Sylvie Michel

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4307700/publications.pdf

Version: 2024-02-01

155 papers 3,051 citations

147801 31 h-index 223800 46 g-index

178 all docs

178 docs citations

times ranked

178

3698 citing authors

#	Article	IF	CITATIONS
1	Pistacia lentiscus L. Distilled Leaves as a Potential Cosmeceutical Ingredient: Phytochemical Characterization, Transdermal Diffusion, and Anti-Elastase and Anti-Tyrosinase Activities. Molecules, 2022, 27, 855.	3.8	8
2	Potent Antiplasmodial Derivatives of Dextromethorphan Reveal the Ent-Morphinan Pharmacophore of Tazopsine-Type Alkaloids. Pharmaceutics, 2022, 14, 372.	4.5	1
3	A kaleidoscope of photosynthetic antenna proteins and their emerging roles. Plant Physiology, 2022, 189, 1204-1219.	4.8	14
4	Bryophyllum pinnatum markers: CPC isolation, simultaneous quantification by a validated UPLC-DAD method and biological evaluations. Journal of Pharmaceutical and Biomedical Analysis, 2021, 193, 113682.	2.8	13
5	Identification of alkylsalicylic acids in Lentisk oil (<i>Pistacia lentiscus</i> L.) and viability assay on Human Normal Dermal Fibroblasts. OCL - Oilseeds and Fats, Crops and Lipids, 2021, 28, 22.	1.4	1
6	Comparison of extraction methods for chemical composition, antibacterial, depigmenting and antioxidant activities of <i>Eryngium maritimum</i> . International Journal of Cosmetic Science, 2020, 42, 127-135.	2.6	12
7	Correlation study on methoxylation pattern of flavonoids and their heme-targeted antiplasmodial activity. Bioorganic Chemistry, 2020, 104, 104243.	4.1	8
8	A Photoalkylative Fluorogenic Probe of Guttiferone A for Live Cell Imaging and Proteome Labeling in Plasmodium falciparum. Molecules, 2020, 25, 5139.	3.8	6
9	Cytotoxic compounds from the leaves and stems of the endemic Thai plant <i>Mitrephora sirikitiae</i> Pharmaceutical Biology, 2020, 58, 490-497.	2.9	8
10	Membrane-Interactive Compounds From Pistacia lentiscus L. Thwart Pseudomonas aeruginosa Virulence. Frontiers in Microbiology, 2020, 11, 1068.	3.5	30
11	Polymethoxyflavones from Gardenia oudiepe (Rubiaceae) induce cytoskeleton disruption-mediated apoptosis and sensitize BRAF-mutated melanoma cells to chemotherapy. Chemico-Biological Interactions, 2020, 325, 109109.	4.0	7
12	Health risk associated with the oral consumption of "Chiniy-trefâ€, a traditional medicinal preparation used in Martinique (French West Indies): Qualitative and quantitative analyses of aristolochic acids contained therein. Toxicon, 2019, 172, 53-60.	1.6	9
13	Collected mass spectrometry data on monoterpene indole alkaloids from natural product chemistry research. Scientific Data, 2019, 6, 15.	5 . 3	37
14	Chemical composition and biological activity of essential oils from <i>Artemisia copa</i> Phil. var. copa (Asteraceae) and <i>Aloysia deserticola</i> (Phil.) Lu-Irving & O'Leary (Verbenaceae), used in the Chilean Atacama's Taira Community (Antofagasta, Chile). Journal of Essential Oil Research, 2019, 31, 425-431.	2.7	5
15	Bioguided identification of triterpenoids and neolignans as bioactive compounds from anti-infectious medicinal plants of the Taira Atacama's community (Calama, Chile). Journal of Ethnopharmacology, 2019, 231, 217-229.	4.1	15
16	Chemical composition and biological properties of Ipomoea procumbens. Revista Brasileira De Farmacognosia, 2019, 29, 191-197.	1.4	7
17	In vitro biological evaluation and molecular docking studies of natural and semisynthetic flavones from Gardenia oudiepe (Rubiaceae) as tyrosinase inhibitors. Bioorganic Chemistry, 2019, 82, 241-245.	4.1	12
18	How light photoperiod and medium composition could increase the production of a potent anticancer metabolite by Nostoc. Planta Medica, 2019, 85, .	1.3	0

