

# Maria-Christina Zennaro

## List of Publications by Year in descending order

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121  
papers

6,504  
citations

57758

44  
h-index

69250

77  
g-index

124  
all docs

124  
docs citations

124  
times ranked

4837  
citing authors

#	ARTICLE	IF	CITATIONS
1	Colocalization of Wnt/ $\beta$ -Catenin and ACTH Signaling Pathways and Paracrine Regulation in Aldosterone-producing Adenoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 419-434.	3.6	5
2	Somatic mutations of GNA11 and GNAQ in CTNNB1-mutant aldosterone-producing adenomas presenting in puberty, pregnancy or menopause. <i>Nature Genetics</i> , 2021, 53, 1360-1372.	21.4	37
3	Circulating microRNAs as Diagnostic Markers in Primary Aldosteronism. <i>Cancers</i> , 2021, 13, 5312.	3.7	6
4	Renin-aldosterone system evaluation over four decades in an extended family with autosomal dominant pseudohypoaldosteronism due to a deletion in the NR3C2 gene. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2020, 204, 105755.	2.5	8
5	Pathogenesis and treatment of primary aldosteronism. <i>Nature Reviews Endocrinology</i> , 2020, 16, 578-589.	9.6	65
6	Genetics, prevalence, screening and confirmation of primary aldosteronism: a position statement and consensus of the Working Group on Endocrine Hypertension of The European Society of Hypertension. <i>Journal of Hypertension</i> , 2020, 38, 1919-1928.	0.5	151
7	Genetic, Cellular, and Molecular Heterogeneity in Adrenals With Aldosterone-Producing Adenoma. <i>Hypertension</i> , 2020, 75, 1034-1044.	2.7	89
8	Glucocorticoid Excess in Patients with Pheochromocytoma Compared with Paraganglioma and Other Forms of Hypertension. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e3374-e3383.	3.6	17
9	Genetic and Genomic Mechanisms of Primary Aldosteronism. <i>Trends in Molecular Medicine</i> , 2020, 26, 819-832.	6.7	20
10	Old and new genes in primary aldosteronism. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2020, 34, 101375.	4.7	13
11	Pathogenesis of hypertension in a mouse model for human CLCN2 related hyperaldosteronism. <i>Nature Communications</i> , 2019, 10, 4678.	12.8	33
12	Retinoic acid receptor $\beta$ as a novel contributor to adrenal cortex structure and function through interactions with Wnt and Vegfa signalling. <i>Scientific Reports</i> , 2019, 9, 14677.	3.3	10
13	Germline and somatic genetic basis of primary aldosteronism. <i>Current Opinion in Endocrine and Metabolic Research</i> , 2019, 8, 160-166.	1.4	0
14	From Transcripts to Proteins. <i>Hypertension</i> , 2019, 73, 284-285.	2.7	2
15	Molecular mechanisms in primary aldosteronism. <i>Journal of Endocrinology</i> , 2019, 242, R67-R79.	2.6	4
16	Simultaneous sequencing of 37 genes identified causative mutations in the majority of children with renal tubulopathies. <i>Kidney International</i> , 2018, 93, 961-967.	5.2	77
17	A gain-of-function mutation in the CLCN2 chloride channel gene causes primary aldosteronism. <i>Nature Genetics</i> , 2018, 50, 355-361.	21.4	154
18	Overview of aldosterone-related genetic syndromes and recent advances. <i>Current Opinion in Endocrinology, Diabetes and Obesity</i> , 2018, 25, 147-154.	2.3	6

