

Haijun Chen

List of Publications by Year in descending order

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77
papers

2,568
citations

185998

28
h-index

197535

49
g-index

78
all docs

78
docs citations

78
times ranked

3976
citing authors

#	ARTICLE	IF	CITATIONS
1	Recent progress in development of new sonosensitizers for sonodynamic cancer therapy. <i>Drug Discovery Today</i> , 2014, 19, 502-509.	3.2	280
2	Nanotechnology-based intelligent drug design for cancer metastasis treatment. <i>Biotechnology Advances</i> , 2014, 32, 761-777.	6.0	151
3	Evolution in medicinal chemistry of ursolic acid derivatives as anticancer agents. <i>European Journal of Medicinal Chemistry</i> , 2015, 92, 648-655.	2.6	116
4	Discovery of <i>N</i> -Alkylamino-Tethered Niclosamide Derivatives as Potent and Orally Bioavailable Anticancer Agents. <i>ACS Medicinal Chemistry Letters</i> , 2013, 4, 180-185.	1.3	108
5	Mifepristone Suppresses Basal Triple-Negative Breast Cancer Stem Cells by Down-regulating KLF5 Expression. <i>Theranostics</i> , 2016, 6, 533-544.	4.6	103
6	Exploring therapeutic potentials of baicalin and its aglycone baicalein for hematological malignancies. <i>Cancer Letters</i> , 2014, 354, 5-11.	3.2	102
7	Evolutions in fragment-based drug design: the deconstruction–reconstruction approach. <i>Drug Discovery Today</i> , 2015, 20, 105-113.	3.2	99
8	Fragment-based drug design and identification of HJC0123, a novel orally bioavailable STAT3 inhibitor for cancer therapy. <i>European Journal of Medicinal Chemistry</i> , 2013, 62, 498-507.	2.6	91
9	Biochemical and Pharmacological Characterizations of ESI-09 Based EPAC Inhibitors: Defining the ESI-09 –Therapeutic Window–. <i>Scientific Reports</i> , 2015, 5, 9344.	1.6	90
10	Dendrimeric anticancer prodrugs for targeted delivery of ursolic acid to folate receptor-expressing cancer cells: Synthesis and biological evaluation. <i>European Journal of Pharmaceutical Sciences</i> , 2015, 70, 55-63.	1.9	64
11	Targeting Kruppel-Like Factor 5 (KLF5) for Cancer Therapy. <i>Current Topics in Medicinal Chemistry</i> , 2015, 15, 699-713.	1.0	63
12	Identification and Characterization of Small Molecules as Potent and Specific EPAC2 Antagonists. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 952-962.	2.9	59
13	Chloroquine in combination with aptamer-modified nanocomplexes for tumor vessel normalization and efficient erlotinib/Survivin shRNA co-delivery to overcome drug resistance in EGFR-mutated non-small cell lung cancer. <i>Acta Biomaterialia</i> , 2018, 76, 257-274.	4.1	58
14	Chitosan-based nanoparticles for improved anticancer efficacy and bioavailability of mifepristone. <i>Beilstein Journal of Nanotechnology</i> , 2016, 7, 1861-1870.	1.5	57
15	5-Cyano-6-oxo-1,6-dihydro-pyrimidines as potent antagonists targeting exchange proteins directly activated by cAMP. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 4038-4043.	1.0	52
16	Discovery of potent anticancer agent HJC0416, an orally bioavailable small molecule inhibitor of signal transducer and activator of transcription 3 (STAT3). <i>European Journal of Medicinal Chemistry</i> , 2014, 82, 195-203.	2.6	52
17	Recent Advances in the Discovery of Small Molecules Targeting Exchange Proteins Directly Activated by cAMP (EPAC). <i>Journal of Medicinal Chemistry</i> , 2014, 57, 3651-3665.	2.9	46
18	Near-infrared/pH dual-responsive nanocomplexes for targeted imaging and chemo/gene/photothermal tri-therapies of non-small cell lung cancer. <i>Acta Biomaterialia</i> , 2020, 107, 242-259.	4.1	45

