## Azita Zadeh-Vakili

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/849650/publications.pdf

Version: 2024-02-01

		471509	580821
39	718	17	25
papers	citations	h-index	g-index
20	20	20	1000
39	39	39	1028
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Association of miR-34a and miR-143 levels with PPAR $\hat{I}^3$ gene expression in adipose tissues of non-diabetic adults. Journal of Physiological Anthropology, 2022, 41, 13.	2.6	9
2	Maternal Exposure to D-galactose Reduces Ovarian Reserve in Female Rat Offspring Later in Life. International Journal of Endocrinology and Metabolism, 2022, 20, .	1.0	2
3	Clinical and Laboratory Characteristics of a Large Iranian Kindred Afflicted with Von Hippel Lindau Disease. International Journal of Endocrinology and Metabolism, 2021, 19, e105189.	1.0	1
4	Biosimilar Gene Therapy: Investigational Assessment of Secukinumab Gene Therapy. Cell Journal, 2020, 21, 433-443.	0.2	0
5	Induced premature ovarian insufficiency by using D galactose and its effects on reproductive profiles in small laboratory animals: a systematic review. Journal of Ovarian Research, 2019, 12, 96.	3.0	9
6	The Principles of Biomedical Scientific Writing: Discussion. International Journal of Endocrinology and Metabolism, 2019, 17, e95415.	1.0	15
7	BRAF V600E mutation and microRNAs are helpful in distinguishing papillary thyroid malignant lesions: Tissues and fine needle aspiration cytology cases. Life Sciences, 2019, 223, 166-173.	4.3	14
8	The Principles of Biomedical Scientific Writing: Materials and Methods. International Journal of Endocrinology and Metabolism, 2019, In Press, e88155.	1.0	8
9	The Principles of Biomedical Scientific Writing: Results. International Journal of Endocrinology and Metabolism, 2019, In Press, e92113.	1.0	10
10	Altered Epigenetic Mechanisms in Thyroid Cancer Subtypes. Molecular Diagnosis and Therapy, 2018, 22, 41-56.	3.8	33
11	The role of matrix metalloproteinase-9 as a prognostic biomarker in papillary thyroid cancer. BMC Cancer, 2018, 18, 1199.	2.6	27
12	The Association of BRAF V600E Mutation With Tissue Inhibitor of Metalloproteinase-3 Expression and Clinicopathological Features in Papillary Thyroid Cancer. International Journal of Endocrinology and Metabolism, 2018, 16, e56120.	1.0	15
13	Hypomethylation of specific CpG sites in the promoter region of steroidogeneic genes (GATA6 and) Tj ETQq $1\ 1\ 0$ .	.784314 rş	gBT/Overloc
14	Effect of long-term nitrite administration on browning of white adipose tissue in type 2 diabetic rats: A stereological study. Life Sciences, 2018, 207, 219-226.	4.3	20
15	Cardio-Metabolic Disease Genetic Risk Factors in Iran: Twenty Years of Tehran Lipid and Glucose Study. International Journal of Endocrinology and Metabolism, 2018, In Press, e84744.	1.0	6
16	Outcomes in the Tehran Lipid and Glucose Study (TLGS) as a Longitudinal Population-Based Cohort Study and a Pragmatic Community Trial. International Journal of Endocrinology and Metabolism, 2018, In Press, e84748.	1.0	31
17	Review of Rationale, Design, and Initial Findings: Tehran Lipid and Glucose Study. International Journal of Endocrinology and Metabolism, 2018, In Press, e84777.	1.0	52
18	Contributions and Implications of the Tehran Lipid and Glucose Study. International Journal of Endocrinology and Metabolism, 2018, 16, e84792.	1.0	29