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19	MicroRNA-204 Is Necessary for Aldosterone-Stimulated T-Type Calcium Channel Expression in Cardiomyocytes. <i>International Journal of Molecular Sciences</i> , 2018, 19, 2941.	4.1	11
20	Molecular genetics of Conn adenomas in the era of exome analysis. <i>Presse Medicale</i> , 2018, 47, e151-e158.	1.9	5
21	Somatic and inherited mutations in primary aldosteronism. <i>Journal of Molecular Endocrinology</i> , 2017, 59, R47-R63.	2.5	42
22	30 YEARS OF THE MINERALOCORTICOID RECEPTOR: Mineralocorticoid receptor mutations. <i>Journal of Endocrinology</i> , 2017, 234, T93-T106.	2.6	39
23	Genetic Causes of Functional Adrenocortical Adenomas. <i>Endocrine Reviews</i> , 2017, 38, 516-537.	20.1	72
24	30 YEARS OF THE MINERALOCORTICOID RECEPTOR: The scientific impact of cloning the mineralocorticoid receptor: 30 years on. <i>Journal of Endocrinology</i> , 2017, 234, E3-E6.	2.6	4
25	Pseudohypoaldosteronism types I and II: little more than a name in common. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2017, 30, 597-601.	0.9	17
26	CACNA1H Mutations Are Associated With Different Forms of Primary Aldosteronism. <i>EBioMedicine</i> , 2016, 13, 225-236.	6.1	119
27	Aldosterone-Producing Adenoma With a Somatic KCNJ5 Mutation Revealing APC-Dependent Familial Adenomatous Polyposis. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 3874-3878.	3.6	32
28	[OP.3A.02] RETINOIC ACID RECEPTOR SIGNALING CONTRIBUTES TO ADRENAL CORTEX MORPHOLOGY AND FUNCTIONAL ZONATION. <i>Journal of Hypertension</i> , 2016, 34, e26.	0.5	0
29	[OP.LB01.12] CACNA1H MUTATIONS ARE ASSOCIATED WITH YOUNG ONSET AND FAMILIAL FORMS OF PRIMARY ALDOSTERONISM. <i>Journal of Hypertension</i> , 2016, 34, e39.	0.5	0
30	SFE/SFHTA/AFCE consensus on primary aldosteronism, part 5: Genetic diagnosis of primary aldosteronism. <i>Annales D'Endocrinologie</i> , 2016, 77, 214-219.	1.4	17
31	SFE/SFHTA/AFCE primary aldosteronism consensus: Introduction and handbook. <i>Annales D'Endocrinologie</i> , 2016, 77, 179-186.	1.4	50
32	Local Control of Aldosterone Production and Primary Aldosteronism. <i>Trends in Endocrinology and Metabolism</i> , 2016, 27, 123-131.	7.1	29
33	Renal Tubular Disorders of Electrolyte Regulation in Children. , 2016, , 1201-1271.		2
34	CO-33: Different somatic mutations in multinodular adrenals with aldosterone-producing adenoma. <i>Annales De Cardiologie Et D'Angiologie</i> , 2015, 64, S16.	0.6	0
35	Overweight Is a Major Contributor to Atherosclerosis in Systemic Lupus Erythematosus Patients at Apparent Low Risk for Cardiovascular Disease. <i>Medicine (United States)</i> , 2015, 94, e2177.	1.0	21
36	Molecular and Cellular Mechanisms of Aldosterone Producing Adenoma Development. <i>Frontiers in Endocrinology</i> , 2015, 6, 95.	3.5	20

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37	Bilateral Idiopathic Adrenal Hyperplasia: Genetics and Beyond. <i>Hormone and Metabolic Research</i> , 2015, 47, 947-952.	1.5	19
38	An update on novel mechanisms of primary aldosteronism. <i>Journal of Endocrinology</i> , 2015, 224, R63-R77.	2.6	56
39	Paracrine control of steroidogenesis by serotonin in adrenocortical neoplasms. <i>Molecular and Cellular Endocrinology</i> , 2015, 408, 198-204.	3.2	21
40	Inherited forms of mineralocorticoid hypertension. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2015, 29, 633-645.	4.7	32
41	Functional histopathological markers of aldosterone producing adenoma and somatic KCNJ5 mutations. <i>Molecular and Cellular Endocrinology</i> , 2015, 408, 220-226.	3.2	23
42	Different Somatic Mutations in Multinodular Adrenals With Aldosterone-Producing Adenoma. <i>Hypertension</i> , 2015, 66, 1014-1022.	2.7	55
43	Mast Cell Hyperplasia Is Associated With Aldosterone Hypersecretion in a Subset of Aldosterone-Producing Adenomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, E550-E560.	3.6	32
44	Renal Tubular Disorders of Electrolyte Regulation in Children. , 2015, , 1-80.		0
45	Genetic Spectrum and Clinical Correlates of Somatic Mutations in Aldosterone-Producing Adenoma. <i>Hypertension</i> , 2014, 64, 354-361.	2.7	248
46	WNT/ $\beta$ -catenin signalling is activated in aldosterone-producing adenomas and controls aldosterone production. <i>Human Molecular Genetics</i> , 2014, 23, 889-905.	2.9	157
47	Increased Arterial Stiffness in Systemic Lupus Erythematosus (SLE) Patients at Low Risk for Cardiovascular Disease: A Cross-Sectional Controlled Study. <i>PLoS ONE</i> , 2014, 9, e94511.	2.5	64
48	Diastrophic Dysplasia Sulfate Transporter (SLC26A2) Is Expressed in the Adrenal Cortex and Regulates Aldosterone Secretion. <i>Hypertension</i> , 2014, 63, 1102-1109.	2.7	21
49	Activation of the Hypothalamic-Pituitary-Adrenal Axis in Adults With Mineralocorticoid Receptor Haploinsufficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, E1586-E1591.	3.6	10
50	From Genetic Abnormalities to Pathophysiological Mechanisms. , 2014, , 53-74.		0
51	Pseudohypoaldosteronism type 1: Management issues. <i>Indian Pediatrics</i> , 2013, 50, 331-333.	0.4	7
52	Cardiovascular Effects of Aldosterone. <i>Circulation: Cardiovascular Genetics</i> , 2013, 6, 381-390.	5.1	27
53	KCNJ5 mutations in aldosterone producing adenoma and relationship with adrenal cortex remodeling. <i>Molecular and Cellular Endocrinology</i> , 2013, 371, 221-227.	3.2	38
54	Somatic mutations in ATP1A1 and ATP2B3 lead to aldosterone-producing adenomas and secondary hypertension. <i>Nature Genetics</i> , 2013, 45, 440-444.	21.4	460

