

Alberto Falchetti

List of Publications by Year in descending order

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163
papers

10,197
citations

50276

46
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34986

98
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194
all docs

194
docs citations

194
times ranked

7524
citing authors

#	ARTICLE	IF	CITATIONS
1	CONSENSUS: Guidelines for Diagnosis and Therapy of MEN Type 1 and Type 2. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5658-5671.	3.6	1,782
2	TNM staging of foregut (neuro)endocrine tumors: a consensus proposal including a grading system. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2006, 449, 395-401.	2.8	1,403
3	CONSENSUS: Guidelines for Diagnosis and Therapy of MEN Type 1 and Type 2. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 5658-5671.	3.6	574
4	Clonality of Parathyroid Tumors in Familial Multiple Endocrine Neoplasia Type 1. New England Journal of Medicine, 1989, 321, 213-218.	27.0	315
5	Gastrinoma (Duodenal and Pancreatic). Neuroendocrinology, 2006, 84, 173-182.	2.5	268
6	Well-Differentiated Pancreatic Tumor/Carcinoma: Insulinoma. Neuroendocrinology, 2006, 84, 183-188.	2.5	248
7	Well-Differentiated Pancreatic Nonfunctioning Tumors/Carcinoma. Neuroendocrinology, 2006, 84, 196-211.	2.5	241
8	Evidence of a linkage disequilibrium between polymorphisms in the human estrogen receptor alpha gene and their relationship to bone mass variation in postmenopausal Italian women. Human Molecular Genetics, 2000, 9, 2043-2050.	2.9	164
9	Thymic Neuroendocrine Carcinoma (Carcinoid) in Multiple Endocrine Neoplasia Type 1 Syndrome: The Italian Series. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 2603-2609.	3.6	160
10	Large-Scale Evidence for the Effect of the COL1A1 Sp1 Polymorphism on Osteoporosis Outcomes: The GENOMOS Study. PLoS Medicine, 2006, 3, e90.	8.4	160
11	Genome-wide association identifies three new susceptibility loci for Paget's disease of bone. Nature Genetics, 2011, 43, 685-689.	21.4	158
12	A Polymorphic CYP19 TTTA Repeat Influences Aromatase Activity and Estrogen Levels in Elderly Men: Effects on Bone Metabolism. Journal of Clinical Endocrinology and Metabolism, 2004, 89, 2803-2810.	3.6	151
13	Influence of Calcium-Sensing Receptor Gene on Urinary Calcium Excretion in Stone-Forming Patients. Journal of the American Society of Nephrology: JASN, 2002, 13, 2517-2523.	6.1	135
14	Well-Differentiated Gastric Tumors/Carcinomas. Neuroendocrinology, 2006, 84, 158-164.	2.5	133
15	Pancreatectomy in Multiple Endocrine Neoplasia Type 1-Related Gastrinomas and Pancreatic Endocrine Neoplasias. Annals of Surgery, 2006, 244, 61-70.	4.2	131
16	Multiple endocrine neoplasia type 1. Orphanet Journal of Rare Diseases, 2006, 1, 38.	2.7	129
17	Rare Functioning Pancreatic Endocrine Tumors. Neuroendocrinology, 2006, 84, 189-195.	2.5	124
18	Genetics of Osteoporosis. , 1999, 20, 788-804.		115

