes by a Pathway Involving 5 α -Pregnane-3 β ,17 β -Diol-20-One as a Key Intermediate. <i>Endocrinology</i> , 2003, 144, 575-580.	1.4	166
200	The 17, 20-Lyase Activity of Cytochrome P450c17 from Human Fetal Testis Favors the Δ^5 Steroidogenic Pathway. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2003, 88, 3762-3766.	1.8	148
201	Aldo is back: recent advances and unresolved controversies in hyperaldosteronism. <i>Current Opinion in Nephrology and Hypertension</i> , 2003, 12, 153-158.	1.0	12
202	TOWARDS A UNIFYING MECHANISM FOR CYP17 MUTATIONS THAT CAUSE ISOLATED 17, 20-LYASE DEFICIENCY. <i>Endocrine Research</i> , 2002, 28, 443-447.	0.6	12
203	THE GENETICS, PATHOPHYSIOLOGX, AND MANAGEMENT OF HUMAN DEFICIENCIES OF P450c17. <i>Endocrinology and Metabolism Clinics of North America</i> , 2001, 30, 101-119.	1.2	239
204	Molecular Modeling of the Hamster Adrenala P450C17. <i>Endocrine Research</i> , 2000, 26, 723-728.	0.6	1
205	Estrogen: Consequences and Implications of Human Mutations in Synthesis and Action1. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1999, 84, 4677-4694.	1.8	334
206	Molecular Modeling of Human P450c17 (17 β -Hydroxylase/ 17,20-Lyase): Insights into Reaction Mechanisms and Effects of Mutations. <i>Molecular Endocrinology</i> , 1999, 13, 1169-1182.	3.7	185
207	P450c17 Mutations R347H and R358Q Selectively Disrupt 17,20-Lyase Activity by Disrupting Interactions with P450 Oxidoreductase and Cytochrome b ₅ . <i>Molecular Endocrinology</i> , 1999, 13, 167-175.	3.7	173
208	Cytochrome b 5 Augments the 17,20-Lyase Activity of Human P450c17 without Direct Electron Transfer. <i>Journal of Biological Chemistry</i> , 1998, 273, 3158-3165.	1.6	490
209	The regulation of 17,20 lyase activity. <i>Steroids</i> , 1997, 62, 133-142.	0.8	218
210	The genetic and functional basis of isolated 17,20 α -lyase deficiency. <i>Nature Genetics</i> , 1997, 17, 201-205.	9.4	306