A Ganesan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9509264/publications.pdf

Version: 2024-02-01

135 papers	6,266 citations	57758 44 h-index	76900 74 g-index
163 all docs	163 docs citations	163 times ranked	7340 citing authors

#	Article	IF	Citations
1	Synthesis of [1â€8â€Nî±C]â€zanriorb A1, alanineâ€containing analogues, and their cytotoxic and antiâ€inflammatory activity. Journal of Peptide Science, 2022, 28, e3405.	1.4	1
2	Euglenatides, Potent Antiproliferative Cyclic Peptides Isolated from the Freshwater Photosynthetic Microalga Euglena gracilis. Angewandte Chemie - International Edition, 2022, 61, .	13.8	9
3	Euglenatides, Potent Antiproliferative Cyclic Peptides Isolated from the Freshwater Photosynthetic Microalga Euglena gracilis. Angewandte Chemie, 2022, 134, .	2.0	1
4	Synthesis of Carboxamideâ€Containing Tranylcypromine Analogues as LSD1 (KDM1A) Inhibitors Targeting Acute Myeloid Leukemia. ChemMedChem, 2021, 16, 1316-1324.	3.2	5
5	From Hit Seeking to Magic Bullets: The Successful Union of Epigenetic and Fragment Based Drug Discovery (EPIDD + FBDD). Journal of Medicinal Chemistry, 2021, 64, 13980-14010.	6.4	12
6	The clinical landscape of HDAC inhibitors. , 2021, , 885-899.		0
7	Insights into the Structure-Activity Relationship of Glycosides as Positive Allosteric Modulators Acting on P2X7 Receptors. Molecular Pharmacology, 2021, 99, 163-174.	2.3	8
8	Stereoselective Synthesis of Protected l-allo-Enduracididine and l-Enduracididine via Asymmetric Nitroaldol Reaction. Synthesis, 2020, 52, 942-948.	2.3	1
9	Editorial overview: Epigenetics equals chemical biology. Current Opinion in Chemical Biology, 2020, 57, A1-A4.	6.1	1
10	Two-hit wonders: The expanding universe of multitargeting epigenetic agents. Current Opinion in Chemical Biology, 2020, 57, 135-154.	6.1	30
11	Thirty Years of HDAC Inhibitors: 2020 Insight and Hindsight. Journal of Medicinal Chemistry, 2020, 63, 12460-12484.	6.4	381
12	Pharmacological inhibition of lysine-specific demethylase 1 (LSD1) induces global transcriptional deregulation and ultrastructural alterations that impair viability in Schistosoma mansoni. PLoS Neglected Tropical Diseases, 2020, 14, e0008332.	3.0	11
13	Epigenetic modulation of secondary metabolite profiles in Aspergillus calidoustus and Aspergillus westerdijkiae through histone deacetylase (HDAC) inhibition by vorinostat. Journal of Antibiotics, 2020, 73, 410-413.	2.0	16
14	New tranylcypromine derivatives containing sulfonamide motif as potent LSD1 inhibitors to target acute myeloid leukemia: Design, synthesis and biological evaluation. Bioorganic Chemistry, 2020, 99, 103808.	4.1	20
15	Three cheers for nitrogen: aza-DKPs, the aza analogues of 2,5-diketopiperazines. RSC Advances, 2020, 10, 43358-43370.	3.6	3
16	HDAC inhibitors in cancer therapy. , 2020, , 19-49.		1
17	Title is missing!. , 2020, 14, e0008332.		0
18	Title is missing!. , 2020, 14, e0008332.		0

#	Article	IF	Citations
19	Title is missing!. , 2020, 14, e0008332.		0
20	Title is missing!. , 2020, 14, e0008332.		0
21	Targeting the Zinc-Dependent Histone Deacetylases (HDACs) for Drug Discovery. Topics in Medicinal Chemistry, 2019, , 1-27.	0.8	2
22	The timeline of epigenetic drug discovery: from reality to dreams. Clinical Epigenetics, 2019, 11, 174.	4.1	275
23	A Decade of Antifungal Leads from Natural Products: 2010â€“2019. Pharmaceuticals, 2019, 12, 182.	3.8	51