#	Article	IF	Citations
19	Alteration in follistatin gene expression detected in prenatally androgenized rats. Gynecological Endocrinology, 2017, 33, 433-437.	1.7	3
20	Transient Congenital Hypothyroidism Alters Gene Expression of Glucose Transporters and Impairs Glucose Sensing Apparatus in Young and Aged Offspring Rats. Cellular Physiology and Biochemistry, 2017, 43, 2338-2352.	1.6	21
21	Hormone-induced rat model of polycystic ovary syndrome: A systematic review. Life Sciences, 2017, 191, 259-272.	4.3	33
22	The Role of Kisspeptin in Female Reproduction. International Journal of Endocrinology and Metabolism, 2017, In Press, e44337.	1.0	38
23	Hereditary Vitamin D Resistant Rickets: Clinical, Laboratory, and Genetic Characteristics of 2 Iranian Siblings. International Journal of Endocrinology and Metabolism, 2017, In Press, e12384.	1.0	4
24	Elevated expression of steroidogenesis pathway genes; CYP17, GATA6 and StAR in prenatally androgenized rats. Gene, 2016, 593, 167-171.	2.2	15
25	Sugarâ€sweetened beverage consumption and risk of incident chronic kidney disease: Tehran lipid and glucose study. Nephrology, 2016, 21, 608-616.	1.6	29
26	Involvement of inducible nitric oxide synthase in the loss of cardioprotection by ischemic postconditioning in hypothyroid rats. Gene, 2016, 580, 169-176.	2.2	18
27	Consumption of nitrate-containing vegetables is inversely associated with hypertension in adults: a prospective investigation from the Tehran Lipid and Glucose Study. Journal of Nephrology, 2016, 29, 377-384.	2.0	25
28	Genetic polymorphism of vitamin D receptor gene affects the phenotype of PCOS. Gene, 2013, 515, 193-196.	2.2	44
29	Sistani Population: a Different Spectrum oF $\hat{I}^2$ -Thalassemia Mutations From other Ethnic Groups of Iran. Hemoglobin, 2013, 37, 138-147.	0.8	17
30	Association of <i>CD36 </i> Gene Variants and Metabolic Syndrome in Iranians. Genetic Testing and Molecular Biomarkers, 2012, 16, 234-238.	0.7	2
31	Profile of $\hat{I}^2$ -Thalassemia and its Prenatal Diagnosis in Khorasan-E-Jonobi Province, Iran. Hemoglobin, 2012, 36, 456-463.	0.8	2
32	Chromosome abnormality rate among Iranian patients with idiopathic mental retardation from consanguineous marriages. Archives of Medical Science, 2011, 2, 321-325.	0.9	11
33	Waist circumference and insulin resistance: a community based cross sectional study on reproductive aged Iranian women. Diabetology and Metabolic Syndrome, 2011, 3, 18.	2.7	40
34	Detection of a Rare Mutation in An Iranian Family: Codons 37/38/39 (7 bp Deletion). Hemoglobin, 2009, 33, 523-527.	0.8	1
35	Molecular characterization of thalassemia intermedia, due to co-inheritance of homozygous alpha triplication and IVSI-5 β-thalassemia. Blood Cells, Molecules, and Diseases, 2009, 43, 158-160.	1.4	12
36	Molecular analysis of sixteen unrelated factor XIIIA deficient families from south-east of Iran. British Journal of Haematology, 2008, 140, 581-584.	2.5	24

#	Article	IF	CITATIONS
37	An Unusually Frequent Î <sup>2</sup> -Thalassemia Mutation in an Iranian Province. Hemoglobin, 2008, 32, 387-392.	0.8	16
38	Hematological Phenotype of the IVS-I-5 (G > C) β-Thalassemia Mutation and Assessment of Iran's Nation Screening Criteria. Hemoglobin, 2008, 32, 440-445.	na  0.8	5
39	Immunization with the hybrid protein vaccine, consisting of Leishmania major cysteine proteinases Type I (CPB) and Type II (CPA), partially protects against leishmaniasis. Vaccine, 2004, 22, 1930-1940.	3.8	59