Massimo Mannelli

List of Publications by Year in Descending Order

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

118 191 14,370 54 h-index g-index citations papers 16,622 6.8 5.58 194 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
191	SDHB and SDHD silenced pheochromocytoma spheroids respond differently to tumour microenvironment and their aggressiveness is inhibited by impairing stroma metabolism <i>Molecular and Cellular Endocrinology</i> , 2022 , 547, 111594	4.4	O
190	A Multicenter Epidemiological Study on Second Malignancy in Non-Syndromic Pheochromocytoma/Paraganglioma Patients in Italy. <i>Cancers</i> , 2021 , 13,	6.6	1
189	International consensus on initial screening and follow-up of asymptomatic SDHx mutation carriers. <i>Nature Reviews Endocrinology</i> , 2021 , 17, 435-444	15.2	12
188	Circulating Fascin 1 as a Promising Prognostic Marker in Adrenocortical Cancer. <i>Frontiers in Endocrinology</i> , 2021 , 12, 698862	5.7	1
187	Adrenocortical carcinoma: current treatment options. <i>Current Opinion in Oncology</i> , 2021 , 33, 16-22	4.2	5
186	Functional significance of germline EPAS1 variants. Endocrine-Related Cancer, 2021, 28, 97-109	5.7	1
185	Analysis of Telomere Maintenance Related Genes Reveals as a New Metastatic-Risk Marker in Pheochromocytoma/Paraganglioma. <i>Cancers</i> , 2021 , 13,	6.6	2
184	Prognostic and Monitoring Value of Circulating Tumor Cells in Adrenocortical Carcinoma: A Preliminary Monocentric Study. <i>Cancers</i> , 2020 , 12,	6.6	6
183	The Role of Metabolic Changes in Shaping the Fate of Cancer-Associated Adipose Stem Cells. <i>Frontiers in Cell and Developmental Biology</i> , 2020 , 8, 332	5.7	3
182	Metabolomics, machine learning and immunohistochemistry to predict succinate dehydrogenase mutational status in phaeochromocytomas and paragangliomas. <i>Journal of Pathology</i> , 2020 , 251, 378-3	8 9 ·4	11
181	Germline Mutation in Gene Associated with Loss of Heterozygosity: Usefulness of Next-Generation Sequencing in the Genetic Screening of Patients with Pheochromocytoma. <i>International Journal of Endocrinology</i> , 2020 , 2020, 3671396	2.7	O
180	Genetics, diagnosis, management and future directions of research of phaeochromocytoma and paraganglioma: 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, 1443-1456	1.9	62
179	Effects of Germline CYP2W1*6 and CYP2B6*6 Single Nucleotide Polymorphisms on Mitotane Treatment in Adrenocortical Carcinoma: A Multicenter ENSAT Study. <i>Cancers</i> , 2020 , 12,	6.6	14
178	Urine steroid metabolomics for the differential diagnosis of adrenal incidentalomas in the EURINE-ACT study: a prospective test validation study. <i>Lancet Diabetes and Endocrinology,the</i> , 2020 , 8, 773-781	18.1	56
177	and Non-Chromaffin Tumors: A Mediastinal Germ Cell Tumor Occurring in a Young Man with Germline Mutation. <i>Medicina (Lithuania)</i> , 2020 , 56,	3.1	1
176	Le crisi di un anziano signore. <i>L Endocrinologo</i> , 2020 , 21, 303-304	O	
175	Pheochromocytomas and Paragangliomas as Causes of Endocrine Hypertension. <i>Frontiers in Endocrinology</i> , 2019 , 10, 333	5.7	7

