Lvks Bhaskar

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8606708/publications.pdf

Version: 2024-02-01

144	1,413	394286	454834
papers	citations	19 h-index	30 g-index

150	150	150	2012
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	NOS3 gene intron 4 a/b polymorphism is associated with ESRD in autosomal dominant polycystic kidney disease patients. Jornal Brasileiro De Nefrologia: Orgao Oficial De Sociedades Brasileira E Latino-Americana De Nefrologia, 2022, 44, 224-231.	0.4	2
2	Risk factors and pathogenic mechanismâ€“associated hepatocellular carcinoma. , 2022, , 33-49.		0
3	Multidrug resistance, a major obstacle in hepatocellular carcinoma treatment: challenges and future perspectives. , 2022, , 227-253.		0
4	Biomarkers of Oxidative Stress Tethered to Cardiovascular Diseases. Oxidative Medicine and Cellular Longevity, 2022, 2022, 1-15.	1.9	29
5	Erythrocyte microRNAs: a tiny magic bullet with great potential for sickle cell disease therapy. Annals of Hematology, 2021, 100, 607-614.	0.8	6
6	Diagnostic and Prognostic Implications of Cardiac Markers for Hepatocellular Carcinoma. Critical Reviews in Oncogenesis, 2021, 26, 1-10.	0.2	0
7	Molecular mechanism, diagnosis, and potential treatment for novel coronavirus (COVID-19): a current literature review and perspective. 3 Biotech, 2021, 11, 94.	1.1	7
8	CYP1B1 rs1056836 polymorphism and endometrial cancer risk: A meta-analysis. , 2021, , 263-270.		0
9	A Review on Hematopoietic Stem Cell Treatment for Epilepsy. CNS and Neurological Disorders - Drug Targets, 2021, 20, 644-656.	0.8	2
10	Genetic variations at 10q26 regions near FGFR2 gene and its association with non-syndromic cleft lip with or without cleft palate. International Journal of Pediatric Otorhinolaryngology, 2021, 143, 110648.	0.4	2
11	Susceptibility to vascular complications in sickle cell anemia patients is associated with intron 4a/b polymorphism of the NOS3 gene: A meta-analysis. Meta Gene, 2021, 28, 100870.	0.3	0
12	Nanotechnology based drug delivery system: Current strategies and emerging therapeutic potential for medical science. Journal of Drug Delivery Science and Technology, 2021, 63, 102487.	1.4	117
13	Association of hypercoagulation with severe acute respiratory syndrome coronavirus 2 infection. Blood Research, 2021, 56, 61-64.	0.5	9
14	World Kidney Day 2021 with the theme of living well with kidney disease; a review of current concepts. Journal of Preventive Epidemiology, 2021, 6, e08-e08.	0.1	0
15	Interleukin-6 gene -174G>C promoter polymorphism reduces the risk of periodontitis in Brazilian populations: A meta-analysis. Journal of Oral Biosciences, 2021, 63, 388-393.	0.8	2
16	Sodium-glucose co-transporter 2 inhibitors (SGLT2i); as a preventive factor of kidney failure in patients with type 2 diabetes; a meta-analysis of randomized controlled trials. Journal of Renal Injury Prevention, 2021, 10, e35-e35.	0.6	2
17	Pharmacoeconomics and cost-effectiveness of treatments related to breast and cervix cancers. , 2021, , 235-244.		0
18	Herbal antioxidants and renal ischemic-reperfusion injury; an updated review. Journal of Nephropharmacology, 2021, 10, e03-e03.	0.2	7

#	Article	IF	Citations
19	Potential of renin-angiotensin system inhibition to improve metabolic bone disorders. Journal of Nephropharmacology, 2021, 10, e16-e16.	0.2	5
20	Reply to the comments on letter to the editor on review article â€œcurrent updates and treatment strategy of the European and WHO registered clinical trials of coronavirus disease 2019â€• Biomedical Journal, 2021, , .	1.4	1