#	Article	lF	Citations
19	Exploring the traditional medicine of Atacama people from Northern Chile as in inestimable source of bioactive compounds. Planta Medica, 2019, 85, .	1.3	0
20	Spirokermeline: A Macrocyclic Spirolactone from <i>Kermadecia elliptica</i> Brongn. & Brongn. & European Journal of Organic Chemistry, 2018, 2018, 5819-5822.	2.4	6
21	Clerodane furanoditerpenoids as the probable cause of toxic hepatitis induced by Tinospora crispa. Scientific Reports, 2018, 8, 13520.	3.3	14
22	Polar lipids in cosmetics: recent trends in extraction, separation, analysis and main applications. Phytochemistry Reviews, 2018, 17, 1179-1210.	6.5	29
23	Chemical constituents of Anthospermum perrieri (Rubiaceae). Biochemical Systematics and Ecology, 2018, 80, 29-31.	1.3	2
24	Comparative metabolomic study between African and Amazonian Symphonia globulifera by tandem LC–HRMS. Phytochemistry Letters, 2017, 20, 309-315.	1.2	6
25	Heme-binding activity of methoxyflavones from Pentzia monodiana Maire (Asteraceae). Fìtoterapìâ, 2017, 118, 1-5.	2.2	10
26	Assessment of two centrifugal partition chromatography devices. Application to the purification of Centaurium erythraea methanolic extract. Phytochemistry Letters, 2017, 20, 401-405.	1.2	6
27	Chemical study of Anthospermum emirnense (Rubiaceae). Biochemical Systematics and Ecology, 2017, 70, 186-191.	1.3	3
28	A Nitrile Glucoside and Biflavones from the Leaves of <i>Campylospermum excavatum</i> (Ochnaceae). Chemistry and Biodiversity, 2017, 14, e1700241.	2.1	9
29	Off-line coupling of new generation centrifugal partition chromatography device with preparative high pressure liquid chromatography-mass spectrometry triggering fraction collection applied to the recovery of secoiridoid glycosides from Centaurium erythraea Rafn. (Gentianaceae). Journal of Chromatography A, 2017, 1513, 149-156.	3.7	10
30	Synthetic Analogue of the Natural Product Piperlongumine as a Potent Inhibitor of Breast Cancer Cell Line Migration. Journal of the Brazilian Chemical Society, 2017, 28, 475-484.	0.6	9
31	Three new trixane glycosides obtained from the leaves of <i>Jungia sellowii</i> Less. using centrifugal partition chromatography. Beilstein Journal of Organic Chemistry, 2016, 12, 674-683.	2.2	13
32	Phytochemical study and biological evaluation of chemical constituents of Platanus orientalis and PlatanusÂ×Âacerifolia buds. Phytochemistry, 2016, 130, 170-181.	2.9	21
33	Guttiferone A Aggregates Modulate Silent Information Regulator 1 (SIRT1) Activity. Journal of Medicinal Chemistry, 2016, 59, 9560-9566.	6.4	6
34	Triterpenes from the exudate of Gardenia urvillei. Phytochemistry, 2016, 122, 193-202.	2.9	14
35	Purification of bioactive compounds from Centaurium erythraea by conventional and new generation designed Centrifugal Partition Chromatography column. Planta Medica, 2016, 81, S1-S381.	1.3	0
36	Isolation of Guttiferones from Renewable Parts of Symphonia globulifera by Centrifugal Partition Chromatography. Planta Medica, 2015, 81, 1604-1608.	1.3	6