174 Malignant Pheochromocytoma **2019**, 460-468

173	Prognosis of Malignant Pheochromocytoma and Paraganglioma (MAPP-Prono Study): A European Network for the Study of Adrenal Tumors Retrospective Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 2367-2374	5.6	57
172	Metabolome-guided genomics to identify pathogenic variants in isocitrate dehydrogenase, fumarate hydratase, and succinate dehydrogenase genes in pheochromocytoma and paraganglioma. <i>Genetics in Medicine</i> , 2019 , 21, 705-717	8.1	36
171	Heat Shock Protein 90 as a Prognostic Marker and Therapeutic Target for Adrenocortical Carcinoma. <i>Frontiers in Endocrinology</i> , 2019 , 10, 487	5.7	4
170	Value of Molecular Classification for Prognostic Assessment of Adrenocortical Carcinoma. <i>JAMA Oncology</i> , 2019 , 5, 1440-1447	13.4	31
169	Integrative multi-omics analysis identifies a prognostic miRNA signature and a targetable miR-21-3p/TSC2/mTOR axis in metastatic pheochromocytoma/paraganglioma. <i>Theranostics</i> , 2019 , 9, 4946-4958	12.1	30
168	La Fascina-1 ในท nuovo biomarker prognostico associato all [hvasivit[del carcinoma corticosurrenalico. <i>L Endocrinologo</i> , 2019 , 20, 392-393	O	
167	The Adipose Stem Cell as a Novel Metabolic Actor in Adrenocortical Carcinoma Progression: Evidence from an In Vitro Tumor Microenvironment Crosstalk Model. <i>Cancers</i> , 2019 , 11,	6.6	8
166	CT Characteristics of Pheochromocytoma: Relevance for the Evaluation of Adrenal Incidentaloma. Journal of Clinical Endocrinology and Metabolism, 2019 , 104, 312-318	5.6	59
165	Fascin-1 Is a Novel Prognostic Biomarker Associated With Tumor Invasiveness in Adrenocortical Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019 , 104, 1712-1724	5.6	17
164	Human fetal adrenal cells retain age-related stem- and endocrine-differentiation potential in culture. <i>FASEB Journal</i> , 2019 , 33, 2263-2277	0.9	16
163	Adrenalectomy Lowers Incident Atrial Fibrillation in Primary Aldosteronism Patients at Long Term. <i>Hypertension</i> , 2018 , 71, 585-591	8.5	95
162	DIAGNOSIS of ENDOCRINE DISEASE: SDHx mutations: beyond pheochromocytomas and paragangliomas. <i>European Journal of Endocrinology</i> , 2018 , 178, R11-R17	6.5	18
161	Role of MDH2 pathogenic variant in pheochromocytoma and paraganglioma patients. <i>Genetics in Medicine</i> , 2018 , 20, 1652-1662	8.1	33
160	The Endocrine Regulation of Blood Pressure. <i>Endocrinology</i> , 2018 , 611-625	0.1	
159	A case of malignant insulinoma responsive to somatostatin analogs treatment. <i>BMC Endocrine Disorders</i> , 2018 , 18, 98	3.3	3
158	Primary fibroblast co-culture stimulates growth and metabolism in Sdhb-impaired mouse pheochromocytoma MTT cells. <i>Cell and Tissue Research</i> , 2018 , 374, 473-485	4.2	12
157	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. <i>Cancer Cell</i> , 2017 , 31, 181-193	24.3	350