24	Combined Ligand and Fragmentâ€based Drug Design of Selective Histone Deacetylase â€“ 6 Inhibitors. Molecular Informatics, 2019, 38, e1800083.	2.5	17
25	Epigenetic drug discovery: a success story for cofactor interference. Philosophical Transactions of the Royal Society B: Biological Sciences, 2018, 373, 20170069.	4.0	39
26	Isoform-selective HDAC1/6/8 inhibitors with an imidazo-ketopiperazine cap containing stereochemical diversity. Philosophical Transactions of the Royal Society B: Biological Sciences, 2018, 373, 20170364.	4.0	6
27	Epigenetics: the first 25 centuries. Philosophical Transactions of the Royal Society B: Biological Sciences, 2018, 373, 20170067.	4.0	14
28	Î²-amino alcohols and their respective 2-phenyl-N-alkyl aziridines as potential DNA minor groove binders. European Journal of Medicinal Chemistry, 2018, 157, 657-664.	5.5	16
29	Synthesis of breast cancer targeting conjugate of temporin-SHa analog and its effect on pro- and anti-apoptotic protein expression in MCF-7 cells. Peptides, 2018, 106, 68-82.	2.4	13
30	LSD1 inhibition attenuates androgen receptor V7 splice variant activation in castration resistant prostate cancer models. Cancer Cell International, 2018, 18, 71.	4.1	19
31	Cellular analysis of the action of epigenetic drugs and probes. Epigenetics, 2017, 12, 308-322.	2.7	5
32	Comparative Study of the Synthesis and Structural and Physicochemical Properties of Diketopiperazines vs Aza-diketopiperazines. Journal of Organic Chemistry, 2017, 82, 3239-3244.	3.2	7
33	Synthesis of Fluorenes with an Allâ€Carbon Quaternary Center via Palladiumâ€Catalyzed Dual Arylation using Cyclic Diaryliodonium Triflates. Advanced Synthesis and Catalysis, 2017, 359, 1152-1156.	4.3	24
34	Fluorinated tranylcypromine analogues as inhibitors of lysine-specific demethylase 1 (LSD1, KDM1A). Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2099-2101.	2.2	22
35	Systematic Analysis of the Relationship among 3D Structure, Bioactivity, and Membrane Permeability of PF1171F, a Cyclic Hexapeptide with Paralyzing Effects on Silkworms. Journal of Organic Chemistry, 2017, 82, 11447-11463.	3.2	9
36	The histone deacetylase inhibitor, romidepsin, as a potential treatment for pulmonary fibrosis. Oncotarget, 2017, 8, 48737-48754.	1.8	48

#	Article	IF	Citations
37	Epigenetic polypharmacology: from combination therapy to multitargeted drugs. Clinical Epigenetics, 2016, 8, 105.	4.1	113
38	Oral Administration of Peptideâ€Based Drugs: Beyond Lipinski's Rule. ChemMedChem, 2016, 11, 2245-2251.	3.2	104
39	Total synthesis, structural, and biological evaluation of stylissatin <scp>A</scp> and related analogs. Journal of Peptide Science, 2016, 22, 607-617.	1.4	13
40	Multitarget Drugs: an Epigenetic Epiphany. ChemMedChem, 2016, 11, 1227-1241.	3.2	76
41	Synthesis of two â€˜heteroaromatic rings of the futureâ€™ for applications in medicinal chemistry. RSC Advances, 2016, 6, 22777-22780.	3.6	9
42	cisâ€cyclopropylamines as mechanismâ€based inhibitors of monoamine oxidases. FEBS Journal, 2015, 282, 3190-3198.	4.7	31
43	Inhibition of NAADP signalling on reperfusion protects the heart by preventing lethal calcium oscillations via two-pore channel 1 and opening of the mitochondrial permeability transition pore. Cardiovascular Research, 2015, 108, 357-366.	3.8	44
44	Hologram quantitative structureâ€“activity relationship and comparative molecular interaction field analysis of aminothiazole and thiazolesulfonamide as reversible LSD1 inhibitors. Future Medicinal Chemistry, 2015, 7, 1381-1394.	2.3	12
