

Nicola Brownlow

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

256
citations

1163117

8
h-index

1199594

12
g-index

13
all docs

13
docs citations

13
times ranked

511
citing authors

#	ARTICLE	IF	CITATIONS
1	The Aurora B specificity switch is required to protect from non-disjunction at the metaphase/anaphase transition. <i>Nature Communications</i> , 2020, 11, 1396.	12.8	12
2	STAT4 expression and activation is increased during mitosis <i>in vitro</i> and <i>in vivo</i> in skin- and mucosa-derived cell types: implications in neoplastic and inflammatory skin diseases. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2017, 31, 1663-1673.	2.4	4
3	DNA replication stress: NoCut to the rescue. <i>Cell Cycle</i> , 2017, 16, 233-234.	2.6	7
4	PKC ϵ switches Aurora B specificity to exit the abscission checkpoint. <i>Nature Communications</i> , 2016, 7, 13853.	12.8	21
5	Mitotic catenation is monitored and resolved by a PKC μ -regulated pathway. <i>Nature Communications</i> , 2014, 5, 5685.	12.8	21
6	Regulation of the cytokinesis cleavage furrow by PKC μ . <i>Biochemical Society Transactions</i> , 2014, 42, 1534-1537.	3.4	5
7	Novel Imatinib Derivatives with Altered Specificity between Bcr-Abl and FMS, KIT, and PDGF Receptors. <i>ChemMedChem</i> , 2010, 5, 130-139.	3.2	17
8	Protein kinase C epsilon in cell division: Control of abscission. <i>Cell Cycle</i> , 2009, 8, 549-555.	2.6	16
9	Dasatinib is a potent inhibitor of tumour-associated macrophages, osteoclasts and the FMS receptor. <i>Leukemia</i> , 2009, 23, 590-594.	7.2	59
10	Comparison of nilotinib and imatinib inhibition of FMS receptor signaling, macrophage production and osteoclastogenesis. <i>Leukemia</i> , 2008, 22, 649-652.	7.2	32
11	Tandutinib inhibits FMS receptor signalling, and macrophage and osteoclast formation <i>in vitro</i> . <i>Leukemia</i> , 2008, 22, 1452-1453.	7.2	6
12	FMS receptor for M-CSF (CSF-1) is sensitive to the kinase inhibitor imatinib and mutation of Asp-802 to Val confers resistance. <i>Oncogene</i> , 2006, 25, 147-151.	5.9	56