#	ARTICLE	IF	CITATIONS
19	Multiple endocrine neoplasia type 2 syndromes (MEN 2): results from the ItaMEN network analysis on the prevalence of different genotypes and phenotypes. <i>European Journal of Endocrinology</i> , 2010, 163, 301-308.	3.7	111
20	Vitamin D Status and SARS-CoV-2 Infection and COVID-19 Clinical Outcomes. <i>Frontiers in Public Health</i> , 2021, 9, 736665.	2.7	108
21	Poorly Differentiated Carcinomas of the Foregut (Gastric, Duodenal and Pancreatic). <i>Neuroendocrinology</i> , 2006, 84, 212-215.	2.5	106
22	Italian Association of Clinical Endocrinologists (AME) and Italian Chapter of the American Association of Clinical Endocrinologists (AACE) Position Statement: Clinical Management of Vitamin D Deficiency in Adults. <i>Nutrients</i> , 2018, 10, 546.	4.1	103
23	Aggressive Forms of Gastric Neuroendocrine Tumors in Multiple Endocrine Neoplasia Type I. <i>American Journal of Surgical Pathology</i> , 1997, 21, 1075-1082.	3.7	102
24	Progression of uremic hyperparathyroidism involves allelic loss on chromosome 11.. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1993, 76, 139-144.	3.6	95
25	<i>Fok</i> Polymorphism at Translation Initiation Site of the Vitamin D Receptor Gene Predicts Bone Mineral Density and Vertebral Fractures in Postmenopausal Italian Women. <i>Journal of Bone and Mineral Research</i> , 1999, 14, 1379-1386.	2.8	94
26	Polymorphism of the Aromatase Gene in Postmenopausal Italian Women: Distribution and Correlation with Bone Mass and Fracture Risk1. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 2263-2269.	3.6	93
27	Large-scale analysis of association between polymorphisms in the transforming growth factor beta 1 gene (TGFB1) and osteoporosis: The GENOMOS study. <i>Bone</i> , 2008, 42, 969-981.	2.9	91
28	Polymorphisms of the Calcitonin Receptor Gene Are Associated with Bone Mineral Density in Postmenopausal Italian Women. <i>Biochemical and Biophysical Research Communications</i> , 1998, 248, 190-195.	2.1	90
29	ERÎ ² is a potent inhibitor of cell proliferation in the HCT8 human colon cancer cell line through regulation of cell cycle components. <i>Endocrine-Related Cancer</i> , 2005, 12, 455-469.	3.1	90
30	Multiple endocrine neoplasia type 1 (MEN1): Not only inherited endocrine tumors. <i>Genetics in Medicine</i> , 2009, 11, 825-835.	2.4	89
31	Progression of uremic hyperparathyroidism involves allelic loss on chromosome 11. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1993, 76, 139-144.	3.6	86
32	Genetics of osteoporosis: role of steroid hormone receptor gene polymorphisms. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2002, 81, 1-24.	2.5	83
33	Is Total Parathyroidectomy the Treatment of Choice for Hyperparathyroidism in Multiple Endocrine Neoplasia Type 1?. <i>Annals of Surgery</i> , 2007, 246, 1075-1082.	4.2	78
34	Two Novel Mutations at Exon 8 of the Sequestosome 1 (SQSTM1) Gene in an Italian Series of Patients Affected by Paget's Disease of Bone (PDB). <i>Journal of Bone and Mineral Research</i> , 2004, 19, 1013-1017.	2.8	74
35	Polymorphism of the Aromatase Gene in Postmenopausal Italian Women: Distribution and Correlation with Bone Mass and Fracture Risk. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2001, 86, 2263-2269.	3.6	74
36	Well-Differentiated Duodenal Tumor/Carcinoma (Excluding Gastrinomas). <i>Neuroendocrinology</i> , 2006, 84, 165-172.	2.5	70

