Marina Ciullo

List of Publications by Year in descending order

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74163 81900 9,505 75 39 75 citations h-index g-index papers 79 79 79 17360 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Genetic analysis of over 1 million people identifies 535 new loci associated with blood pressure traits. Nature Genetics, 2018, 50, 1412-1425.	21.4	924
2	Genome-wide association analyses identify 18 new loci associated with serum urate concentrations. Nature Genetics, 2013, 45, 145-154.	21.4	675
3	A catalog of genetic loci associated with kidney function from analyses of a million individuals. Nature Genetics, 2019, 51, 957-972.	21.4	549
4	Genome-wide association analysis identifies novel blood pressure loci and offers biological insights into cardiovascular risk. Nature Genetics, 2017, 49, 403-415.	21.4	492
5	Genomic analyses identify hundreds of variants associated with age at menarche and support a role for puberty timing in cancer risk. Nature Genetics, 2017, 49, 834-841.	21.4	426
6	Genetic associations at 53 loci highlight cell types and biological pathways relevant for kidney function. Nature Communications, 2016, 7, 10023.	12.8	412
7	New gene functions in megakaryopoiesis and platelet formation. Nature, 2011, 480, 201-208.	27.8	401
8	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. Nature Genetics, 2015, 47, 1294-1303.	21.4	357
9	Meta-analysis of genome-wide association studies for personality. Molecular Psychiatry, 2012, 17, 337-349.	7.9	340
10	Genome Analyses of >200,000 Individuals Identify 58 Loci for Chronic Inflammation and Highlight Pathways that Link Inflammation and Complex Disorders. American Journal of Human Genetics, 2018, 103, 691-706.	6.2	326
11	Seventy-five genetic loci influencing the human red blood cell. Nature, 2012, 492, 369-375.	27.8	320
12	Trans-ancestry genome-wide association study identifies 12 genetic loci influencing blood pressure and implicates a role for DNA methylation. Nature Genetics, 2015, 47, 1282-1293.	21.4	294
13	Meta-analysis of Genome-wide Association Studies for Neuroticism, and the Polygenic Association With Major Depressive Disorder. JAMA Psychiatry, 2015, 72, 642.	11.0	289
14	Genome-wide analysis identifies 12 loci influencing human reproductive behavior. Nature Genetics, 2016, 48, 1462-1472.	21.4	284
15	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. Nature Genetics, 2019, 51, 1459-1474.	21.4	251
16	Age- And Sex-Related Variations in Platelet Count in Italy: A Proposal of Reference Ranges Based on 40987 Subjects' Data. PLoS ONE, 2013, 8, e54289.	2.5	190
17	Genetic insights into biological mechanisms governing human ovarian ageing. Nature, 2021, 596, 393-397.	27.8	183
18	Meta-analysis of Genome-Wide Association Studies for Extraversion: Findings from the Genetics of Personality Consortium. Behavior Genetics, 2016, 46, 170-182.	2.1	178