21	Physical Activity and Nutritional Influence on Immune Function: An Important Strategy to Improve Immunity and Health Status. Frontiers in Physiology, 2021, 12, 751374.	1.3	21
22	Meta-Analysis on the Association of Neuropeptide Y rs16139 Variant With the Risk of Alcoholism. Frontiers in Psychiatry, 2021, 12, 737440.	1.3	0
23	The M235T polymorphism in the angiotensinogen gene is not a major risk factor for diabetic nephropathy; a meta-analysis. Journal of Preventive Epidemiology, 2021, 7, e15-e15.	0.1	0
24	Glomerulonephritis associated with SARS-CoV-2 infection. Journal of Nephropharmacology, 2021, 10, e07-e07.	0.2	0
25	SARS-CoV-2 infection in people with pre-existing liver disease: Further research is warranted. World Journal of Gastroenterology, 2021, 27, 7855-7858.	1.4	2
26	Metformin induced acute kidney injury; a systematic review. Journal of Nephropharmacology, 2021, 10, e13-e13.	0.2	1
27	Strive for kidney health for everyone during COVID-19; the possible theme for the world kidney day 2021. Journal of Nephropharmacology, 2021, 10, e12-e12.	0.2	1
28	Interleukin-10 gene promoter variants and susceptibility to diabetic nephropathy; a meta-analysis. Journal of Nephropathology, 2021, 10, e38-e38.	0.1	0
29	Understanding kidney injury in COVID-19; a pressing priority. Journal of Nephropharmacology, 2021, 10, e19-e19.	0.2	0
30	COVID-19 and the kidney; mechanisms of tubular injury by SARS-CoV-2. Journal of Renal Injury Prevention, 2021, 10, e08-e08.	0.6	1
31	Association of Clinical and Hematological variables with the disease severity in Indian Sickle cell anemia patients. Research Journal of Pharmacy and Technology, 2021, , 5254-5257.	0.2	1
32	Association Between MTHFD1 1958Gâ€‰›â€‰A Variant and non-Syndromic Cleft lip and Palate: An Updated Meta-Analysis. Cleft Palate-Craniofacial Journal, 2021, , 105566562110464.	0.5	1
33	Influence of Neuropeptide Y and Neuropeptide Y 2 Receptor Variants in Acute Coronary Syndrome. International Journal of Cardiovascular Sciences, 2021, , .	0.0	0
34	Gender differences in the relationship between alcohol consumption and gastric cancer risk are uncertain and not well-delineated. World Journal of Gastrointestinal Oncology, 2021, 13, 2216-2218.	0.8	0
35	Maternal and infant MTHFR gene polymorphisms and non-syndromic oral cleft susceptibility: An updated meta-analysis. Process Biochemistry, 2020, 89, 81-88.	1.8	1
36	Genetic association of GSTM1, GSTT1, and GSTP1 polymorphisms with sickle cell disease complications: A systematic review and meta-analysis. Meta Gene, 2020, 26, 100815.	0.3	0

#	Article	IF	Citations
37	Micro RNA facilitated chemoresistance in gastric cancer: a novel biomarkers and potential therapeutics. Alexandria Journal of Medicine, 2020, 56, 81-92.	0.4	5
38	Current updates on the European and WHO registered clinical trials of coronavirus disease 2019 (COVID-19). Biomedical Journal, 2020, 43, 424-433.	1.4	22
39	Dynamic Propagation and Impact of Pandemic Influenza A (2009 H1N1) in Children: A Detailed Review. Current Microbiology, 2020, 77, 3809-3820.	1.0	19
40	HIF1A C1772T genetic variation is associated with the elevated risk of breast cancer among Asians: An updated meta-analysis. Meta Gene, 2020, 25, 100722.	0.3	0
41	The Fuzzy Connection between SARS-CoV-2 Infection and Loss of Renal Function. American Journal of Nephrology, 2020, 51, 572-573.	1.4	3