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55	Genetics in endocrinology: Genetics of mineralocorticoid excess: an update for clinicians. <i>European Journal of Endocrinology</i> , 2013, 169, R15-R25.	3.7	26
56	Progress in Primary Aldosteronism 2. <i>Hormone and Metabolic Research</i> , 2012, 44, 155-156.	1.5	0
57	<i>KCNJ5</i> Mutations in European Families With Nonglucocorticoid Remediable Familial Hyperaldosteronism. <i>Hypertension</i> , 2012, 59, 235-240.	2.7	176
58	Primary Aldosteronism Takes (KCNJ)Five!. <i>Endocrinology</i> , 2012, 153, 1575-1577.	2.8	2
59	Task3 Potassium Channel Gene Inactivation Causes Low Renin and Salt-Sensitive Arterial Hypertension. <i>Endocrinology</i> , 2012, 153, 4740-4748.	2.8	63
60	<i>Dkk3</i> is a component of the genetic circuitry regulating aldosterone biosynthesis in the adrenal cortex. <i>Human Molecular Genetics</i> , 2012, 21, 4922-4929.	2.9	22
61	Integrating Genetics and Genomics in Primary Aldosteronism. <i>Hypertension</i> , 2012, 60, 580-588.	2.7	22
62	A network perspective on metabolic inconsistency. <i>BMC Systems Biology</i> , 2012, 6, 41.	3.0	26
63	Prevalence, Clinical, and Molecular Correlates of <i>KCNJ5</i> Mutations in Primary Aldosteronism. <i>Hypertension</i> , 2012, 59, 592-598.	2.7	246
64	Aldosterone resistance: Structural and functional considerations and new perspectives. <i>Molecular and Cellular Endocrinology</i> , 2012, 350, 206-215.	3.2	60
65	The role of the mineralocorticoid receptor in adipocyte biology and fat metabolism. <i>Molecular and Cellular Endocrinology</i> , 2012, 350, 281-288.	3.2	109
66	Pseudohypoaldosteronism type 1: the index case revisited. <i>Clinical Endocrinology</i> , 2011, 74, 408-410.	2.4	4
67	Progesterone increase counteracts aldosterone action in a pregnant woman with primary aldosteronism. <i>Clinical Endocrinology</i> , 2011, 74, 278-279.	2.4	25
68	Mineralocorticoid Receptor Mutations Differentially Affect Individual Gene Expression Profiles in Pseudohypoaldosteronism Type 1. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011, 96, E519-E527.	3.6	30
69	Antiadipogenic Effects of the Mineralocorticoid Receptor Antagonist Drosiprenone: Potential Implications for the Treatment of Metabolic Syndrome. <i>Endocrinology</i> , 2011, 152, 113-125.	2.8	124
70	Aldosterone-Producing Adenoma Formation in the Adrenal Cortex Involves Expression of Stem/Progenitor Cell Markers. <i>Endocrinology</i> , 2011, 152, 4753-4763.	2.8	85
71	Mutations in <i>KCNJ5</i> Gene Cause Hyperaldosteronism. <i>Circulation Research</i> , 2011, 108, 1417-1418.	4.5	16
72	Mineralocorticoid Receptor Mutations and a Severe Recessive Pseudohypoaldosteronism Type 1. <i>Journal of the American Society of Nephrology: JASN</i> , 2011, 22, 1997-2003.	6.1	33

