

# James R Fuchs

## List of Publications by Year in descending order

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88  
papers

5,397  
citations

57758

44  
h-index

88630

70  
g-index

97  
all docs

97  
docs citations

97  
times ranked

6650  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pancreatic Cancer-Associated Stellate Cells Promote Differentiation of Myeloid-Derived Suppressor Cells in a STAT3-Dependent Manner. <i>Cancer Research</i> , 2013, 73, 3007-3018.	0.9	340
2	Novel STAT3 Phosphorylation Inhibitors Exhibit Potent Growth-Suppressive Activity in Pancreatic and Breast Cancer Cells. <i>Cancer Research</i> , 2010, 70, 2445-2454.	0.9	211
3	Allosteric integrase inhibitor potency is determined through the inhibition of HIV-1 particle maturation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 8690-8695.	7.1	178
4	Total Synthesis of (±)-Perophoramidine. <i>Journal of the American Chemical Society</i> , 2004, 126, 5068-5069.	13.7	175
5	Reactions of SmI <sub>2</sub> with Alkyl Halides and Ketones: An Inner-Sphere vs Outer-Sphere Electron Transfer in Reactions of Sm(II) Reductants. <i>Journal of the American Chemical Society</i> , 2000, 122, 7718-7722.	13.7	170
6	Multimode, Cooperative Mechanism of Action of Allosteric HIV-1 Integrase Inhibitors. <i>Journal of Biological Chemistry</i> , 2012, 287, 16801-16811.	3.4	167
7	Targeting colon cancer stem cells using a new curcumin analogue, GO-Y030. <i>British Journal of Cancer</i> , 2011, 105, 212-220.	6.4	137
8	Modulation of DNA Methylation by a Sesquiterpene Lactone Parthenolide. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2009, 329, 505-514.	2.5	133
9	Structural and mechanistic bases for a potent HIV-1 capsid inhibitor. <i>Science</i> , 2020, 370, 360-364.	12.6	114
10	Indol-2-one Intermediates: Mechanistic Evidence and Synthetic Utility. Total Syntheses of (±)-Flustramines A and C. <i>Organic Letters</i> , 2005, 7, 677-680.	4.6	112
11	STAT3 signaling pathway is necessary for cell survival and tumorsphere forming capacity in ALDH <sup>+</sup> /CD133 <sup>+</sup> stem cell-like human colon cancer cells. <i>Biochemical and Biophysical Research Communications</i> , 2011, 416, 246-251.	2.1	112
12	A New Class of Multimerization Selective Inhibitors of HIV-1 Integrase. <i>PLoS Pathogens</i> , 2014, 10, e1004171.	4.7	112
13	Intramolecular Diels-Alder/1,3-Dipolar Cycloaddition Cascade of 1,3,4-Oxadiazoles. <i>Journal of the American Chemical Society</i> , 2006, 128, 10589-10595.	13.7	111
14	HIV-1 Integrase Binds the Viral RNA Genome and Is Essential during Virion Morphogenesis. <i>Cell</i> , 2016, 166, 1257-1268.e12.	28.9	110
15	Rocaglamide, silvestrol and structurally related bioactive compounds from <i>Aglaia</i> species. <i>Natural Product Reports</i> , 2014, 31, 924-939.	10.3	108
16	The small molecule curcumin analog FLLL32 induces apoptosis in melanoma cells via STAT3 inhibition and retains the cellular response to cytokines with anti-tumor activity. <i>Molecular Cancer</i> , 2010, 9, 165.	19.2	106
17	Structure-activity relationship studies of curcumin analogues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 2065-2069.	2.2	105
18	Total Synthesis of (±)-Lennoxamine and (±)-Aphanorphine by Intramolecular Electrophilic Aromatic Substitution Reactions of 2-Amidoacroleins. <i>Organic Letters</i> , 2001, 3, 3923-3925.	4.6	104

