

Neill J Liptrott

List of Publications by Year in descending order

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Version: 2024-02-01

47
papers

1,280
citations

361296

20
h-index

377752

34
g-index

49
all docs

49
docs citations

49
times ranked

2213
citing authors

#	ARTICLE	IF	CITATIONS
1	Firefly luciferase offers superior performance to AkaLuc for tracking the fate of administered cell therapies. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2022, 49, 796-808.	3.3	16
2	Exposure of human immune cells, to the antiretrovirals efavirenz and lopinavir, leads to lower glucose uptake and altered bioenergetic cell profiles through interactions with SLC2A1. <i>Biomedicine and Pharmacotherapy</i> , 2022, 150, 112999.	2.5	3
3	Physiologically based pharmacokinetic modeling of intravenously administered nanoformulated substances. <i>Drug Delivery and Translational Research</i> , 2022, 12, 2132-2144.	3.0	3
4	Linear and branched polymer prodrugs of the water-soluble nucleoside reverse-transcriptase inhibitor emtricitabine as structural materials for long-acting implants. <i>Journal of Materials Chemistry B</i> , 2022, 10, 4395-4404.	2.9	3
5	Dose prediction for repurposing nitazoxanide in SARS-CoV-2 treatment or chemoprophylaxis. <i>British Journal of Clinical Pharmacology</i> , 2021, 87, 2078-2088.	1.1	46
6	Induction of Cytokines by Nucleic Acid Nanoparticles (NANPs) Depends on the Type of Delivery Carrier. <i>Molecules</i> , 2021, 26, 652.	1.7	26
7	AUTOSTERE: Systematic Search for Scaffold Replacement Opportunities within Structural Databases. <i>Journal of Chemical Information and Modeling</i> , 2021, 61, 1778-1788.	2.5	0
8	Drug delivery systems as immunomodulators for therapy of infectious disease: Relevance to COVID-19. <i>Advanced Drug Delivery Reviews</i> , 2021, 178, 113848.	6.6	6
9	Assessment of changes in autophagic vesicles in human immune cell lines exposed to nano particles. <i>Cell and Bioscience</i> , 2021, 11, 133.	2.1	3
10	Scalable nanoprecipitation of niclosamide and <i>in vivo</i> demonstration of long-acting delivery after intramuscular injection. <i>Nanoscale</i> , 2021, 13, 6410-6416.	2.8	11
11	Critical considerations for targeting colorectal liver metastases with nanotechnology. <i>Wiley Interdisciplinary Reviews: Nanomedicine and Nanobiotechnology</i> , 2020, 12, e1588.	3.3	14
12	In Vitro Determination of the Immunogenic Impact of Nanomaterials on Primary Peripheral Blood Mononuclear Cells. <i>International Journal of Molecular Sciences</i> , 2020, 21, 5610.	1.8	7
13	Prioritization of Anti-SARS-CoV-2 Drug Repurposing Opportunities Based on Plasma and Target Site Concentrations Derived from their Established Human Pharmacokinetics. <i>Clinical Pharmacology and Therapeutics</i> , 2020, 108, 775-790.	2.3	118
14	Safety assessment of a new nanoemulsion-based drug-delivery system reveals unexpected, drug-free anticoagulant activity. <i>Nanomedicine</i> , 2020, 15, 1361-1373.	1.7	0
15	Immunotoxicity Considerations for Next Generation Cancer Nanomedicines. <i>Advanced Science</i> , 2019, 6, 1900133.	5.6	54
16	Long-Acting Injectable Statins "Is It Time for a Paradigm Shift?". <i>Molecules</i> , 2019, 24, 2685.	1.7	7
17	Bridging communities in the field of nanomedicine. <i>Regulatory Toxicology and Pharmacology</i> , 2019, 106, 187-196.	1.3	32
18	Overview of the blood compatibility of nanomedicines: A trend analysis of <i>in vitro</i> and <i>in vivo</i> studies. <i>Wiley Interdisciplinary Reviews: Nanomedicine and Nanobiotechnology</i> , 2019, 11, e1546.	3.3	29

