Sriram Ravindran

List of Publications by Year in descending order

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1039880 839398 19 322 9 18 citations h-index g-index papers 19 19 19 409 citing authors docs citations times ranked all docs

#	Article	lF	Citations
1	Fisetin Confers Cardioprotection against Myocardial Ischemia Reperfusion Injury by Suppressing Mitochondrial Oxidative Stress and Mitochondrial Dysfunction and Inhibiting Glycogen Synthase Kinase $3 < i > \hat{l}^2 < i>$ Activity. Oxidative Medicine and Cellular Longevity, 2018, 2018, 1-16.	1.9	64
2	Sodium Thiosulfate Preconditioning Ameliorates Ischemia/Reperfusion Injury in Rat Hearts Via Reduction of Oxidative Stress and Apoptosis. Cardiovascular Drugs and Therapy, 2017, 31, 511-524.	1.3	41
3	Sodium thiosulfate post-conditioning protects rat hearts against ischemia reperfusion injury via reduction of apoptosis and oxidative stress. Chemico-Biological Interactions, 2017, 274, 24-34.	1.7	41
4	Hydrogen sulfide post-conditioning preserves interfibrillar mitochondria of rat heart during ischemia reperfusion injury. Cell Stress and Chaperones, 2016, 21, 571-582.	1.2	29
5	Role of endogenous hydrogen sulfide in cardiac mitochondrial preservation during ischemia reperfusion injury. Biomedicine and Pharmacotherapy, 2018, 97, 271-279.	2.5	20
6	Effect of Sodium Thiosulfate Postconditioning on Ischemia-Reperfusion Injury Induced Mitochondrial Dysfunction in Rat Heart. Journal of Cardiovascular Translational Research, 2018, 11, 246-258.	1.1	18
7	Preconditioning the rat heart with sodium thiosulfate preserved the mitochondria in response to ischemia-reperfusion injury. Journal of Bioenergetics and Biomembranes, 2019, 51, 189-201.	1.0	15
8	Hydrogen sulfide preconditioning shows differential protection towards interfibrillar and subsarcolemmal mitochondria from isolated rat heart subjected to revascularization injury. Cardiovascular Pathology, 2016, 25, 306-315.	0.7	13
9	Nicorandil attenuates neuronal mitochondrial dysfunction and oxidative stress associated with murine model of vascular calcification. Acta Neurobiologiae Experimentalis, 2017, 77, 57-67.	0.4	12
10	Eventual analysis of global cerebral ischemia-reperfusion injury in rat brain: a paradigm of a shift in stress and its influence on cognitive functions. Cell Stress and Chaperones, 2019, 24, 581-594.	1.2	11
11	Vascular calcification abrogates the nicorandil mediated cardio-protection in ischemia reperfusion injury of rat heart. Vascular Pharmacology, 2017, 89, 31-38.	1.0	9
12	The role of secretory phospholipases as therapeutic targets for the treatment of myocardial ischemia reperfusion injury. Biomedicine and Pharmacotherapy, 2017, 92, 7-16.	2.5	9
13	Renal mitochondria can withstand hypoxic/ischemic injury secondary to renal failure in uremic rats pretreated with sodium thiosulfate. Indian Journal of Pharmacology, 2017, 49, 317.	0.4	9
14	The renal mitochondrial dysfunction in patients with vascular calcification is prevented by sodium thiosulfate. International Urology and Nephrology, 2016, 48, 1927-1935.	0.6	7
15	Beneficial effect of sodium thiosulfate extends beyond myocardial tissue in isoproterenol model of infarction: Implication for nootropic effects. Journal of Biochemical and Molecular Toxicology, 2020, 34, e22606.	1.4	7
16	Sodium thiosulfate mediated cardioprotection against myocardial ischemia-reperfusion injury is defunct in rat heart with co-morbidity of vascular calcification. Biochimie, 2018, 147, 80-88.	1.3	6
17	Addressing the alterations in cerebral ischemia-reperfusion injury on the brain mitochondrial activity: A possible link to cognitive decline. Biochemical and Biophysical Research Communications, 2019, 518, 100-106.	1.0	6
18	Hydrogen sulfide-mediated cardioprotection against ischemia reperfusion is linked to KATP channel for mitochondrial preservation but not for its distinct preference on interfibrillar mitochondria. Bangladesh Journal of Pharmacology, 2019, 14, 107-115.	0.1	3

#	Article	IF	CITATIONS
19	Erythrocyte Membrane Bound ATPase and Antioxidant Enzyme Changes Associated with Vascular Calcification is Reduced by Sodium Thiosulfate. Indian Journal of Clinical Biochemistry, 2017, 32, 487-492.	0.9	2