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19	The effect of cosolvent on the reducing power of SmI2 in tetrahydrofuran. <i>Tetrahedron Letters</i> , 1998, 39, 4429-4432.	1.4	103
20	Discovery of Anticancer Agents of Diverse Natural Origin. <i>Anticancer Research</i> , 2016, 36, 5623-5638.	1.1	94
21	Distribution and Redistribution of HIV-1 Nucleocapsid Protein in Immature, Mature, and Integrase-Inhibited Virions: a Role for Integrase in Maturation. <i>Journal of Virology</i> , 2015, 89, 9765-9780.	3.4	91
22	Curcumin induces proapoptotic effects against human melanoma cells and modulates the cellular response to immunotherapeutic cytokines. <i>Molecular Cancer Therapeutics</i> , 2009, 8, 2726-2735.	4.1	90
23	The effect of lithium bromide and lithium chloride on the reactivity of SmI2 in THF. <i>Tetrahedron Letters</i> , 1997, 38, 8157-8158.	1.4	88
24	HRP2 determines the efficiency and specificity of HIV-1 integration in LEDGF/p75 knockout cells but does not contribute to the antiviral activity of a potent LEDGF/p75-binding site integrase inhibitor. <i>Nucleic Acids Research</i> , 2012, 40, 11518-11530.	14.5	86
25	The A128T Resistance Mutation Reveals Aberrant Protein Multimerization as the Primary Mechanism of Action of Allosteric HIV-1 Integrase Inhibitors. <i>Journal of Biological Chemistry</i> , 2013, 288, 15813-15820.	3.4	85
26	The novel curcumin analog FLLL32 decreases STAT3 DNA binding activity and expression, and induces apoptosis in osteosarcoma cell lines. <i>BMC Cancer</i> , 2011, 11, 112.	2.6	83
27	New structural analogues of curcumin exhibit potent growth suppressive activity in human colorectal carcinoma cells. <i>BMC Cancer</i> , 2009, 9, 99.	2.6	82
28	New curcumin analogues exhibit enhanced growth suppressive activity and inhibit AKT and signal transducer and activator of transcription 3 phosphorylation in breast and prostate cancer cells. <i>Cancer Science</i> , 2009, 100, 1719-1727.	3.9	82
29	Site-Selective C-H Functionalization of (Hetero)Arenes via Transient, Non-symmetric Iodanes. <i>CheM</i> , 2019, 5, 417-428.	11.7	80
30	Structurally Modified Curcumin Analogs Inhibit STAT3 Phosphorylation and Promote Apoptosis of Human Renal Cell Carcinoma and Melanoma Cell Lines. <i>PLoS ONE</i> , 2012, 7, e40724.	2.5	80
31	EGFR-STAT3 signaling promotes formation of malignant peripheral nerve sheath tumors. <i>Oncogene</i> , 2014, 33, 173-180.	5.9	75
32	Small molecules, LLL12 and FLLL32, inhibit STAT3 and exhibit potent growth suppressive activity in osteosarcoma cells and tumor growth in mice. <i>Investigational New Drugs</i> , 2012, 30, 916-926.	2.6	67
33	IL-6, a risk factor for hepatocellular carcinoma: FLLL32 inhibits IL-6-Induced STAT3 phosphorylation in human hepatocellular cancer cells. <i>Cell Cycle</i> , 2010, 9, 3423-3427.	2.6	65
34	HAb18G/CD147 Promotes pSTAT3-Mediated Pancreatic Cancer Development via CD44s. <i>Clinical Cancer Research</i> , 2013, 19, 6703-6715.	7.0	65
35	Potent Cytotoxic Arylnaphthalene Lignan Lactones from <i>Phyllanthus poilanei</i> . <i>Journal of Natural Products</i> , 2014, 77, 1494-1504.	3.0	65
36	Bioactive Flavaglines and Other Constituents Isolated from <i>Aglaia perviridis</i> . <i>Journal of Natural Products</i> , 2013, 76, 394-404.	3.0	63