156	Prognostic factors in ectopic Cushing's syndrome due to neuroendocrine tumors: a multicenter study. <i>European Journal of Endocrinology</i> , 2017 , 176, 453-461	6.5	39
155	Long-Term Outcomes of Adjuvant Mitotane Therapy in Patients With Radically Resected Adrenocortical Carcinoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 1358-1365	5.6	75
154	Quantitative Value of Aldosterone-Renin Ratio for Detection of Aldosterone-Producing Adenoma: The Aldosterone-Renin Ratio for Primary Aldosteronism (AQUARR) Study. <i>Journal of the American Heart Association</i> , 2017 , 6,	6	41
153	The mTORC1 Complex Is Significantly Overactivated in SDHX-Mutated Paragangliomas. <i>Neuroendocrinology</i> , 2017 , 105, 384-393	5.6	6
152	Sunitinib in the therapy of malignant paragangliomas: report on the efficacy in a SDHB mutation carrier and review of the literature. <i>Archives of Endocrinology and Metabolism</i> , 2017 , 61, 90-97	2.2	12
151	The microenvironment induces collective migration in -silenced mouse pheochromocytoma spheroids. <i>Endocrine-Related Cancer</i> , 2017 , 24, 555-564	5.7	19
150	Evaluation and diagnostic potential of circulating extracellular vesicle-associated microRNAs in adrenocortical tumors. <i>Scientific Reports</i> , 2017 , 7, 5474	4.9	33
149	Assessment of VAV2 Expression Refines Prognostic Prediction in Adrenocortical Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2017, 102, 3491-3498	5.6	24
148	PheoSeq: A Targeted Next-Generation Sequencing Assay for Pheochromocytoma and Paraganglioma Diagnostics. <i>Journal of Molecular Diagnostics</i> , 2017 , 19, 575-588	5.1	51
147	Consensus Statement on next-generation-sequencing-based diagnostic testing of hereditary phaeochromocytomas and paragangliomas. <i>Nature Reviews Endocrinology</i> , 2017 , 13, 233-247	15.2	140
146	New insights in the clinical and translational relevance of miR483-5p in adrenocortical cancer. <i>Oncotarget</i> , 2017 , 8, 65525-65533	3.3	21
145	Potential Pitfalls of SDH Immunohistochemical Detection in Paragangliomas and Phaeochromocytomas Harbouring Germline Gene Mutation. <i>Anticancer Research</i> , 2017 , 37, 805-812	2.3	20
144	DNA Methylation Is an Independent Prognostic Marker of Survival in Adrenocortical Cancer. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017 , 102, 923-932	5.6	35
143	Genetic Landscape of Sporadic Unilateral Adrenocortical Adenomas Without PRKACA p.Leu206Arg Mutation. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 3526-38	5.6	46
142	Metodiche di dosaggio di metanefrine e catecolamine: rischi e variabili interpretative per i dosaggi effettuati in centri non specializzati. <i>L Endocrinologo</i> , 2016 , 17, 107-108	О	
141	Epigenetic Mutation of the Succinate Dehydrogenase C Promoter in a Patient With Two Paragangliomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016 , 101, 359-63	5.6	36
140	Metformin as a new anti-cancer drug in adrenocortical carcinoma. <i>Oncotarget</i> , 2016 , 7, 49636-49648	3.3	30
139	The Endocrine Regulation of Blood Pressure. <i>Endocrinology</i> , 2016 , 1-15	0.1	

138	Commentary. Clinical Chemistry, 2016 , 62, 929	5.5	
137	Caratterizzazione genomica del carcinoma surrenalico. <i>L Endocrinologo</i> , 2016 , 17, 293-299	0	
136	Functional and in silico assessment of MAX variants of unknown significance. <i>Journal of Molecular Medicine</i> , 2015 , 93, 1247-55	5.5	17
135	DNA Methylation Profiling in Pheochromocytoma and Paraganglioma Reveals Diagnostic and Prognostic Markers. <i>Clinical Cancer Research</i> , 2015 , 21, 3020-30	12.9	44
134	Cushing's syndrome in pregnancy. <i>Gynecological Endocrinology</i> , 2015 , 31, 102-4	2.4	11
133	Prognostic factors in stage III-IV adrenocortical carcinomas (ACC): an European Network for the Study of Adrenal Tumor (ENSAT) study. <i>Annals of Oncology</i> , 2015 , 26, 2119-25	10.3	134
132	Rare diseases in clinical endocrinology: a taxonomic classification system. <i>Journal of Endocrinological Investigation</i> , 2015 , 38, 193-259	5.2	9
131	SDHB/SDHA immunohistochemistry in pheochromocytomas and paragangliomas: a multicenter interobserver variation analysis using virtual microscopy: a Multinational Study of the European Network for the Study of Adrenal Tumors (ENS@T). <i>Modern Pathology</i> , 2015 , 28, 807-21	9.8	142
130	Immunohistochemical expression of stem cell markers in pheochromocytomas/paragangliomas is associated with SDHx mutations. <i>European Journal of Endocrinology</i> , 2015 , 173, 43-52	6.5	14
129	Role of microenvironment on neuroblastoma SK-N-AS SDHB-silenced cell metabolism and function. <i>Endocrine-Related Cancer</i> , 2015 , 22, 409-17	5.7	18
128	15 YEARS OF PARAGANGLIOMA: Metabolism and pheochromocytoma/paraganglioma. Endocrine-Related Cancer, 2015 , 22, T83-90	5.7	5
127	2D-DIGE proteomic analysis identifies new potential therapeutic targets for adrenocortical carcinoma. <i>Oncotarget</i> , 2015 , 6, 5695-706	3.3	22
126	Oncogenic features of the bone morphogenic protein 7 (BMP7) in pheochromocytoma. <i>Oncotarget</i> , 2015 , 6, 39111-26	3.3	11
125	Analysis of circulating microRNAs in adrenocortical tumors. <i>Laboratory Investigation</i> , 2014 , 94, 331-9	5.9	79
124	Patient affected by neurofibromatosis type 1 and thyroid C-cell hyperplasia harboring pathogenic germ-line mutations in both NF1 and RET genes. <i>Gene</i> , 2014 , 536, 332-5	3.8	16
123	Opposing effects of HIF1[and HIF2[bn chromaffin cell phenotypic features and tumor cell proliferation: Insights from MYC-associated factor X. <i>International Journal of Cancer</i> , 2014 , 135, 2054-6.	4 ^{7·5}	54
122	Integrated genomic characterization of adrenocortical carcinoma. <i>Nature Genetics</i> , 2014 , 46, 607-12	36.3	423
121	Succinate dehydrogenase subunit B mutations modify human neuroblastoma cell metabolism and proliferation. <i>Hormones and Cancer</i> , 2014 , 5, 174-84	5	17

