

Chao Xuan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1568083/publications.pdf>

Version: 2024-02-01

39
papers

734
citations

471371
17
h-index

526166
27
g-index

40
all docs

40
docs citations

40
times ranked

1268
citing authors

#	ARTICLE	IF	CITATIONS
1	Mitochondrial membrane potential and reactive oxygen species in cancer stem cells. <i>Familial Cancer</i> , 2015, 14, 19-23.	0.9	105
2	Association Between Polymorphism of Methylenetetrahydrofolate Reductase (MTHFR) C677T and Risk of Myocardial Infarction: A Meta-analysis for 8,140 Cases and 10,522 Controls. <i>Archives of Medical Research</i> , 2011, 42, 677-685.	1.5	67
3	Levels of asymmetric dimethylarginine (ADMA), an endogenous nitric oxide synthase inhibitor, and risk of coronary artery disease: A meta-analysis based on 4713 participants. <i>European Journal of Preventive Cardiology</i> , 2016, 23, 502-510.	0.8	48
4	Association Between MTHFR Polymorphisms and Congenital Heart Disease: A Meta-analysis based on 9,329 cases and 15,076 controls. <i>Scientific Reports</i> , 2014, 4, 7311.	1.6	39
5	Identification of Altered Plasma Proteins by Proteomic Study in Valvular Heart Diseases and the Potential Clinical Significance. <i>PLoS ONE</i> , 2013, 8, e72111.	1.1	33
6	L-citrulline for protection of endothelial function from ADMA-induced injury in porcine coronary artery. <i>Scientific Reports</i> , 2015, 5, 10987.	1.6	32
7	Association between OCTN1/2 gene polymorphisms (1672C-T, 207G-C) and susceptibility of Crohn's disease: a meta-analysis. <i>International Journal of Colorectal Disease</i> , 2012, 27, 11-19.	1.0	30
8	<i>PTPN22</i> Gene Polymorphism (C1858T) Is Associated with Susceptibility to Type 1 Diabetes: A Meta-Analysis of 19,495 Cases and 25,341 Controls. <i>Annals of Human Genetics</i> , 2013, 77, 191-203.	0.3	29
9	Zebrafish xenotransplantation as a tool for in vivo cancer study. <i>Familial Cancer</i> , 2015, 14, 487-493.	0.9	29
10	Endothelial nitric oxide synthase enhancer for protection of endothelial function from asymmetric dimethylarginine-induced injury in human internal thoracic artery. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2012, 144, 697-703.	0.4	25
11	Genetic 135G/C polymorphism of RAD51 gene and risk of cancer: a meta-analysis of 28,956 cases and 28,372 controls. <i>Familial Cancer</i> , 2014, 13, 515-526.	0.9	24
12	No association between APOE epsilon 4 allele and multiple sclerosis susceptibility: A meta-analysis from 5472 cases and 4727 controls. <i>Journal of the Neurological Sciences</i> , 2011, 308, 110-116.	0.3	22
13	Serum Uric Acid as an Independent Risk Factor for the Presence and Severity of Early-Onset Coronary Artery Disease: A Case-Control Study. <i>Disease Markers</i> , 2018, 2018, 1-8.	0.6	22
14	SELDI-TOF MS profiling of serum for detection of nasopharyngeal carcinoma. <i>Journal of Experimental and Clinical Cancer Research</i> , 2009, 28, 85.	3.5	21
15	Role of TRPC3 Channel in Human Internal Mammary Artery. <i>Archives of Medical Research</i> , 2012, 43, 431-437.	1.5	21
16	Proteomic Study Reveals Plasma Protein Changes in Congenital Heart Diseases. <i>Annals of Thoracic Surgery</i> , 2014, 97, 1414-1419.	0.7	21
17	Increased serum concentrations of asymmetric dimethylarginine (ADMA) in patients with early-onset coronary artery disease. <i>Clinica Chimica Acta</i> , 2017, 464, 195-199.	0.5	20
18	Acute phase proteins altered in the plasma of patients with congenital ventricular septal defect. <i>Proteomics - Clinical Applications</i> , 2015, 9, 1087-1096.	0.8	19