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19	Role of generation on folic acid-modified poly(amidoamine) dendrimers for targeted delivery of baicalin to cancer cells. <i>Materials Science and Engineering C</i> , 2017, 75, 182-190.	3.8	39
20	Structure-Activity Relationship Studies of Substituted 2-(Isoxazol-3-yl)-2-oxo-1-phenyl-acetohydrazonoyl Cyanide Analogues: Identification of Potent Exchange Proteins Directly Activated by cAMP (EPAC) Antagonists. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6033-6047.	2.9	38
21	Biomimetic Oxidative Coupling Cyclization Enabling Rapid Construction of Isochromanoindolenines. <i>Organic Letters</i> , 2018, 20, 5457-5460.	2.4	37
22	Oxidative Rearrangement Coupling Reaction for the Functionalization of Tetrahydrocarbolines with Aromatic Amines. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 14968-14972.	7.2	36
23	Efficient synthesis of ESI-09, a novel non-cyclic nucleotide EPAC antagonist. <i>Tetrahedron Letters</i> , 2013, 54, 1546-1549.	0.7	33
24	Design, synthesis, and characterization of novel apigenin analogues that suppress pancreatic stellate cell proliferation in vitro and associated pancreatic fibrosis in vivo. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 3393-3404.	1.4	33
25	Folate and Heptamethine Cyanine Modified Chitosan-Based Nanotheranostics for Tumor Targeted Near-Infrared Fluorescence Imaging and Photodynamic Therapy. <i>Biomacromolecules</i> , 2017, 18, 2146-2160.	2.6	33
26	Monascus Pigment Rubropunctatin: A Potential Dual Agent for Cancer Chemotherapy and Phototherapy. <i>Journal of Agricultural and Food Chemistry</i> , 2016, 64, 2541-2548.	2.4	32
27	Erlotinib-Guided Self-Assembled Trifunctional Click Nanotheranostics for Distinguishing Druggable Mutations and Synergistic Therapy of Nonsmall Cell Lung Cancer. <i>Molecular Pharmaceutics</i> , 2018, 15, 5146-5161.	2.3	32
28	Dual-responsive nanosystem for precise molecular subtyping and resistant reversal of EGFR targeted therapy. <i>Chemical Engineering Journal</i> , 2019, 372, 483-495.	6.6	32
29	STAT3 inhibition suppresses hepatic stellate cell fibrogenesis: HJC0123, a potential therapeutic agent for liver fibrosis. <i>RSC Advances</i> , 2016, 6, 100652-100663.	1.7	28
30	Synthesis and potent cytotoxic activity of a novel diosgenin derivative and its phytosomes against lung cancer cells. <i>Beilstein Journal of Nanotechnology</i> , 2019, 10, 1933-1942.	1.5	27
31	Acetic Acid Accelerated Visible-Light Photoredox Catalyzed Demethylation of Dimethylaminophenyl Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 687-692.	2.1	26
32	STAT3 Modulation to Enhance Motor Neuron Differentiation in Human Neural Stem Cells. <i>PLoS ONE</i> , 2014, 9, e100405.	1.1	25
33	Ursolic acid derivative FZU-03,010 inhibits STAT3 and induces cell cycle arrest and apoptosis in renal and breast cancer cells. <i>Acta Biochimica Et Biophysica Sinica</i> , 2017, 49, 367-373.	0.9	25
34	A Combined Bioinformatics and Chemoinformatics Approach for Developing Asymmetric Bivalent AMPA Receptor Positive Allosteric Modulators as Neuroprotective Agents. <i>ChemMedChem</i> , 2013, 8, 226-230.	1.6	24
35	Chemical Modification of Chitosan for Developing Cancer Nanotheranostics. <i>Biomacromolecules</i> , 2022, 23, 2197-2218.	2.6	24
36	Indocyanine green-encapsulated erlotinib modified chitosan nanoparticles for targeted chemo-photodynamic therapy of lung cancer cells. <i>Dyes and Pigments</i> , 2019, 170, 107588.	2.0	21