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73	A homozygous missense mutation in SCNN1A is responsible for a transient neonatal form of pseudohypoaldosteronism type 1. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2011, 301, E467-E473.	3.5	35
74	A Homozygous Missense Mutation in SCNN1A Is Responsible for a Transient Form of Pseudohypoaldosteronism Type 1. , 2011, , P2-740-P2-740.		1
75	Asymptomatic myocardial ischemic disease in antiphospholipid syndrome: A controlled cardiac magnetic resonance imaging study. <i>Arthritis and Rheumatism</i> , 2010, 62, 2093-2100.	6.7	43
76	The Functional c.-2G>C Variant of the Mineralocorticoid Receptor Modulates Blood Pressure, Renin, and Aldosterone Levels. <i>Hypertension</i> , 2010, 56, 995-1002.	2.7	46
77	Adrenal Cortex Remodeling and Functional Zona Glomerulosa Hyperplasia in Primary Aldosteronism. <i>Hypertension</i> , 2010, 56, 885-892.	2.7	128
78	Analysis of Insulin Sensitivity in Adipose Tissue of Patients with Primary Aldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 4037-4042.	3.6	40
79	Mineralocorticoid receptors in the metabolic syndrome. <i>Trends in Endocrinology and Metabolism</i> , 2009, 20, 444-451.	7.1	69
80	Tubular Disorders of Electrolyte Regulation. , 2009, , 929-977.		9
81	Potential role of progestogens in the control of adipose tissue and salt sensitivity via interaction with the mineralocorticoid receptor. <i>Climacteric</i> , 2008, 11, 258-264.	2.4	6
82	Pseudohypoaldosteronisms, report on a 10-patient series. <i>Nephrology Dialysis Transplantation</i> , 2008, 23, 1636-1641.	0.7	69
83	The Mineralocorticoid Receptor in Endothelial Physiology and Disease: Novel Concepts in the Understanding of Erectile Dysfunction. <i>Current Pharmaceutical Design</i> , 2008, 14, 3749-3757.	1.9	19
84	Pivotal role of the mineralocorticoid receptor in corticosteroid-induced adipogenesis. <i>FASEB Journal</i> , 2007, 21, 2185-2194.	0.5	277
85	Mineralocorticoid receptor mutations are the principal cause of renal type 1 pseudohypoaldosteronism. <i>Human Mutation</i> , 2007, 28, 33-40.	2.5	79
86	A Common Polymorphism in the Mineralocorticoid Receptor Modulates Stress Responsiveness. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2006, 91, 5083-5089.	3.6	188
87	Autosomal Dominant Pseudohypoaldosteronism Type 1. <i>Journal of the American Society of Nephrology: JASN</i> , 2006, 17, 1429-1436.	6.1	118
88	Enhancement of $\beta_2$ -adrenergic cAMP-signaling by the mineralocorticoid receptor. <i>Molecular and Cellular Endocrinology</i> , 2005, 231, 23-31.	3.2	15
89	Prolactin potentiates insulin-stimulated leptin expression and release from differentiated brown adipocytes. <i>Journal of Molecular Endocrinology</i> , 2004, 33, 679-691.	2.5	32
90	New Naturally Occurring Missense Mutations of the Human Mineralocorticoid Receptor Disclose Important Residues Involved in Dynamic Interactions with Deoxyribonucleic Acid, Intracellular Trafficking, and Ligand Binding. <i>Molecular Endocrinology</i> , 2004, 18, 2151-2165.	3.7	37

