Che-Ming Teng

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

 431
 13,697
 60
 81

 papers
 citations
 h-index
 g-index

 443
 14,490
 5.1
 5.81

 ext. papers
 ext. citations
 avg, IF
 L-index

#	Paper	IF	Citations
431	Ring-opening of five-membered heterocycles conjugated 4-isopropylresorcinol scaffold-based benzamides as HSP90 inhibitors suppressing tumor growth in vitro and in vivo. <i>European Journal of Medicinal Chemistry</i> , 2021 , 219, 113428	6.8	2
430	Retraction: Non-epigenetic function of HDAC8 in regulating breast cancer stem cells by maintaining Notch1 protein stability. <i>Oncotarget</i> , 2020 , 11, 1096	3.3	
429	Combination treatment strategy for pancreatic cancer involving the novel HDAC inhibitor MPT0E028 with a MEK inhibitor beyond K-Ras status. <i>Clinical Epigenetics</i> , 2019 , 11, 85	7.7	10
428	Lanatoside C, a cardiac glycoside, acts through protein kinase CIto cause apoptosis of human hepatocellular carcinoma cells. <i>Scientific Reports</i> , 2017 , 7, 46134	4.9	23
427	The apoptotic mechanisms of MT-6, a mitotic arrest inducer, in human ovarian cancer cells. <i>Scientific Reports</i> , 2017 , 7, 46149	4.9	
426	An oral quinoline derivative, MPT0B392, causes leukemic cells mitotic arrest and overcomes drug resistant cancer cells. <i>Oncotarget</i> , 2017 , 8, 27772-27785	3.3	4
425	ERK Activation Globally Downregulates miRNAs through Phosphorylating Exportin-5. <i>Cancer Cell</i> , 2016 , 30, 723-736	24.3	96
424	MPT0G066, a novel anti-mitotic drug, induces JNK-independent mitotic arrest, JNK-mediated apoptosis, and potentiates antineoplastic effect of cisplatin in ovarian cancer. <i>Scientific Reports</i> , 2016 , 6, 31664	4.9	10
423	LTP-1, a novel antimitotic agent and Stat3 inhibitor, inhibits human pancreatic carcinomas in vitro and in vivo. <i>Scientific Reports</i> , 2016 , 6, 27794	4.9	8
422	TW-01, a piperazinedione-derived compound, inhibits Ras-mediated cell proliferation and angioplasty-induced vascular restenosis. <i>Toxicology and Applied Pharmacology</i> , 2016 , 305, 194-202	4.6	
421	Novel histone deacetylase inhibitor MPT0G009 induces cell apoptosis and synergistic anticancer activity with tumor necrosis factor-related apoptosis-inducing ligand against human hepatocellular carcinoma. <i>Oncotarget</i> , 2016 , 7, 402-17	3.3	17
420	Non-epigenetic function of HDAC8 in regulating breast cancer stem cells by maintaining Notch1 protein stability. <i>Oncotarget</i> , 2016 , 7, 1796-807	3.3	21
419	Integrin-linked kinase as a novel molecular switch of the IL-6-NF- B signaling loop in breast cancer. <i>Carcinogenesis</i> , 2016 , 37, 430-442	4.6	16
418	N-Sulfonyl-aminobiaryls as Antitubulin Agents and Inhibitors of Signal Transducers and Activators of Transcription 3 (STAT3) Signaling. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 6549-58	8.3	22
417	The synergic effect of vincristine and vorinostat in leukemia in vitro and in vivo. <i>Journal of Hematology and Oncology</i> , 2015 , 8, 82	22.4	45
416	Heteronemin, a Spongean Sesterterpene, Induces Cell Apoptosis and Autophagy in Human Renal Carcinoma Cells. <i>BioMed Research International</i> , 2015 , 2015, 738241	3	23
415	Dual phosphoinositide 3-kinase/mammalian target of rapamycin inhibitor is an effective radiosensitizer for colorectal cancer. <i>Cancer Letters</i> , 2015 , 357, 582-90	9.9	35

