Seyed Alireza Dastgheib

List of Publications by Year
in descending order

77 papers	749 citations	840119 11 h-index	610482 24 g-index
78 all docs	78 docs citations	78 times ranked	1117 citing authors

#	Article	IF	Citations
1	Vertical Transmission of Coronavirus Disease 19 (COVID-19) from Infected Pregnant Mothers to Neonates: A Review. Fetal and Pediatric Pathology, 2020, 39, 246-250.	0.4	304
2	Association of MTHFR 677C>T and 1298A>C polymorphisms with susceptibility to autism: A systematic review and meta-analysis. Asian Journal of Psychiatry, 2019, 46, 54-61.	0.9	27
3	The Risk and Prevalence of COVID-19 Infection in Colorectal Cancer Patients: a Systematic Review and Meta-analysis. Journal of Gastrointestinal Cancer, 2021, 52, 73-79.	0.6	22
4	Association of Endothelial Nitric Oxide Synthase 894Gâ€‰>â€‰T Polymorphism with Preeclampsia Risk: A Systematic Review and Meta-Analysis based on 35 Studies. Fetal and Pediatric Pathology, 2021, 40, 455-470.	0.4	20
5	Proportion and mortality of Iranian diabetes mellitus, chronic kidney disease, hypertension and cardiovascular disease patients with COVID-19: a meta-analysis. Journal of Diabetes and Metabolic Disorders, 2021, 20, 905-917.	0.8	18
6	Association of eNOS and ACE Polymorphisms with Retinopathy of Prematurity: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2020, 39, 334-345.	0.4	15
7	Association of BMP4 rs17563 Polymorphism with Nonsyndromic Cleft Lip with or without Cleft Palate Risk: Literature Review and Comprehensive Meta-Analysis. Fetal and Pediatric Pathology, 2021, 40, 305-319.	0.4	15
8	Evidence from a meta-analysis for association of MC4R rs17782313 and FTO rs9939609 polymorphisms with susceptibility to obesity in children. Diabetes and Metabolic Syndrome: Clinical Research and Reviews, 2021, 15, 102234.	1.8	15
9	Association of Promoter Region Polymorphisms of IL-6 and IL-18 Genes with Risk of Recurrent Pregnancy Loss: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2020, 39, 346-359.	0.4	13
10	Phenotype of ST3GAL3 deficient patients: A case and review of the literature. European Journal of Medical Genetics, 2021, 64, 104250.	0.7	12
11	Association of IL-6 â^'174 G>C Polymorphism with Susceptibility to Colorectal Cancer and Gastric Cancer: a Systematic Review and Meta-Analysis. Acta Medica (Hradec Kralove), 2019, 62, 137-146.	0.2	12
12	Association of MTHFR 1298Aâ€‰>â€‰C Polymorphism with Susceptibility to Non-Syndromic Cleft Lip with or without Palate: A Case-Control Study and Meta-Analysis. Fetal and Pediatric Pathology, 2021, 40, 1-17.	0.4	11
13	Association of TNF-Î± rs1800629, CASP3 rs72689236 and FCGR2A rs1801274 Polymorphisms with Susceptibility to Kawasaki Disease: A Comprehensive Meta-Analysis. Fetal and Pediatric Pathology, 2019, 40, 1-17.	0.4	10
14	Association of Interleukin-10 -1082Gâ€‰>â€‰A Polymorphism with Susceptibility to Preeclampsia: A Systematic Review and Meta-Analysis Based on 21 Studies. Fetal and Pediatric Pathology, 2020, 39, 518-532.	0.4	10
15	The Impact of Methylenetetrahydrofolate Reductase (MTHFR) Sperm Methylation and Variants on Semen Parameters and the Chance of Recurrent Pregnancy Loss in the Couple. Clinical Laboratory, 2018, 64, 1121-1128.	0.2	10
16	A novel mutation in SEPN1 causing rigid spine muscular dystrophy 1: a Case report. BMC Medical Genetics, 2019, 20, 13.	2.1	9
17	Association of plasminogen activator inhibitor-1 4G5G Polymorphism with risk of diabetic nephropathy:Âa systematic review and meta-analysis. Journal of Diabetes and Metabolic Disorders, 2020, 19, 2005-2016.	0.8	9
18	An analysis of inhibition of the severe acute respiratory syndrome coronavirus 2 RNA-dependent RNA polymerase by zinc ion: an inÂsilico approach. Future Virology, 2021, 16, 331-339.	0.9	9

