

Heidi M Kieler-Ferguson

List of Publications by Year in descending order

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Version: 2024-02-01

23
papers

563
citations

687220

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h-index

642610

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23
all docs

23
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23
times ranked

1124
citing authors

#	ARTICLE	IF	CITATIONS
1	STimulator of INterferon Genes Agonism Accelerates Antitumor Activity in Poorly Immunogenic Tumors. <i>Molecular Cancer Therapeutics</i> , 2022, 21, 282-293.	1.9	6
2	Oxetane Promise Delivered: Discovery of Long-Acting IDO1 Inhibitors Suitable for Q3W Oral or Parenteral Dosing. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 6001-6016.	2.9	8
3	Development of High-Throughput Assays for Evaluation of Hematopoietic Progenitor Kinase 1 Inhibitors. <i>SLAS Discovery</i> , 2021, 26, 88-99.	1.4	15
4	Combination of EP ⁴ antagonist MF-766 and anti-PD-1 promotes anti-tumor efficacy by modulating both lymphocytes and myeloid cells. <i>Onc Immunology</i> , 2021, 10, 1896643.	2.1	28
5	Carbamate and <i>N</i> -Pyrimidine Mitigate Amide Hydrolysis: Structure-Based Drug Design of Tetrahydroquinoline IDO1 Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 389-396.	1.3	14
6	Flexibility in Drug Product Development: A Perspective. <i>Molecular Pharmaceutics</i> , 2021, 18, 2455-2469.	2.3	4
7	Hierarchical Particle Approach for Co-Precipitated Amorphous Solid Dispersions for Use in Preclinical In Vivo Studies. <i>Pharmaceutics</i> , 2021, 13, 1034.	2.0	12
8	Sonoporation-Enhanced Delivery of STING Agonist Induced Robust Immune Modulation and Tumor Regression. <i>Advanced Therapeutics</i> , 2021, 4, 2100154.	1.6	3
9	SAR towards indoline and 3-aza-indoline classes of IDO1 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 47, 128214.	1.0	4
10	Discovery of IDO1 inhibitors containing a decahydroquinoline, decahydro-1,6-naphthyridine, or octahydro-1H-pyrrolo[3,2-c]pyridine scaffold. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 49, 128314.	1.0	7
11	Discovery of <i>N</i> -(Indazol-3-yl)piperidine-4-carboxylic Acids as ROR ^γ t Allosteric Inhibitors for Autoimmune Diseases. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 114-119.	1.3	18
12	Discovery of Potent and Orally Available Bicyclo[1.1.1]pentane-Derived Indoleamine-2,3-dioxygenase 1 (IDO1) Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1548-1554.	1.3	44
13	Discovery of Amino-cyclobutane-derived Indoleamine-2,3-dioxygenase 1 (IDO1) Inhibitors for Cancer Immunotherapy. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 1530-1536.	1.3	38
14	The NLRP3 inflammasome mediates DSS-induced intestinal inflammation in <i>Nod2</i> knockout mice. <i>Innate Immunity</i> , 2019, 25, 132-143.	1.1	27
15	Encapsulation, controlled release, and antitumor efficacy of cisplatin delivered in liposomes composed of sterol-modified phospholipids. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 103, 85-93.	1.9	33
16	Identification of quinazoline based inhibitors of IRAK4 for the treatment of inflammation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 2721-2726.	1.0	18
17	Inhibition of ROR ^γ T Skews TCR ^α Gene Rearrangement and Limits T Cell Repertoire Diversity. <i>Cell Reports</i> , 2016, 17, 3206-3218.	2.9	51
18	A robust and quantitative method for tracking liposome contents after intravenous administration. <i>Journal of Controlled Release</i> , 2014, 176, 86-93.	4.8	11

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19	Clinical developments of chemotherapeutic nanomedicines: polymers and liposomes for delivery of camptothecins and platinum (II) drugs. Wiley Interdisciplinary Reviews: Nanomedicine and Nanobiotechnology, 2013, 5, 130-138.	3.3	41
20	Resolution of enantiomers of a series of chiral alkoxy-modified phthalocyaninato nickel(ii) complexes by enantioselective HPLC. Dalton Transactions, 2011, 40, 11809.	1.6	5
21	Anti-tumor activity of liposome encapsulated fluoroorotic acid as a single agent and in combination with liposome irinotecan. Journal of Controlled Release, 2011, 153, 288-296.	4.8	50
22	Chemotherapeutic Evaluation of a Synthetic Tubulysin Analogueâ€“Dendrimer Conjugate in C26 Tumor Bearing Mice. ChemMedChem, 2011, 6, 49-53.	1.6	31
23	Design, Synthesis, and Biological Evaluation of a Robust, Biodegradable Dendrimer. Bioconjugate Chemistry, 2010, 21, 764-773.	1.8	95