Filomena Cetani

List of Publications by Year
in descending order

146	6,305	87723	74018
papers	citations	38 h-index	75 g-index

162	162	162	5284
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Chronic mucocutaneous candidiasis in APECED or thymoma patients correlates with autoimmunity to Th17-associated cytokines. Journal of Experimental Medicine, 2010, 207, 299-308.	4.2	593
2	Surgery or Surveillance for Mild Asymptomatic Primary Hyperparathyroidism: A Prospective, Randomized Clinical Trial. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 3114-3121.	1.8	347
3	Primary Hyperparathyroidism. New England Journal of Medicine, 2011, 365, 2389-2397.	13.9	312
4	Thyroid Ultrasonography Helps to Identify Patients with Diffuse Lymphocytic Thyroiditis who Are Prone to Develop Hypothyroidism*. Journal of Clinical Endocrinology and Metabolism, 1991, 72, 209-213.	1.8	267
5	Parathyroid Carcinoma. Journal of Bone and Mineral Research, 2008, 23, 1869-1880.	3.1	243
6	Genetic Analyses of theHRPT2Gene in Primary Hyperparathyroidism: Germline and Somatic Mutations in Familial and Sporadic Parathyroid Tumors. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 5583-5591.	1.8	234
7	Autoantibodies against Type I Interferons as an Additional Diagnostic Criterion for Autoimmune Polyendocrine Syndrome Type I. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 4389-4397.	1.8	176
8	A Novel Mutation of the Autoimmune Regulator Gene in an Italian Kindred with Autoimmune Polyendocrinopathy-Candidiasis-Ectodermal Dystrophy, Acting in a Dominant Fashion and Strongly Cosegregating with Hypothyroid Autoimmune Thyroiditis. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 4747-4752.	1.8	170
9	Morphometric Vertebral Fractures in Postmenopausal Women with Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 2306-2312.	1.8	170
10	Outcome of 309 patients with metastatic differentiated thyroid carcinoma treated with radioiodine. World Journal of Surgery, 1994, 18, 600-604.	0.8	164
11	Should parafibromin staining replace HRTP2 gene analysis as an additional tool for histologic diagnosis of parathyroid carcinoma?. European Journal of Endocrinology, 2007, 156, 547-554.	1.9	144
12	Functional characteristics of three new germline mutations of the thyrotropin receptor gene causing autosomal dominant toxic thyroid hyperplasia.. Journal of Clinical Endocrinology and Metabolism, 1996, 81, 547-554.	1.8	128
13	A Novel Mutation in the Upstream Open Reading Frame of the CDKN1B Gene Causes a MEN4 Phenotype. PLoS Genetics, 2013, 9, e1003350.	1.5	125
14	No Evidence for Mutations in the Calcium-Sensing Receptor Gene in Sporadic Parathyroid Adenomas. Journal of Bone and Mineral Research, 1999, 14, 878-882.	3.1	98
15	Update on parathyroid carcinoma. Journal of Endocrinological Investigation, 2016, 39, 595-606.	1.8	98
16	A quick intraoperative parathyroid hormone assay in the surgical management of patients with primary hyperparathyroidism: a study of 206 consecutive cases. European Journal of Endocrinology, 2002, 146, 783-788.	1.9	95
17	The microRNA cluster C19MC is deregulated in parathyroid tumours. Journal of Molecular Endocrinology, 2012, 49, 115-124.	1.1	89
18	Atypical parathyroid adenomas: challenging lesions in the differential diagnosis of endocrine tumors. Endocrine-Related Cancer, 2019, 26, R441-R464.	1.6	87

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19	Hyperfunctioning Thyroid Nodules in Toxic Multinodular Goiter Share Activating Thyrotropin Receptor Mutations with Solitary Toxic Adenoma¹. Journal of Clinical Endocrinology and Metabolism, 1998, 83, 492-498.	1.8	82