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19	Association of PAI-1 rs1799889 Polymorphism with Susceptibility to Ischemic Stroke: a Huge Meta-Analysis based on 44 Studies. Acta Medica (Hradec Kralove), 2020, 63, 31-42.	0.2	9
20	Association of MTHFR 677C>T, 1298A>C and MTR 2756A>G Polymorphisms with Risk of Retinoblastoma. Klinicka Onkologie, 2019, 32, 375-379.	0.1	9
21	AssociaÃ§Ã£o dos polimorfismos ESRÎ± XbaI Aâ€‰>â€‰G, ESRÎ± PvuII Tâ€‰>â€‰C e ESRÎ² AlwNI Tâ€‰>â€‰C com o risco desenvolver escoliose idiopÃ¡tica da adolescÃªncia: RevisÃ£o sistemÃ¡tica e metanÃ¡lise genÃ©tica. Revista Brasileira De Ortopedia, 2020, 55, 008-016.	0.2	8
22	A Candidate Gene Association Study of Bone Mineral Density in an Iranian Population. Frontiers in Endocrinology, 2016, 7, 141.	1.5	7
23	Clinical and molecular characterization of three patients with Hepatocerebral form of mitochondrial DNA depletion syndrome: a case series. BMC Medical Genetics, 2019, 20, 167.	2.1	7
24	Association of Fetal MTHFR 677Câ€‰>â€‰T Polymorphism with Non-Syndromic Cleft Lip with or without Palate Risk: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2021, 40, 337-353.	0.4	7
25	Association of MTHFR 677Câ€‰>â€‰T and 1298Aâ€‰>â€‰C polymorphisms with susceptibility to attention deficit and hyperactivity disorder. Fetal and Pediatric Pathology, 2020, 39, 422-429.	0.4	7
26	Association of Axis Inhibition Protein 2 Polymorphisms with Non-Syndromic Cleft Lip with or without Cleft Palate in Iranian Children. Fetal and Pediatric Pathology, 2020, 39, 29-37.	0.4	7
27	MCM2 mutation causes autosomal dominant nonsyndromic hearing loss (DFNA70): novel variant in the second family. Journal of Genetics, 2022, 101, 1.	0.4	7
28	Association of IL-6 -176Gâ€‰>â€‰C Polymorphism with Susceptibility to Preeclampsia: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2020, 39, 491-502.	0.4	6
29	Exome sequencing identified a de novo frameshift pathogenic variant of CTBP1 in an extremely rare case of HADDTS. Journal of Genetics, 2021, 100, 1.	0.4	6
30	A meta-analysis for the risk and prevalence of preeclampsia among pregnant women with COVID-19. TâˆšÂªrk Jinekoloji Ve Obstetrik Dernei Dergisi, 2021, 18, 224-235.	0.3	6
31	A novel non-sense mutation in TDP2 causes spinocerebellar ataxia autosomal recessive 23 accompanied by bilateral upward gaze; report of a case and review of the literature. European Journal of Medical Genetics, 2021, 64, 104348.	0.7	6
32	ASSOCIATION OF IL-8 -251T>A (RS4073) POLYMORPHISM WITH SUSCEPTIBILITY TO GASTRIC CANCER: A SYSTEMATIC REVIEW AND META-ANALYSIS BASED ON 33 CASE-CONTROL STUDIES. Arquivos De Gastroenterologia, 2020, 57, 91-99.	0.3	6
33	Clinical features of patients with Yin Yang 1 deficiency causing Gabrieleâ€de Vries syndrome: A new case and review of the literature. Annals of Human Genetics, 2022, 86, 52-62.	0.3	6
34	A Meta-Analysis for Association of XRCC3 rs861539, MTHFR rs1801133, IL-6 rs1800795, IL-12B rs3212227, TNF-Î± rs1800629, and TLR9 rs352140 Polymorphisms with Susceptibility to Cervical Carcinoma. Asian Pacific Journal of Cancer Prevention, 2021, 22, 3419-3431.	0.5	6
35	Association of MMP-2, MMP-3, and MMP-9 Polymorphisms with Susceptibility to Recurrent Pregnancy Loss. Fetal and Pediatric Pathology, 2021, 40, 378-386.	0.4	5
36	Cumulative Evidence for Association between IL-10 Polymorphisms and Kawasaki Disease Susceptibility: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2021, 40, 153-165.	0.4	5