#	Article	IF	Citations
37	Antileishmanial activity of fucosterol recovered from Lessonia vadosa Searles (Lessoniaceae) by SFE, PSE and CPC. Phytochemistry Letters, 2015, 11, 418-423.	1.2	39
38	Antivascular and anti-parasite activities of natural and hemisynthetic flavonoids from New Caledonian Gardenia species (Rubiaceae). European Journal of Medicinal Chemistry, 2015, 93, 93-100.	5.5	32
39	Ion tree-based structure elucidation of acetophenone dimers (AtA) from <i>Acronychia pedunculata</i> and their identification in extracts by liquid chromatography electrospray ionization LTQ-Orbitrap mass spectrometry. Journal of Mass Spectrometry, 2015, 50, 495-512.	1.6	8
40	Polycyclic Polyprenylated Xanthones from <i>Symphonia globulifera</i> Electrosynthesis. Journal of Natural Products, 2015, 78, 2136-2140.	3.0	10
41	Cymoside, a monoterpene indole alkaloid with a hexacyclic fused skeleton from Chimarrhis cymosa. Tetrahedron Letters, 2015, 56, 5377-5380.	1.4	16
42	Viability of a [2 + 2 + 1] Hetero-Pauson–Khand Cycloaddition Strategy towardSecurinegaAlkaloids: Synthesis of the BCD-Ring Core of Securinine and Related Alkaloids. Journal of Organic Chemistry, 2015, 80, 6525-6528.	3.2	10
43	Symphonia globulifera, a Widespread Source of Complex Metabolites with Potent Biological Activities. Planta Medica, 2015, 81, 95-107.	1.3	16
44	Phytochemical study of Capraria biflora L. aerial parts (Scrophulariaceae) from Martinique island (French West Indies). Phytochemistry Letters, 2015, 13, 194-199.	1.2	2
45	Chiroptical study and absolute configuration of securinine oxidation products. Natural Product Research, 2015, 29, 1235-1242.	1.8	3
46	A new 3,4-seco-cycloartane from the leaves of Hopea odorata Roxb Natural Product Research, 2015, 29, 1820-1827.	1.8	8
47	Rapid Identification of Antioxidant Compounds of Genista saharae Coss. & DPPH Scavenging Assay and HPTLC-MS. Molecules, 2014, 19, 4369-4379.	3.8	25
48	Synthesis, Antitumor Activity, and Mechanism of Action of Benzo[$<$ i> $>$ b $<$ /i $>$]chromeno[6 ,5- $<$ i $>$ g $<$ /i $>$][1 ,8]naphthyridin-7-one Analogs of Acronycine. Journal of Medicinal Chemistry, 2014, 57, 10329-10342.	6.4	18
49	Comparative LC–MS-based metabolite profiling of the ancient tropical rainforest tree Symphonia globulifera. Phytochemistry, 2014, 108, 102-108.	2.9	13
50	One-Step Semisynthesis of Oleacein and the Determination as a 5-Lipoxygenase Inhibitor. Journal of Natural Products, 2014, 77, 441-445.	3.0	60
51	Antifungal ether diglycosides from Matayba guianensis Aublet. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 1414-1416.	2.2	7
52	Geranylated homogentisic acid derivatives and flavonols from Miliusa umpangensis. Biochemical Systematics and Ecology, 2014, 54, 179-181.	1.3	10
53	Toxic hepatitis induced by a herbal medicine: Tinospora crispa. Phytomedicine, 2014, 21, 1120-1123.	5.3	37
54	Cytotoxic turrianes from Kermadecia elliptica: Hemisynthesis and biological activities of kermadecin A derivatives. Phytochemistry Letters, 2014, 10, 249-254.	1.2	5