(2013-2015)

414	Novel oral histone deacetylase inhibitor, MPT0E028, displays potent growth-inhibitory activity against human B-cell lymphoma in vitro and in vivo. <i>Oncotarget</i> , 2015 , 6, 4976-91	3.3	13
413	eIF4E binding protein 1 expression is associated with clinical survival outcomes in colorectal cancer. <i>Oncotarget</i> , 2015 , 6, 24092-104	3.3	15
412	Molecular mechanisms underlying the antitumor activity of (E)-N-hydroxy-3-(1-(4-methoxyphenylsulfonyl)-1,2,3,4-tetrahydroquinolin-6-yl)acrylamide in human colorectal cancer cells in vitro and in vivo. <i>Oncotarget</i> , 2015 , 6, 35991-6002	3.3	5
411	YC-1 inhibits proliferation of breast cancer cells by down-regulating EZH2 expression via activation of c-Cbl and ERK. <i>British Journal of Pharmacology</i> , 2014 , 171, 4010-25	8.6	22
410	Effect of C7-substitution of 1-arylsulfonyl-5-(N-hydroxyacrylamide)indolines on the selectivity towards a subclass of histone deacetylases. <i>Organic and Biomolecular Chemistry</i> , 2014 , 12, 8966-76	3.9	7
409	Design, synthesis, mechanisms of action, and toxicity of novel 20(s)-sulfonylamidine derivatives of camptothecin as potent antitumor agents. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 6008-18	8.3	54
408	Indole-3-ethylsulfamoylphenylacrylamides: potent histone deacetylase inhibitors with anti-inflammatory activity. <i>European Journal of Medicinal Chemistry</i> , 2014 , 85, 468-79	6.8	35
407	Synthesis of analogues of gingerol and shogaol, the active pungent principles from the rhizomes of Zingiber officinale and evaluation of their anti-platelet aggregation effects. <i>International Journal of Molecular Sciences</i> , 2014 , 15, 3926-51	6.3	30
406	In vitro and in vivo anti-tumour effects of MPT0B014, a novel derivative aroylquinoline, and in combination with erlotinib in human non-small-cell lung cancer cells. <i>British Journal of Pharmacology</i> , 2014 , 171, 122-33	8.6	6
405	Radiation-induced VEGF-C expression and endothelial cell proliferation in lung cancer. <i>Strahlentherapie Und Onkologie</i> , 2014 , 190, 1154-62	4.3	27
404	Synergistic interaction between the HDAC inhibitor, MPT0E028, and sorafenib in liver cancer cells in vitro and in vivo. <i>Clinical Cancer Research</i> , 2014 , 20, 1274-1287	12.9	37
403	NPRL-Z-1, as a new topoisomerase II poison, induces cell apoptosis and ROS generation in human renal carcinoma cells. <i>PLoS ONE</i> , 2014 , 9, e112220	3.7	8
402	A novel class I HDAC inhibitor, MPT0G030, induces cell apoptosis and differentiation in human colorectal cancer cells via HDAC1/PKCland E-cadherin. <i>Oncotarget</i> , 2014 , 5, 5651-62	3.3	26
401	A novel action mechanism for MPT0G013, a derivative of arylsulfonamide, inhibits tumor angiogenesis through up-regulation of TIMP3 expression. <i>Oncotarget</i> , 2014 , 5, 9838-50	3.3	10
400	Moscatilin inhibits migration and metastasis of human breast cancer MDA-MB-231 cells through inhibition of Akt and Twist signaling pathway. <i>Journal of Molecular Medicine</i> , 2013 , 91, 347-56	5.5	35
399	Denbinobin induces human glioblastoma multiforme cell apoptosis through the IKKEAkt-FKHR signaling cascade. <i>European Journal of Pharmacology</i> , 2013 , 698, 103-9	5.3	8
398	Activated PAR-2 regulates pancreatic cancer progression through ILK/HIF-Enduced TGF-Expression and MEK/VEGF-A-mediated angiogenesis. <i>American Journal of Pathology</i> , 2013 , 183, 566-75	5.8	35
397	TW01001, a novel piperazinedione compound, induces mitotic arrest and autophagy in non-small cell lung cancer A549 cells. <i>Cancer Letters</i> , 2013 , 336, 370-8	9.9	12