42	Parental transmission effect of PDGF-C gene variants on non-syndromic cleft lip with or without cleft palate. Meta Gene, 2020, 24, 100669.	0.3	1
43	Molecular Signaling Pathways Involved in Gastric Cancer Chemoresistance. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 117-134.	0.2	5
44	Administration of metformin in type 2 diabetes mellitus patients with chronic kidney disease; facts and myths. Journal of Nephropathology, 2020, 9, e04-e04.	0.1	2
45	Tumor Necrosis Factor-Alpha Gene Promoter (TNF-Î± G-308A) Polymorphisms Increase the Risk of Hepatocellular Carcinoma in Asians: A Meta-Analysis. Critical Reviews in Oncogenesis, 2020, 25, 11-20.	0.2	5
46	A Retrospective Look at Anti-EGFR Agents in Pancreatic Cancer Therapy. Current Drug Metabolism, 2020, 20, 958-966.	0.7	9
47	Coronavirus-nephropathy; renal involvement in COVID-19. Journal of Renal Injury Prevention, 2020, 9, e18-e18.	0.6	37
48	Perspectives on the relationship of renal disease and coronavirus disease 2019. Journal of Nephropharmacology, 2020, 9, e22-e22.	0.2	17
49	The K469E genetic variant in the ICAM1 gene is associated with type 2 diabetes but not with its vascular complications: a meta-analysis. Journal of Nephropharmacology, 2020, 9, e16-e16.	0.2	0
50	Association Between IL6 Gene Polymorphisms and Gastric Cancer Risk: A Meta-Analysis of Case-Control Studies. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 171-182.	0.2	0
51	Meta-analysis of NFKB1-94 ATTG Ins/Del Polymorphism and Risk of Breast Cancer. Current Drug Metabolism, 2020, 21, 221-225.	0.7	1
52	MicroRNA a small magic bullet for gastric cancer. Gene, 2020, 753, 144801.	1.0	1
53	Understanding Colorectal Cancer: The Basics. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 93-115.	0.2	0
54	An Intergenic Variant rs4779584 Between SCG5 and GREM1 Contributes to the Increased Risk of Colorectal Cancer: A Meta-Analysis. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 159-169.	0.2	0

#	Article	IF	Citations
55	Meta-Analysis Reveals no Significant Association of EPHX1 Tyr113His and His139Arg Polymorphisms with the Colorectal Cancer Risk. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 135-150.	0.2	1
56	Meta-Analysis of Genetic Variants in Alcohol Metabolizing Enzymes and their Association with Colorectal Cancer Risk. Diagnostics and Therapeutic Advances in GI Malignancies, 2020, , 151-166.	0.2	0
57	Current Therapeutic Strategies and Perspectives for Neuroprotection in Parkinsonâ€™s Disease. Current Pharmaceutical Design, 2020, 26, 4738-4746.	0.9	3
58	Association between MTHFR 677C>T polymorphism and vascular complications in sickle cell disease: A meta-analysis. Transfusion Clinique Et Biologique, 2019, 26, 284-288.	0.2	16
59	Molecular Markers for Treatment Response and Toxicity of Gemcitabine. , 2019, , 175-195.		2
60	Meta-analysis of MTHFR polymorphisms and pancreatic cancer susceptibility. , 2019, , 263-274.		0
61	Small molecule tyrosine kinase inhibitors and pancreatic cancerâ€”Trials and troubles. Seminars in Cancer Biology, 2019, 56, 149-167.	4.3	23
62	Genetic association of ACE gene I/D polymorphism with the risk of diabetic kidney disease; a meta-analysis. Journal of Nephropathology, 2019, 8, 44-44.	0.1	2
63	Factor V Leiden and prothrombin G20210A mutations and risk of vaso-occlusive complications in sickle cell disease: A meta-analysis through the lens of nephrology. Journal of Nephropharmacology, 2019, 8, 16-16.	0.2	2