45	Structure Revision of Similanamide to PF1171C by Total Synthesis. Journal of Natural Products, 2015, 78, 2286-2291.	3.0	12
46	Synthesis and biological evaluation of santacruzamate A and analogs as potential anticancer agents. RSC Advances, 2015, 5, 1109-1112.	3.6	7
47	Macrocyclic Inhibitors of Zinc-dependent Histone Deacetylases (HDACs). RSC Drug Discovery Series, 2014, , 109-140.	0.3	3
48	Intra- and Intermolecular Alkylation ofN,O-Acetals and Ï€-Activated Alcohols Catalyzed by in Situ Generated Acid. Journal of Organic Chemistry, 2014, 79, 1900-1912.	3.2	33
49	Synthesis of N-Acyl-N,O-acetals Mediated by Titanium Ethoxide. Organic Letters, 2014, 16, 10-13.	4.6	33
50	Oneâ€Pot Î±â€Amidosulfoneâ€Mediated Variation of the Pictetâ€“Spengler Tetrahydroisoquinoline Synthesis, Suitable for Amideâ€Type Substrates. European Journal of Organic Chemistry, 2014, 2014, 5720-5727.	2.4	11
51	Total Synthesis and Biological Evaluation of PF1171A, C, F, and G, Cyclic Hexapeptides with Insecticidal Activity. Journal of Organic Chemistry, 2014, 79, 7844-7853.	3.2	22
52	Three-Component Pd/Cu-Catalyzed Cascade Reactions of Cyclic Iodoniums, Alkynes, and Boronic Acids: An Approach to Methylidenefluorenes. Organic Letters, 2014, 16, 2350-2353.	4.6	68
53	Protein Recognition by Short Peptide Reversible Inhibitors of the Chromatin-Modifying LSD1/CoREST Lysine Demethylase. ACS Chemical Biology, 2013, 8, 1677-1682.	3.4	60
54	Expanding the Druggable Space of the LSD1/CoREST Epigenetic Target: New Potential Binding Regions for Drug-Like Molecules, Peptides, Protein Partners, and Chromatin. PLoS Computational Biology, 2013, 9, e1003158.	3.2	27

#	Article	IF	Citations
55	The Impact of Natural Products Upon Cancer Chemotherapy. , 2013, , 3-15.		0
56	Spiruchostatin A Inhibits Proliferation and Differentiation of Fibroblasts from Patients with Pulmonary Fibrosis. American Journal of Respiratory Cell and Molecular Biology, 2012, 46, 687-694.	2.9	57
57	Solid-Phase Total Synthesis of Cherimolacyclopeptide E and Discovery of More Potent Analogues by Alanine Screening. Journal of Natural Products, 2012, 75, 1882-1887.	3.0	19
58	Combinatorial Aid for Underprivileged Scaffolds: Solution and Solid-phase Strategies for a Rapid and Efficient Access To Novel Aza-diketopiperazines (Aza-DKP). ACS Combinatorial Science, 2012, 14, 323-334.	3.8	26
59	Biologic activity of LSD1 inhibition by novel tranylcypromine structural analogues in prostate cancer cells.. Journal of Clinical Oncology, 2012, 30, 82-82.	1.6	0
60	Pictet-Spengler Reaction Using Ion-Exchange Resin as a Catalyst and Support for â€˜Catch and Releaseâ€™ Purification. Bioscience, Biotechnology and Biochemistry, 2011, 75, 391-392.	1.3	3
61	The Depsipeptide HDAC Inhibitor FK228 (Romidepsin) Has Anti-Fibrotic Properties In Fibrotic Primary Pulmonary Fibroblasts. , 2011, , .		0
62	The histone deacetylase inhibitors vorinostat and romidepsin downmodulate ILâ€10 expression in cutaneous Tâ€cell lymphoma cells. British Journal of Pharmacology, 2011, 162, 1590-1602.	5.4	78
63	Total Synthesis and Stereochemical Assignment of Burkholdac B, a Depsipeptide HDAC Inhibitor. Organic Letters, 2011, 13, 6334-6337.	4.6	39
64	Enantioselective synthesis of tranylcypromine analogues as lysine demethylase (LSD1) inhibitors. Bioorganic and Medicinal Chemistry, 2011, 19, 3709-3716.	3.0	87
65	Total synthesis of largazole and analogues: HDAC inhibition, antiproliferative activity and metabolic stability. Bioorganic and Medicinal Chemistry, 2011, 19, 3650-3658.	3.0	56