396	1-arylsulfonyl-5-(N-hydroxyacrylamide)indolines histone deacetylase inhibitors are potent cytokine release suppressors. <i>ChemBioChem</i> , 2013 , 14, 1248-54	3.8	18
395	Depletion of 4E-BP1 and regulation of autophagy lead to YXM110-induced anticancer effects. <i>Carcinogenesis</i> , 2013 , 34, 2050-60	4.6	7
394	Azatyrosinamides: novel RAS-related anticancer agents. <i>Anticancer Research</i> , 2013 , 33, 425-32	2.3	1
393	Synthesis and biological evaluation of 1-arylsulfonyl-5-(N-hydroxyacrylamide)indoles as potent histone deacetylase inhibitors with antitumor activity in vivo. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 3777-91	8.3	45
392	Cytotoxic apigenin derivatives from Chrysopogon aciculatis. <i>Journal of Natural Products</i> , 2012 , 75, 198	-2015	21
391	Protopine, a novel microtubule-stabilizing agent, causes mitotic arrest and apoptotic cell death in human hormone-refractory prostate cancer cell lines. <i>Cancer Letters</i> , 2012 , 315, 1-11	9.9	42
390	Aciculatin induces p53-dependent apoptosis via MDM2 depletion in human cancer cells in vitro and in vivo. <i>PLoS ONE</i> , 2012 , 7, e42192	3.7	24
389	Anticancer activity of MPT0E028, a novel potent histone deacetylase inhibitor, in human colorectal cancer HCT116 cells in vitro and in vivo. <i>PLoS ONE</i> , 2012 , 7, e43645	3.7	23
388	Dehydrocostuslactone suppresses angiogenesis in vitro and in vivo through inhibition of Akt/GSK-3land mTOR signaling pathways. <i>PLoS ONE</i> , 2012 , 7, e31195	3.7	33
387	Antitumor agents 295. E-ring hydroxylated antofine and cryptopleurine analogues as antiproliferative agents: design, synthesis, and mechanistic studies. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 6751-61	8.3	21
386	DYZ-2-90, a novel neo-tanshinlactone ring-opened compound, induces ERK-mediated mitotic arrest and subsequent apoptosis by activating JNK in human colorectal cancer cells. <i>ChemBioChem</i> , 2012 , 13, 1663-72	3.8	7
385	QS-ZYX-1-61 induces apoptosis through topoisomerase II in human non-small-cell lung cancer A549 cells. <i>Cancer Science</i> , 2012 , 103, 80-7	6.9	13
384	Physalin F induces cell apoptosis in human renal carcinoma cells by targeting NF-kappaB and generating reactive oxygen species. <i>PLoS ONE</i> , 2012 , 7, e40727	3.7	37
383	Aciculatin inhibits granulocyte colony-stimulating factor production by human interleukin 1卧timulated fibroblast-like synoviocytes. <i>PLoS ONE</i> , 2012 , 7, e42389	3.7	8
382	From tine-mediated anti-cancer activity in vitro and in vivo through cell cycle- and caspase-independent pathways. <i>PLoS ONE</i> , 2012 , 7, e44093	3.7	24
381	Histone deacetylase inhibitors stimulate histone H3 lysine 4 methylation in part via transcriptional repression of histone H3 lysine 4 demethylases. <i>Molecular Pharmacology</i> , 2011 , 79, 197-206	4.3	97
380	Combined blockade of thrombin anion binding exosite-1 and PAR4 produces synergistic antiplatelet effect in human platelets. <i>Thrombosis and Haemostasis</i> , 2011 , 105, 88-95	7	14
379	The inhibition of angiogenesis and tumor growth by denbinobin is associated with the blocking of insulin-like growth factor-1 receptor signaling. <i>Journal of Nutritional Biochemistry</i> , 2011 , 22, 625-33	6.3	9

