Charles F Reese

List of Publications by Year in descending order

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Version: 2024-02-01

1163117 1281871 11 221 8 11 citations h-index g-index papers 11 11 11 364 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Bleomycin delivery by osmotic minipump: similarity to human scleroderma interstitial lung disease. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2014, 306, L736-L748.	2.9	50
2	Enhanced chemokine-receptor expression, function, and signaling in healthy African American and scleroderma-patient monocytes are regulated by caveolin-1. Fibrogenesis and Tissue Repair, 2015, 8, 11.	3.4	32
3	Caveolinâ€1 Deficiency May Predispose African Americans to Systemic Sclerosis–Related Interstitial Lung Disease. Arthritis and Rheumatology, 2014, 66, 1909-1919.	5.6	27
4	Caveolin-1 regulates chemokine receptor 5-mediated contribution of bone marrow-derived cells to dermal fibrosis. Frontiers in Pharmacology, 2014, 5, 140.	3.5	24
5	Fibrocytes in the fibrotic lung: altered phenotype detected by flow cytometry. Frontiers in Pharmacology, 2014, 5, 141.	3.5	23
6	Suppression of angiotensin II-induced pathological changes in heart and kidney by the caveolin-1 scaffolding domain peptide. PLoS ONE, 2018, 13, e0207844.	2.5	19
7	Reversal of maladaptive fibrosis and compromised ventricular function in the pressure overloaded heart by a caveolin-1 surrogate peptide. Laboratory Investigation, 2017, 97, 370-382.	3.7	16
8	Adipose-derived mesenchymal stromal/stem cells in systemic sclerosis: Alterations in function and beneficial effect on lung fibrosis are regulated by caveolin-1. Journal of Scleroderma and Related Disorders, 2019, 4, 127-136.	1.7	11
9	The Caveolin-1 Scaffolding Domain Peptide Reverses Aging-Associated Deleterious Changes in Multiple Organs. Journal of Pharmacology and Experimental Therapeutics, 2021, 378, 1-9.	2.5	8
10	Differential regulation of cell functions by CSD peptide subdomains. Respiratory Research, 2013, 14, 90.	3.6	7
11	Multiple subregions within the caveolin-1 scaffolding domain inhibit fibrosis, microvascular leakage, and monocyte migration. PLoS ONE, 2022, 17, e0264413.	2.5	4