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37	Multiple endocrine neoplasia type 2. Orphanet Journal of Rare Diseases, 2006, 1, 45.	2.7	67
38	Genetics of multiple endocrine neoplasia type 1 syndrome: what's new and what's old. F1000Research, 2017, 6, 73.	1.6	62
39	Paget's Disease of Bone. Calcified Tissue International, 2019, 104, 483-500.	3.1	59
40	Relaxin influences growth, differentiation and cell-cell adhesion of human breast-cancer cells in culture. International Journal of Cancer, 1994, 57, 129-134.	5.1	58
41	Multiple Endocrine Neoplasia Type 1. Frontiers of Hormone Research, 2013, 41, 1-15.	1.0	55
42	Pathophysiology and Management of Type 2 Diabetes Mellitus Bone Fragility. Journal of Diabetes Research, 2020, 2020, 1-18.	2.3	55
43	Azidothymidine Induces Apoptosis and Inhibits Cell Growth and Telomerase Activity of Human Parathyroid Cancer Cells in Culture. Journal of Bone and Mineral Research, 2005, 20, 410-418.	2.8	54
44	Modulatory effect of farnesyl pyrophosphate synthase (<i>FDPS</i>) <i>rs2297480</i> polymorphism on the response to long-term amino-bisphosphonate treatment in postmenopausal osteoporosis. Current Medical Research and Opinion, 2008, 24, 2609-2615.	1.9	52
45	Production of basic fibroblast growth factor by gastric carcinoid tumors and their putative cells of origin. Human Pathology, 1994, 25, 175-180.	2.0	51
46	Allelic Variants of Human Calcitonin Receptor: Distribution and Association with Bone Mass in Postmenopausal Italian Women. Biochemical and Biophysical Research Communications, 1998, 245, 622-626.	2.1	51
47	Protective Effect of Denosumab on Bone in Older Women with Primary Hyperparathyroidism. Journal of the American Geriatrics Society, 2018, 66, 518-524.	2.6	51
48	Characterization of a Non-UBA Domain Missense Mutation of Sequestosome 1 (SQSTM1) in Paget's Disease of Bone. Journal of Bone and Mineral Research, 2009, 24, 632-642.	2.8	48
49	Surgery for gastroenteropancreatic tumours in multiple endocrine neoplasia type 1: review and personal experience. Journal of Internal Medicine, 2005, 257, 38-49.	6.0	47
50	Characterization and function of the receptor for IGF-I in human preosteoclastic cells. Bone, 1996, 18, 269-276.	2.9	46
51	Genetic Epidemiology of Paget's Disease of Bone in Italy: sequestosome1/p62 Gene Mutational Test and Haplotype Analysis at 5q35 in a Large Representative Series of Sporadic and Familial Italian Cases of Paget's Disease of Bone. Calcified Tissue International, 2009, 84, 20-37.	3.1	46
52	Pharmacogenetics of bisphosphonate-associated osteonecrosis of the jaw. Frontiers in Bioscience - Elite, 2011, E3, 364-370.	1.8	44
53	MEN1 gene mutation analysis in Italian patients with multiple endocrine neoplasia type 1. European Journal of Endocrinology, 2000, 142, 131-137.	3.7	42
54	Surgical Approach in Hereditary Hyperparathyroidism. Endocrine Journal, 2009, 56, 827-841.	1.6	41

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55	DIAGNOSIS OF ENDOCRINE DISEASE: Evaluation of bone fragility in endocrine disorders. <i>European Journal of Endocrinology</i> , 2019, 180, R213-R232.	3.7	40
56	A patient with MEN1-associated hyperparathyroidism, responsive to cinacalcet. <i>Nature Clinical Practice Endocrinology and Metabolism</i> , 2008, 4, 351-357.	2.8	38
57	Calcium Bioavailability From a Calcium-Rich Mineral Water, With Some Observations on Method. <i>Journal of Clinical Gastroenterology</i> , 2004, 38, 761-766.	2.2	37
58	Multiple endocrine neoplasms. <i>Best Practice and Research in Clinical Rheumatology</i> , 2008, 22, 149-163.	3.3	37
59	The Regulatory Network Menin-MicroRNA 26a As a Possible Target for RNA-Based Therapy of Bone Diseases. <i>Nucleic Acid Therapeutics</i> , 2012, 22, 103-108.	3.6	35
60	Management of familial hyperparathyroidism syndromes: MEN1, MEN2, MEN4, HPT-Jaw tumour, Familial isolated hyperparathyroidism, FHH, and neonatal severe hyperparathyroidism. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2018, 32, 861-875.	4.7	35
61	Hypovitaminosis D: Is It Time to Consider the Use of Calcifediol?. <i>Nutrients</i> , 2019, 11, 1016.	4.1	34
62	Genetic markers of osteoarticular disorders: facts and hopes. <i>Arthritis Research</i> , 2001, 3, 270.	2.0	31
63	A Novel Recessive Mutation of Fibroblast Growth Factor-23 in Tumoral Calcinosis. <i>Journal of Bone and Joint Surgery - Series A</i> , 2009, 91, 1190-1198.	3.0	31
64	Echocardiographic Evaluation of Children with and without Family History of Essential Hypertension. <i>American Journal of Hypertension</i> , 1988, 1, 305-308.	2.0	30
65	RFLP Analysis of Human Chromosome 11 Region q13 in Multiple Symmetrical Lipomatosis and Multiple Endocrine Neoplasia Type 1-Associated Lipomas. <i>Biochemical and Biophysical Research Communications</i> , 1995, 207, 363-368.	2.1	29
66	DNA-based test: when and why to apply it to primary hyperparathyroidism clinical phenotypes. <i>Journal of Internal Medicine</i> , 2009, 266, 69-83.	6.0	27
67	Homozygotes for the autosomal dominant neoplasia syndrome (MEN1). <i>American Journal of Human Genetics</i> , 1993, 53, 1167-72.	6.2	27
68	Management of Osteoporosis in Men: A Narrative Review. <i>International Journal of Molecular Sciences</i> , 2021, 22, 13640.	4.1	26
69	Treatment options for glucocorticoid-induced osteoporosis. <i>Expert Opinion on Pharmacotherapy</i> , 2020, 21, 721-732.	1.8	25
70	Steroid 5 α -reductase 2 deficiency: virilization in early infancy may be due to partial function of mutant enzyme. <i>Clinical Endocrinology</i> , 1996, 44, 477-482.	2.4	24
71	Microencapsulation of Human Parathyroid Cells: An <i>in Vitro</i> Study. <i>Journal of Surgical Research</i> , 2001, 96, 81-89.	1.6	24
72	Calcium citrate: from biochemistry and physiology to clinical applications. <i>Reviews in Endocrine and Metabolic Disorders</i> , 2019, 20, 353-364.	5.7	24