(2009-2011)

378	Aciculatin inhibits lipopolysaccharide-mediated inducible nitric oxide synthase and cyclooxygenase-2 expression via suppressing NF- B and JNK/p38 MAPK activation pathways. Journal of Biomedical Science, 2011 , 18, 28	13.3	54
377	Thrombin induces expression of twist and cell motility via the hypoxia-inducible factor-1 translational pathway in colorectal cancer cells. <i>Journal of Cellular Physiology</i> , 2011 , 226, 1060-8	7	31
376	Novel mechanism by which histone deacetylase inhibitors facilitate topoisomerase III degradation in hepatocellular carcinoma cells. <i>Hepatology</i> , 2011 , 53, 148-59	11.2	33
375	Antitumor agents 288: design, synthesis, SAR, and biological studies of novel heteroatom-incorporated antofine and cryptopleurine analogues as potent and selective antitumor agents. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 5097-107	8.3	32
374	Denbinobin suppresses breast cancer metastasis through the inhibition of Src-mediated signaling pathways. <i>Journal of Nutritional Biochemistry</i> , 2011 , 22, 732-40	6.3	14
373	Total synthesis of moniliformediquinone and calanquinone A as potent inhibitors for breast cancer. <i>Tetrahedron</i> , 2011 , 67, 6166-6172	2.4	17
372	Energy restriction-mimetic agents induce apoptosis in prostate cancer cells in part through epigenetic activation of KLF6 tumor suppressor gene expression. <i>Journal of Biological Chemistry</i> , 2011 , 286, 9968-76	5.4	13
371	Potent anti-inflammatory effects of denbinobin mediated by dual inhibition of expression of inducible no synthase and cyclooxygenase 2. <i>Shock</i> , 2011 , 35, 191-7	3.4	23
370	The roles and mechanisms of PAR4 and P2Y12/phosphatidylinositol 3-kinase pathway in maintaining thrombin-induced platelet aggregation. <i>British Journal of Pharmacology</i> , 2010 , 161, 643-58	8.6	39
369	CHM-1, a new vascular targeting agent, induces apoptosis of human umbilical vein endothelial cells via p53-mediated death receptor 5 up-regulation. <i>Journal of Biological Chemistry</i> , 2010 , 285, 5497-506	5.4	22
368	The indazole derivative YD-3 specifically inhibits thrombin-induced angiogenesis in vitro and in vivo. <i>Shock</i> , 2010 , 34, 580-5	3.4	5
367	Antitumor agents. 272. Structure-activity relationships and in vivo selective anti-breast cancer activity of novel neo-tanshinlactone analogues. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 2299-308	8.3	56
366	Moscatilin, a bibenzyl derivative from the India orchid Dendrobrium loddigesii, suppresses tumor angiogenesis and growth in vitro and in vivo. <i>Cancer Letters</i> , 2010 , 292, 163-70	9.9	50
365	Moscatilin repressed lipopolysaccharide-induced HIF-1alpha accumulation and NF-kappaB activation in murine RAW264.7 cells. <i>Shock</i> , 2010 , 33, 70-5	3.4	22
364	Synthesis and preclinical evaluations of 2-(2-fluorophenyl)-6,7-methylenedioxyquinolin-4-one monosodium phosphate (CHM-1-P-Na) as a potent antitumor agent. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 1616-26	8.3	26
363	Synthesis of 1-benzyl-3-(5-hydroxymethyl-2-furyl)selenolo[3,2-c]pyrazole derivatives as new anticancer agents. <i>European Journal of Medicinal Chemistry</i> , 2010 , 45, 1395-402	6.8	34
362	EPOX inhibits angiogenesis by degradation of Mcl-1 through ERK inactivation. <i>Clinical Cancer Research</i> , 2009 , 15, 4904-14	12.9	19
361	Synthesis, antiplatelet and vasorelaxing activities of xanthone derivatives. <i>Archiv Der Pharmazie</i> , 2009 , 342, 19-26	4.3	8