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19	Directional dominance on stature and cognition inÂdiverse human populations. Nature, 2015, 523, 459-462.	27.8	173
20	Genome-Wide Association and Functional Follow-Up Reveals New Loci for Kidney Function. PLoS Genetics, 2012, 8, e1002584.	3.5	166
21	Meta-Analysis of Genome-Wide Association Studies Identifies Six New Loci for Serum Calcium Concentrations. PLoS Genetics, 2013, 9, e1003796.	3.5	142
22	Initiation of the breakage-fusion-bridge mechanism through common fragile site activation in human breast cancer cells: the model of PIP gene duplication from a break at FRA7I. Human Molecular Genetics, 2002, 11, 2887-2894.	2.9	124
23	Novel Blood Pressure Locus and Gene Discovery Using Genome-Wide Association Study and Expression Data Sets From Blood and the Kidney. Hypertension, 2017, 70, .	2.7	123
24	Genome-wide meta-analysis associates HLA-DQA1/DRB1 and LPA and lifestyle factors with human longevity. Nature Communications, 2017, 8, 910.	12.8	118
25	52 Genetic Loci Influencing MyocardialÂMass. Journal of the American College of Cardiology, 2016, 68, 1435-1448.	2.8	113
26	Large-scale genome-wide analysis identifies genetic variants associated with cardiac structure and function. Journal of Clinical Investigation, 2017, 127, 1798-1812.	8.2	106
27	Harmonization of Neuroticism and Extraversion phenotypes across inventories and cohorts in the Genetics of Personality Consortium: an application of Item Response Theory. Behavior Genetics, 2014, 44, 295-313.	2.1	103
28	1000 Genomes-based meta-analysis identifies 10 novel loci for kidney function. Scientific Reports, 2017, 7, 45040.	3.3	98
29	Meta-analysis of genome-wide association studies identifies common variants in CTNNA2 associated with excitement-seeking. Translational Psychiatry, 2011, 1, e49-e49.	4.8	97
30	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	12.8	84
31	Hearing function and thresholds: a genome-wide association study in European isolated populations identifies new loci and pathways. Journal of Medical Genetics, 2011, 48, 369-374.	3.2	71
32	Genetics of VEGF Serum Variation in Human Isolated Populations of Cilento: Importance of VEGF Polymorphisms. PLoS ONE, 2011, 6, e16982.	2.5	68
33	Body mass index is directly associated with biomarkers of angiogenesis and inflammation in children and adolescents. Nutrition, 2012, 28, 262-266.	2.4	67
34	Integration of genome-wide association studies with biological knowledge identifies six novel genes related to kidney function. Human Molecular Genetics, 2012, 21, 5329-5343.	2.9	64
35	Modulation of Genetic Associations with Serum Urate Levels by Body-Mass-Index in Humans. PLoS ONE, 2015, 10, e0119752.	2.5	64
36	Meta-Analysis of the INSIG2 Association with Obesity Including 74,345 Individuals: Does Heterogeneity of Estimates Relate to Study Design?. PLoS Genetics, 2009, 5, e1000694.	3.5	62

3

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37	Six Novel Loci Associated with Circulating VEGF Levels Identified by a Meta-analysis of Genome-Wide Association Studies. PLoS Genetics, 2016, 12, e1005874.	3.5	56
38	Large-scale whole-exome sequencing association studies identify rare functional variants influencing serum urate levels. Nature Communications, 2018, 9, 4228.	12.8	43
39	Angiogenesis and biomarkers of cardiovascular risk in adults with metabolic syndrome. Journal of Internal Medicine, 2010, 268, 338-347.	6.0	40
40	New susceptibility locus for hypertension on chromosome 8q by efficient pedigree-breaking in an Italian isolate. Human Molecular Genetics, 2006, 15, 1735-1743.	2.9	39
41	Expression and molecular characterization of alternative transcripts of the ARHGEF5/TIM oncogene specific for human breast cancer. Human Molecular Genetics, 2003, 13, 323-334.	2.9	37
42	Heritability and Demographic Analyses in the Large Isolated Population of Val Borbera Suggest Advantages in Mapping Complex Traits Genes. PLoS ONE, 2009, 4, e7554.	2.5	37
43	Genome-wide association analysis on normal hearing function identifies <i>PCDH20</i> and <i>SLC28A3</i> as candidates for hearing function and loss. Human Molecular Genetics, 2015, 24, 5655-5664.	2.9	37
44	Influence of age, sex and ethnicity on platelet count in five Italian geographic isolates: mild thrombocytopenia may be physiological. British Journal of Haematology, 2012, 157, 384-387.	2.5	33
45	Common Variants in Mendelian Kidney Disease Genes and Their Association with Renal Function. Journal of the American Society of Nephrology: JASN, 2013, 24, 2105-2117.	6.1	33
46	Comparing population structure as inferred from genealogical versus genetic information. European Journal of Human Genetics, 2009, 17, 1635-1641.	2.8	31
47	Salt-inducible kinase 3, SIK3, is a new gene associated with hearing. Human Molecular Genetics, 2014, 23, 6407-6418.	2.9	30
48	Campora: A Young Genetic Isolate in South Italy. Human Heredity, 2007, 64, 123-135.	0.8	28
49	Strategies for phasing and imputation in a population isolate. Genetic Epidemiology, 2018, 42, 201-213.	1.3	27
50	Variation of hemoglobin levels in normal Italian populations from genetic isolates. Haematologica, 2008, 93, 1372-1375.	3.5	25
51	Biosynthesis and immunobiochemical characterization of gp17/GCDFP-15. A glycoprotein from seminal vesicles and from breast tumors, in HeLa cells and in Pichia pastoris yeast. FEBS Journal, 1999, 265, 664-670.	0.2	24
52	Intragenic amplification and formation of extrachromosomal small circular DNA molecules from thePIP gene on chromosome 7 in primary breast carcinomas. International Journal of Cancer, 2002, 99, 370-377.	5.1	21
53	Genetic loci and prioritization of genes for kidney function decline derived from a meta-analysis of 62 longitudinal genome-wide association studies. Kidney International, 2022, 102, 624-639.	5.2	18
54	Age-related hearing loss in four Italian genetic isolates: An epidemiological study. International Journal of Audiology, 2009, 48, 465-472.	1.7	17

