

# Teresa Pasqua

## List of Publications by Year in descending order

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Version: 2024-02-01

23  
papers

609  
citations

567281

15  
h-index

642732

23  
g-index

23  
all docs

23  
docs citations

23  
times ranked

874  
citing authors

#	ARTICLE	IF	CITATIONS
1	The chromogranin A 1-73 fragment reveals how a single change in the protein sequence exerts strong cardioregulatory effects by engaging neuropilin-1. <i>Acta Physiologica</i> , 2021, 231, e13570.	3.8	14
2	Cateslytin abrogates lipopolysaccharide-induced cardiomyocyte injury by reducing inflammation and oxidative stress through toll like receptor 4 interaction. <i>International Immunopharmacology</i> , 2021, 94, 107487.	3.8	16
3	Immunosuppression of Macrophages Underlies the Cardioprotective Effects of CST (Catestatin). <i>Hypertension</i> , 2021, 77, 1670-1682.	2.7	31
4	PI3K $\gamma$ Inhibition as a Potential Therapeutic Target in COVID-19. <i>Frontiers in Immunology</i> , 2020, 11, 2094.	4.8	23
5	Cardiac and Metabolic Impact of Functional Foods with Antioxidant Properties Based on Whey Derived Proteins Enriched with Hemp Seed Oil. <i>Antioxidants</i> , 2020, 9, 1066.	5.1	13
6	Nesfatin-1 in cardiovascular orchestration: From bench to bedside. <i>Pharmacological Research</i> , 2020, 156, 104766.	7.1	11
7	Mechanisms and Pathophysiology of Obesity: Upgrading a Complex Scenario. <i>Current Medicinal Chemistry</i> , 2020, 27, 172-173.	2.4	2
8	Cardiac Damage in Anthracyclines Therapy: Focus on Oxidative Stress and Inflammation. <i>Antioxidants and Redox Signaling</i> , 2020, 32, 1081-1097.	5.4	40
9	Modulation of the coronary tone in the expanding scenario of Chromogranin-A and its derived peptides. <i>Future Medicinal Chemistry</i> , 2019, 11, 1501-1511.	2.3	7
10	Progress in the emerging role of selenoproteins in cardiovascular disease: focus on endoplasmic reticulum-resident selenoproteins. <i>Cellular and Molecular Life Sciences</i> , 2019, 76, 3969-3985.	5.4	53
11	Physiological levels of chromogranin A prevent doxorubicin-induced cardiotoxicity without impairing its anticancer activity. <i>FASEB Journal</i> , 2019, 33, 7734-7747.	0.5	20
12	Role of Brain Neuroinflammatory Factors on Hypertension in the Spontaneously Hypertensive Rat. <i>Neuroscience</i> , 2018, 375, 158-168.	2.3	17
13	Notch1 Mediates Preconditioning Protection Induced by GPER in Normotensive and Hypertensive Female Rat Hearts. <i>Frontiers in Physiology</i> , 2018, 9, 521.	2.8	32
14	Role of NLRP-3 Inflammasome in Hypertension: A Potential Therapeutic Target. <i>Current Pharmaceutical Biotechnology</i> , 2018, 19, 708-714.	1.6	44
15	Granin-derived peptides. <i>Progress in Neurobiology</i> , 2017, 154, 37-61.	5.7	65
16	Protective Role of GPER Agonist G6615 on Cardiotoxicity Induced by Doxorubicin. <i>Journal of Cellular Physiology</i> , 2017, 232, 1640-1649.	4.1	46
17	Biological Roles of the Eclectic Chromogranin-A-derived Peptide Catestatin. <i>Current Medicinal Chemistry</i> , 2017, 24, 3356-3372.	2.4	8
18	Cardiac and hepatic role of HSP70: basal effects and protection against ischemic and sepsis conditions. <i>Journal of Cellular and Molecular Medicine</i> , 2015, 19, 1492-1503.	3.6	13

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19	Indenopyrazole oxime ethers: Synthesis and $\beta$ 1-adrenergic blocking activity. <i>European Journal of Medicinal Chemistry</i> , 2015, 92, 672-681.	5.5	21
20	Chromofungin, CgA47-66-derived peptide, produces basal cardiac effects and postconditioning cardioprotective action during ischemia/reperfusion injury. <i>Peptides</i> , 2015, 71, 40-48.	2.4	26
21	Catestatin Increases the Expression of Anti-Apoptotic and Pro-Angiogenetic Factors in the Post-Ischemic Hypertrophied Heart of SHR. <i>PLoS ONE</i> , 2014, 9, e102536.	2.5	29
22	Full-Length Human Chromogranin-A Cardioactivity: Myocardial, Coronary, and Stimulus-Induced Processing Evidence in Normotensive and Hypertensive Male Rat Hearts. <i>Endocrinology</i> , 2013, 154, 3353-3365.	2.8	41
23	Phosphodiesterase type-2 and NO-dependent S-nitrosylation mediate the cardioinhibition of the antihypertensive catestatin. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2012, 302, H431-H442.	3.2	37