66	Spiruchostatin A Inhibits Proliferation And Differentiation Of Primary Fibroblasts From Patients With Interstitial Lung Disease. , 2010, , .		0
67	TPC2 Is a Novel NAADP-sensitive Ca2+ Release Channel, Operating as a Dual Sensor of Luminal pH and Ca2+. Journal of Biological Chemistry, 2010, 285, 35039-35046.	3.4	197
68	Determination of Molecular Torsion Angles Using Nuclear Singlet Relaxation. Journal of the American Chemical Society, 2010, 132, 8225-8227.	13.7	40
69	Analogues of the Nicotinic Acid Adenine Dinucleotide Phosphate (NAADP) Antagonist Ned-19 Indicate Two Binding Sites on the NAADP Receptor. Journal of Biological Chemistry, 2009, 284, 34930-34934.	3.4	40
70	Thioflavin S (NSC71948) Interferes with Bcl-2-Associated Athanogene (BAG-1)-Mediated Protein-Protein Interactions. Journal of Pharmacology and Experimental Therapeutics, 2009, 331, 680-689.	2.5	30
71	Identification of a chemical probe for NAADP by virtual screening. Nature Chemical Biology, 2009, 5, 220-226.	8.0	274
72	A solid-phase total synthesis of the cyclic depsipeptide HDAC inhibitor spiruchostatin A. Tetrahedron Letters, 2009, 50, 2970-2972.	1.4	24

#	Article	IF	Citations
73	Epigenetic Therapy: Histone Acetylation, DNA Methylation and Anti-Cancer Drug Discovery. Current Cancer Drug Targets, 2009, 9, 963-981.	1.6	94
74	The impact of natural products upon modern drug discovery. Current Opinion in Chemical Biology, 2008, 12, 306-317.	6.1	505
75	Will histone deacetylase inhibitors require combination with other agents to fulfil their therapeutic potential?. British Journal of Cancer, 2008, 99, 689-694.	6.4	82
76	Characterisation of the in vitro activity of the depsipeptide histone deacetylase inhibitor spiruchostatin A. Biochemical Pharmacology, 2008, 76, 463-475.	4.4	67
77	Macrolactamization versus Macrolactonization: Total Synthesis of FK228, the Depsipeptide Histone Deacetylase Inhibitor. Journal of Organic Chemistry, 2008, 73, 9353-9361.	3.2	56
78	The First Biologically Active Synthetic Analogues of FK228, the Depsipeptide Histone Deacetylase Inhibitor. Journal of Medicinal Chemistry, 2007, 50, 5720-5726.	6.4	89
79	Total Synthesis of Azumamide A and Azumamide E, Evaluation as Histone Deacetylase Inhibitors, and Design of a More Potent Analogue. Organic Letters, 2007, 9, 1105-1108.	4.6	57
80	A total synthesis of spiruchostatin A. Tetrahedron Letters, 2006, 47, 1177-1180.	1.4	53
81	Azumamides Aâ€“E: Histone Deacetylase Inhibitory Cyclic Tetrapeptides from the Marine SpongeMycale izuensis. Angewandte Chemie - International Edition, 2006, 45, 7553-7557.	13.8	105
82	Cover Picture: Azumamides Aâ€“E: Histone Deacetylase Inhibitory Cyclic Tetrapeptides from the Marine SpongeMycale izuensis / Total Synthesis of Azumamides A and E Z602047 Z602033 (Angew. Chem. Int. Ed.) Tj ETQq0 0 0 rgBT /Overloc		
83	Solid-Phase Synthesis in the Twenty-First Century. Mini-Reviews in Medicinal Chemistry, 2006, 6, 3-10.	2.4	21
84	The Transcriptional Coactivator p300 Plays a Critical Role in the Hypertrophic and Protective Pathways Induced by Phenylephrine in Cardiac Cells but Is Specific to the Hypertrophic Effect of Urocortin. ChemBioChem, 2005, 6, 162-170.	2.6	40
85	Rapid deprotection of N-Boc amines by TFA combined with freebase generation using basic ion-exchange resins. Molecular Diversity, 2005, 9, 291-293.	3.9	52
86	Literature Highlights in Combinatorial Science. QSAR and Combinatorial Science, 2005, 24, 189-196.	1.4	1