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37	Challenges and Opportunities from Basic Cancer Biology for Nanomedicine for Targeted Drug Delivery. <i>Current Cancer Drug Targets</i> , 2019, 19, 257-276.	0.8	21
38	Scaffold Repurposing of Old Drugs Towards New Cancer Drug Discovery. <i>Current Topics in Medicinal Chemistry</i> , 2016, 16, 2107-2114.	1.0	21
39	Design, synthesis and biological evaluation of a novel Cu ²⁺ -selective fluorescence sensor for bio-detection and chelation. <i>RSC Advances</i> , 2015, 5, 80110-80117.	1.7	20
40	An intelligent hypoxia-relieving chitosan-based nanoplatfrom for enhanced targeted chemo-sonodynamic combination therapy on lung cancer. <i>Carbohydrate Polymers</i> , 2021, 274, 118655.	5.1	20
41	Self-assembled chitosan/rose bengal derivative nanoparticles for targeted sonodynamic therapy: preparation and tumor accumulation. <i>RSC Advances</i> , 2015, 5, 17915-17923.	1.7	19
42	One-pot synthesis of tricyclo-1,4-benzoxazines via visible-light photoredox catalysis in continuous flow. <i>Tetrahedron Letters</i> , 2017, 58, 1395-1398.	0.7	17
43	Tri-component programmable nanoregulator with Three-pronged penetration boosts immunotherapy of Triple-Negative breast cancer. <i>Chemical Engineering Journal</i> , 2022, 439, 135712.	6.6	17
44	Discovery of novel mifepristone derivatives via suppressing KLF5 expression for the treatment of triple-negative breast cancer. <i>European Journal of Medicinal Chemistry</i> , 2018, 146, 354-367.	2.6	16
45	Catalytic Oxidative Coupling Cyclization for Construction of Benzofuroindolenines under Mild Reaction Conditions. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 432-435.	2.1	16
46	Hypoxia/pH dual-responsive nitroimidazole-modified chitosan/rose bengal derivative nanoparticles for enhanced photodynamic anticancer therapy. <i>Dyes and Pigments</i> , 2020, 179, 108395.	2.0	16
47	Co-delivery of gefitinib and hematoporphyrin by aptamer-modified fluorinated dendrimer for hypoxia alleviation and enhanced synergistic chemo-photodynamic therapy of NSCLC. <i>European Journal of Pharmaceutical Sciences</i> , 2021, 167, 106004.	1.9	15
48	Mifepristone Derivative FZU-00,003 Suppresses Triple-negative Breast Cancer Cell Growth partially via miR-153-KLF5 axis. <i>International Journal of Biological Sciences</i> , 2020, 16, 611-619.	2.6	14
49	Synthesis and structure-activity relationship studies of MI-2 analogues as MALT1 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 3321-3344.	1.4	13
50	Manipulation of Water for Diversified Functionalization of Tetrahydro- β -carbolines (TH β Cs) with Indoles. <i>Organic Letters</i> , 2019, 21, 6160-6163.	2.4	13
51	A novel synthetic ursolic acid derivative inhibits growth and induces apoptosis in breast cancer cell lines. <i>Oncology Letters</i> , 2018, 15, 2323-2329.	0.8	11
52	AMPA receptor positive allosteric modulators attenuate morphine tolerance and dependence. <i>Neuropharmacology</i> , 2018, 137, 50-58.	2.0	11
53	Oxidation of Tetrahydro- β -carbolines by Persulfate. <i>Organic Letters</i> , 2019, 21, 7475-7477.	2.4	11
54	Diverse Functionalization of Tetrahydro- β -carbolines or Tetrahydro- β -carbolines via Oxidative Coupling Rearrangement. <i>Journal of Organic Chemistry</i> , 2021, 86, 794-812.	1.7	11

