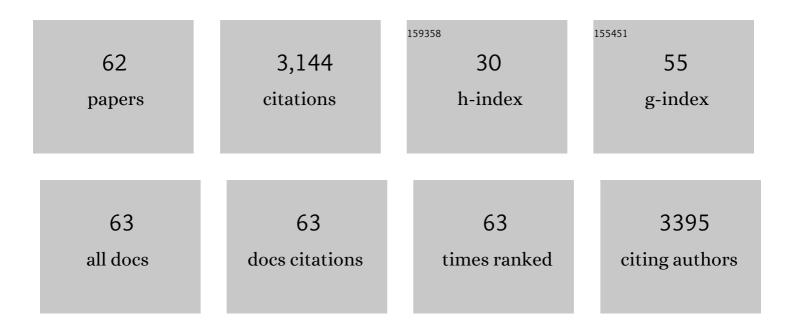
Paolo Emidio Macchia

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Quantitative but not qualitative flavor recognition impairments in COVID-19 patients. Irish Journal of Medical Science, 2022, 191, 1759-1766.	0.8	1
2	Obesity and Thyroid Cancer Risk: An Update. International Journal of Environmental Research and Public Health, 2022, 19, 1116.	1.2	32
3	Relationship between salt consumption and iodine intake in a pediatric population. European Journal of Nutrition, 2021, 60, 2193-2202.	1.8	7
4	Iodine Intake Estimated by 24 h Urine Collection in the Italian Adult Population: 2008–2012 Survey. Nutrients, 2021, 13, 1529.	1.7	5
5	lodine Intake from Food and Iodized Salt as Related to Dietary Salt Consumption in the Italian Adult General Population. Nutrients, 2021, 13, 3486.	1.7	7
6	Epigenetic Mechanisms of Endocrine-Disrupting Chemicals in Obesity. Biomedicines, 2021, 9, 1716.	1.4	17
7	Flavor identification inversely correlates with body mass index (BMI). Nutrition, Metabolism and Cardiovascular Diseases, 2020, 30, 1299-1305.	1.1	7
8	Influences of Age, Sex and Smoking Habit on Flavor Recognition in Healthy Population. International Journal of Environmental Research and Public Health, 2020, 17, 959.	1.2	15
9	Food and Nutrition as Prime Environmental Factors. , 2020, , 3-16.		1
10	Reply to A Olivieri et al American Journal of Clinical Nutrition, 2019, 110, 1267.	2.2	0
11	The flavor test is a sensitive tool in identifying the flavor sensorineural dysfunction in Parkinson's disease. Neurological Sciences, 2019, 40, 1351-1356.	0.9	11
12	Quercetin and its derivative Q2 modulate chromatin dynamics in adipogenesis and Q2 prevents obesity and metabolic disorders in rats. Journal of Nutritional Biochemistry, 2019, 69, 151-162.	1.9	40
13	Similarities and differences in the reproductive phenotypes of women with congenital hypogonadotrophic hypogonadism caused byGNRHRmutations and women with polycystic ovary syndrome. Human Reproduction, 2019, 34, 137-147.	0.4	10
14	Epigenetic and Metabolism: Glucose and Homeotic Transcription Factor PREP1 VRP Suggested Epigenetics and Metabolism. , 2019, , 761-776.		0
15	High-resolution melting analysis (HRM) for mutational screening of Dnajc17 gene in patients affected by thyroid dysgenesis. Journal of Endocrinological Investigation, 2018, 41, 711-717.	1.8	2
16	Nutritional and Environmental Factors in Thyroid Carcinogenesis. International Journal of Environmental Research and Public Health, 2018, 15, 1735.	1.2	50
17	Germline polymorphisms of the VEGF-pathway predict recurrence in non-advanced differentiated thyroid cancer. Journal of Clinical Endocrinology and Metabolism, 2017, 102, jc.2016-2555.	1.8	23
18	Influence of nutrition on somatotropic axis: Milk consumption inÂadult individuals with moderate-severe obesity. Clinical Nutrition, 2017, 36, 293-301.	2.3	47