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91	Mineralocorticoid resistance. Trends in Endocrinology and Metabolism, 2004, 15, 264-270.	7.1	75
92	Inactivating mutations of the mineralocorticoid receptor in Type I pseudohypoaldosteronism. Molecular and Cellular Endocrinology, 2004, 217, 119-125.	3.2	61
93	Expression and function of the human mineralocorticoid receptor: lessons from transgenic mouse models. Molecular and Cellular Endocrinology, 2004, 217, 127-136.	3.2	41
94	GILZ, a new target for the transcription factor FoxO3, protects T lymphocytes from interleukin-2 withdrawal-induced apoptosis. Blood, 2004, 104, 215-223.	1.4	139
95	Aldosterone Receptors. , 2004, , 158-163.		0
96	Different Inactivating Mutations of the Mineralocorticoid Receptor in Fourteen Families Affected by Type I Pseudohypoaldosteronism. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 2508-2517.	3.6	81
97	Protein Inhibitor of Activated Signal Transducer and Activator of Transcription 1 Interacts with the N-Terminal Domain of Mineralocorticoid Receptor and Represses Its Transcriptional Activity: Implication of Small Ubiquitin-Related Modifier 1 Modification. Molecular Endocrinology, 2003, 17, 2529-2542.	3.7	109
98	Mineralocorticoid Receptor, Natural Mutations of. , 2003, , 691-695.		0
99	Characterization of Rat NDRG2 (N-Myc Downstream Regulated Gene 2), a Novel Early Mineralocorticoid-specific Induced Gene. Journal of Biological Chemistry, 2002, 277, 31506-31515.	3.4	131
100	Brown adipocytes are novel sites of expression and regulation of adiponectin and resistin. FEBS Letters, 2002, 532, 345-350.	2.8	103
101	Mineralocorticoid and glucocorticoid receptors inhibit UCP expression and function in brown adipocytes. American Journal of Physiology - Endocrinology and Metabolism, 2001, 280, E640-E649.	3.5	90
102	Alteration of Cardiac and Renal Functions in Transgenic Mice Overexpressing Human Mineralocorticoid Receptor. Journal of Biological Chemistry, 2001, 276, 38911-38920.	3.4	106
103	A New Human MR Splice Variant Is a Ligand-Independent Transactivator Modulating Corticosteroid Action. Molecular Endocrinology, 2001, 15, 1586-1598.	3.7	94
104	A New Human MR Splice Variant Is a Ligand-Independent Transactivator Modulating Corticosteroid Action. Molecular Endocrinology, 2001, 15, 1586-1598.	3.7	32
105	Transgenic mouse models to study human mineralocorticoid receptor function in vivo. Kidney International, 2000, 57, 1299-1306.	5.2	15
106	The mineralocorticoid receptor mediates aldosterone-induced differentiation of T37i cells into brown adipocytes. American Journal of Physiology - Endocrinology and Metabolism, 2000, 279, E386-E394.	3.5	70
107	Targeted Oncogenesis Reveals a Distinct Tissue-specific Utilization of Alternative Promoters of the Human Mineralocorticoid Receptor Gene in Transgenic Mice. Journal of Biological Chemistry, 2000, 275, 7878-7886.	3.4	44
108	Mineralocorticoid receptor isoforms. Current Opinion in Endocrinology, Diabetes and Obesity, 1998, 5, 183-188.	0.6	5

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109	Syndromes of glucocorticoid and mineralocorticoid resistance. <i>European Journal of Endocrinology</i> , 1998, 139, 127-138.	3.7	14
110	Hibernoma development in transgenic mice identifies brown adipose tissue as a novel target of aldosterone action.. <i>Journal of Clinical Investigation</i> , 1998, 101, 1254-1260.	8.2	118
111	Tissue-Specific Expression of $\hat{1}\pm$ and $\hat{1}^2$ Messenger Ribonucleic Acid Isoforms of the Human Mineralocorticoid Receptor in Normal and Pathological States. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1997, 82, 1345-1352.	3.6	65
112	Tissue-Specific Expression of $\hat{A}$ and $\hat{A}$ Messenger Ribonucleic Acid Isoforms of the Human Mineralocorticoid Receptor in Normal and Pathological States. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1997, 82, 1345-1352.	3.6	59
113	Mineralocorticoid resistance. <i>Steroids</i> , 1996, 61, 189-192.	1.8	13
114	Characterization of the human mineralocorticoid receptor gene 5'- regulatory region: evidence for differential hormonal regulation of two alternative promoters via nonclassical mechanisms. <i>Molecular Endocrinology</i> , 1996, 10, 1549-1560.	3.7	61
115	Human Mineralocorticoid Receptor Genomic Structure and Identification of Expressed Isoforms. <i>Journal of Biological Chemistry</i> , 1995, 270, 21016-21020.	3.4	131
116	Pseudohypoaldosteronism: Evaluation of type I receptors by radioreceptor assay and by antireceptor antibodies. <i>Steroids</i> , 1995, 60, 161-163.	1.8	5
117	Molecular characterization of the mineralocorticoid receptor in pseudohypoaldosteronism. <i>Steroids</i> , 1995, 60, 164-167.	1.8	12
118	No alteration in the primary structure of the mineralocorticoid receptor in a family with pseudohypoaldosteronism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1994, 79, 32-38.	3.6	40
119	Regulation of aldosterone receptors in hypertension. <i>Steroids</i> , 1993, 58, 611-613.	1.8	9
120	Corticosteroid receptors and aging. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1993, 45, 191-194.	2.5	17
121	Pseudohypoaldosteronism and mineralocorticoid receptor abnormalities. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1991, 40, 363-365.	2.5	18