20	Genetic and Clinical Features of Multiple Endocrine Neoplasia Types 1 and 2. Journal of Oncology, 2012, 2012, 1-15.	0.6	79
21	TSH receptor and disease. Clinical Endocrinology, 1996, 44, 621-633.	1.2	78
22	Multiple endocrine neoplasia syndrome type 1: institution, management, and data analysis of a nationwide multicenter patient database. Endocrine, 2017, 58, 349-359.	1.1	77
23	CDC73 mutational status and loss of parafibromin in the outcome of parathyroid cancer. Endocrine Connections, 2013, 2, 186-195.	0.8	76
24	Differential effects of NaCl concentration on the constitutive activity of the thyrotropin and the luteinizing hormone/chorionic gonadotropin receptors. FEBS Letters, 1996, 378, 27-31.	1.3	75
25	Specific activation of the thyrotropin receptor by trypsin. Molecular and Cellular Endocrinology, 1996, 119, 161-168.	1.6	73
26	European expert consensus on practical management of specific aspects of parathyroid disorders in adults and in pregnancy: recommendations of the ESE Educational Program of Parathyroid Disorders (PARAT 2021). European Journal of Endocrinology, 2022, 186, R33-R63.	1.9	73
27	Parathyroid Expression of Calcium-Sensing Receptor Protein andin VivoParathyroid Hormone-Ca²⁺Set-Point in Patients with Primary Hyperparathyroidism¹. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 4789-4794.	1.8	70
28	Parathyroid carcinoma. Best Practice and Research in Clinical Endocrinology and Metabolism, 2018, 32, 877-889.	2.2	70
29	A reappraisal of the Rb1 gene abnormalities in the diagnosis of parathyroid cancer. Clinical Endocrinology, 2004, 60, 99-106.	1.2	67
30	Parathyroid Carcinoma. Frontiers of Hormone Research, 2019, 51, 63-76.	1.0	63
31	Unmet therapeutic, educational and scientific needs in parathyroid disorders: Consensus Statement from the first European Society of Endocrinology Workshop (PARAT). European Journal of Endocrinology, 2019, 181, P1-P19.	1.9	61
32	Parathyroid Expression of Calcium-Sensing Receptor Protein and in Vivo Parathyroid Hormone-Ca2+ Set-Point in Patients with Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 4789-4794.	1.8	61
33	Genetic analyses in familial isolated hyperparathyroidism: implication for clinical assessment and surgical management. Clinical Endocrinology, 2006, 64, 146-152.	1.2	59
34	Î²-catenin activation is not involved in sporadic parathyroid carcinomas and adenomas. Endocrine-Related Cancer, 2010, 17, 1-6.	1.6	54
35	Persistent secondary hyperparathyroidism and vertebral fractures in kidney transplantation: Role of calcium-sensing receptor polymorphisms and vitamin D deficiency. Journal of Bone and Mineral Research, 2010, 25, 841-848.	3.1	51
36	Functional characterization of a CDKN1B mutation in a Sardinian kindred with multiple endocrine neoplasia type 4. Endocrine Connections, 2015, 4, 1-8.	0.8	44

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37	Bone disease in primary hyperparathyrodism. Therapeutic Advances in Musculoskeletal Disease, 2012, 4, 357-368.	1.2	43
38	Impact of vitamin D deficiency on the clinical and biochemical phenotype in women with sporadic primary hyperparathyroidism. Endocrine, 2017, 55, 256-265.	1.1	42
39	Loss of p27 expression is associated with MEN1 gene mutations in sporadic parathyroid adenomas. Endocrine, 2017, 55, 386-397.	1.1	42
40	Efficacy and safety of long-term management of patients with chronic post-surgical hypoparathyroidism. Journal of Endocrinological Investigation, 2018, 41, 1221-1226.	1.8	40