87	Novel sulfasalazine analogues with enhanced NF-ĸB inhibitory and apoptosis promoting activity. Apoptosis: an International Journal on Programmed Cell Death, 2005, 10, 481-491.	4.9	63
88	PS-COD and PS-9-BBN:â€‰ Polymer-Supported Reagents for Solution-Phase Parallel Synthesisâ€. Organic Letters, 2005, 7, 831-833.	4.6	18
89	Solid-Phase Total Synthesis of Kahalalide A and Related Analogues. Journal of Medicinal Chemistry, 2005, 48, 1330-1335.	6.4	42
90	Current Strategies to Target the Anti-Apoptotic Bcl-2 Protein in Cancer Cells. Current Medicinal Chemistry, 2004, 11, 1031-1040.	2.4	40

#	Article	IF	Citations
91	Natural products as a hunting ground for combinatorial chemistry. Current Opinion in Biotechnology, 2004, 15, 584-590.	6.6	94
92	Natural products and combinatorial chemistry: back to the future. Current Opinion in Chemical Biology, 2004, 8, 271-280.	6.1	296
93	A â€˜triflate-likeâ€™ tetrafluoroarylsulfonate linker for multifunctional solid-phase organic synthesis. Chemical Communications, 2004, , 1916-1917.	4.1	27
94	Modular Three-Component Solid-Phase Synthesis of Unsymmetrical Guanidines via Resin Capture of Carbodiimidesâ€. ACS Combinatorial Science, 2004, 6, 32-34.	3.3	13
95	Total Synthesis of Spiruchostatin A, a Potent Histone Deacetylase Inhibitor. Journal of the American Chemical Society, 2004, 126, 1030-1031.	13.7	99
96	Radical Reactions in Combinatorial Chemistry. , 2004, , 225-246.		0
97	Synthesis of Functionalized 1,5-Cyclooctadienes by LICKOR Metalation.. ChemInform, 2003, 34, no.	0.0	0
98	Total Synthesis of (+)-Okaramine J Featuring an Exceptionally Facile N-Reverse-prenyl to C-Prenyl Aza-Claisen Rearrangement. Organic Letters, 2003, 5, 2825-2827.	4.6	63
99	Enzymatic Synthesis of an Indole Diterpene by an Oxidosqualene Cyclase:â€‰ Mechanistic, Biosynthetic, and Phylogenetic Implications. Journal of the American Chemical Society, 2003, 125, 9002-9003.	13.7	32
100	Cyclative Cleavage Strategies for the Solid-Phase Synthesis of Heterocycles and Natural Products. Methods in Enzymology, 2003, 369, 415-434.	1.0	15
101	Total Synthesis of Debromoflustramine B via Biomimetic Alkylative Cyclization. Organic Letters, 2003, 5, 1801-1803.	4.6	50
102	Synthesis of Functionalized 1,5-Cyclooctadienes by LICKOR Metalation. Journal of Organic Chemistry, 2002, 67, 6250-6252.	3.2	9
103	Solid-Phase Synthesis of Tetrahydro-Î²-carbolinehydantoins via the N-Acyliminium Pictetâ€“Spengler Reaction and Cyclative Cleavage. ACS Combinatorial Science, 2002, 4, 546-548.	3.3	50
104	Ionic Liquid Acceleration of Solid-Phase Suzukiâ€“Miyaura Cross-Coupling Reactions. Organic Letters, 2002, 4, 3071-3073.	4.6	93
105	Recent developments in combinatorial organic synthesis. Drug Discovery Today, 2002, 7, 47-55.	6.4	41
106	Recent developments in combinatorial organic synthesis. Drug Discovery Today, 2002, 7, 47-55.	6.4	39
107	Regioselective Synthesis of 3-Alkylindoles Mediated by Zinc Triflate. Journal of Organic Chemistry, 2002, 67, 2705-2708.	3.2	78
108	Integrating natural product synthesis and combinatorial chemistry. Pure and Applied Chemistry, 2001, 73, 1033-1039.	1.9	24

#	Article	IF	Citations
109	Parallel modification of tropane alkaloids. Tetrahedron Letters, 2001, 42, 1975-1977.	1.4	16
110	New tools for parallel automated chemistry. Drug Discovery Today, 2001, 6, 238-241.	6.4	10
111	A selenide linker for â€œtracelessâ€•solid-phase organic synthesis. Biotechnology and Bioengineering, 2000, 71, 104-106.	3.3	8