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73	Updates in epidemiology, pathophysiology and management strategies of glucocorticoid-induced osteoporosis. <i>Expert Review of Endocrinology and Metabolism</i> , 2020, 15, 283-298.	2.4	24
74	Telomerase Repeat Amplification Protocol (TRAP): A New Molecular Marker for Parathyroid Carcinoma. <i>Biochemical and Biophysical Research Communications</i> , 1999, 265, 252-255.	2.1	23
75	Thyroid Cancer: Current Molecular Perspectives. <i>Journal of Oncology</i> , 2010, 2010, 1-17.	1.3	23
76	Phenotypes Associated With MEN1 Syndrome: A Focus on Genotype-Phenotype Correlations. <i>Frontiers in Endocrinology</i> , 2020, 11, 591501.	3.5	23
77	Segregation of a M404V mutation of the p62/sequestosome 1 (p62/SQSTM1) gene with polyostotic Paget's disease of bone in an Italian family. <i>Arthritis Research and Therapy</i> , 2005, 7, R1289.	3.5	21
78	Cardiovascular complications of mild autonomous cortisol secretion. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2021, 35, 101494.	4.7	21
79	FokI Polymorphism of the Vitamin D Receptor Gene Correlates with Parameters of Bone Mass and Turnover in a Female Population of the Italian Island of Lampedusa. <i>Calcified Tissue International</i> , 2007, 80, 15-20.	3.1	20
80	Paget's disease of bone: there's more than the affected skeletal "a" a clinical review and suggestions for the clinical practice. <i>Current Opinion in Rheumatology</i> , 2010, 22, 410-423.	4.3	19
81	Genetics of Primary Hyperparathyroidism. <i>Urologia Internationalis</i> , 2004, 72, 11-16.	1.3	18
82	Ribozyme-mediated compensatory induction of menin-oncosuppressor function in primary fibroblasts from MEN1 patients. <i>Cancer Gene Therapy</i> , 2010, 17, 814-825.	4.6	18
83	Cortisol Secretion, Sensitivity, and Activity Are Associated With Hypertension in Postmenopausal Eucortisolemic Women. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 4441-4448.	3.6	18
84	Energy Metabolism and Ketogenic Diets: What about the Skeletal Health? A Narrative Review and a Prospective Vision for Planning Clinical Trials on this Issue. <i>International Journal of Molecular Sciences</i> , 2021, 22, 435.	4.1	18
85	Genetic screening for multiple endocrine neoplasia syndrome type 1 (MEN-1): when and how. <i>F1000 Medicine Reports</i> , 2010, 2, .	2.9	18
86	Genetics of parathyroids disorders: Overview. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2018, 32, 781-790.	4.7	17
87	Multiple Endocrine Neoplasia Type I Variants and Phenocopies: More than a Nosological Issue?. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 1518-1520.	3.6	16
88	Efficacy and safety of abaloparatide for the treatment of post-menopausal osteoporosis. <i>Expert Opinion on Pharmacotherapy</i> , 2019, 20, 805-811.	1.8	16
89	Emerging therapeutic targets for osteoporosis. <i>Expert Opinion on Therapeutic Targets</i> , 2020, 24, 115-130.	3.4	16
90	Silent Familial Isolated Pituitary Adenomas: Histopathological and Clinical Case Report. <i>Endocrine Pathology</i> , 2008, 19, 40-46.	9.0	15