41	Different somatic alterations of the HRPT2 gene in a patient with recurrent sporadic primary hyperparathyroidism carrying an HRPT2 germline mutation. Endocrine-Related Cancer, 2007, 14, 493-499.	1.6	38
42	Normocalcemic primary hyperparathyroidism: a survey in a small village of Southern Italy. Endocrine Connections, 2015, 4, 172-178.	0.8	37
43	Familial and Hereditary Forms of Primary Hyperparathyroidism. Frontiers of Hormone Research, 2019, 51, 40-51.	1.0	36
44	Mapping thyroid peroxidase epitopes using recombinant protein fragments. European Journal of Endocrinology, 1995, 132, 53-61.	1.9	35
45	A Novel Mutation of the Autoimmune Regulator Gene in an Italian Kindred with Autoimmune Polyendocrinopathy-Candidiasis-Ectodermal Dystrophy, Acting in a Dominant Fashion and Strongly Cosegregating with Hypothyroid Autoimmune Thyroiditis. , 0, .		33
46	Genetic analysis of the MEN1 gene and HPRT2 locus in two Italian kindreds with familial isolated hyperparathyroidism. Clinical Endocrinology, 2002, 56, 457-464.	1.2	32
47	Cinacalcet in the management of primary hyperparathyroidism: post marketing experience of an Italian multicentre group. Clinical Endocrinology, 2013, 79, 20-26.	1.2	32
48	Clinical presentation and management of patients with primary hyperparathyroidism in Italy. Journal of Endocrinological Investigation, 2018, 41, 1339-1348.	1.8	32
49	Calcium-sensing receptor gene polymorphisms in primary hyperparathyroidism. Journal of Endocrinological Investigation, 2002, 25, 614-619.	1.8	31
50	Mutational and large deletion study of genes implicated in hereditary forms of primary hyperparathyroidism and correlation with clinical features. PLoS ONE, 2017, 12, e0186485.	1.1	31
51	Aryl Hydrocarbon Receptor Interacting Protein (AIP) Mutations Occur Rarely in Sporadic Parathyroid Adenomas. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 2800-2810.	1.8	29
52	Genetic analysis of the TSH receptor gene in differentiated human thyroid carcinomas. Journal of Endocrinological Investigation, 1999, 22, 273-278.	1.8	27
53	Two Italian kindreds with familial hypocalciuric hypercalcaemia caused by loss-of-function mutations in the calcium-sensing receptor (CaR) gene: functional characterization of a novel CaR missense mutation. Clinical Endocrinology, 2003, 58, 199-206.	1.2	27
54	Activating Antibodies to The Calcium-sensing Receptor in Immunotherapy-induced Hypoparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1581-1588.	1.8	27

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55	Evaluation of formalin-fixed paraffin-embedded tissues in the proteomic analysis of parathyroid glands. Proteome Science, 2011, 9, 29.	0.7	25
56	Expression, function, and regulation of the embryonic transcription factor TBX1 in parathyroid tumors. Laboratory Investigation, 2017, 97, 1488-1499.	1.7	25
57	Cinacalcet efficacy in patients with moderately severe primary hyperparathyroidism according to the European Medicine Agency prescription labeling. Journal of Endocrinological Investigation, 2012, 35, 655-60.	1.8	25
58	A proteomic approach to study parathyroid glands. Molecular BioSystems, 2011, 7, 687-699.	2.9	24
59	HypoparaNet: A Database of Chronic Hypoparathyroidism Based on Expert Medical-Surgical Centers in Italy. Calcified Tissue International, 2018, 103, 151-163.	1.5	23
60	Re: Familial hyperparathyroidism: Surgical outcome after 30 years of follow-up in three families with germline HRPT2 mutations. Surgery, 2008, 144, 839-840.	1.0	22