360	Apoptosis signal-regulating kinase 1 mediates denbinobin-induced apoptosis in human lung adenocarcinoma cells. <i>Journal of Biomedical Science</i> , 2009 , 16, 43	13.3	31
359	Combined treatment with denbinobin and Fas ligand has a synergistic cytotoxic effect in human pancreatic adenocarcinoma BxPC-3 cells. <i>British Journal of Pharmacology</i> , 2009 , 157, 1175-85	8.6	14
358	Evodiamine represses hypoxia-induced inflammatory proteins expression and hypoxia-inducible factor 1alpha accumulation in RAW264.7. <i>Shock</i> , 2009 , 32, 263-9	3.4	45
357	A New and Facile Synthesis of Rutaecarpine Alkaloids. <i>Heterocycles</i> , 2009 , 78, 1047	0.8	15
356	YC-1 induces apoptosis of human renal carcinoma A498 cells in vitro and in vivo through activation of the JNK pathway. <i>British Journal of Pharmacology</i> , 2008 , 155, 505-13	8.6	22
355	Anticancer mechanisms of YC-1 in human lung cancer cell line, NCI-H226. <i>Biochemical Pharmacology</i> , 2008 , 75, 360-8	6	44
354	Denbinobin induces apoptosis in human lung adenocarcinoma cells via Akt inactivation, Bad activation, and mitochondrial dysfunction. <i>Toxicology Letters</i> , 2008 , 177, 48-58	4.4	54
353	Molecular mechanism of the inhibitory effect of KS-5 on bFGF-induced angiogenesis in vitro and in vivo. <i>Cancer Letters</i> , 2008 , 263, 114-21	9.9	11
352	CHM-1, a novel synthetic quinolone with potent and selective antimitotic antitumor activity against human hepatocellular carcinoma in vitro and in vivo. <i>Molecular Cancer Therapeutics</i> , 2008 , 7, 350-60	6.1	45
351	Moscatilin induces apoptosis in human colorectal cancer cells: a crucial role of c-Jun NH2-terminal protein kinase activation caused by tubulin depolymerization and DNA damage. <i>Clinical Cancer Research</i> , 2008 , 14, 4250-8	12.9	44
350	Antiplatelet effect and selective binding to cyclooxygenase by molecular docking analysis of 3-alkylaminopropoxy-9,10-anthraquinone derivatives. <i>Biological and Pharmaceutical Bulletin</i> , 2008 , 31, 1547-51	2.3	14
349	Denbinobin induces apoptosis by apoptosis-inducing factor releasing and DNA damage in human colorectal cancer HCT-116 cells. <i>Naunyn-Schmiedebergn Archives of Pharmacology</i> , 2008 , 378, 447-57	3.4	18
348	Baicalein attenuates intimal hyperplasia after rat carotid balloon injury through arresting cell-cycle progression and inhibiting ERK, Akt, and NF-kappaB activity in vascular smooth-muscle cells. <i>Naunyn-Schmiedeberg Archives of Pharmacology</i> , 2008 , 378, 579-88	3.4	38
347	Synthesis and antiplatelet activity of ethyl 4-(1-benzyl-1H-indazol-3-yl)benzoate (YD-3) derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 1262-78	3.4	31
346	YC-1 induces heat shock protein 70 expression and prevents oxidized LDL-mediated apoptosis in vascular smooth muscle cells. <i>Shock</i> , 2008 , 30, 274-9	3.4	13
345	Synthesis of furopyrazole analogs of 1-benzyl-3-(5-hydroxymethyl-2-furyl)indazole (YC-1) as novel anti-leukemia agents. <i>Bioorganic and Medicinal Chemistry,</i> 2007 , 15, 1732-40	3.4	70
344	Proteomic approach to studying the cytotoxicity of YC-1 on U937 leukemia cells and antileukemia activity in orthotopic model of leukemia mice. <i>Proteomics</i> , 2007 , 7, 3305-17	4.8	43
343	Quinolone analogue inhibits tubulin polymerization and induces apoptosis via Cdk1-involved signaling pathways. <i>Biochemical Pharmacology</i> , 2007 , 74, 10-9	6	50