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55	Discovery of FZU-03,010 as a self-assembling anticancer amphiphile for acute myeloid leukemia. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 1007-1011.	1.0	9
56	Discovery of novel negletein derivatives as potent anticancer agents for acute myeloid leukemia. <i>Chemical Biology and Drug Design</i> , 2018, 91, 924-932.	1.5	9
57	Construction of Bisindolines via Oxidative Coupling Cyclization. <i>Organic Letters</i> , 2020, 22, 116-119.	2.4	9
58	Doxorubicin/Nucleophosmin Binding Protein-Conjugated Nanoparticle Enhances Anti-leukemia Activity in Acute Lymphoblastic Leukemia Cells in vitro and in vivo. <i>Frontiers in Pharmacology</i> , 2021, 12, 607755.	1.6	9
59	E35 ablates acute leukemia stem and progenitor cells in vitro and in vivo. <i>Journal of Cellular Physiology</i> , 2020, 235, 8023-8034.	2.0	8
60	Convenient Tuning of the Elasticity of Self-Assembled Nano-Sized Triterpenoids to Regulate Their Biological Activities. <i>ACS Applied Materials & Interfaces</i> , 2021, 13, 44065-44078.	4.0	8
61	Development of a concise synthetic approach to access oroxin A. <i>RSC Advances</i> , 2014, 4, 45151-45154.	1.7	7
62	Nickel-catalyzed C–O Cross-Coupling Reaction at Low Catalytic Loading with Weak Base Participation. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 519-522.	1.2	7
63	Oxidative Rearrangement Coupling Reaction for the Functionalization of Tetrahydrocarbolines with Aromatic Amines. <i>Angewandte Chemie</i> , 2017, 129, 15164-15168.	1.6	6
64	Stabilization of Transient 3-Chloroindolenines Enables Diverse Functionalization. <i>Organic Letters</i> , 2019, 21, 8884-8887.	2.4	6
65	Synthesis and structure–activity relationship studies of LLY-507 analogues as SMYD2 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127598.	1.0	5
66	Synthesis of metapristone through an efficient N-demethylation of mifepristone. <i>RSC Advances</i> , 2016, 6, 7195-7197.	1.7	4
67	<i>Monascus</i> pigment rubropunctatin derivative FZU-H reduces A β (1-42)-induced neurotoxicity in Neuro-2A cells. <i>RSC Advances</i> , 2018, 8, 17389-17398.	1.7	4
68	Facile access to evodiakine enabled by aerobic copper-catalyzed oxidative rearrangement. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 8811-8815.	1.5	3
69	Construction and biological evaluation of different self-assembled nanoarchitectures of FZU-03,010. <i>European Journal of Pharmaceutical Sciences</i> , 2018, 121, 382-391.	1.9	2
70	Isochromanoindolenines suppress triple-negative breast cancer cell proliferation partially via inhibiting Akt activation. <i>International Journal of Biological Sciences</i> , 2021, 17, 986-994.	2.6	2
71	Pyrrolo [3,4-b]-quinolin-9-amine compound FZU-0038-056 suppresses triple-negative breast cancer partially through inhibiting the expression of Bcl-2. <i>Aging</i> , 2020, 12, 9621-9632.	1.4	2
72	A Direct Approach to 3-Azo-Substituted 2-Oxindoles at Room Temperature by Nickel-catalyzed Oxidative Coupling Reaction. <i>Asian Journal of Organic Chemistry</i> , 2019, 8, 475-478.	1.3	1

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73	Cu ^{II} -Catalyzed Aerobic Oxidative Coupling of Tetrahydro- β -carbolines with Indoles. <i>ChemistrySelect</i> , 2021, 6, 6272-6274.	0.7	1
74	Adriamycin/Nucleophosmin Binding Protein-Conjugated Nanoparticle (ADR-PMs-NPMBP) Enhances Anti-Leukemia Activities of Adriamycin in Acute Lymphoblastic Leukemia Cells. <i>Blood</i> , 2020, 136, 16-16.	0.6	1
75	Induction of Genes Implicated in Stress Response and Autophagy by a Novel Quinolin-8-yl-nicotinamide QN523 in Pancreatic Cancer. <i>Journal of Medicinal Chemistry</i> , 2022, , .	2.9	1
76	Direct C-H functionalization of tetrahydro- β -carbolines at the β -position. <i>New Journal of Chemistry</i> , 2022, 46, 9511-9514.	1.4	1
77	Biochemical and Pharmacological Characterizations of ESI ⁻ based EPAC inhibitors. <i>FASEB Journal</i> , 2015, 29, 1022.4.	0.2	0