#	ARTICLE	IF	CITATIONS
19	Dimethylarginine Dimethylaminohydrolase 2 (DDAH 2) Gene Polymorphism, Asymmetric Dimethylarginine (ADMA) Concentrations, and Risk of Coronary Artery Disease: A Case-Control Study. <i>Scientific Reports</i> , 2016, 6, 33934.	1.6	15
20	Screening and Identification of Pregnancy Zone Protein and Leucine-Rich Alpha ₂ -Glycoprotein as Potential Serum Biomarkers for Early-Onset Myocardial Infarction using Protein Profile Analysis. <i>Proteomics - Clinical Applications</i> , 2019, 13, e1800079.	0.8	14
21	Identification of Two Novel Mutations of the <i>HOMEZ</i> Gene in Chinese Patients with Isolated Ventricular Septal Defect. <i>Genetic Testing and Molecular Biomarkers</i> , 2013, 17, 390-394.	0.3	13
22	Association Between MTHFR Gene Common Variants, Serum Homocysteine, and Risk of Early-Onset Coronary Artery Disease: A Case-Control Study. <i>Biochemical Genetics</i> , 2020, 58, 245-256.	0.8	13
23	A Novel Variation of <i>PLAGL1</i> in Chinese Patients with Isolated Ventricular Septal Defect. <i>Genetic Testing and Molecular Biomarkers</i> , 2012, 16, 984-987.	0.3	12
24	Diagnostic Accuracy of Glycosylated Hemoglobin in Chinese Patients with Gestational Diabetes Mellitus: A Meta-Analysis Based on 2,812 Patients and 5,918 Controls. <i>Genetic Testing and Molecular Biomarkers</i> , 2013, 17, 687-695.	0.3	11
25	Serum adenosine deaminase activity and coronary artery disease: a retrospective case-control study based on 9929 participants. <i>Therapeutic Advances in Chronic Disease</i> , 2019, 10, 204062231989153.	1.1	11
26	Quantitative Assessment of Serum Amino Acids and Association with Early-Onset Coronary Artery Disease. <i>Clinical Interventions in Aging</i> , 2021, Volume 16, 465-474.	1.3	9
27	BIIB021: A novel inhibitor to heat shock protein 90α-addicted oncology. <i>Tumor Biology</i> , 2017, 39, 101042831769835.	0.8	8
28	Serum fatty acids profile and association with early-onset coronary artery disease. <i>Therapeutic Advances in Chronic Disease</i> , 2021, 12, 204062232110331.	1.1	5
29	Association between the methylenetetrahydrofolate reductase C677T polymorphism and susceptibility to preeclampsia: the need for data clarification in a recent meta-analysis. <i>Hypertension Research</i> , 2013, 36, 463-464.	1.5	4
30	Association between <i>ALMS1</i> variants and early-onset coronary artery disease: a case-control study in Chinese population. <i>Bioscience Reports</i> , 2020, 40, .	1.1	3
31	Corrective surgery alters plasma protein profiling in congenital heart diseases and clinical perspectives. <i>American Journal of Translational Research (discontinued)</i> , 2020, 12, 1319-1337.	0.0	3
32	Association between interleukin-10 gene polymorphisms and risk of oral carcinoma: A meta-analysis. <i>Histology and Histopathology</i> , 2020, 35, 1329-1336.	0.5	3
33	Comparison of protein profiles between acetonitrile- and non-acetonitrile-treated sera from patients with nasopharyngeal carcinomas. <i>Oncology Letters</i> , 2011, 2, 477-481.	0.8	1
34	Reply: Association Between Methylenetetrahydrofolate Reductase (MTHFR) C677T Polymorphism and Risk of Myocardial Infarction: Need for Clarification of Data in a Recent Meta-analysis. <i>Archives of Medical Research</i> , 2012, 43, 490.	1.5	1
35	A rapid and highly effective approach for SARS-CoV-2 nucleic acid daily testing in more than four thousand single-tube samples. <i>Journal of Clinical Laboratory Analysis</i> , 2022, 36, e24211.	0.9	1
36	Mutations of <i>HOMEZ</i> gene in Congenital Heart Diseases. <i>FASEB Journal</i> , 2012, 26, 1134.8.	0.2	0

#	ARTICLE	IF	CITATIONS
37	Effect of Azelnidipine in Human Internal Mammary Artery and Clinical Implications. FASEB Journal, 2013, 27, 1101.2.	0.2	0
38	Identification of Altered Plasma Proteins by Proteomic Study in Valvular Heart Diseases and the Potential Clinical Significance. FASEB Journal, 2013, 27, 1189.3.	0.2	0
39	Proteomic Changes After Surgical Repair in Congenital Heart Diseases. FASEB Journal, 2018, 32, 675.3.	0.2	0