

Kepa B Uribe

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

34
papers

592
citations

12
h-index

24
g-index

44
ext. papers

1,025
ext. citations

5.7
avg, IF

4.09
L-index

| # | Paper | IF | Citations |
|----|--|-----|-----------|
| 34 | Boosting Cholesterol Efflux from Foam Cells by Sequential Administration of rHDL to Deliver MicroRNA and to Remove Cholesterol in a Triple-Cell 2D Atherosclerosis Model.. <i>Small</i> , 2022 , e2105915 | 11 | 1 |
| 33 | Familial hypercholesterolemia 2022 , 501-524 | | |
| 32 | MLb-LDLr: A Machine Learning Model for Predicting the Pathogenicity of Missense Variants. <i>JACC Basic To Translational Science</i> , 2021 , 6, 815-827 | 8.7 | 0 |
| 31 | Evaluation of Multifunctional Gold Nanorods for Boron Neutron Capture and Photothermal Therapies. <i>ACS Applied Materials & Interfaces</i> , 2021 , 13, 49589-49601 | 9.5 | 4 |
| 30 | Cholesterol stimulates the lytic activity of Adenylate Cyclase Toxin on lipid membranes by promoting toxin oligomerization and formation of pores with a greater effective size. <i>FEBS Journal</i> , 2021 , 288, 6795-6814 | 5.7 | 0 |
| 29 | Novel PCSK9 (Proprotein Convertase Subtilisin Kexin Type 9) Variants in Patients With Familial Hypercholesterolemia From Cape Town. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2021 , 41, 934-943 | 9.4 | 3 |
| 28 | Pre-targeting with ultra-small nanoparticles: boron carbon dots as drug candidates for boron neutron capture therapy. <i>Journal of Materials Chemistry B</i> , 2021 , 9, 410-420 | 7.3 | 7 |
| 27 | Molecular mechanisms of lipotoxicity-induced pancreatic β cell dysfunction. <i>International Review of Cell and Molecular Biology</i> , 2021 , 359, 357-402 | 6 | 6 |
| 26 | (r)HDL in theranostics: how do we apply HDL biology for precision medicine in atherosclerosis management?. <i>Biomaterials Science</i> , 2021 , 9, 3185-3208 | 7.4 | 1 |
| 25 | A Systematic Approach to Assess the Activity and Classification of PCSK9 Variants.. <i>International Journal of Molecular Sciences</i> , 2021 , 22, | 6.3 | 4 |
| 24 | Statin Treatment-Induced Development of Type 2 Diabetes: From Clinical Evidence to Mechanistic Insights. <i>International Journal of Molecular Sciences</i> , 2020 , 21, | 6.3 | 16 |
| 23 | Mutation type classification and pathogenicity assignment of sixteen missense variants located in the EGF-precursor homology domain of the LDLR. <i>Scientific Reports</i> , 2020 , 10, 1727 | 4.9 | 12 |
| 22 | Cholesterol Efflux Efficiency of Reconstituted HDL Is Affected by Nanoparticle Lipid Composition. <i>Biomedicines</i> , 2020 , 8, | 4.8 | 5 |
| 21 | miR-27b Modulates Insulin Signaling in Hepatocytes by Regulating Insulin Receptor Expression. <i>International Journal of Molecular Sciences</i> , 2020 , 21, | 6.3 | 7 |
| 20 | Pathophysiology of Type 2 Diabetes Mellitus. <i>International Journal of Molecular Sciences</i> , 2020 , 21, | 6.3 | 222 |
| 19 | Functional Analysis of LDLR (Low-Density Lipoprotein Receptor) Variants in Patient Lymphocytes to Assess the Effect of Evinacumab in Homozygous Familial Hypercholesterolemia Patients With a Spectrum of LDLR Activity. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2019 , 39, 2248-2260 | 9.4 | 33 |
| 18 | Membrane Permeabilization by Pore-Forming RTX Toxins: What Kind of Lesions Do These Toxins Form?. <i>Toxins</i> , 2019 , 11, | 4.9 | 16 |