#	Article	IF	Citations
55	Dereplication and metabolomics strategies for the discovery of bioactive natural products: The Acronychia example. Planta Medica, 2014, 80, .	1.3	3
56	Towards the first SAR study on the Securinega alkaloids. Planta Medica, 2014, 80, .	1.3	1
57	Influence of solvents and catalysts on the formation and hydrolysis of polyfunctional enoxysilanes derived from aucubin. Arkivoc, 2014, 2014, 184-196.	0.5	2
58	Neolignans from leaves of Miliusa mollis. Fìtoterapìâ, 2013, 85, 49-56.	2.2	25
59	Synthesis of novel guttiferone A derivatives: In-vitro evaluation toward Plasmodium falciparum, Trypanosoma brucei and Leishmania donovani. European Journal of Medicinal Chemistry, 2013, 65, 284-294.	5.5	25
60	New neolignans and a lignan from Miliusa fragrans, and their anti-herpetic and cytotoxic activities. Tetrahedron Letters, 2013, 54, 4259-4263.	1.4	30
61	New triterpenoids from the stem bark ofHypodaphnis zenkeri. Natural Product Research, 2013, 27, 137-145.	1.8	7
62	Synthesis of a new bis(indolyl)methane that inhibits growth and induces apoptosis in human prostate cancer cells. Natural Product Research, 2013, 27, 2039-2045.	1.8	44
63	Tetrahydroalstonine. Acta Crystallographica Section E: Structure Reports Online, 2013, 69, o1389-o1390.	0.2	0
64	New neolignans from leaves of Miliusa mollis. Planta Medica, 2013, 79, .	1.3	0
65	A New Sphingolipid and Furanocoumarins with Antimicrobial Activity from <i>Ficus exasperata</i> . Chemical and Pharmaceutical Bulletin, 2012, 60, 1072-1075.	1.3	30
66	Lupane triterpenes from the leaves of the tropical rain forest tree Hopea odorata Roxb. and their cytotoxic activities. Biochemical Systematics and Ecology, 2012, 44, 407-412.	1.3	12
67	Synthesis and Antimicrobial Activities of Some Sulphur Containing Chromene Derivatives. Natural Product Communications, 2012, 7, 1934578X1200700.	0.5	2
68	Semisythesis of Guttiferone A analogs. Planta Medica, 2012, 78, .	1.3	1
69	Methodology for the preparation of olive oil open ring secoiridoids. Planta Medica, 2012, 78, .	1.3	13
70	Natural products as models for the conception of new active products: Benzopyran, a privileged structure. Planta Medica, 2012, 78, .	1.3	0
71	Cymoside, an original hexacyclic monoterpene indole alkaloid and others compounds from Chimarrhis cymosa (Rubiaceae). Planta Medica, 2012, 78, .	1.3	0
72	Synthesis and biological activity of some C(9)-hydroxymethyl-5,11-dimethylellipticine derivatives. Planta Medica, 2012, 78, .	1.3	0