61	The Sulfaphenazole-Sensitive Pathway Acts as a Compensatory Mechanism for Impaired Nitric Oxide Availability in Patients with Primary Hyperparathyroidism. Effect of Surgical Treatment. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 920-927.	1.8	22
62	Patient with monoclonal gammopathy, thyrotoxicosis, pretibial myxedema and thyroid-associated ophthalmopathy; demonstration of direct binding of autoantibodies to the thyrotropin receptor. European Journal of Endocrinology, 1996, 134, 97-103.	1.9	21
63	Hyperparathyroidism 2 gene (HRPT2, CDC73) and parafibromin studies in two patients with primary hyperparathyroidism and uncertain pathological assessment. Journal of Endocrinological Investigation, 2008, 31, 900-904.	1.8	21
64	Multiple endocrine neoplasia type 1: analysis of germline MEN1 mutations in the Italian multicenter MEN1 patient database. Endocrine, 2018, 62, 215-233.	1.1	21
65	Assays of TSH-Receptor Antibodies in 576 Patients with Various Thyroid Disorders: Their Incidence, Significance and Clinical Usefulness. Autoimmunity, 1989, 3, 103-112.	1.2	20
66	Parathyroid carcinoma: a clinical and genetic perspective. Minerva Endocrinology, 2018, 43, 144-155.	0.6	20
67	A new mutation of the MEN1 gene in an italian kindred with multiple endocrine neoplasia type 1. European Journal of Endocrinology, 1999, 140, 429-433.	1.9	19
68	High-intensity focused ultrasound as an alternative to the surgical approach in primary hyperparathyroidism: A preliminary experience. Journal of Endocrinological Investigation, 2011, 34, 655-659.	1.8	19
69	Non-surgical management of primary hyperparathyroidism. Best Practice and Research in Clinical Endocrinology and Metabolism, 2018, 32, 821-835.	2.2	19
70	Increased Prevalence of theGCM2Polymorphism, Y282D, in Primary Hyperparathyroidism: Analysis of Three Italian Cohorts. Journal of Clinical Endocrinology and Metabolism, 2014, 99, E2794-E2798.	1.8	18
71	A nonfunctioning parathyroid carcinoma misdiagnosed as a follicular thyroid nodule. World Journal of Surgical Oncology, 2015, 13, 270.	0.8	18
72	Whole exome sequencing in familial isolated primary hyperparathyroidism. Journal of Endocrinological Investigation, 2020, 43, 231-245.	1.8	18

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73	The aberrantly expressed miR-372 partly impairs sensitivity to apoptosis in parathyroid tumor cells. Endocrine-Related Cancer, 2018, 25, 761-771.	1.6	17
74	Hereditary Hypercalcemia Caused by a Homozygous Pathogenic Variant in theCYP24A1Gene: A Case Report and Review of the Literature. Case Reports in Endocrinology, 2019, 2019, 1-7.	0.2	17
75	Hypercalcemia due to CYP24A1 mutations: a systematic descriptive review. European Journal of Endocrinology, 2022, 186, 137-149.	1.9	17
76	Calcium-sensing receptor gene polymorphism is not associated with bone mineral density in Italian postmenopausal women. European Journal of Endocrinology, 2003, 148, 603-607.	1.9	16
77	Spontaneous short-term remission of primary hyperparathyroidism from infarction of a parathyroid adenoma. Journal of Endocrinological Investigation, 2004, 27, 687-690.	1.8	16
78	Novel association of MEN1 gene mutations with parathyroid carcinoma. Oncology Letters, 2017, 14, 23-30.	0.8	16
79	Hypercalciuria: its value as a predictive risk factor for nephrolithiasis in asymptomatic primary hyperparathyroidism?. Journal of Endocrinological Investigation, 2020, 43, 677-682.	1.8	16
80	Oncogenic mutations in thyroid adenoma: methodological criteria. European Journal of Endocrinology, 1996, 135, 444-446.	1.9	15