342	2-(3-Fluorophenyl)-6-methoxyl-4-oxo-1,4-dihydroquinoline-3-carboxylic acid (YJC-1) induces mitotic phase arrest in A549 cells. <i>European Journal of Pharmacology</i> , 2007 , 559, 14-20	5.3	19
341	CHM-1 inhibits hepatocyte growth factor-induced invasion of SK-Hep-1 human hepatocellular carcinoma cells by suppressing matrix metalloproteinase-9 expression. <i>Cancer Letters</i> , 2007 , 257, 87-96	9.9	20
340	Antitumor agents 253. Design, synthesis, and antitumor evaluation of novel 9-substituted phenanthrene-based tylophorine derivatives as potential anticancer agents. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3674-80	8.3	68
339	Selective COX-2 inhibitors. Part 1: synthesis and biological evaluation of phenylazobenzenesulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 4440-3	2.9	39
338	Comparison of the effects of PAR1 antagonists, PAR4 antagonists, and their combinations on thrombin-induced human platelet activation. <i>European Journal of Pharmacology</i> , 2006 , 546, 142-7	5.3	41
337	Effect of isoquinoline alkaloids of different structural types on antiplatelet aggregation in vitro. <i>Planta Medica</i> , 2006 , 72, 1238-41	3.1	16
336	YC-1 [3-(5Phydroxymethyl-2Pfuryl)-1-benzyl indazole] inhibits neointima formation in balloon-injured rat carotid through suppression of expressions and activities of matrix metalloproteinases 2 and 9. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006 , 316, 35-41	4.7	37
335	Alisol B acetate, a triterpene from Alismatis rhizoma, induces Bax nuclear translocation and apoptosis in human hormone-resistant prostate cancer PC-3 cells. <i>Cancer Letters</i> , 2006 , 231, 270-8	9.9	82
334	Synthesis of N2-(substituted benzyl)-3-(4-methylphenyl)indazoles as novel anti-angiogenic agents. <i>Bioorganic and Medicinal Chemistry</i> , 2006 , 14, 528-36	3.4	67
333	Induction of endoplasmic reticulum stress and apoptosis by a marine prostanoid in human hepatocellular carcinoma. <i>Journal of Hepatology</i> , 2005 , 43, 679-86	13.4	41
332	Induction of mitotic arrest and apoptosis in human prostate cancer pc-3 cells by evodiamine. Journal of Urology, 2005 , 173, 256-61	2.5	41
331	Three new taxane diterpenoids from Taxus sumatrana. <i>Journal of Natural Products</i> , 2005 , 68, 90-3	4.9	16
330	Chemical and Bioactive Constituents from Formosan Zanthoxylum Species. <i>Frontiers in Natural Product Chemistry</i> , 2005 , 1, 201-208	2	1
329	Enhancement of learning behaviour by a potent nitric oxide-guanylate cyclase activator YC-1. <i>European Journal of Neuroscience</i> , 2005 , 21, 1679-88	3.5	61
328	YC-1-inhibited proliferation of rat mesangial cells through suppression of cyclin D1-independent of cGMP pathway and partially reversed by p38 MAPK inhibitor. <i>European Journal of Pharmacology</i> , 2005 , 517, 1-10	5.3	14
327	Interference of neutrophil-platelet interaction by YC-1: a cGMP-dependent manner on heterotypic cell-cell interaction. <i>European Journal of Pharmacology</i> , 2005 , 519, 158-67	5.3	10
326	Goniothalamin induces cell cycle-specific apoptosis by modulating the redox status in MDA-MB-231 cells. <i>European Journal of Pharmacology</i> , 2005 , 522, 20-9	5.3	58
325	Denbinobin-mediated anticancer effect in human K562 leukemia cells: role in tubulin polymerization and Bcr-Abl activity. <i>Journal of Biomedical Science</i> , 2005 , 12, 113-21	13.3	28