120	Dissecting the origin of inducible brown fat in adult humans through a novel adipose stem cell model from adipose tissue surrounding pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E1903-12	5.6	15
119	Outcomes of adrenal-sparing surgery or total adrenalectomy in phaeochromocytoma associated with multiple endocrine neoplasia type 2: an international retrospective population-based study. <i>Lancet Oncology, The</i> , 2014 , 15, 648-55	21.7	110
118	Detection of circulating tumor cells in adrenocortical neoplasms. <i>Pathology</i> , 2014 , 46, S13-S14	1.6	
117	Novel somatic mutations in the catalytic subunit of the protein kinase A as a cause of adrenal Cushing's syndrome: a European multicentric study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E2093-100	5.6	67
116	Krebs cycle metabolite profiling for identification and stratification of pheochromocytomas/paragangliomas due to succinate dehydrogenase deficiency. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, 3903-11	5.6	83
115	Pitfalls in genetic analysis of pheochromocytomas/paragangliomas-case report. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, 2321-6	5.6	7
114	H-RAS mutations are restricted to sporadic pheochromocytomas lacking specific clinical or pathological features: data from a multi-institutional series. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E1376-80	5.6	39
113	Integrative genetic, epigenetic and pathological analysis of paraganglioma reveals complex dysregulation of NOTCH signaling. <i>Acta Neuropathologica</i> , 2013 , 126, 575-94	14.3	22
112	Mitotane therapy in adrenocortical cancer induces CYP3A4 and inhibits 5E eductase, explaining the need for personalized glucocorticoid and androgen replacement. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2013 , 98, 161-71	5.6	104
	Endournology and Metabolism, 2010, 10111		
111	Diagnosis and Clinical Pictures 2013 , 35-53		1
111		4.2	4
	Diagnosis and Clinical Pictures 2013 , 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective,	4.2 5.6	
110	Diagnosis and Clinical Pictures 2013 , 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective, single-center study. <i>Head and Neck</i> , 2013 , 35, 1345-8 Detection of circulating tumor cells in patients with adrenocortical carcinoma: a monocentric	ŕ	4
110	Diagnosis and Clinical Pictures 2013, 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective, single-center study. Head and Neck, 2013, 35, 1345-8 Detection of circulating tumor cells in patients with adrenocortical carcinoma: a monocentric preliminary study. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 3731-8 Yeast model for evaluating the pathogenic significance of SDHB, SDHC and SDHD mutations in	5.6	30
110	Diagnosis and Clinical Pictures 2013, 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective, single-center study. Head and Neck, 2013, 35, 1345-8 Detection of circulating tumor cells in patients with adrenocortical carcinoma: a monocentric preliminary study. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 3731-8 Yeast model for evaluating the pathogenic significance of SDHB, SDHC and SDHD mutations in PHEO-PGL syndrome. Human Molecular Genetics, 2013, 22, 804-15 The reticulin algorithm for adrenocortical tumor diagnosis: a multicentric validation study on 245	5.6	4 30 22
110 109 108	Diagnosis and Clinical Pictures 2013, 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective, single-center study. Head and Neck, 2013, 35, 1345-8 Detection of circulating tumor cells in patients with adrenocortical carcinoma: a monocentric preliminary study. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 3731-8 Yeast model for evaluating the pathogenic significance of SDHB, SDHC and SDHD mutations in PHEO-PGL syndrome. Human Molecular Genetics, 2013, 22, 804-15 The reticulin algorithm for adrenocortical tumor diagnosis: a multicentric validation study on 245 unpublished cases. American Journal of Surgical Pathology, 2013, 37, 1433-40 Tumoral EPAS1 (HIF2A) mutations explain sporadic pheochromocytoma and paraganglioma in the	5.6 5.6 6.7	4 30 22 54
110 109 108 107	Diagnosis and Clinical Pictures 2013, 35-53 Feasibility and safety of minimal-incision thyroidectomy for Graves' disease: a prospective, single-center study. Head and Neck, 2013, 35, 1345-8 Detection of circulating tumor cells in patients with adrenocortical carcinoma: a monocentric preliminary study. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 3731-8 Yeast model for evaluating the pathogenic significance of SDHB, SDHC and SDHD mutations in PHEO-PGL syndrome. Human Molecular Genetics, 2013, 22, 804-15 The reticulin algorithm for adrenocortical tumor diagnosis: a multicentric validation study on 245 unpublished cases. American Journal of Surgical Pathology, 2013, 37, 1433-40 Tumoral EPAS1 (HIF2A) mutations explain sporadic pheochromocytoma and paraganglioma in the absence of erythrocytosis. Human Molecular Genetics, 2013, 22, 2169-76 Morphofunctional effects of mitotane on mitochondria in human adrenocortical cancer cells.	5.6 5.6 6.7 5.6	4 30 22 54 120