#	Article	IF	CITATIONS
73	UHPLC-LTQ-ORBITRAP based identification and HSCCC isolation of antifungal components from Platanus SP. (Platanaceae). Planta Medica, 2012, 78, .	1.3	O
74	Selective antiproliferative activity of spinasterol from Physospemum verticillatum against A549 and COR-L23 cancer cells. Planta Medica, 2012, 78, .	1.3	0
75	Evaluation of the antiangiogenic and anti-parasitic activities of flavonoids from gardenia species and their modified analogues. Planta Medica, 2012, 78, .	1.3	0
76	New 3,4-secocycloartane and lupane triterpenes from the leaves of the tropical rain forest tree Hopea odorata Roxb. Planta Medica, 2012, 78, .	1.3	0
77	A one-pot synthesis of 7-phenylindolo[3,2-a]carbazoles from indoles and \hat{l}^2 -nitrostyrenes, via an unprecedented reaction sequence. Organic and Biomolecular Chemistry, 2011, 9, 7780.	2.8	39
78	Dammarane Triterpenes from <i>Gardenia aubryi</i> <scp>Vieill</scp> Helvetica Chimica Acta, 2011, 94, 656-661.	1.6	6
79	Solvent/Base Effects in the Selective Domino Synthesis of Phenanthridinones That Involves Highâ€Valent Palladium Species: Experimental and Theoretical Studies. Chemistry - A European Journal, 2011, 17, 12809-12819.	3.3	19
80	Synthesis and cytotoxic activity of benzo[a]acronycine and benzo[b]acronycine substituted on the A ring. European Journal of Medicinal Chemistry, 2011, 46, 1861-1873.	5.5	10
81	Synthesis and biological evaluation of N-substituted benzo[c]phenanthrolines and benzo[c]phenanthrolinones as antiproliferative agents. European Journal of Medicinal Chemistry, 2011, 46, 2117-2131.	5.5	13
82	Tröger's bases in the acronycine, benzo[a]acronycine, and benzo[b]acronycine series. Tetrahedron Letters, 2011, 52, 4426-4429.	1.4	15
83	Biological Potential and Structure-Activity Relationships of Most Recently Developed Vascular Disrupting Agents: An Overview of New Derivatives of Natural Combretastatin A-4. Current Medicinal Chemistry, 2011, 18, 3035-3081.	2.4	64
84	Synthesis and cytotoxic activity of psorospermin and acronycine analogues in the 3-propyloxy-acridin- $9(10H)$ -one and -benzo[b]acridin- $12(5H)$ -one series. European Journal of Medicinal Chemistry, 2010, 45, 581-587.	5.5	16
85	Synthesis and biological evaluation of dialkylaminoalkylamino benzo $[c][1,7]$ and $[1,8]$ phenanthrolines as antiproliferative agents. European Journal of Medicinal Chemistry, 2010, 45, 2547-2558.	5.5	14
86	Synthesis, biological activity, and evaluation of the mode of action of novel antitubercular benzofurobenzopyrans substituted on A ring. European Journal of Medicinal Chemistry, 2010, 45, 5833-5847.	5.5	33
87	The synthesis and Angiotensin Converting Enzyme (ACE) inhibitory activity of chalcones and their pyrazole derivatives. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 1990-1993.	2.2	77
88	Structureâ€"activity relationships of indole compounds derived from combretastatin A4: Synthesis and biological screening of 5-phenylpyrrolo[3,4-a]carbazole-1,3-diones as potential antivascular agents. European Journal of Medicinal Chemistry, 2010, 45, 3726-3739.	5.5	23
89	Selective Unusual Pd-Mediated Biaryl Coupling Reactions: Solvent Effects with Carbonate Bases. Organic Letters, 2010, 12, 156-158.	4.6	28
90	Tri-ionizable calix[4]arene ligands: synthesis and lanthanide ion complexation. Arkivoc, 2010, 2010, 191-202.	0.5	6