81	Vitamin D measurement and effect on outcome in a cohort of patients with heart failure. Endocrine Connections, 2018, 7, 957-964.	0.8	15
82	Six novelMEN1 gene mutations in sporadic parathyroid tumors. Human Mutation, 2000, 16, 445-445.	1.1	14
83	Gene expression profile in metastatic and non-metastatic parathyroid carcinoma. Endocrine-Related Cancer, 2021, 28, 111-134.	1.6	14
84	Functional Characteristics of a Variant Thyrotropin Receptor. FEBS Journal, 1996, 238, 490-494.	0.2	13
85	Familial Hypocalciuric Hypercalcemia in a Woman with Metastatic Breast Cancer: A Case Report of Mistaken Identity. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 5132-5136.	1.8	13
86	The impact of thyroidectomy on parathyroid glands: A biochemical and clinical profile. Journal of Endocrinological Investigation, 2007, 30, 666-671.	1.8	13
87	Molecular pathogenesis of primary hyperparathyroidism. Journal of Endocrinological Investigation, 2011, 34, 35-9.	1.8	13
88	Post-surgical follow-up of differentiated thyroid cancer. Journal of Endocrinological Investigation, 1995, 18, 165-166.	1.8	12
89	First evidence of <scp>TRPV</scp>5 and <scp>TRPV</scp>6 channels in human parathyroid glands: possible involvement in neoplastic transformation. Journal of Cellular and Molecular Medicine, 2014, 18, 1944-1952.	1.6	12
90	Diagnostic, therapeutic and healthcare management protocols in parathyroid surgery: II Consensus Conference of the Italian Association of Endocrine Surgery Units (U.E.C. CLUB). Journal of Endocrinological Investigation, 2014, 37, 149-165.	1.8	12

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91	Hypomagnesuria is Associated With Nephrolithiasis in Patients With Asymptomatic Primary Hyperparathyroidism. Journal of Clinical Endocrinology and Metabolism, 2020, 105, e2789-e2795.	1.8	12
92	Renal complications and quality of life in postsurgical hypoparathyroidism: a caseâ€“control study. Journal of Endocrinological Investigation, 2022, 45, 573-582.	1.8	12
93	Novel Glial Cells Missing-2 (GCM2) variants in parathyroid disorders. European Journal of Endocrinology, 2022, 186, 351-366.	1.9	12
94	Rare diseases in clinical endocrinology: a taxonomic classification system. Journal of Endocrinological Investigation, 2015, 38, 193-259.	1.8	11
95	The Oncosuppressors <scp>MEN1</scp> and <scp>CDC73</scp> Are Involved in <scp>lncRNA</scp> Deregulation in Human Parathyroid Tumors. Journal of Bone and Mineral Research, 2020, 35, 2423-2431.	3.1	11
96	Do the Heterozygous Carriers of a CYP24A1 Mutation Display a Different Biochemical Phenotype Than Wild Types?. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 708-717.	1.8	11
97	Long-term Efficacy and Safety of Rifampin in the Treatment of a Patient Carrying a CYP24A1 Loss-of-Function Variant. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e3159-e3166.	1.8	11
98	The thyroperoxidase doublet is not produced by alternative splicing. Molecular and Cellular Endocrinology, 1995, 115, 125-132.	1.6	10
99	Parathyroid Carcinoma and Adenoma Co-existing in One Patient: Case Report and Comparative Proteomic Analysis. Cancer Genomics and Proteomics, 2021, 18, 781-796.	1.0	9
100	Update on the use of cinacalcet in the management of primary hyperparathyroidism. Journal of Endocrinological Investigation, 2012, 35, 90-5.	1.8	9
101	MEN1 gene alterations do not correlate with the phenotype of sporadic primary hyperparathyroidism. Journal of Endocrinological Investigation, 2002, 25, 508-512.	1.8	8
102	Microvessel Density in Human Normal and Neoplastic Parathyroids. Endocrine Pathology, 2006, 17, 175-182.	5.2	8