64	Perspectives and Molecular Understanding of Pancreatic Cancer Stem Cells. , 2019, , 157-172.		0
65	Association between the methylenetetrahydrofolate reductase (<i>MTHFR</i>) gene 677C>T and 1298A>C polymorphisms and the risk of diabetic nephropathy; a meta-analysis. Journal of Renal Injury Prevention, 2019, 8, 175-184.	0.6	2
66	Apolipoprotein L1 associated nephropathy; an overview. Journal of Renal Injury Prevention, 2019, 8, 311-315.	0.6	0
67	Relationship between matrilin-1 gene polymorphisms and mandibular retrognathism. American Journal of Orthodontics and Dentofacial Orthopedics, 2018, 153, 255-261.e1.	0.8	19
68	IRF6 rs2235375 single nucleotide polymorphism is associated with isolated non-syndromic cleft palate but not with cleft lip with or without palate in South Indian population. Brazilian Journal of Otorhinolaryngology, 2018, 84, 473-477.	0.4	10
69	Compound Heterozygosity of Î²-Thalassemia and the Sickle Cell Hemoglobin in Various Populations of Chhattisgarh State, India. Hemoglobin, 2018, 42, 84-90.	0.4	7
70	5-Lipoxygenase: Its involvement in gastrointestinal malignancies. Critical Reviews in Oncology/Hematology, 2018, 127, 50-55.	2.0	26
71	Pain Management Issues as Part of the Comprehensive Care of Patients with Sickle Cell Disease. Pain Management Nursing, 2018, 19, 558-572.	0.4	17
72	Retrospection of the effect of hydroxyurea treatment in patients with sickle cell disease. Acta Haematologica Polonica, 2018, 49, 1-8.	0.1	13

#	Article	IF	Citations
73	Cyclic Guanosine Monophosphate-Dependent Protein Kinase I Stimulators and Activators Are Therapeutic Alternatives for Sickle Cell Disease. Turkish Journal of Haematology, 2018, 35, 77-78.	0.2	4
74	Drug Resistance Against Tyrosine Kinase Inhibitor in Gastrointestinal Malignancies. , 2018, , 191-224.		0
75	Molecular docking analysis of phytoconstituent from Momordica charantia with Guanylate Cyclase catalytic domain. Bioinformation, 2018, 14, 378-383.	0.2	2
76	Transmission analysis of TGFB1 gene polymorphisms in non-syndromic cleft lip with or without cleft palate. International Journal of Pediatric Otorhinolaryngology, 2017, 100, 14-17.	0.4	6
77	Attitudes and beliefs among high- and low-risk population groups towards Î²-thalassemia prevention: a cross-sectional descriptive study from India. Journal of Community Genetics, 2017, 8, 159-166.	0.5	8
78	Assessment of renal function in Indian patients with sickle cell disease. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2017, 28, 524.	0.4	9
79	Aldosterone synthase gene is not a major susceptibility gene for progression of chronic kidney disease in patients with autosomal dominant polycystic kidney disease. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2017, 28, 552.	0.4	1
80	Ocular manifestations of sickle cell disease and genetic susceptibility for refractive errors. Taiwan Journal of Ophthalmology, 2017, 7, 89.	0.3	7
81	ECE1 gene variant shows tendency toward chronic kidney disease advancement among autosomal polycystic kidney disease patients. Hong Kong Journal of Nephrology, 2016, 18, 20-25.	0.0	2
82	Polymorphic Regions in Fc Gamma Receptor and Tumor Necrosis Factor-Î± Genes and Susceptibility to Chronic Periodontitis in a Cohort From South India. Journal of Periodontology, 2016, 87, 914-922.	1.7	8
83	Role of myosin 1H gene polymorphisms in mandibular retrognathism. American Journal of Orthodontics and Dentofacial Orthopedics, 2016, 149, 699-704.	0.8	26