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55	Body composition, leg length and blood pressure in a rural Italian population: A test of the capacity-load model. Nutrition, Metabolism and Cardiovascular Diseases, 2014, 24, 1204-1212.	2.6	17
56	Differential and shared genetic effects on kidney function between diabetic and non-diabetic individuals. Communications Biology, 2022, 5, .	4.4	17
57	Identification and Replication of a Novel Obesity Locus on Chromosome 1q24 in Isolated Populations of Cilento. Diabetes, 2008, 57, 783-790.	0.6	16
58	Differential expression of insulin-dependent diabetes mellitus-associated HLA-DQA1 allelesin vivo. European Journal of Immunology, 1997, 27, 1549-1556.	2.9	15
59	Association of a variant in the CHRNA5-A3-B4 gene cluster region to heavy smoking in the Italian population. European Journal of Human Genetics, 2011, 19, 593-596.	2.8	13
60	Genetic Variants Modulating CRIPTO Serum Levels Identified by Genome-Wide Association Study in Cilento Isolates. PLoS Genetics, 2015, 11, e1004976.	3.5	13
61	Longevity candidate genes and their association with personality traits in the elderly. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 192-200.	1.7	12
62	Genetic and Environmental Factors Influencing the Placental Growth Factor (PGF) Variation in Two Populations. PLoS ONE, 2012, 7, e42537.	2.5	11
63	Control of Nucleo-cytoplasmic HLA-DRA MRNA Partitioning by Interaction of a Retention Signal with Compartmentalized Proteins. Journal of Molecular Biology, 1994, 240, 193-204.	4.2	10
64	Unbalanced expression of HLA-A and -B antigens: A specific feature of cutaneous melanoma and other non-hemopoietic malignancies reverted by IFN-?. International Journal of Cancer, 2001, 91, 500-507.	5.1	10
65	Whole-Exome Sequencing in the Isolated Populations of Cilento from South Italy. Scientific Reports, 2019, 9, 4059.	3.3	7
66	Genetics of PIGF plasma levels highlights a role of its receptors and supports the link between angiogenesis and immunity. Scientific Reports, 2021, 11, 16821.	3.3	6
67	Detecting the dominance component of heritability in isolated and outbred human populations. Scientific Reports, 2018, 8, 18048.	3.3	3
68	Polymorphism in the $5\hat{a}\in^2$ terminal region of the mRNA of HLA-DQA1 gene: Identification of four groups of transcripts and their association with polymorphism in the a 1 domain. Human Immunology, 1997, 53, 167-173.	2.4	2
69	Regulation of HLA class II gene expression: the case for posttranscriptional control levels. Microbes and Infection, 1999, 1, 943-948.	1.9	2
70	Downstream Sequence Adjacent to AUG Affects Translation of Chloramphenicol Acetyl Transferase in Eukaryotic Cells. DNA and Cell Biology, 2000, 19, 39-46.	1.9	2
71	SNP-Based Linkage Analysis in Extended Pedigrees: Comparison between Two Alternative Approaches. Human Heredity, 2014, 78, 27-37.	0.8	1
72	A transnational collaborative network dedicated to the study and applications of the vascular endothelial growth factor-A in medical practice: the VEGF Consortium. Clinical Chemistry and Laboratory Medicine, 2018, 56, 83-86.	2.3	1

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73	Unbalanced expression of HLAâ∈A and â∈B antigens: A specific feature of cutaneous melanoma and other nonâ∈hemopoietic malignancies reverted by IFNâ∈γ. International Journal of Cancer, 2001, 91, 500-507.	5.1	1
74	Moment estimators of relatedness from low-depth whole-genome sequencing data. BMC Bioinformatics, 2022, 23, .	2.6	1
7 5	Large-Scale Genomic Analyses Link Reproductive Aging to Hypothalamic Signaling, Breast Cancer Susceptibility, and BRCA1-Mediated DNA Repair. Obstetrical and Gynecological Survey, 2015, 70, 758-762.	0.4	0