(2010-2012)

Plasma methoxytyramine: a novel biomarker of metastatic pheochromocytoma and paraganglioma in relation to established risk factors of tumour size, location and SDHB mutation status. <i>European Journal of Cancer</i> , 2012 , 48, 1739-49	7·5	228
Updated and new perspectives on diagnosis, prognosis, and therapy of malignant pheochromocytoma/paraganglioma. <i>Journal of Oncology</i> , 2012 , 2012, 872713	4.5	88
Genetic-clinical profile of subjects with apparently sporadic extra-adrenal paragangliomas. <i>Frontiers in Endocrinology</i> , 2012 , 3, 65	5.7	2
Mitochondrial function and content in pheochromocytoma/paraganglioma of succinate dehydrogenase mutation carriers. <i>Endocrine-Related Cancer</i> , 2012 , 19, 261-9	5.7	17
Head and neck paragangliomas: genetic spectrum and clinical variability in 79 consecutive patients. <i>Endocrine-Related Cancer</i> , 2012 , 19, 149-55	5.7	61
Oral lixivaptan effectively increases serum sodium concentrations in outpatients with euvolemic hyponatremia. <i>Kidney International</i> , 2012 , 82, 1215-22	9.9	27
Perioperative management of pheochromocytoma/paraganglioma: is there a state of the art?. <i>Hormone and Metabolic Research</i> , 2012 , 44, 373-8	3.1	23
MAX mutations cause hereditary and sporadic pheochromocytoma and paraganglioma. <i>Clinical Cancer Research</i> , 2012 , 18, 2828-37	12.9	226
Catecholamine metabolomic and secretory phenotypes in phaeochromocytoma. <i>Endocrine-Related Cancer</i> , 2011 , 18, 97-111	5.7	127
Exome sequencing identifies MAX mutations as a cause of hereditary pheochromocytoma. <i>Nature Genetics</i> , 2011 , 43, 663-7	36.3	409
Measurements of plasma methoxytyramine, normetanephrine, and metanephrine as discriminators of different hereditary forms of pheochromocytoma. <i>Clinical Chemistry</i> , 2011 , 57, 411-20	5.5	225
Age at diagnosis of pheochromocytoma differs according to catecholamine phenotype and tumor location. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2011 , 96, 375-84	5.6	65
Within-patient reproducibility of the aldosterone: renin ratio in primary aldosteronism. <i>Hypertension</i> , 2010 , 55, 83-9	8.5	54
Pheochromocytoma in rats with multiple endocrine neoplasia (MENX) shares gene expression patterns with human pheochromocytoma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 18493-8	11.5	32
Rosiglitazone impairs proliferation of human adrenocortical cancer: preclinical study in a xenograft mouse model. <i>Endocrine-Related Cancer</i> , 2010 , 17, 169-77	5.7	27
Role of the PPAR-Bystem in normal and tumoral pituitary corticotropic cells and adrenal cells. <i>Neuroendocrinology</i> , 2010 , 92 Suppl 1, 23-7	5.6	16
Progress in primary aldosteronism: present challenges and perspectives. <i>Hormone and Metabolic Research</i> , 2010 , 42, 374-81	3.1	42
Low sensitivity of glucagon provocative testing for diagnosis of pheochromocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 238-45	5.6	21