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|----|--|-----|----|
| 17 | Membrane Permeabilization by Adenylate Cyclase Toxin Involves Pores of Tunable Size. <i>Biomolecules</i> , 2019 , 9, | 5.9 | 6 |
| 16 | The Arg499His gain-of-function mutation in the C-terminal domain of PCSK9. <i>Atherosclerosis</i> , 2019 , 289, 162-172 | 3.1 | 12 |
| 15 | Site-specific -glycosylation of members of the low-density lipoprotein receptor superfamily enhances ligand interactions. <i>Journal of Biological Chemistry</i> , 2018 , 293, 7408-7422 | 5.4 | 38 |
| 14 | Validation of LDLr Activity as a Tool to Improve Genetic Diagnosis of Familial Hypercholesterolemia: A Retrospective on Functional Characterization of LDLr Variants. <i>International Journal of Molecular Sciences</i> , 2018 , 19, | 6.3 | 22 |
| 13 | Familial Hypercholesterolemia: The Most Frequent Cholesterol Metabolism Disorder Caused Disease. <i>International Journal of Molecular Sciences</i> , 2018 , 19, | 6.3 | 36 |
| 12 | p.(Asp47Asn) and p.(Thr62Met): non deleterious LDL receptor missense variants functionally characterized in vitro. <i>Scientific Reports</i> , 2018 , 8, 16614 | 4.9 | 3 |
| 11 | Replacement of cysteine at position 46 in the first cysteine-rich repeat of the LDL receptor impairs apolipoprotein recognition. <i>PLoS ONE</i> , 2018 , 13, e0204771 | 3.7 | |
| 10 | Analysis of LDLR variants from homozygous FH patients carrying multiple mutations in the LDLR gene. <i>Atherosclerosis</i> , 2017 , 263, 163-170 | 3.1 | 4 |
| 9 | The leucine stretch length of PCSK9 signal peptide and its role in development of autosomal dominant hypercholesterolaemia: Unravelling the activities of P.LEU23DEL and P.LEU22_LEU23DUP variants. <i>Atherosclerosis</i> , 2017 , 263, e37 | 3.1 | 3 |
| 8 | Phospholipase A activity of adenylate cyclase toxin mediates translocation of its adenylate cyclase domain. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, E6784-E6793 ^{11,5,18} | | |
| 7 | Identification and in vitro characterization of two new PCSK9 Gain of Function variants found in patients with Familial Hypercholesterolemia. <i>Scientific Reports</i> , 2017 , 7, 15282 | 4.9 | 24 |
| 6 | Understanding the Mechanism of Translocation of Adenylate Cyclase Toxin across Biological Membranes. <i>Toxins</i> , 2017 , 9, | 4.9 | 7 |
| 5 | Adenylate Cyclase Toxin promotes bacterial internalisation into non phagocytic cells. <i>Scientific Reports</i> , 2015 , 5, 13774 | 4.9 | 8 |
| 4 | Calpain-Mediated Processing of Adenylate Cyclase Toxin Generates a Cytosolic Soluble Catalytically Active N-Terminal Domain. <i>PLoS ONE</i> , 2013 , 8, e67648 | 3.7 | 15 |
| 3 | Ca ²⁺ influx and tyrosine kinases trigger Bordetella adenylate cyclase toxin (ACT) endocytosis. Cell physiology and expression of the CD11b/CD18 integrin major determinants of the entry route. <i>PLoS ONE</i> , 2013 , 8, e74248 | 3.7 | 5 |
| 2 | Functional characterization of splicing and ligand-binding domain variants in the LDL receptor. <i>Human Mutation</i> , 2012 , 33, 232-43 | 4.7 | 33 |
| 1 | Adenylate cyclase toxin promotes internalisation of integrins and raft components and decreases macrophage adhesion capacity. <i>PLoS ONE</i> , 2011 , 6, e17383 | 3.7 | 12 |