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37	TALEN Knockout of the <i>PSIP1</i> Gene in Human Cells: Analyses of HIV-1 Replication and Allosteric Integrase Inhibitor Mechanism. <i>Journal of Virology</i> , 2014, 88, 9704-9717.	3.4	63
38	Overcoming chemo/radio-resistance of pancreatic cancer by inhibiting STAT3 signaling. <i>Oncotarget</i> , 2016, 7, 11708-11723.	1.8	58
39	Curcumin analogues exhibit enhanced growth suppressive activity in human pancreatic cancer cells. <i>Anti-Cancer Drugs</i> , 2009, 20, 444-449.	1.4	57
40	Cytotoxic and NF- $\kappa$ B Inhibitory Constituents of the Stems of <i>Cratoxylum cochinchinense</i> and Their Semisynthetic Analogues. <i>Journal of Natural Products</i> , 2011, 74, 1117-1125.	3.0	57
41	Insertional Mutagenesis Identifies a STAT3/Arid1b/ $\beta$ -catenin Pathway Driving Neurofibroma Initiation. <i>Cell Reports</i> , 2016, 14, 1979-1990.	6.4	55
42	Sterols with antileishmanial activity isolated from the roots of <i>Pentalinon andrieuxii</i> . <i>Phytochemistry</i> , 2012, 82, 128-135.	2.9	49
43	A liquid chromatography-tandem mass spectrometric method for quantification of curcuminoids in cell medium and mouse plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2010, 878, 3045-3051.	2.3	48
44	Potential Anticancer Agents Characterized from Selected Tropical Plants. <i>Journal of Natural Products</i> , 2019, 82, 657-679.	3.0	48
45	HIV-1 integrase tetramers are the antiviral target of pyridine-based allosteric integrase inhibitors. <i>ELife</i> , 2019, 8, .	6.0	41
46	Sensitization of Head and Neck Cancer to Cisplatin Through the Use of a Novel Curcumin Analog. <i>JAMA Otolaryngology</i> , 2011, 137, 499.	1.2	40
47	Intramolecular Electrophilic Aromatic Substitution Reactions of 2-Amidoacroleins: A New Method for the Preparation of Tetrahydroisoquinolines, Tetrahydro-3-benzazepines, and Hexahydro-3-benzazocines. <i>Organic Letters</i> , 2001, 3, 3349-3351.	4.6	39
48	The mechanism of H171T resistance reveals the importance of N <sup>+</sup> -protonated His171 for the binding of allosteric inhibitor BI-D to HIV-1 integrase. <i>Retrovirology</i> , 2014, 11, 100.	2.0	39
49	Pro-growth role of the JMJD2C histone demethylase in HCT-116 colon cancer cells and identification of curcuminoids as JMJD2 inhibitors. <i>American Journal of Translational Research (discontinued)</i> , 2014, 6, 236-47.	0.0	39
50	Inhibition of the JAK2/STAT3 signaling pathway exerts a therapeutic effect on osteosarcoma. <i>Molecular Medicine Reports</i> , 2015, 12, 498-502.	2.4	37
51	Cytotoxic and natural killer cell stimulatory constituents of <i>Phyllanthus songboiensis</i> . <i>Phytochemistry</i> , 2015, 111, 132-140.	2.9	36
52	Phosphoproteomic analysis of anaplastic lymphoma kinase (ALK) downstream signaling pathways identifies signal transducer and activator of transcription 3 as a functional target of activated ALK in neuroblastoma cells. <i>FEBS Journal</i> , 2013, 280, 5269-5282.	4.7	35
53	Trichormamides A and B with Antiproliferative Activity from the Cultured Freshwater Cyanobacterium <i>Trichormus</i> sp. UIC 10339. <i>Journal of Natural Products</i> , 2014, 77, 1871-1880.	3.0	31
54	Resistance to pyridine-based inhibitor KF116 reveals an unexpected role of integrase in HIV-1 Gag-Pol polyprotein proteolytic processing. <i>Journal of Biological Chemistry</i> , 2017, 292, 19814-19825.	3.4	31

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55	Allosteric HIV-1 integrase inhibitors promote aberrant protein multimerization by directly mediating inter-subunit interactions: Structural and thermodynamic modeling studies. <i>Protein Science</i> , 2016, 25, 1911-1917.	7.6	30
56	The Competitive Interplay between Allosteric HIV-1 Integrase Inhibitor BI/D and LEDGF/p75 during the Early Stage of HIV-1 Replication Adversely Affects Inhibitor Potency. <i>ACS Chemical Biology</i> , 2016, 11, 1313-1321.	3.4	29
57	A Critical Role of the C-terminal Segment for Allosteric Inhibitor-induced Aberrant Multimerization of HIV-1 Integrase. <i>Journal of Biological Chemistry</i> , 2014, 289, 26430-26440.	3.4	28
58	Indole-based allosteric inhibitors of HIV-1 integrase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 4748-4752.	2.2	28
59	STarFish: A Stacked Ensemble Target Fishing Approach and its Application to Natural Products. <i>Journal of Chemical Information and Modeling</i> , 2019, 59, 4906-4920.	5.4	27
60	Novel Anti-Campylobacter Compounds Identified Using High Throughput Screening of a Pre-selected Enriched Small Molecules Library. <i>Frontiers in Microbiology</i> , 2016, 7, 405.	3.5	24
61	Phyllanthusmin Derivatives Induce Apoptosis and Reduce Tumor Burden in High-Grade Serous Ovarian Cancer by Late-Stage Autophagy Inhibition. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 2123-2135.	4.1	24
62	Identification of a Small Molecule Anti-biofilm Agent Against <i>Salmonella enterica</i> . <i>Frontiers in Microbiology</i> , 2018, 9, 2804.	3.5	23
63	Trichormamides C and D, antiproliferative cyclic lipopeptides from the cultured freshwater cyanobacterium cf. <i>Oscillatoria</i> sp. UIC 10045. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 3153-3162.	3.0	22
64	Synthesis and antiproliferative activity of derivatives of the phyllanthusmin class of aryl-naphthalene lignan lactones. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 2354-2364.	3.0	21
65	<sup>15</sup> N Stable Isotope Labeling and Comparative Metabolomics Facilitates Genome Mining in Cultured Cyanobacteria. <i>ACS Chemical Biology</i> , 2020, 15, 758-765.	3.4	21
66	A New Class of Allosteric HIV-1 Integrase Inhibitors Identified by Crystallographic Fragment Screening of the Catalytic Core Domain. <i>Journal of Biological Chemistry</i> , 2016, 291, 23569-23577.	3.4	20
67	Novel small molecule IL-6 inhibitor suppresses autoreactive Th17 development and promotes Treg development. <i>Clinical and Experimental Immunology</i> , 2019, 196, 215-225.	2.6	19
68	Discovery of Anticancer Agents of Diverse Natural Origin. <i>Journal of Natural Products</i> , 2022, 85, 702-719.	3.0	19
69	Discovery of novel small molecule modulators of <i>Clavibacter michiganensis</i> subsp. <i>michiganensis</i> . <i>Frontiers in Microbiology</i> , 2015, 6, 1127.	3.5	18
70	A Novel Sterol Isolated from a Plant Used by Mayan Traditional Healers Is Effective in Treatment of Visceral Leishmaniasis Caused by <i>Leishmania donovani</i> . <i>ACS Infectious Diseases</i> , 2015, 1, 497-506.	3.8	18
71	An Isoquinoline Scaffold as a Novel Class of Allosteric HIV-1 Integrase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 215-220.	2.8	18
72	Two small molecule compounds, LLL12 and FLLL32, exhibit potent inhibitory activity on STAT3 in human rhabdomyosarcoma cells. <i>International Journal of Oncology</i> , 2011, 38, 279-85.	3.9	18