	in relation to established risk factors of tumour size, location and SDHB mutation status. European Journal of Cancer, 2012, 48, 1739-49 Journal of Cancer, 2012, 2012, 872713 Genetic-clinical profile of subjects with apparently sporadic extra-adrenal paragangliomas. Frontiers in Endocrinology, 2012, 3, 65 Mitochondrial function and content in pheochromocytoma/paraganglioma of succinate dehydrogenase mutation carriers. Endocrine-Related Cancer, 2012, 19, 261-9 Head and neck paragangliomas: genetic spectrum and clinical variability in 79 consecutive patients. Endocrine-Related Cancer, 2012, 19, 149-55 Oral lixivaptan effectively increases serum sodium concentrations in outpatients with euvolemic hyponatremia. Kidney International, 2012, 82, 1215-22 Perioperative management of pheochromocytoma/paraganglioma: is there a state of the art?. Hormone and Metabolic Research, 2012, 44, 373-8 MAX mutations cause hereditary and sporadic pheochromocytoma and paraganglioma. Clinical Cancer Research, 2012, 18, 2828-37 Catecholamine metabolomic and secretory phenotypes in phaeochromocytoma. Endocrine-Related Cancer, 2011, 18, 97-111 Exome sequencing identifies MAX mutations as a cause of hereditary pheochromocytoma. Nature Genetics, 2011, 43, 663-7 Measurements of plasma methoxytyramine, normetanephrine, and metanephrine as discriminators of different hereditary forms of pheochromocytoma. Clinical Chemistry, 2011, 57, 411-20 Age at diagnosis of pheochromocytoma differs according to catecholamine phenotype and tumor location. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 375-84 Within-patient reproducibility of the aldosterone: renin ratio in primary aldosteronism. Hypertension, 2010, 55, 83-9 Pheochromocytoma in rats with multiple endocrine neoplasia (MENX) shares gene expression patterns with human pheochromocytoma. Proceedings of the National Academy of Sciences of the Uniked Scates of	in relation to established risk factors of tumour size, location and SDHB mutation status. European Journal of Cancer, 2012, 48, 1739-49 Updated and new perspectives on diagnosis, prognosis, and therapy of malignant pheochromocytoma/paraganglioma. Journal of Oncology, 2012, 2012, 872713 Genetic-clinical profile of subjects with apparently sporadic extra-adrenal paragangliomas. Frontiers in Endocrinology, 2012, 3, 65 Mitochondrial function and content in pheochromocytoma/paraganglioma of succinate dehydrogenase mutation carriers. Endocrine-Related Cancer, 2012, 19, 261-9 Head and neck paragangliomas: genetic spectrum and clinical variability in 79 consecutive patients. Endocrine-Related Cancer, 2012, 19, 149-55 Oral lixivaptan effectively increases serum sodium concentrations in outpatients with euvolemic hyponatremia. Kidney international, 2012, 82, 1215-22 Perioperative management of pheochromocytoma/paraganglioma: is there a state of the art?. Hormone and Metabolic Research, 2012, 44, 373-8 MAX mutations cause hereditary and sporadic pheochromocytoma and paraganglioma. Clinical Cancer Research, 2012, 18, 2828-37 Catecholamine metabolomic and secretory phenotypes in phaeochromocytoma. Endocrine-Related Cancer, 2011, 18, 97-111 Exome sequencing identifies MAX mutations as a cause of hereditary pheochromocytoma. Nature Genetics, 2011, 43, 663-7 Measurements of plasma methoxytyramine, normetanephrine, and metanephrine as discriminators of different hereditary forms of pheochromocytoma. Clinical Chemistry, 2011, 57, 411-20 Age at diagnosis of sheochromocytoma differs according to catecholamine phenotype and tumor location. Journal of Clinical Endocrinology and Metabolism, 2011, 96, 375-84 Within-patient reproducibility of the aldosterone: renin ratio in primary aldosteronism. Hypertension, 2010, 55, 83-9 Pheochromocytoma in rats with multiple endocrine neoplasia (MENX) shares gene expression patterns with human pheochromocytoma. Proceedings of the National Academy of Sciences of the United States of A