324	Investigation of anticancer mechanism of clavulone II, a coral cyclopentenone prostaglandin analog, in human acute promyelocytic leukemia. <i>Journal of Biomedical Science</i> , 2005 , 12, 335-45	13.3	11
323	A potential role of YC-1 on the inhibition of cytokine release in peripheral blood mononuclear leukocytes and endotoxemic mouse models. <i>Thrombosis and Haemostasis</i> , 2005 , 93, 940-8	7	31
322	YC-1 [3-(5Phydroxymethyl-2Pfuryl)-1-benzyl indazole] inhibits endothelial cell functions induced by angiogenic factors in vitro and angiogenesis in vivo models. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2005 , 314, 35-42	4.7	55
321	YC-1 suppresses constitutive nuclear factor-kappaB activation and induces apoptosis in human prostate cancer cells. <i>Molecular Cancer Therapeutics</i> , 2005 , 4, 1628-35	6.1	47
320	CIL-102 interacts with microtubule polymerization and causes mitotic arrest following apoptosis in the human prostate cancer PC-3 cell line. <i>Journal of Biological Chemistry</i> , 2005 , 280, 2771-9	5.4	40
319	Antitumor mechanism of evodiamine, a constituent from Chinese herb Evodiae fructus, in human multiple-drug resistant breast cancer NCI/ADR-RES cells in vitro and in vivo. <i>Carcinogenesis</i> , 2005 , 26, 968-75	4.6	102
318	YC-1 [3-(5PHydroxymethyl-2Pfuryl)-1-benzyl Indazole] exhibits a novel antiproliferative effect and arrests the cell cycle in G0-G1 in human hepatocellular carcinoma cells. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2005 , 312, 917-25	4.7	48
317	Structure-activity relationship studies on chalcone derivatives: potent inhibition of platelet aggregation. <i>Journal of Pharmacy and Pharmacology</i> , 2004 , 56, 1333-7	4.8	19
316	The indazole derivative YD-3 inhibits thrombin-induced vascular smooth muscle cell proliferation and attenuates intimal thickening after balloon injury. <i>Thrombosis and Haemostasis</i> , 2004 , 92, 1232-9	7	22
315	Soluble decoy receptor 3 induces angiogenesis by neutralization of TL1A, a cytokine belonging to tumor necrosis factor superfamily and exhibiting angiostatic action. <i>Cancer Research</i> , 2004 , 64, 1122-9	10.1	96
314	A new aristolactam alkaloid and anti-platelet aggregation constituents from Piper taiwanense. <i>Planta Medica</i> , 2004 , 70, 174-7	3.1	32
313	Peptidoglycan induces nuclear factor-kappaB activation and cyclooxygenase-2 expression via Ras, Raf-1, and ERK in RAW 264.7 macrophages. <i>Journal of Biological Chemistry</i> , 2004 , 279, 20889-97	5.4	76
312	Investigation of extrinsic and intrinsic apoptosis pathways of new clerodane diterpenoids in human prostate cancer PC-3 cells. <i>European Journal of Pharmacology</i> , 2004 , 503, 17-24	5.3	37
311	Investigation of ouabain-induced anticancer effect in human androgen-independent prostate cancer PC-3 cells. <i>Biochemical Pharmacology</i> , 2004 , 67, 727-33	6	91
310	Genistein inversely affects tubulin-binding agent-induced apoptosis in human breast cancer cells. <i>Biochemical Pharmacology</i> , 2004 , 67, 2031-8	6	34
309	Modulation of anti-adhesion molecule MUC-1 is associated with arctiin-induced growth inhibition in PC-3 cells. <i>Prostate</i> , 2004 , 59, 260-7	4.2	19
308	New prostanoids with cytotoxic activity from Taiwanese octocoral Clavularia viridis. <i>Journal of Natural Products</i> , 2004 , 67, 542-6	4.9	39
307	Induction of mitotic arrest and apoptosis by evodiamine in human leukemic T-lymphocytes. <i>Life Sciences</i> , 2004 , 75, 35-49	6.8	53