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91	Management and Medical Therapy of Mild Hypercortisolism. International Journal of Molecular Sciences, 2021, 22, 11521.	4.1	15
92	Germline mutations in MEN1 and BRCA1 genes in a woman with familial multiple endocrine neoplasia type 1 and inherited breast-ovarian cancer syndromes: a case report. Cancer Genetics and Cytogenetics, 2009, 195, 75-79.	1.0	14
93	What Is the Relationship Between Paget's Disease of Bone and Hyperparathyroidism?. Journal of Bone and Mineral Research, 2006, 21, P69-P74.	2.8	13
94	Prediction of hypertension, diabetes and fractures in eucortisolemic women by measuring parameters of cortisol milieu. Endocrine, 2020, 68, 411-419.	2.3	13
95	Selenium: A Trace Element for a Healthy Skeleton - A Narrative Review. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2021, 21, 577-585.	1.2	13
96	Lessons from genes mutated in multiple endocrine neoplasia (MEN) syndromes. Annales D'Endocrinologie, 2005, 66, 195-205.	1.4	12
97	Hidden hypercortisolism: a too frequently neglected clinical condition. Journal of Endocrinological Investigation, 2021, 44, 1581-1596.	3.3	12
98	Mazabraud's Syndrome: A Case Report and Up-To-Date Literature Review. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2019, 19, 885-893.	1.2	10
99	Allelic Loss in Parathyroid Tumors from Individuals Homozygous for Multiple Endocrine Neoplasia Type 11. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2278-2282.	3.6	9
100	MEN1 family with a novel frameshift mutation. Journal of Endocrinological Investigation, 2006, 29, 450-456.	3.3	9
101	Genetic aspects of the Paget's disease of bone: concerns on the introduction of DNA-based tests in the clinical practice. Advantages and disadvantages of its application. European Journal of Clinical Investigation, 2010, 40, 655-667.	3.4	9
102	Defining Nonfunctioning Adrenal Adenomas on the Basis of the Occurrence of Hypocortisolism after Adrenalectomy. Journal of the Endocrine Society, 2020, 4, bvaa079.	0.2	9
103	Idiopathic Osteoporosis and Nephrolithiasis: Two Sides of the Same Coin?. International Journal of Molecular Sciences, 2020, 21, 8183.	4.1	9
104	Pathophysiology of Mild Hypercortisolism: From the Bench to the Bedside. International Journal of Molecular Sciences, 2022, 23, 673.	4.1	9
105	Clonal Analysis by Chromosome 11 Microsatellite-PCR of Microdissected Parathyroid Tumors from MEN 1 Patients. Biochemical and Biophysical Research Communications, 1996, 227, 736-742.	2.1	8
106	Allelic Loss in Parathyroid Tumors from Individuals Homozygous for Multiple Endocrine Neoplasia Type 1. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2278-2282.	3.6	8
107	Lack of allelic loss at the multiple endocrine neoplasia type 1 (MEN-1) gene locus in a pancreatic ductal (non-endocrine) adenocarcinoma of a patient with the MEN-1 syndrome. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 1995, 426, 203-8.	2.8	7
108	Relationship of Volumetric Bone Mineral Density and Structural Parameters with ERÎ± Gene Polymorphisms. Calcified Tissue International, 2007, 80, 307-315.	3.1	7