84	Il dosaggio delle metanefrine nella diagnosi del feocromocitoma: vantaggi clinici e problematiche di laboratorio. <i>L Endocrinologo</i> , 2010 , 11, 65-74	О	1
83	Spectrum and prevalence of FP/TMEM127 gene mutations in pheochromocytomas and paragangliomas. <i>JAMA - Journal of the American Medical Association</i> , 2010 , 304, 2611-9	27.4	144
82	Incidental and metastatic adrenal masses. Seminars in Oncology, 2010, 37, 649-61	5.5	14
81	Seladin-1 expression is regulated by promoter methylation in adrenal cancer. <i>BMC Cancer</i> , 2010 , 10, 20	14.8	11
80	Clinically guided genetic screening in a large cohort of italian patients with pheochromocytomas and/or functional or nonfunctional paragangliomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009 , 94, 1541-7	5.6	227
79	Functional study in a yeast model of a novel succinate dehydrogenase subunit B gene germline missense mutation (C191Y) diagnosed in a patient affected by a glomus tumor. <i>Human Molecular Genetics</i> , 2009 , 18, 1860-8	5.6	50
78	Clinical aspects of SDHx-related pheochromocytoma and paraganglioma. <i>Endocrine-Related Cancer</i> , 2009 , 16, 391-400	5.7	101
77	Sindromi feocromocitoma/paraganglioma familiari. <i>L Endocrinologo</i> , 2009 , 10, 26-31	О	
76	Biochemistry, genetics and therapy of malignant pheochromocytomas. <i>Annales Dl</i> Endocrinologie, 2009 , 70, 166-7	1.7	3
75	An immunohistochemical procedure to detect patients with paraganglioma and phaeochromocytoma with germline SDHB, SDHC, or SDHD gene mutations: a retrospective and prospective analysis. <i>Lancet Oncology, The</i> , 2009 , 10, 764-71	21.7	405
74	Uncommon clinical presentations of pheochromocytoma and paraganglioma in two different patients affected by two distinct novel VHL germline mutations. <i>Clinical Endocrinology</i> , 2008 , 68, 762-8	3.4	20
73	The Y606C RET mutation causes a receptor gain of function. <i>Clinical Endocrinology</i> , 2008 , 69, 253-8	3.4	15
72	Differential expression of the regulated catecholamine secretory pathway in different hereditary forms of pheochromocytoma. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2008 , 295, E1223-33	6	47
71	Body mass index predicts plasma aldosterone concentrations in overweight-obese primary hypertensive patients. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2008 , 93, 2566-71	5.6	141
70	Rosiglitazone inhibits adrenocortical cancer cell proliferation by interfering with the IGF-IR intracellular signaling. <i>PPAR Research</i> , 2008 , 2008, 904041	4.3	37
69	Pheochromocytoma: recommendations for clinical practice from the First International Symposium. October 2005. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2007 , 3, 92-102		467
68	Comparison of the captopril and the saline infusion test for excluding aldosterone-producing adenoma. <i>Hypertension</i> , 2007 , 50, 424-31	8.5	120
67	A case of hyponatremia caused by central hypocortisolism. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2007 , 3, 369-75		7

(2005-2007)

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