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37	Association of IL-10 -1082A>G, -819C>T, and -592C>A polymorphisms with susceptibility to chronic and aggressive periodontitis: a systematic review and meta-analysis. Inflammation Research, 2021, 70, 509-524.	1.6	5
38	Association of PON1, LEP and LEPR Polymorphisms with Susceptibility to Breast Cancer: A Meta-Analysis. Asian Pacific Journal of Cancer Prevention, 2021, 22, 2323-2334.	0.5	5
39	Meta-analysis of the frequency of intrauterine growth restriction and preterm premature rupture of the membranes in pregnant women with COVID-19. TâˆšÂºrk Jinekoloji Ve Obstetrik Dernei Dergisi, 2021, 18, 236-244.	0.3	5
40	Association of -753C>T and -1562C>T Polymorphisms with Chronic/Aggressive Periodontitis Risk: A Systematic Review and Meta-Analysis. Iranian Journal of Public Health, 2019, 48, 1227-1238.	0.3	5
41	Phenotypic spectrum of autosomal recessive Keratitis-Ichthyosis-Deafness Syndrome (KIDAR) due to mutations in AP1B1. European Journal of Medical Genetics, 2022, 65, 104449.	0.7	5
42	Association of MTHFR 677C>T polymorphism with IUGR and placental abruption risk: A systematic review and meta-analysis. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 256, 130-139.	0.5	4
43	Association of IL-6 -174Gâ€‰>â€‰C and -572Gâ€‰>â€‰C Polymorphisms with Risk of Legg-Calve-Perthes Disease in Iranian Children. Fetal and Pediatric Pathology, 2021, 40, 206-213.	0.4	4
44	A novel frameshift pathogenic variant in ST3GAL5 causing salt and pepper developmental regression syndrome (SPDRS): A case report. Human Genome Variation, 2021, 8, 33.	0.4	4
45	Proportion of hematological cancer patients with SARS-CoV-2 infection during the COVID-19 pandemic: A systematic review and meta-analysis. Hematology, Transfusion and Cell Therapy, 2022, 44, 225-234.	0.1	4
46	Association of Endothelial Nitric Oxide Synthase Gene Polymorphisms with Susceptibility to Prostate Cancer: a Comprehensive Systematic Review and Meta-Analysis. Urology Journal, 2020, 17, 329-337.	0.3	4
47	An Updated and Comprehensive Meta-Analysis of Association between VEGA -634Gâ€‰>â€‰C, -460Tâ€‰>â€‰C, +405Gâ€‰>â€‰C and +936Câ€‰>â€‰T Polymorphisms and Retinopathy of Prematurity Risk. Fetal and Pediatric Pathology, 2021, 40, 233-249.	Ped	3
48	A novel PTRH2 missense mutation causing IMNEPD: a case report. Human Genome Variation, 2021, 8, 23.	0.4	3
49	Association of Transforming Growth Factor-Î²1 rs1982073 Polymorphism with Susceptibility to Acute Renal Rejection: a Systematic Review and Meta-Analysis. Urology Journal, 2020, 18, 1-10.	0.3	3
50	Association of catechol-O-methyltranferase 472G>A (Val158Met) polymorphism with susceptibility to fibromyalgia syndrome. Journal of Orthopaedics, 2020, 20, 257-260.	0.6	2
51	Association of +1923Câ€‰>â€‰T, -1112Câ€‰>â€‰T and +2044Aâ€‰>â€‰G Polymorphisms in IL-13 Gene with Susce to Pediatric Asthma: A Systematic Review and Meta-Analysis. Fetal and Pediatric Pathology, 2022, 41, 259-277.	0.4	2
52	Association Between the hOGG1 1245C>G (rs1052133) Polymorphism and Susceptibility to Colorectal Cancer: a Meta-analysis Based on 7010 Cases and 10,674 Controls. Journal of Gastrointestinal Cancer, 2021, 52, 389-398.	0.6	2
53	Association of Neuregulin 1 rs7835688â€‰Gâ€‰>â€‰C, rs16879552 Tâ€‰>â€‰C and rs2439302â€‰Gâ€‰>â€‰C Po with Susceptibility to Non-Syndromic Hirschsprungâ€™s Disease. Fetal and Pediatric Pathology, 2021, 40, 198-205.	0.4	2
54	Gender difference in determinant factors of being overweight among the 40â€“70-year-old population of Kharameh cohort study, Iran. BMC Public Health, 2021, 21, 746.	1.2	2