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109	The genetic ascertainment of multiple endocrine neoplasia type 1 syndrome by ancient DNA analysis. <i>Journal of Endocrinological Investigation</i> , 2008, 31, 905-909.	3.3	7
110	Management of bone fragility in type 2 diabetes: Perspective from an interdisciplinary expert panel. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2021, 31, 2210-2233.	2.6	7
111	Thiazolidinediones and bone. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2007, 4, 103-7.	1.0	7
112	Evidence for Bioeffects of LY 139478 on the Human Preosteoclastic Cell Line FLG 29.1. <i>Biochemical and Biophysical Research Communications</i> , 1995, 211, 857-863.	2.1	6
113	Genetic screening to identify the gene carrier in Italian and German kindreds affected by multiple endocrine neoplasia type 1 (MEN 1) syndrome. <i>Journal of Endocrinological Investigation</i> , 1995, 18, 329-335.	3.3	6
114	Infiltrating giant cell tumor in a case of Paget's disease of bone. <i>Archives of Osteoporosis</i> , 2009, 4, 91-94.	2.4	6
115	Secondary prevention of fragility fractures: where do we stand during the COVID-19 pandemic?. <i>Journal of Endocrinological Investigation</i> , 2021, 44, 2521-2524.	3.3	6
116	Giant cell tumor in a case of Paget's disease of bone: an aggressive benign tumor exhibiting a quick response to an innovative therapeutic agent. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2010, 7, 145-52.	1.0	6
117	Calcium Citrate Versus Calcium Carbonate in the Management of Chronic Hypoparathyroidism: A Randomized, Double-Blind, Crossover Clinical Trial. <i>Journal of Bone and Mineral Research</i> , 2020, 37, 1251-1259.	2.8	6
118	Calcium agonists in hyperparathyroidism. <i>Expert Opinion on Investigational Drugs</i> , 2004, 13, 229-244.	4.1	5
119	Vitamin K and bone metabolism: the myth and the truth. <i>Expert Review of Precision Medicine and Drug Development</i> , 2016, 1, 301-317.	0.7	5
120	Mitochondriopathies and bone health. <i>Trends in Biomedical Research</i> , 2018, 1, .	0.2	5
121	Bone metabolism, bone mass and structural integrity profile in professional male football players. <i>Journal of Sports Medicine and Physical Fitness</i> , 2020, 60, 912-918.	0.7	5
122	The role of osteoprotegerin (OPG) and estrogen receptor (ER- α) gene polymorphisms in rheumatoid arthritis. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2007, 4, 156-60.	1.0	5
123	Polyostotic form of fibrous dysplasia in a 13 years old Colombian girl showing clinical and biochemical response to neridronate intravenous therapy. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2009, 6, 264-5.	1.0	5
124	HPE Cells: A Clonal Endothelial Cell Line Established From Human Parathyroid Tissue (Human) Tj ETQq0 0 0 rgBT /Oyerlock 10 Tf 50 142	1.7	4
125	Multiple Endocrine Neoplasia Type 1. , 2008. , 1345-1374.		4
126	A novel germline inactivating mutation in the CASR gene in an Italian kindred affected by familial hypocalciuric hypercalcemia. <i>European Journal of Endocrinology</i> , 2012, 166, 933-940.	3.7	4