103	Parafibromin as a Tool for the Diagnosis of Parathyroid Tumors. Advances in Anatomic Pathology, 2008, 15, 179.	2.4	8
104	Identification and functional characterization of loss-of-function mutations of the calcium-sensing receptor in four Italian kindreds with familial hypocalciuric hypercalcemia. European Journal of Endocrinology, 2009, 160, 481-489.	1.9	8
105	Time for Revival of Bone Biopsy with Histomorphometric Analysis in Chronic Kidney Disease (CKD): Moving from Skepticism to Pragmatism. Nutrients, 2022, 14, 1742.	1.7	8
106	TSH receptor antibodies do not alter the function of gonadotropin receptors stably expressed in eukaryotic cells. European Journal of Endocrinology, 2004, 150, 381-387.	1.9	7
107	Development of an algorithm to predict serum vitamin D levels using a simple questionnaire based on sunlight exposure. Endocrine, 2017, 55, 85-92.	1.1	7
108	Do Patients With Atypical Parathyroid Adenoma Need Close Follow-up?. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e4565-e4579.	1.8	7

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109	Serum calcium levels are associated with cognitive function in hypoparathyroidism: a neuropsychological and biochemical study in an Italian cohort of patients with chronic post-surgical hypoparathyroidism. Journal of Endocrinological Investigation, 2022, 45, 1909-1918.	1.8	7
110	Impaired GH secretion to provocative stimuli in two families with hypocalciuric hypercalcaemia. Clinical Endocrinology, 2003, 59, 604-606.	1.2	6
111	Cinacalcet in the management of primary hyperparathyroidism. Expert Review of Endocrinology and Metabolism, 2012, 7, 45-53.	1.2	6
112	Parathyroid Carcinoma. , 2015, , 409-421.		6
113	A large functioning parathyroid lipoadenoma. Endocrine, 2016, 53, 615-616.	1.1	6
114	Clinical profile of juvenile primary hyperparathyroidism: a prospective study. Endocrine, 2018, 59, 344-352.	1.1	6
115	Pseudohypoparathyroidism: Focus on Cerebral and Renal Calcifications. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3005-e3020.	1.8	6
116	Six novel MEN1 gene mutations in sporadic parathyroid tumors Communicated by: Richard G.H. Cotton Online Citation: Human Mutation, Mutation in Brief #373 (2000) Online http://journals.wiley.com/1059-7794/pdf/mutation/373.pdf. Human Mutation, 2000, 16, 445.	1.1	6
117	The Long Non-Coding BC200 Is a Novel Circulating Biomarker of Parathyroid Carcinoma. Frontiers in Endocrinology, 2022, 13, 869006.	1.5	6
118	Functioning glucagonoma associated with primary hyperparathyroidism: multiple endocrine neoplasia type 1 or incidental association?. BMC Cancer, 2012, 12, 614.	1.1	5
119	The use of positron emission tomography with 11C-methionine in patients with primary hyperparathyroidism. Endocrine, 2013, 43, 251-252.	1.1	5
120	Effect of neridronate in osteopenic patients after heart, liver or lung transplant: a multicenter, randomized, double-blind, placebo-controlled study. Panminerva Medica, 2021, 63, 214-223.	0.2	5
121	Hypoparathyroidism and pseudohypoparathyroidism in pregnancy: an Italian retrospective observational study. Orphanet Journal of Rare Diseases, 2021, 16, 421.	1.2	5
122	Parathyroid Carcinoma and Ectopic Secretion of Parathyroid hormone. Endocrinology and Metabolism Clinics of North America, 2021, 50, 683-709.	1.2	5
123	Multicenter retro-prospective observational study on chronic hypoparathyroidism and rhPTH (1â€“84) treatment. Journal of Endocrinological Investigation, 2022, 45, 1653-1662.	1.8	5