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55	A meta-analysis of the association of the ACE I/D and PAI-1 4G/5G polymorphisms with recurrent pregnancy loss in Iranian women: Are the investigations adequate?. TâˆšÂºrk Jinekoloji Ve Obstetrik Dernei Dergisi, 2021, 18, 139-150.	0.3	2
56	Association of ACE I/D and PAI-1 4G/5G polymorphisms with susceptibility to type 2 diabetes mellitus. Journal of Diabetes and Metabolic Disorders, 2021, 20, 1191-1197.	0.8	2
57	A meta-analysis for association of TNF-Î± -308G>A polymorphism with susceptibility to Ankylosing Spondylitis. Journal of Orthopaedics, 2021, 26, 79-87.	0.6	2
58	A meta-analysis for association of eNOS VNTR 4b/a, â€‰â€“â€‰786ÂTâ€‰>â€‰C andâ€‰+â€‰894Gâ€‰>â€‰T polymorphisms recurrent pregnancy loss. Archives of Gynecology and Obstetrics, 2021, 304, 1135-1151.	0.8	2
59	Association of Interleukin-10 Polymorphisms with Susceptibility to Colorectal Cancer and Gastric Cancer: an Updated Meta-analysis Based on 106 Studies. Journal of Gastrointestinal Cancer, 2022, 53, 1066-1082.	0.6	2
60	Association of interlukine-18 polymorphisms with susceptibility to prostate cancer in Iranian population. Neoplasma, 2020, 67, 644-649.	0.7	2
61	Genotypic and phenotypic spectrum of Myofibrillar Myopathy 7 as a result of Kyphoscoliosis Peptidase deficiency: The first description of a missense mutation in KY and literature review. European Journal of Medical Genetics, 2022, 65, 104552.	0.7	2
62	Peripheral Arterial Disease in Patients with Diabetes Mellitus. Southern Medical Journal, 2006, 99, 707-708.	0.3	1
63	Association of Epidermal Growth Factor 61A>G, Survivin -31G>C, and EFNA1 -1732G>A Polymorphisms with Susceptibility to Colorectal Cancer. Journal of Gastrointestinal Cancer, 2022, 53, 78-83.	0.6	1
64	AssociaÃ§Ã£o do polimorfismo IL-6 -174Gâ€‰>â€‰C (rs1800795) com escoliose idiopÃ¡tica da adolescÃªncia: EvidÃªncias de um estudo de caso-controle e metanÃ¡lise. Revista Brasileira De Ortopedia, 2020, 55, 017-026.	0.2	1
65	Association of ACE I/D, -240Aâ€‰>â€‰T and AT1R A1166C polymorphisms with susceptibility to breast cancer: a systematic review and meta-analysis based on 35 case-control studies. Nucleosides, Nucleotides and Nucleic Acids, 2021, 40, 117-135.	0.4	1
66	Cumulative Evidence for Association Between IL-8 -251T>A and IL-18 -607C>A Polymorphisms and Colorectal Cancer Susceptibility: a Systematic Review and Meta-analysis. Journal of Gastrointestinal Cancer, 2021, 52, 31-40.	0.6	1
67	Association of SERPINE1 rs1799889 polymorphism with arterial ischemic stroke in children: a systematic review and meta-analysis. Nucleosides, Nucleotides and Nucleic Acids, 2021, 40, 1018-1035.	0.4	1
68	Two novel Warburg micro syndrome 1 cases caused by pathogenic variants in RAB3GAP1. Human Genome Variation, 2021, 8, 39.	0.4	1
69	Chronic Renal Failure and Diabetes Mellitus: Are They Comparable Risk Factors of Coronary Artery Disease?. Southern Medical Journal, 2007, 100, 6-7.	0.3	1
70	Recurrent Infections and Immunodeficiency Caused by Severe Pancytopenia Associated with a Novel Life-Threatening Mutation in Hypoxia-Upregulated Protein 1. Immunological Investigations, 2022, , 1-12.	1.0	1
71	Associations of MTHFR rs1801133 (677C>T) and rs180113 (1298A>C) Polymorphisms with Susceptibility to Bladder Cancer: A Systematic Review and Meta-Analysis. Asian Pacific Journal of Cancer Prevention, 2022, 23, 1465-1482.	0.5	1
72	A single-amino-acid in-frame deletion in CYP17A1 results in combined 17-hydroxylase and 17,20-lyase deficiency in an Iranian family despite the protein mutation site. Human Genome Variation, 2021, 8, 31.	0.4	0

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73	A Meta-Analysis for Association of XRCC1, XRCC2 and XRCC3 Polymorphisms with Susceptibility to Thyroid Cancer. Asian Pacific Journal of Cancer Prevention, 2021, 22, 2221-2236.	0.5	0
74	Association of AXIN2 s2240308 C>T, rs1133683 C>T, rs7224837 A>G Polymorphisms with Susceptibility to Breast Cancer. Asian Pacific Journal of Cancer Prevention, 2021, 22, 2717-2722.	0.5	0
75	Association of NAD(P)H Quinine Oxidoreductase 1 rs1800566 Polymorphism with Bladder and Prostate Cancers â€“ a Systematic Review and Meta-Analysis. Klinicka Onkologie, 2020, 33, 92-100.	0.1	0
76	Exome sequencing identified a frameshift pathogenic variant of in an extremely rare case of HADDTS. Journal of Genetics, 2021, 100, .	0.4	0
77	Association of XPG rs17655G>C and XPF rs1799801T>C Polymorphisms with Susceptibility to Cutaneous Malignant Melanoma: Evidence from a Case-Control Study, Systematic Review and Meta-Analysis. Klinicka Onkologie, 2020, 33, 184-194.	0.1	0