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127	Molecular diagnosis of parathyroid carcinoma: a reality in the near future. <i>Expert Opinion on Medical Diagnostics</i> , 2012, 6, 27-37.	1.6	4
128	The interaction between vitamin C and bone health: a narrative review. <i>Expert Review of Precision Medicine and Drug Development</i> , 2018, 3, 215-223.	0.7	4
129	Early post-natal life stress induces permanent adrenocorticotropin-dependent hypercortisolism in male mice. <i>Endocrine</i> , 2021, 73, 186-195.	2.3	4
130	Metabolic Syndromes and Dismobility. <i>American Journal of Physical Medicine and Rehabilitation</i> , 2021, 100, e32-e33.	1.4	4
131	Italian guidelines for the diagnosis and treatment of Paget's disease of bone. <i>Reumatismo</i> , 2011, 59, .	0.9	3
132	A novel germline mutation at exon 10 of MEN1 gene: a clinical survey and positive genotype-phenotype analysis of a MEN1 Italian family, including monozygotic twins. <i>Hormones</i> , 2018, 17, 427-435.	1.9	3
133	A novel polymorphism at the GNAS1 gene associated with low circulating calcium levels. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2007, 4, 139-45.	1.0	2
134	Genetic testing for multiple endocrine neoplasias. <i>Endocrine-Related Cancer</i> , 1998, 5, 37-44.	3.1	1
135	6. MEN syndromes. <i>Tumori</i> , 2010, 96, 823-826.	1.1	1
136	Multiple endocrine neoplasia type 2 syndromes (MEN 2): results from the ItaMEN network analysis on the prevalence of different genotypes and phenotypes. <i>European Journal of Endocrinology</i> , 2010, 163, 963.	3.7	1
137	LRP5 gene polymorphism and cortical bone. <i>Ageing Clinical and Experimental Research</i> , 2010, 22, 281-288.	2.9	1
138	Response to Letter to the Editor: "Methodological Issues Regarding Cortisol Secretion, Sensitivity, and Activity are Associated With Hypertension in Postmenopausal Eucortisolemic Women". <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 376-377.	3.6	1
139	A Novel Germline Mutation of ADA2 Gene in Two "Discordant" Homozygous Female Twins Affected by Adenosine Deaminase 2 Deficiency: Description of the Bone-Related Phenotype. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8331.	4.1	1
140	Vitamin D Status and SARS-CoV2 Clinical Outcomes: A Systematic Review and Meta-Analysis. <i>SSRN Electronic Journal</i> , 0, , .	0.4	1
141	Construction of a database for the evaluation and the clinical management of patients with breast cancer treated with antiestrogens and/or aromatase inhibitors. <i>Clinical Cases in Mineral and Bone Metabolism</i> , 2011, 8, 37-50.	1.0	1
142	Fragility fractures: clinical and therapeutic aspects. <i>Journal of Biological Regulators and Homeostatic Agents</i> , 2015, 29, 761-9.	0.7	1
143	Metabolic syndrome and fragility fracture risk. <i>Minerva Obstetrics and Gynecology</i> , 2021, 73, 744-753.	1.0	1
144	Are allelic losses at 11q13 universal in MEN 1 tumors?. <i>Journal of Clinical Endocrinology and Metabolism</i> , 1996, 81, 3162-3163.	3.6	1

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145	Somatic Mutation Analysis of the MEN1 Gene in Adrenocortical Tumors Using Denaturing Gradient Gel Electrophoresis (DGGE). International Journal on Disability and Human Development, 2001, 2, .	0.2	0
146	Fisiopatologia e trattamento nelle forme familiari dell'iperparatiroidismo primitivo. Giornale De Tecniche Nefrologiche & Dialitiche, 2009, 21, 18-24.	0.1	0
147	Aspetti epidemiologici della calcolosi di calcio in Italia: distribuzione dei fenotipi intermedi nella popolazione italiana. Giornale De Tecniche Nefrologiche & Dialitiche, 2009, 21, 74-77.	0.1	0
148	SAT0284...A Self-Administered Questionnaire May Allow General Practitioners to Perform a Rapid Screening to Identify Patients at Risk for Fragility Fractures and to Address Them to Bone Metabolism Specialized Clinical Centers. Annals of the Rheumatic Diseases, 2015, 74, 761.1-761.	0.9	0
149	SAT0285...Application of Criteria for the Diagnosis of "Clinical Osteoporosis" in a Population of Postmenopausal Women from Emilia-Romagna Region. Annals of the Rheumatic Diseases, 2015, 74, 761.2-762.	0.9	0
150	SUN-614 Prediction of Hypertension, Diabetes and Fractures in Eucortisolemic Women by Measuring Parameters of Cortisol Milieu. Journal of the Endocrine Society, 2020, 4, .	0.2	0
151	Adenosine Deaminase 2 Deficiency: Its pleiotropic manifestations may also affect bone metabolism? A case study on two homozygous female twins. Bone Reports, 2021, 14, 101021.	0.4	0
152	What Do We Talk About When We Talk About Frailty?. Frontiers in Rehabilitation Sciences, 2021, 2, .	1.2	0
153	Genetics of Osteoporosis. Medical Science Symposia Series, 2002, , 31-42.	0.0	0
154	Multiple Endocrine Neoplasia Type 1. , 2002, , 1047-1066.		0
155	Phytoestrogens: Food or Drug?. , 2006, , 219-231.		0
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