84	Association of TFAP2A gene polymorphism with susceptibility to non-syndromic cleft lip with or without palate risk in south Indian population. Meta Gene, 2016, 9, 181-184.	0.3	2
85	Renin gene rs1464816 polymorphism contributes to chronic kidney disease progression in ADPKD. Journal of Biomedical Science, 2016, 23, 1.	2.6	61
86	Insights on the structural perturbations in human MTHFR Ala222Val mutant by protein modeling and molecular dynamics. Journal of Biomolecular Structure and Dynamics, 2016, 34, 892-905.	2.0	9
87	Angiotensinogen gene polymorphisms and progression of chronic kidney disease in ADPKD patients. Clinical and Experimental Nephrology, 2016, 20, 561-568.	0.7	3
88	NOS3 27-bp and IL4 70-bp VNTR Polymorphisms Do Not Contribute to the Risk of Sickle Cell Crisis. Turkish Journal of Haematology, 2016, 33, 365-366.	0.2	6
89	Two promoter polymorphisms in TBX22 are associated with the risk of NSCLP in Indian women. Clinical Dysmorphology, 2015, 24, 140-143.	0.1	4
90	Burden among Parkinson's disease care givers for a community based study from India. Journal of the Neurological Sciences, 2015, 358, 276-281.	0.3	30

#	Article	IF	Citations
91	CBS c.844ins68 Polymorphism Frequencies in Control Populations: Implications on Nonsyndromic Cleft Lip with or without Cleft Palate. Cleft Palate-Craniofacial Journal, 2015, 52, 49-53.	0.5	2
92	Polymorphic Regions in the Interleukin-1 Gene and Susceptibility to Chronic Periodontitis: A Genetic Association Study. Genetic Testing and Molecular Biomarkers, 2015, 19, 175-181.	0.3	20
93	Relationship between reduced folate carrier gene polymorphism and non-syndromic cleft lip and palate in Indian population. Journal of Maternal-Fetal and Neonatal Medicine, 2015, 28, 329-332.	0.7	6
94	Evidence of the involvement of the polymorphisms near MSX1 gene in non-syndromic cleft lip with or without cleft palate. International Journal of Pediatric Otorhinolaryngology, 2015, 79, 1081-1084.	0.4	7
95	Genetic variant in MTRR A66G, but not MTR A2756G, is associated with risk of non-syndromic cleft lip and palate in Indian population. Journal of Oral and Maxillofacial Surgery, Medicine, and Pathology, 2015, 27, 782-785.	0.2	7
96	Genotyping and meta-analysis of KIF6 Trp719Arg polymorphism in South Indian Coronary Artery Disease patients: A caseâ€“control study. Meta Gene, 2015, 5, 129-134.	0.3	8
97	SATB2 gene variants in non-syndromic cleft lip with or without cleft palate in Indian population. Journal of Oral Biology and Craniofacial Research, 2015, 5, 161-164.	0.8	4
98	Polymorphic variants near 1p22 and 20q11.2 loci and the risk of non-syndromic cleft lip and palate in South Indian population. International Journal of Pediatric Otorhinolaryngology, 2015, 79, 2389-2393.	0.4	11
99	TYMS gene 5'- and 3'-untranslated region polymorphisms and risk of non-syndromic cleft lip and palate in an Indian population. Journal of Biomedical Research, 2015, 29, 337.	0.7	3
100	Significant association of MTHFD1 1958G A single nucleotide polymorphism with nonsyndromic cleft lip and palate in Indian population. Medicina Oral, Patologia Oral Y Cirugia Bucal, 2014, 19, e616-e621.	0.7	8
101	<scp>NOS3 tagSNPs</scp> does not modify the chronic kidney disease progression in autosomal dominant polycystic kidney disease. Nephrology, 2014, 19, 537-541.	0.7	7
102	Polymorphisms in genes encoding dopamine signalling pathway and risk of alcohol dependence: a systematic review. Acta Neuropsychiatrica, 2014, 26, 69-80.	1.0	8