#	Article	IF	CITATIONS
91	Influence of the Stereoisomeric Position of the Reactive Acetate Groups of the Benzo[<i>b</i>)Acronycine derivative S23906-1 on Its DNA Alkylation, Helix-Opening, Cytotoxic, and Antitumor Activities. Molecular Pharmacology, 2009, 76, 1172-1185.	2.3	10
92	Acronycine Derivatives: A Promising Series of Anti-Cancer Agents. Anti-Cancer Agents in Medicinal Chemistry, 2009, 9, 804-815.	1.7	19
93	Isolation and chemistry of the alkaloids from Papaver arachnoideum Kadereit. Biochemical Systematics and Ecology, 2009, 37, 501-503.	1.3	2
94	Synthesis of N-substituted benzo[c][1,7]- and benzo[c][1,8] phenanthrolin-(5H)-6-ones through a Pd-mediated Suzuki–Miyaura heteroaryl-aryl coupling reaction. Tetrahedron, 2009, 65, 10009-10015.	1.9	13
95	Diversity-oriented synthesis of furo [3,2-f] chromanes with antimycobacterial activity. European Journal of Medicinal Chemistry, 2009, 44, 2497-2505.	5.5	81
96	Synthesis, cytotoxic activity, and DNA binding properties of antitumor cis-1,2-dihydroxy-1,2-dihydrobenzo[b]acronycine cinnamoyl esters. Bioorganic and Medicinal Chemistry, 2009, 17, 1918-1927.	3.0	9
97	Cytotoxic activity of Brazilian Cerrado plants used in traditional medicine against cancer cell lines. Journal of Ethnopharmacology, 2009, 123, 439-445.	4.1	122
98	Antitumor <i>Psoropermum</i> Xanthones and <i>Sarcomelicope</i> Acridones: Privileged Structures Implied in DNA Alkylation. Journal of Natural Products, 2009, 72, 527-539.	3.0	67
99	Structure–activity relationships in the acronycine and benzo[b]acronycine series: Role of the pyran ring. European Journal of Medicinal Chemistry, 2008, 43, 2677-2687.	5.5	9
100	Synthesis and biological evaluation of new disubstituted analogues of 6-methoxy-3-(3′,4′,5′-trimethoxybenzoyl)-1H-indole (BPROLO75), as potential antivascular agents. Bioorganic and Medicinal Chemistry, 2008, 16, 7494-7503.	3.0	33
101	A new synthetic access to furo [3,2-f] chromene analogues of an antimycobacterial. Bioorganic and Medicinal Chemistry, 2008, 16, 8264-8272.	3.0	92
102	Synthesis, Cytotoxic Activity, and Mechanism of Action of Furo[2,3- <i>c</i>) acridin-6-one and Benzo[<i>b</i>) furo[3,2- <i>h</i>) acridin-6-one Analogues of Psorospermin and Acronycine. Journal of Medicinal Chemistry, 2008, 51, 7287-7297.	6.4	21
103	Structure-activity relationships in the acronycine and benzo[b]acronycine series: Role of the pyran ring. Planta Medica, 2008, 74, .	1.3	0
104	Natural products as privileged structures for the conception of novel antimycobacterial agents. Planta Medica, 2008, 74, .	1.3	0
105	Design, synthesis and biological evaluation of 13-aza derivatives of benzo[b]acronycine. Planta Medica, 2008, 74, .	1.3	0
106	Novel potential antitumor analogues of fagaronine and nitidine in the Benzo[c]phenanthroline series. Planta Medica, 2008, 74, .	1.3	0
107	Synthesis, cytotoxic activity and mechanism of action of new Psorospermin-Acronycine analogs. Planta Medica, 2008, 74, .	1.3	0
108	Antimycobacterial Benzofuro [3,2-f] chromenes from a 5-Bromochromen-6-ol. Synthesis, 2007, 2007, 1566-1570.	2.3	1