ΡΑΟΙΟ ΕΜΙΟΙΟ ΜΑССΗΙΑ

#	Article	IF	CITATIONS
19	Dietary polyphenols and chromatin remodeling. Critical Reviews in Food Science and Nutrition, 2017, 57, 2589-2599.	5.4	61
20	Sunshine vitamin and thyroid. Reviews in Endocrine and Metabolic Disorders, 2017, 18, 347-354.	2.6	40
21	Selenium supplementation modulates apoptotic processes in thyroid follicular cells. BioFactors, 2017, 43, 415-423.	2.6	22
22	Preliminary results demonstrating the impact of Mediterranean diet on bone health. Journal of Translational Medicine, 2017, 15, 81.	1.8	48
23	Adherence to the Mediterranean Diet and Circulating Levels of Sirtuin 4 in Obese Patients: A Novel Association. Oxidative Medicine and Cellular Longevity, 2017, 2017, 1-14.	1.9	48
24	Mediterranean Diet and Phase Angle in a Sample of Adult Population: Results of a Pilot Study. Nutrients, 2017, 9, 151.	1.7	61
25	Epigenetic and Metabolism: Glucose and Homeotic Transcription Factor PREP1 VRP Suggested Epigenetics and Metabolism. , 2017, , 1-16.		Ο
26	Bioelectrical phase angle and psoriasis: a novel association with psoriasis severity, quality of life and metabolic syndrome. Journal of Translational Medicine, 2016, 14, 130.	1.8	58
27	Long period fiber grating nano-optrode for cancer biomarker detection. Biosensors and Bioelectronics, 2016, 80, 590-600.	5.3	79
28	Flavor perception test: evaluation in patients with Kallmann syndrome. Endocrine, 2016, 52, 236-243.	1.1	16
29	Glucose-induced expression of the homeotic transcription factor Prep1 is associated with histone post-translational modifications in skeletal muscle. Diabetologia, 2016, 59, 176-186.	2.9	27
30	Nutrition: a key environmental dietary factor in clinical severity and cardio-metabolic risk in psoriatic male patients evaluated by 7-day food-frequency questionnaire. Journal of Translational Medicine, 2015, 13, 303.	1.8	63
31	Effects of treatment modalities for Graves' hyperthyroidism on Graves' orbitopathy: a 2015 Italian Society of Endocrinology Consensus Statement. Journal of Endocrinological Investigation, 2015, 38, 481-487.	1.8	44
32	Nutrition and psoriasis: is there any association between the severity of the disease and adherence to the Mediterranean diet?. Journal of Translational Medicine, 2015, 13, 18.	1.8	112
33	Pregnancy outcome in women treated with methimazole or propylthiouracil during pregnancy. Journal of Endocrinological Investigation, 2015, 38, 977-985.	1.8	41
34	Identification and Functional Characterization of a Novel Mutation in the <i>NKX2-1</i> Gene: Comparison with the Data in the Literature. Thyroid, 2013, 23, 675-682.	2.4	29
35	The molecular causes of thyroid dysgenesis: a systematic review. Journal of Endocrinological Investigation, 2013, 36, 654-64.	1.8	40
36	Thyroid Nodules Treated with Percutaneous Radiofrequency Thermal Ablation: A Comparative Study. Journal of Clinical Endocrinology and Metabolism, 2012, 97, 4439-4445.	1.8	107