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73	Novel narrow spectrum benzyl thiophene sulfonamide derivatives to control <i>Campylobacter</i> . <i>Journal of Antibiotics</i> , 2019, 72, 555-565.	2.0	14
74	FLLL12 induces apoptosis in lung cancer cells through a p53/p73-independent but death receptor 5-dependent pathway. <i>Cancer Letters</i> , 2015, 363, 166-175.	7.2	13
75	Evaluation of synergy between host and pathogen-directed therapies against intracellular <i>Leishmania donovani</i> . <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2019, 10, 125-132.	3.4	12
76	Total Synthesis of Scytonemide A Employing Weinreb AM Solid-Phase Resin. <i>Journal of Natural Products</i> , 2018, 81, 534-542.	3.0	11
77	Semisynthetic Derivatives of the Verticillin Class of Natural Products through Acylation of the C11 Hydroxy Group. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 625-630.	2.8	11
78	Evaluation of Novel Quorum Sensing Inhibitors Targeting Auto-Inducer 2 (AI-2) for the Control of Avian Pathogenic <i>Escherichia coli</i> Infections in Chickens. <i>Microbiology Spectrum</i> , 2022, 10, e0028622.	3.0	11
79	Computational and synthetic approaches for developing Lavendustin B derivatives as allosteric inhibitors of HIV-1 integrase. <i>European Journal of Medicinal Chemistry</i> , 2016, 123, 673-683.	5.5	10
80	Pentalinosterol, a Constituent of <i>Pentalinon andrieuxii</i> , Possesses Potent Immunomodulatory Activity and Primes T Cell Immune Responses. <i>Journal of Natural Products</i> , 2017, 80, 2515-2523.	3.0	10
81	PHY34 inhibits autophagy through V-ATPase VOA2 subunit inhibition and CAS/CSE1L nuclear cargo trafficking in high grade serous ovarian cancer. <i>Cell Death and Disease</i> , 2022, 13, 45.	6.3	10
82	Preclinical <i>In Vitro</i> , <i>In Vivo</i> , and Pharmacokinetic Evaluations of FLLL12 for the Prevention and Treatment of Head and Neck Cancers. <i>Cancer Prevention Research</i> , 2016, 9, 63-73.	1.5	9
83	Calothrixamides A and B from the Cultured Cyanobacterium <i>Calothrix</i> sp. UIC 10520. <i>Journal of Natural Products</i> , 2018, 81, 2083-2090.	3.0	9
84	Use of the Hollow Fiber Assay for the Discovery of Novel Anticancer Agents from Fungi. <i>Methods in Molecular Biology</i> , 2012, 944, 267-277.	0.9	6
85	Specialized metabolites of the United States lichen <i>Niebla homalea</i> and their antiproliferative activities. <i>Phytochemistry</i> , 2020, 180, 112521.	2.9	6
86	Identification and Optimization of a Novel HIV-1 Integrase Inhibitor. <i>ACS Omega</i> , 2022, 7, 4482-4491.	3.5	4
87	Pentalinosterol, a Phytosterol from <i>Pentalinon andrieuxii</i> , is Immunomodulatory through Phospholipase A2 in Macrophages toward its Antileishmanial Action. <i>Cell Biochemistry and Biophysics</i> , 2021, , 1.	1.8	2
88	Preparation and Cyclization of 2-Amidoacroleins: Application to the Synthesis of Alkaloid Natural Products. , 2003, , 153.		0