Shuichiro Okamoto

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2047450/publications.pdf

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10 papers	104 citations	1478505 6 h-index	9 g-index
10	10	10	196
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Withaferin A suppresses the growth of myelodysplasia and leukemia cell lines by inhibiting cell cycle progression. Cancer Science, 2016, 107, 1302-1314.	3.9	35
2	Five-aza-2′-deoxycytidine-induced hypomethylation of cholesterol 25-hydroxylase gene is responsible for cell death of myelodysplasia/leukemia cells. Scientific Reports, 2015, 5, 16709.	3.3	20
3	The NADPH oxidase NOX4 promotes the directed migration of endothelial cells by stabilizing vascular endothelial growth factor receptor 2 protein. Journal of Biological Chemistry, 2020, 295, 11877-11890.	3.4	12
4	An MDS-derived cell line and a series of its sublines serve as an in vitro model for the leukemic evolution of MDS. Leukemia, 2018, 32, 1846-1850.	7.2	11
5	The rRNA synthesis inhibitor CX-5461 may induce autophagy that inhibits anticancer drug-induced cell damage to leukemia cells. Bioscience, Biotechnology and Biochemistry, 2020, 84, 2319-2326.	1.3	8
6	Constitutive activity of NADPH oxidase 1 (Nox1) that promotes its own activity suppresses the colon epithelial cell migration. Free Radical Research, 2020, 54, 640-648.	3.3	7
7	Coculture inÂvitro with endothelial cells induces cytarabine resistance of acute myeloid leukemia cells in a VEGF-A/VEGFR-2 signaling–independent manner. Biochemical and Biophysical Research Communications, 2022, 587, 78-84.	2.1	6
8	The downregulation of NADPH oxidase Nox4 during hypoxia in hemangioendothelioma cells: a possible role of p22 <i>^{phox}</i> on Nox4 protein stability. Free Radical Research, 2021, 55, 996-1004.	3.3	3
9	Fine definition of the epitopes on the human gp91 /NOX2 for the monoclonal antibodies CL-5 and 48. Journal of Immunological Methods, 2022, 501, 113213.	1.4	2
10	Malignant Progression of an MDS-Derived Cell Line Serves As an in Vitro Model for the Leukemic Evolution of MDS. Blood, 2018, 132, 5501-5501.	1.4	O