#	Article	IF	CITATIONS
109	Synthesis and Cytotoxic Activity of Dimeric Analogs of Acronycine in the Benzo[b]pyrano[3,2-h]acridin-7-one Series. Chemical and Pharmaceutical Bulletin, 2007, 55, 734-738.	1.3	5
110	Synthesis and antimycobacterial evaluation of benzofurobenzopyran analogues. Bioorganic and Medicinal Chemistry, 2007, 15, 2177-2186.	3.0	47
111	Synthesis and Angiotensin Converting Enzyme (ACE) inhibition activity of chalcone derivatives. Planta Medica, 2007, 73, .	1.3	0
112	Synthesis, Antitumor Activity, and Mechanism of Action of Benzo[a]pyrano[3,2-h]acridin-7-one Analogues of Acronycine. Journal of Medicinal Chemistry, 2006, 49, 3383-3394.	6.4	20
113	seco-Cycloartane Triterpenes fromGardeniaaubryi. Journal of Natural Products, 2006, 69, 1711-1714.	3.0	39
114	Synthesis and Cytotoxic Activity of Benzo[a]pyrano[3,2-h] and [2,3-i]xanthone Analogues of Psorospermine, Acronycine, and Benzo[a]acronycine. Chemical and Pharmaceutical Bulletin, 2006, 54, 1113-1118.	1.3	14
115	Benzofuro[3,2-f][1]benzopyrans: A new class of antitubercular agents. Bioorganic and Medicinal Chemistry, 2006, 14, 5423-5428.	3.0	54
116	Synthesis and biological evaluation of (3,4,5-trimethoxyphenyl)indol-3-ylmethane derivatives as potential antivascular agents. Bioorganic and Medicinal Chemistry, 2006, 14, 4410-4426.	3.0	33
117	Synthesis of benzo[c][1,8]phenanthrolin-6-one through cyclization ofN-(isoquinol-5-yl)-2-bromo-benzamide derivatives. Journal of Heterocyclic Chemistry, 2006, 43, 1261-1265.	2.6	3
118	Design, Synthesis, and Cytotoxic Activity of Michael Acceptors and Enol Esters in the Benzo[b]acronycine Series. Chemical and Pharmaceutical Bulletin, 2005, 53, 919-922.	1.3	6
119	New antitubulin derivatives in the combretastatin A4 series: synthesis and biological evaluation. Bioorganic and Medicinal Chemistry, 2005, 13, 3853-3864.	3.0	46
120	Covalent binding of antitumor benzoacronycines to double-stranded DNA induces helix opening and the formation of single-stranded DNA: unique consequences of a novel DNA-bonding mechanism. Molecular Cancer Therapeutics, 2005, 4, 71-80.	4.1	34
121	Benzo[b]acronycine Derivatives: A Novel Class of Antitumor Agents. ChemInform, 2004, 35, no.	0.0	0
122	A transesterification reaction is implicated in the covalent binding of benzo[b]acronycine anticancer agents with DNA and glutathion. Bioorganic and Medicinal Chemistry, 2004, 12, 23-29.	3.0	24
123	Synthesis and cytotoxic activity of benzo $[c][1,7]$ and $[1,8]$ phenanthrolines analogues of nitidine and fagaronine. Bioorganic and Medicinal Chemistry, 2004, 12, 3943-3953.	3.0	39
124	Benzo[b]acronycine derivatives: a novel class of antitumor agents. European Journal of Medicinal Chemistry, 2004, 39, 649-655.	5.5	38
125	New Diterpenes fromCroton insularis. Journal of Natural Products, 2004, 67, 685-688.	3.0	32
126	Synthesis and Cytotoxic and Antitumor Activity of 1,2-Dihydroxy-1,2-dihydrobenzo[b]acronycine Diacid Hemiesters and Carbamates. Chemical and Pharmaceutical Bulletin, 2004, 52, 293-297.	1.3	15

#	Article	IF	Citations
127	Synthesis and Cytotoxic Activity of Pyranocarbazole Analogues of Ellipticine and Acronycine. Chemical and Pharmaceutical Bulletin, 2004, 52, 540-545.	1.3	16
128	Design of Novel Antitumor DNA Alkylating Agents: The Benzacronycine Series. Anti-Cancer Agents in Medicinal Chemistry, 2004, 4, 83-92.	7.0	12
129	Covalent binding to glutathione of the DNA-alkylating antitumor agent, S23906-1. FEBS Journal, 2003, 270, 2848-2859.	0.2	27
130	Structureâ^'Activity Relationships and Mechanism of Action of Antitumor Benzo[b]pyrano[3,2-h]acridin-7-one Acronycine Analogues. Journal of Medicinal Chemistry, 2003, 46, 3072-3082.	6.4	52
131	A New Pyranoacridone Alkaloid from the Bark of Medicosma subsessilis (Rutaceae). Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences, 2003, 58, 1234-1236.	0.7	6
132	Synthesis of 6-dialkylaminoalkylamino pyrano[2,3-c]acridones and benzo[b]pyrano[3,2-h]acridones: soluble acronycine analogues with increased cytotoxic activity. Oncology Research, 2003, 13, 191-7.	1.5	7
133	A New Phenylpropanoid Ester from the Bark of Zanthoxylum scandens (Rutaceae). Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 2002, 57, 986-989.	1.4	6
134	Structure-Activity Relationships in The Acronycine Series. Current Medicinal Chemistry, 2002, 9, 1689-700.	2.4	17
135	Induction of Cyclin E and Inhibition of DNA Synthesis by the Novel Acronycine Derivative S23906-1 Precede the Irreversible Arrest of Tumor Cells in S Phase Leading to Apoptosis. Molecular Pharmacology, 2001, 60, 1383-1391.	2.3	73
136	Synthesis and Cytotoxic Activity of Benzopyranoxanthone Analogues of Benzo[b]acronycine and Psorospermine Chemical and Pharmaceutical Bulletin, 2001, 49, 675-679.	1.3	13
137	A New Diprenyl Coumarin and Alkaloids from the Bark of Zanthoxylum dimorphophyllum (Rutaceae). Zeitschrift Fur Naturforschung - Section C Journal of Biosciences, 2001, 56, 492-494.	1.4	7
138	Synthesis and Cytotoxic and Antitumor Activity of Benzo[b]pyrano[3,2-h]acridin-7-one Analogues of Acronycine. Journal of Medicinal Chemistry, 2000, 43, 2395-2402.	6.4	78
139	Chiral Dihydroxylation of Acronycine:Â Absolute Configuration of Naturalcis-1,2-Dihydroxy-1,2-dihydroacronycine and Cytotoxicity of (1R,2R)- and (1S,2S)-1,2-Diacetoxy-1,2-dihydroacronycine. Journal of Natural Products, 1999, 62, 490-492.	3.0	27
140	Synthesis and Cytotoxic Activity of 11-Nitro and 11-Amino Derivatives of Acronycine and 6-Demethoxyacronycine Chemical and Pharmaceutical Bulletin, 1999, 47, 1604-1606.	1.3	14
141	Prodrugs of Anthracyclines for Use in Antibody-Directed Enzyme Prodrug Therapy. Journal of Medicinal Chemistry, 1998, 41, 3572-3581.	6.4	84
142	Acronycine-Type Alkaloids : Chemistry and Biology. Alkaloids: Chemical and Biological Perspectives, 1998, 12, 1-102.	0.2	23
143	Synthesis and Aromatization of Dihydropyrimidines Structurally Related to Calcium Channel Modulators of the Nifedipine-Type. Heterocycles, 1997, 45, 1967.	0.7	84
144	Bioactive natural and synthetic acronycine derivatives modified at the pyran ring. Studies in Natural Products Chemistry, 1997, , 789-815.	1.8	3