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19	Derivation of CYP3A4 and CYP2B6 degradation rate constants in primary human hepatocytes: A siRNA-silencing-based approach. <i>Drug Metabolism and Pharmacokinetics</i> , 2018, 33, 179-187.	1.1	11
20	Advances in nanomedicine drug delivery applications for HIV therapy. <i>Future Science OA</i> , 2018, 4, FSO230.	0.9	15
21	Assessment of interactions of efavirenz solid drug nanoparticles with human immunological and haematological systems. <i>Journal of Nanobiotechnology</i> , 2018, 16, 22.	4.2	18
22	Sound understanding of environmental, health and safety, clinical, and market aspects is imperative to clinical translation of nanomedicines. <i>Nanotoxicology</i> , 2017, 11, 147-149.	1.6	29
23	Functionalized superparamagnetic iron oxide nanoparticles provide highly efficient iron-labeling in macrophages for magnetic resonance-based detection in vivo. <i>Cytotherapy</i> , 2017, 19, 555-569.	0.3	44
24	Incompatibility of chemical protein synthesis inhibitors with accurate measurement of extended protein degradation rates. <i>Pharmacology Research and Perspectives</i> , 2017, 5, e00359.	1.1	12
25	Lack of interaction of lopinavir solid drug nanoparticles with cells of the immune system. <i>Nanomedicine</i> , 2017, 12, 2043-2054.	1.7	5
26	In vitro characterisation of solid drug nanoparticle compositions of efavirenz in a brain endothelium cell line. <i>Journal of Interdisciplinary Nanomedicine</i> , 2017, 2, 157-169.	3.6	0
27	Efavirenz Is Predicted To Accumulate in Brain Tissue: an In Silico , In Vitro , and In Vivo Investigation. <i>Antimicrobial Agents and Chemotherapy</i> , 2017, 61, .	1.4	27
28	Determining the relationship between nanoparticle characteristics and immunotoxicity: key challenges and approaches. <i>Nanomedicine</i> , 2016, 11, 1447-1464.	1.7	28
29	Cytotoxic chemotherapy and the evolution of cellular and viral resistance to antiretroviral therapy in HIV- infected individuals with lymphoma. <i>HIV Clinical Trials</i> , 2016, 17, 197-203.	2.0	2
30	Towards a rational design of solid drug nanoparticles with optimised pharmacological properties. <i>Journal of Interdisciplinary Nanomedicine</i> , 2016, 1, 110-123.	3.6	17
31	Accelerated oral nanomedicine discovery from miniaturized screening to clinical production exemplified by paediatric HIV nanotherapies. <i>Nature Communications</i> , 2016, 7, 13184.	5.8	44
32	Toxicity and inflammatory response in Swiss albino mice after intraperitoneal and oral administration of polyurethane nanoparticles. <i>Toxicology Letters</i> , 2016, 246, 17-27.	0.4	16
33	Opportunities and Challenges in Nanotechnology-enabled Antiretroviral Delivery. <i>Frontiers in Nanobiomedical Research</i> , 2016, , 205-239.	0.1	0
34	The Application of Nanotechnology to Drug Delivery in Medicine. , 2015, , 173-223.		12
35	Interactions of antiretroviral drugs with the SLC22A1 (OCT1) drug transporter. <i>Frontiers in Pharmacology</i> , 2015, 6, 78.	1.6	19
36	Flow cytometric analysis of the physical and protein-binding characteristics of solid drug nanoparticle suspensions. <i>Nanomedicine</i> , 2015, 10, 1407-1421.	1.7	9

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37	Antiretroviral Solid Drug Nanoparticles with Enhanced Oral Bioavailability: Production, Characterization, and In Vitro–In Vivo Correlation. <i>Advanced Healthcare Materials</i> , 2014, 3, 400-411.	3.9	73
38	Partial mitigation of gold nanoparticle interactions with human lymphocytes by surface functionalization with a “mixed matrix”™. <i>Nanomedicine</i> , 2014, 9, 2467-2479.	1.7	16
39	High-throughput nanoprecipitation of the organic antimicrobial triclosan and enhancement of activity against <i>Escherichia coli</i> . <i>Journal of Materials Chemistry B</i> , 2013, 1, 4455.	2.9	15
40	Rilpivirine Inhibits Drug Transporters ABCB1, SLC22A1, and SLC22A2 <i>In Vitro</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2013, 57, 5612-5618.	1.4	26
41	Research Spotlight: Nanomedicines for HIV therapy. <i>Therapeutic Delivery</i> , 2013, 4, 153-156.	1.2	23
42	Association of ABCC10 polymorphisms with nevirapine plasma concentrations in the German Competence Network for HIV/AIDS. <i>Pharmacogenetics and Genomics</i> , 2012, 22, 10-19.	0.7	38
43	Plasma and Intracellular (Peripheral Blood Mononuclear Cells) Pharmacokinetics of Once-Daily Raltegravir (800 Milligrams) in HIV-Infected Patients. <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 72-75.	1.4	24
44	Genetic Variants of ABCC10, a Novel Tenofovir Transporter, Are Associated With Kidney Tubular Dysfunction. <i>Journal of Infectious Diseases</i> , 2011, 204, 145-153.	1.9	102
45	Plasma and Intracellular Pharmacokinetics of Darunavir/Ritonavir Once Daily and Raltegravir Once and Twice Daily in HIV-Infected Individuals. <i>Journal of Acquired Immune Deficiency Syndromes (1999)</i> , 2011, 58, 450-457.	0.9	31
46	Raltegravir Is a Substrate for SLC22A6: a Putative Mechanism for the Interaction between Raltegravir and Tenofovir. <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 879-887.	1.4	58
47	HIV protease inhibitors are substrates for OATP1A2, OATP1B1 and OATP1B3 and lopinavir plasma concentrations are influenced by SLCO1B1 polymorphisms. <i>Pharmacogenetics and Genomics</i> , 2010, 20, 112-120.	0.7	160