103	MTHFR C677T and A1298C polymorphisms and risk of nonsyndromic orofacial clefts in a south Indian population. International Journal of Pediatric Otorhinolaryngology, 2014, 78, 339-342.	0.4	15
104	ZwiÄ…zek miÄ™dzy polimorfizmem rs7903146 genu TCF7L2 a nefropatiÄ… cukrzycowÄ… nie jest niezależny od cukrzycy typu 2 â€" badanie populacji Indii PoÅ‚udniowych i metaanaliza. Endokrynologia Polska, 2014, 65, 298-305.	0.3	28
105	Genetic variants associated with insulin signaling and glucose homeostasis in the pathogenesis of insulin resistance in polycystic ovary syndrome: a systematic review. Journal of Assisted Reproduction and Genetics, 2013, 30, 883-895.	1.2	22
106	EPHX1 gene polymorphisms among south Indian populations. Molecular and Cellular Toxicology, 2013, 9, 219-225.	0.8	3
107	EPHX1 Gene Polymorphisms in Alcohol Dependence and their Distribution among the Indian Populations. American Journal of Drug and Alcohol Abuse, 2013, 39, 16-22.	1.1	7
108	HLA class II SNP interactions and the association with type 1 diabetes mellitus in Bengali speaking patients of Eastern India. Journal of Biomedical Science, 2013, 20, 12.	2.6	9

#	Article	IF	Citations
109	Analysis of Microsatellite Polymorphisms in South Indian Patients with Non Syndromic Cleft Lip and Palate. Balkan Journal of Medical Genetics, 2013, 16, 49-54.	0.5	6
110	Association between Neuropeptide Y Gene Polymorphisms and Alcohol Dependence: A Case-Control Study in Two Independent Populations. European Addiction Research, 2013, 19, 307-313.	1.3	14
111	Endothelial nitric oxide synthase VNTR (intron 4 a/b) polymorphism association with nonsyndromic oral clefts. Turkish Journal of Biochemistry, 2013, 38, 308-312.	0.3	0
112	Role of the ACE ID and PPARG P12A Polymorphisms in Genetic Susceptibility of Diabetic Nephropathy in a South Indian Population. Nephro-Urology Monthly, 2013, 5, 813-7.	0.0	13
113	Functional PstI/RsaI Polymorphisms in the CYP2E1 Gene among South Indian Populations. Asian Pacific Journal of Cancer Prevention, 2013, 14, 179-182.	0.5	11
114	Effect of flupirtine on the growth and viability of U373 malignant glioma cells. Cancer Biology and Medicine, 2013, 10, 142-7.	1.4	3
115	Molecular Basis of Î²-Thalassemia in Karnataka, India. Genetic Testing and Molecular Biomarkers, 2012, 16, 138-141.	0.3	14
116	Dopamine Transporter (DAT1) VNTR Polymorphism and Alcoholism in Two Culturally Different Populations of South India. American Journal on Addictions, 2012, 21, 343-347.	1.3	23
117	CYP1A1 genotypes and haplotypes and risk of oral cancer: a case-control study in South Indians. Genetics and Molecular Biology, 2012, 35, 407-412.	0.6	8
118	An unlikely role for the NAT2 genotypes and haplotypes in the oral cancer of south Indians. Archives of Oral Biology, 2012, 57, 513-518.	0.8	5
119	Lack of Association ofEPHX1Genotypes and Haplotypes with Oral Cancer in South Indians. Genetic Testing and Molecular Biomarkers, 2011, 15, 595-599.	0.3	5
120	Lack of Association Between the IL-10 Gene Polymorphisms and Features of the Metabolic Syndrome. Journal of Investigative Medicine, 2011, 59, 267-271.	0.7	2
121	Polymorphisms in genes involved in folate metabolism and orofacial clefts. Archives of Oral Biology, 2011, 56, 723-737.	0.8	39
122	Indian Siddis: African Descendants with Indian Admixture. American Journal of Human Genetics, 2011, 89, 154-161.	2.6	50