112	Total Synthesis of the Fumiquinazoline Alkaloids:Â Solid-Phase Studies1. ACS Combinatorial Science, 2000, 2, 186-194.	3.3	56
113	A Biomimetic Total Synthesis of (â^')-Spirotryprostatin B and Related Studies. Journal of Organic Chemistry, 2000, 65, 4685-4693.	3.2	100
114	Total Synthesis of the Fumiquinazoline Alkaloids:Â Solution-Phase Studies1. Journal of Organic Chemistry, 2000, 65, 1022-1030.	3.2	81
115	Synthesis and Evaluation of Tryprostatin B and Demethoxyfumitremorgin C Analogues. Journal of Medicinal Chemistry, 2000, 43, 1577-1585.	6.4	58
116	A solid-phase equivalent of van Leusen's TosMIC, and its application in oxazole synthesis. Tetrahedron Letters, 1999, 40, 5633-5636.	1.4	47
117	Solution-phase parallel oxazole synthesis with TosMIC. Tetrahedron Letters, 1999, 40, 5637-5638.	1.4	46
118	Targeting proteinÃ¢Â€Â“protein interactions: the HIV protease. Drug Discovery Today, 1999, 4, 387-388.	6.4	0
119	The N-Acyliminium Pictetâ^'Spengler Condensation as a Multicomponent Combinatorial Reaction on Solid Phase and Its Application to the Synthesis of Demethoxyfumitremorgin C Analogues. Organic Letters, 1999, 1, 1647-1649.	4.6	107
120	Solid-Phase Synthesis of Î²-Keto Esters via Sequential Baylisâ^'Hillman and Heck Reactions. ACS Combinatorial Science, 1999, 1, 373-378.	3.3	49
121	Solid-phase C-acylation of active methylene compounds. Tetrahedron Letters, 1998, 39, 2195-2198.	1.4	31
122	Solid-phase synthesis of tetramic acids. Tetrahedron Letters, 1998, 39, 4369-4372.	1.4	41
123	Solid-phase synthesis of N-acyl-Nâ€²-carbamoylguanidines. Tetrahedron Letters, 1998, 39, 9789-9792.	1.4	22
124	Solid-phase combinatorial synthesis of 4-hydroxyquinolin-2(1H)-ones. Tetrahedron Letters, 1998, 39, 6399-6402.	1.4	38
125	Solid-phase synthesis of peptidomimetic oligomers with a phosphodiester backbone. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 511-514.	2.2	19
126	Solid-phase synthesis of potential protein tyrosine phosphatase inhibitors via the Ugi four-component condensation. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 2443-2446.	2.2	27

#	Article	IF	Citations
127	Solution-phase combinatorial synthesis of 4-hydroxyquinolin-2(1H)-ones. Chemical Communications, 1998, , 785-786.	4.1	30
128	Poly(4-Vinylpyridinium P-Toluenesulfonate) as a Polymer-Supported Catalyst for Hydrolysis of Tetrahydropyranyl Ethers. Synthetic Communications, 1998, 28, 3209-3212.	2.1	20
129	Solution-Phase Synthesis of a Combinatorial Thiohydantoin Library1. Journal of Organic Chemistry, 1997, 62, 3230-3235.	3.2	72
130	Solution-phase synthesis of a Î²-amino alcohol combinatorial library. Bioorganic and Medicinal Chemistry Letters, 1997, 7, 1511-1514.	2.2	88
131	Concise synthesis of the cell cycle inhibitor demethoxyfumitremorgin C. Tetrahedron Letters, 1997, 38, 4327-4328.	1.4	39
132	When Two Steroids are Better than One: The Dimeric Steroid-Pyrazine Marine Alkaloids. Studies in Natural Products Chemistry, 1995, 18, 875-906.	1.8	11
133	Specific binding of the DNA repair enzyme AlkA to a pyrrolidine-based inhibitor. Journal of the American Chemical Society, 1995, 117, 6623-6624.	13.7	54
134	A Stereochemical test of the mechanism of electrophilic substitution in 3-substituted indoles. Tetrahedron Letters, 1993, 34, 439-440.	1.4	36
135	Total Synthesis of Altissimacoumarin D, a Small Molecule Sirtuin1 Activator. Journal of the Brazilian Chemical Society, 0, , .	0.6	1