124	A novel germline mutation of MEN 1 gene in a patient with acromegaly and multiple endocrine tumors. Journal of Endocrinological Investigation, 2004, 27, 577-582.	1.8	4
125	HRPT2gene analysis and the diagnosis of parathyroid carcinoma. Expert Review of Endocrinology and Metabolism, 2008, 3, 377-389.	1.2	4
126	A novel mutation in the calcium-sensing receptor in a French family with familial hypocalciuric hypercalcaemia. European Journal of Endocrinology, 2011, 165, 359-363.	1.9	4

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127	Incidental occurrence of metastatic medullary thyroid carcinoma in a patient with multiple endocrine neoplasia type 1 carrying germline MEN1 and somatic RET mutations. Journal of Surgical Oncology, 2017, 116, 1197-1199.	0.8	4
128	Concomitant Primary Hyperparathyroidism in Patients with Multiple Myeloma: A Possible Link?. Acta Haematologica, 2021, 144, 302-307.	0.7	4
129	Familial hypocalciuric hypercalcemia type 1 due to a novel homozygous mutation of the calcium-sensing receptor gene. Journal of Endocrinological Investigation, 2017, 40, 1271-1272.	1.8	4
130	A New MEN2 Syndrome with Clinical Features of Both MEN2A and MEN2B Associated with a New RET Germline Deletion. Case Reports in Endocrinology, 2020, 2020, 1-7.	0.2	3
131	Day case parathyroidectomy: is this the right way for the patients?. Gland Surgery, 2020, 9, S6-S13.	0.5	3
132	Remission of Primary Hyperparathyroidism Following Fine-NeedleÂAspiration Biopsy: A Case Report and Review ofÂtheÂLiterature. AACE Clinical Case Reports, 2021, 7, 75-79.	0.4	3
133	The reduction of bone mineral density in postmenopausal women with primary hyperparathyroidism is higher in the presence of concomitant GH secretion impairment. European Journal of Endocrinology, 2006, 155, 41-45.	1.9	2
134	Acute severe primary hyperparathyroidism: spontaneous remission after 2Âyears follow-up. Journal of Endocrinological Investigation, 2019, 42, 243-244.	1.8	2
135	A severe inactivating PTH/PTHrP signaling disorder type 2 in a patient carrying a novel large deletion of the GNAS gene: a case report and review of the literature. Endocrine, 2020, 67, 466-472.	1.1	2
136	Parathyroid Carcinoma. , 2009, , 321-333.		2
137	Ipocalcemie. L Endocrinologo, 2002, 3, 3-14.	0.0	0
138	Terapia medica dellâ€™iperparatiroidismo primario. L Endocrinologo, 2008, 9, 54-61.	0.0	0
139	Somatostatin Analogues do not Aff ect Calcium Metabolism in Patients with Acromegaly and Primary Hyperparathyroidism due to MEN 1-Like Syndrome. Hormone and Metabolic Research, 2011, 43, e1-e1.	0.7	0
140	Somatostatin Analogues do not Affect Calcium Metabolism in Patients with Acromegaly and Primary Hyperparathyroidism due to MEN 1-Like Syndrome. Hormone and Metabolic Research, 2011, 43, 126-129.	0.7	0
141	A patient with MEN1 and endâ€‘stage chronic kidney disease due to Alport syndrome: Decision making on the eligibility of transplantation. Molecular and Clinical Oncology, 2018, 8, 449-452.	0.4	0
142	Acute pancreatitis as the first manifestation in a young boy with primary hyperparathyroidism. Endocrine, 2018, 62, 267-268.	1.1	0
143	Thyrotoxicosis; Treatment. , 2018, , 673-679.		0
144	Late-onset postsurgical hypoparathyroidism following parathyroidectomy for recurrent primary hyperparathyroidism: a case report and literature review. Endocrine, 2020, 69, 402-409.	1.1	0

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145	Thyrotoxicosis; Systemic Manifestations. , 2018, , 665-672.		0
146	Hypercalcemia. Endocrinology and Metabolism Clinics of North America, 2021, 50, xv-xvi.	1.2	0