(2002-2004)

306	Cerebrosides from the Rhizomes of Gynura Japonica. <i>Journal of the Chinese Chemical Society</i> , 2004 , 51, 1429-1434	1.5	17	
305	The inhibitory mechanism of YC-1, a benzyl indazole, on smooth muscle cell proliferation: an in vitro and in vivo study. <i>Journal of Pharmacological Sciences</i> , 2004 , 94, 252-60	3.7	28	
304	Enhancement of long-term potentiation by a potent nitric oxide-guanylyl cyclase activator, 3-(5-hydroxymethyl-2-furyl)-1-benzyl-indazole. <i>Molecular Pharmacology</i> , 2003 , 63, 1322-8	4.3	69	
303	Anti-platelet aggregation and chemical constituents from the rhizome of Gynura japonica. <i>Planta Medica</i> , 2003 , 69, 757-64	3.1	50	
302	Soluble guanylyl cyclase activator YC-1 inhibits human neutrophil functions through a cGMP-independent but cAMP-dependent pathway. <i>Molecular Pharmacology</i> , 2003 , 64, 1419-27	4.3	87	
301	The role of PAR4 in thrombin-induced thromboxane production in human platelets. <i>Thrombosis and Haemostasis</i> , 2003 , 90, 299-308	7	39	
300	Pharmacological evaluation of several major ingredients of Chinese herbal medicines in human hepatoma Hep3B cells. <i>European Journal of Pharmaceutical Sciences</i> , 2003 , 19, 403-12	5.1	65	
299	Esculetin inhibits Ras-mediated cell proliferation and attenuates vascular restenosis following angioplasty in rats. <i>Biochemical Pharmacology</i> , 2003 , 65, 1897-905	6	59	
298	Potentiation of tumor necrosis factor-alpha expression by YC-1 in alveolar macrophages through a cyclic GMP-independent pathway. <i>Biochemical Pharmacology</i> , 2003 , 66, 149-56	6	17	
297	YC-1 inhibits proliferation of human vascular endothelial cells through a cyclic GMP-independent pathway. <i>Biochemical Pharmacology</i> , 2003 , 66, 263-71	6	43	
296	Identification of Apoptotic and Antiangiogenic Activities of Terazosin in Human Prostate Cancer and Endothelial Cells. <i>Journal of Urology</i> , 2003 , 169, 724-729	2.5	22	
295	Effect of a potent cyclooxygenase inhibitor, 5-ethyl-4-methoxy-2-phenylquinoline (KTC-5), on human platelets. <i>Journal of Pharmacy and Pharmacology</i> , 2002 , 54, 967-74	4.8	2	
294	Characterization of some novel alpha 1-adrenoceptor antagonists in human hyperplastic prostate. <i>European Journal of Pharmacology</i> , 2002 , 445, 125-31	5.3	5	
293	Inhibition of ras-mediated cell proliferation by benzyloxybenzaldehyde. <i>Journal of Biomedical Science</i> , 2002 , 9, 622-630	13.3	5	
292	Cardiac glycosides induce resistance to tubulin-dependent anticancer drugs in androgen-independent human prostate cancer. <i>Journal of Biomedical Science</i> , 2002 , 9, 443-52	13.3	10	
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(2000-2001)

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[1996-1996]

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