#	Article	IF	CITATIONS
145	Synthesis and Cytotoxic and Antitumor Activity of Esters in the 1,2-Dihydroxy-1,2-dihydroacronycine Series. Journal of Medicinal Chemistry, 1996, 39, 4762-4766.	6.4	118
146	Synthesis and Cytotoxic Activity of Acronycine Derivatives Modified at the Pyran Ring Chemical and Pharmaceutical Bulletin, 1996, 44, 2165-2168.	1.3	16
147	A Novel Synthesis of 6-Demethoxyacronycine. Heterocycles, 1992, 34, 799.	0.7	33
148	Synthesis of novel targeted pro-prodrugs of anthracyclines potentially activated by a monoclonal antibody galactosidase conjugate (part 1). Bioorganic and Medicinal Chemistry Letters, 1992, 2, 1093-1096.	2.2	29
149	Strellidimine: the first natural bis-ellipticine alkaloid. Journal of the Chemical Society Chemical Communications, 1987, , 229.	2.0	26
150	Brafouédine et Isobrafouédine: Nouveaux Alcaloïdes Indoliques Mineurs de Strychnos dinklagei. Journal of Natural Products, 1986, 49, 452-455.	3.0	10
151	Plantes de Nouvelle-Calédonie. 94e Communication. AlcaloÃ⁻des monoterpéniques deScaevola racemigeraDÄNIKER. Helvetica Chimica Acta, 1985, 68, 1679-1685.	1.6	19
152	Alcalo $ ilde{A}^-$ des des Feuilles de Strychnos dinklagei. Journal of Natural Products, 1985, 48, 86-92.	3.0	9
153	La dinklageine, alcaloide monoterpenique d'un type nouveau. Tetrahedron Letters, 1984, 25, 2783-2786.	1.4	6
154	AlcaloÃ ⁻ des Des Écorces De Tiges De Strychnos dinklagei. Journal of Natural Products, 1982, 45, 489-494.	3.0	24
155	L'Ellipticine, AlcaloÃ⁻de Majeur Des Écorces de Strychnos dinklagei. Journal of Natural Products, 1980, 43, 294-295.	3.0	26