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37	Screening for mutations in the ISL1 gene in patients with thyroid dysgenesis. Journal of Endocrinological Investigation, 2011, 34, e149-52.	1.8	1
38	Benign hereditary chorea: Clinical and neuroimaging features in an Italian family. Movement Disorders, 2010, 25, 1491-1495.	2.2	32
39	Characterization of a novel lossâ€ofâ€function mutation of PAX8 associated with congenital hypothyroidism. Clinical Endocrinology, 2010, 73, 808-814.	1.2	29
40	Genetic Defects in Thyroid Hormone Synthesis and Action. , 2010, , 1721-1732.		0
41	Thyroid Nodules and Related Symptoms Are Stably Controlled Two Years After Radiofrequency Thermal Ablation. Thyroid, 2009, 19, 219-225.	2.4	239
42	lodine status assessment in Campania (Italy) as determined by urinary iodine excretion. Nutrition, 2009, 25, 926-929.	1.1	25
43	Mutations in TAZ/WWTR1, a co-activator of NKX2.1 and PAX8 are not a frequent cause of thyroid dysgenesis. Journal of Endocrinological Investigation, 2009, 32, 238-241.	1.8	7
44	A Novel <i>NKX2.1</i> Mutation in a Family with Hypothyroidism and Benign Hereditary Chorea. Thyroid, 2008, 18, 1005-1009.	2.4	55
45	A New Case of Familial Nonautoimmune Hyperthyroidism Caused by the M463V Mutation in the TSH Receptor with Anticipation of the Disease Across Generations: A Possible Role of Iodine Supplementation. Thyroid, 2007, 17, 677-680.	2.4	23
46	Missense Mutation in the Transcription Factor NKX2–5: A Novel Molecular Event in the Pathogenesis of Thyroid Dysgenesis. Journal of Clinical Endocrinology and Metabolism, 2006, 91, 1428-1433.	1.8	157
47	Autosomal Dominant Resistance to Thyrotropin as a Distinct Entity in Five Multigenerational Kindreds: Clinical Characterization and Exclusion of Candidate Loci. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4025-4034.	1.8	27
48	A Mouse Model Demonstrates a Multigenic Origin of Congenital Hypothyroidism. Endocrinology, 2005, 146, 5038-5047.	1.4	108
49	Thyroid function and effect of aging in combined hetero/homozygous mice deficient in thyroid hormone receptors alpha and beta genes. Journal of Endocrinology, 2002, 172, 177-185.	1.2	26
50	A Preservation Method That Allows Recovery of Intact RNA from Tissues Dissected by Laser Capture Microdissection. Analytical Biochemistry, 2002, 300, 139-145.	1.1	38
51	High-dose intravenous corticosteroid therapy for Graves' ophthalmopathy. Journal of Endocrinological Investigation, 2001, 24, 152-158.	1.8	88
52	Basi molecolari dell'ipotiroidismo congenito. L Endocrinologo, 2001, 2, 91-98.	0.0	0
53	Increased sensitivity to thyroid hormone in mice with complete deficiency of thyroid hormone receptor alpha. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 349-354.	3.3	82
54	Recent advances in understanding the molecular basis of primary congenital hypothyroidism. Trends in Molecular Medicine, 2000, 6, 36-42.	2.6	78

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#	Article	IF	CITATIONS
55	Molecular genetics of congenital hypothyroidism. Current Opinion in Genetics and Development, 1999, 9, 289-294.	1.5	41
56	Cloning, chromosomal localization and identification of polymorphisms in the human thyroid transcription factor 2 gene (TITF2). Biochimie, 1999, 81, 433-440.	1.3	57
57	Structural defects of a Pax8 mutant that give rise to congenital hypothyroidism. Biochemical Journal, 1999, 341, 89.	1.7	7
58	A mouse model for hereditary thyroid dysgenesis and cleft palate. Nature Genetics, 1998, 19, 395-398.	9.4	302
59	PAX8 mutations associated with congenital hypothyroidism caused by thyroid dysgenesis. Nature Genetics, 1998, 19, 83-86.	9.4	446
60	Mutations in the Gene EncodingThyroid Transcription Factor-1 (TTF-1) Are Not a Frequent Cause of Congenital Hypothyroidism (CH) with Thyroid Dysgenesis. Thyroid, 1997, 7, 383-387.	2.4	68
61	Epidermal growth factor receptor and lipid membrane components in human lung cancers. Journal of Endocrinological Investigation, 1993, 16, 99-107.	1.8	4
62	Epidermal growth factor receptor in human brain tumors. Journal of Endocrinological Investigation, 1992, 15, 31-37.	1.8	28