123	Association Between the M268T Polymorphism in the Angiotensinogen Gene and Essential Hypertension in a South Indian Population. Biochemical Genetics, 2011, 49, 474-482.	0.8	9
124	Apelin gene polymorphism influences apelin expression and obesity phenotypes in Chinese women. American Journal of Clinical Nutrition, 2011, 94, 921-928.	2.2	26
125	A Novel Polymorphism in Codon 25 of the KRAS Gene Associated with Gallbladder Carcinoma Patients of the Eastern Part of India. Genetic Testing and Molecular Biomarkers, 2011, 15, 431-434.	0.3	8
126	Neuropeptide Y gene polymorphisms are not associated with obesity in a South Indian population. European Journal of Clinical Nutrition, 2010, 64, 868-872.	1.3	11

#	Article	IF	Citations
127	Neuropeptide Y gene functional polymorphism influences susceptibility to hypertension in Indian population. Journal of Human Hypertension, 2010, 24, 617-622.	1.0	12
128	Population-Based Case-Control Study of DRD2 Gene Polymorphisms and Alcoholism. Journal of Addictive Diseases, 2010, 29, 475-480.	0.8	13
129	Single Nucleotide Polymorphisms of PARKIN Gene in Ten Indian Populations. , 2010, , 105-114.		0
130	Current concepts in genetics of nonsyndromic clefts. Indian Journal of Plastic Surgery, 2009, 42, 68.	0.2	58
131	Dopamine transporter (DAT1) VNTR polymorphism in 12 Indian populations. Neurological Sciences, 2009, 30, 487-493.	0.9	7
132	Genetic heterogeneity in the Indian stocks of seahorse (HippocampusÂkuda and) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 542 Td (Hippoca	1.0	8
133	Maternal Footprints of Southeast Asians in North India. Human Heredity, 2008, 66, 1-9.	0.4	43
134	Allelic Variation and Haplotype Structure of the Dopamine Receptor GeneDRD2in Nine Indian Populations. Genetic Testing and Molecular Biomarkers, 2008, 12, 153-160.	1.7	11
135	Single nucleotide polymorphisms of the ALDH2 gene in six Indian populations. Annals of Human Biology, 2007, 34, 607-619.	0.4	16
136	Anthropological Perspective of the Single Nucleotide Polymorphisms in the NPY and DRD2 Genes among the Socio-Economically Stratified Populations of Andhra Pradesh, India. International Journal of Human Genetics, 2007, 7, 277-284.	0.1	2
137	Single nucleotide polymorphisms in alcohol dehydrogenase genes among some Indian populations. American Journal of Human Biology, 2007, 19, 338-344.	0.8	16
138	Allelic variation in the NPY gene in 14 Indian populations. Journal of Human Genetics, 2007, 52, 592-598.	1.1	14
139	Single Nucleotide Polymorphisms of the Alcohol Dehydrogenase Genes among the 28 Caste and Tribal Populations of India. International Journal of Human Genetics, 2006, 6, 309-316.	0.1	8
140	Genetic affinities among the lower castes and tribal groups of India: inference from Y chromosome and mitochondrial DNA. BMC Genetics, 2006, 7, 42.	2.7	93
141	The HBG2 rs7482144 (Câ€‰>â€‰T) Polymorphism is Linked to HbF Levels but not to the Severity of Sickle Cell Anemia. Journal of Pediatric Genetics, 0, , .	0.3	1
142	Ameliorative impact of sodiumâ€“glucose cotransporter-2 inhibitors in diabetic kidney disease; a mini-review to current evidence. Journal of Renal Injury Prevention, 0, , .	0.6	0
143	Molecular docking study of Momordica charantia Linn phytoconstituent with caspase 3 and implications for renoprotective actions in diabetes mellitus. Journal of Nephropharmacology, 0, , .	0.2	0
144	IL1RN VNTR Polymorphism and kidney damage in sickle cell anemia patients. Journal of Nephropharmacology, 0, , .	0.2	0