Nina Pereza

List of Publications by Year
in descending order

29	369	1040056 9	794594 19
papers	citations	h-index	g-index

29	29	29	678
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Maternal LINE-1 DNA Methylation in Early Spontaneous Preterm Birth. Biological Research for Nursing, 2022, 24, 85-93.	1.9	2
2	Editorial: The Importance of Genetic Literacy and Education in Medicine. Frontiers in Genetics, 2022, 13, 910530.	2.3	0
3	Genetic variation in the maternal vitamin D receptor FokI gene as a risk factor for recurrent pregnancy loss. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 2221-2226.	1.5	9
4	The impact of needs-based education on the change of knowledge and attitudes towards medical genetics in medical students. European Journal of Human Genetics, 2021, 29, 726-735.	2.8	7
5	Current State of Compulsory Basic and Clinical Courses in Genetics for Medical Students at Medical Faculties in Balkan Countries With Slavic Languages. Frontiers in Genetics, 2021, 12, 793834.	2.3	0
6	Vitamin D receptor polymorphisms in spontaneous preterm birth: a case-control study. Croatian Medical Journal, 2020, 61, 338-345.	0.7	6
7	DNMT3B rs1569686 and rs2424913 gene polymorphisms are associated with positive family history of preterm birth and smoking status. Croatian Medical Journal, 2020, 61, 8-17.	0.7	8
8	Combination of QFâ€PCR and aCGH is an efficient diagnostic strategy for the detection of chromosome aberrations in recurrent miscarriage. Molecular Genetics & Genomic Medicine, 2019, 7, e980.	1.2	7
9	Matrix metalloproteinase and tissue inhibitors of metalloproteinases gene polymorphisms in disorders that influence fertility and pregnancy complications: A systematic review and meta-analysis. Gene, 2018, 647, 48-60.	2.2	22
10	Student section of Medicina Fluminensis â€“ a guarantee for a bright future. Medicina Fluminensis, 2018, 54, 4-5.	0.3	1
11	Functional single nucleotide polymorphisms of matrix metalloproteinase 7 and 12 genes in idiopathic recurrent spontaneous abortion. Journal of Assisted Reproduction and Genetics, 2017, 34, 365-371.	2.5	2
12	A Single Nucleotide Polymorphism of <scp>DNA</scp> methyltransferase 3B gene is a risk factor for recurrent spontaneous abortion. American Journal of Reproductive Immunology, 2017, 78, e12765.	1.2	8
13	Systematic review and meta-analysis of genetic association studies in idiopathic recurrent spontaneous abortion. Fertility and Sterility, 2017, 107, 150-159.e2.	1.0	110
14	Insertion/deletion polymorphism in intron 16 of ACE gene in idiopathic recurrent spontaneous abortion: case-control study, systematic review and meta-analysis. Reproductive BioMedicine Online, 2016, 32, 237-246.	2.4	14
15	The â^'2549 insertion/deletion polymorphism in the promoter region of the VEGFA gene in couples with idiopathic recurrent spontaneous abortion. Journal of Assisted Reproduction and Genetics, 2015, 32, 1789-1794.	2.5	9
16	Cornelia de Lange syndrome caused by heterozygous deletions of chromosome 8q24: Comments on the article by Pereza et al. [2012]. American Journal of Medical Genetics, Part A, 2015, 167, 1426-1427.	1.2	8
17	A critical update on endothelial nitric oxide synthase gene variations in women with idiopathic recurrent spontaneous abortion: genetic association study, systematic review and meta-analyses. Molecular Human Reproduction, 2015, 21, 466-478.	2.8	14
18	Functional Polymorphisms of Matrix Metalloproteinases 1 and 9 Genes in Women with Spontaneous Preterm Birth. Disease Markers, 2014, 2014, 1-7.	1.3	13

#	Article	IF	Citations
19	Genetic variation in tissue inhibitors of metalloproteinases as a risk factorÂfor idiopathic recurrent spontaneous abortion. Fertility and Sterility, 2013, 99, 1923-1929.	1.0	7
20	Y chromosome azoospermia factor region microdeletions are not associated with idiopathic recurrent spontaneous abortion in a Slovenian population: association study and literature review. Fertility and Sterility, 2013, 99, 1663-1667.	1.0	9
21	Insulin-like growth factor 2 and insulin-like growth factor 2 receptor gene polymorphisms in idiopathic male infertility. Journal of reproductive medicine, The, 2013, 58, 132-6.	0.2	0
22	Matrix metalloproteinases 1, 2, 3 and 9 functional single-nucleotide polymorphisms in idiopathic recurrent spontaneous abortion. Reproductive BioMedicine Online, 2012, 24, 567-575.	2.4	33
23	The insulin-like growth factor 2 receptor gene Gly1619Arg polymorphism and idiopathic recurrent spontaneous abortion. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 429-431.	1.5	4
24	Third case of 8q23.3â€q24.13 deletion in a patient with Langerâ€“Giedion syndrome phenotype without TRPS1 gene deletion. American Journal of Medical Genetics, Part A, 2012, 158A, 659-663.	1.2	20
25	Body mass index, waist circumference and waist-to-hip ratio: which anthropometric indicator is better predictor for the hypertension development in women population of the island Cres. Collegium Antropologicum, 2012, 36, 363-8.	0.2	0
26	Recurrent achalasia in a child with Williams-Beuren syndrome. Collegium Antropologicum, 2011, 35, 941-4.	0.2	1
27	Association between genetic polymorphisms in cytokine genes and recurrent miscarriage â€“ a meta-analysis. Reproductive BioMedicine Online, 2009, 19, 406-414.	2.4	24
28	A current genetic and epigenetic view on human aging mechanisms. Collegium Antropologicum, 2009, 33, 687-99.	0.2	7
29	ORIGINAL ARTICLE: Genetic Predisposition to Idiopathic Recurrent Spontaneous Abortion: Contribution of Genetic Variations in IGFâ€2 and H19 Imprinted Genes. American Journal of Reproductive Immunology, 2008, 60, 111-117.	1.2	24