

# Adele M Lehane

## List of Publications by Year in descending order

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Version: 2024-02-01

38  
papers

2,032  
citations

279798

23  
h-index

330143

37  
g-index

41  
all docs

41  
docs citations

41  
times ranked

2898  
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery of spirooxadiazoline oxindoles with dual-stage antimalarial activity. <i>European Journal of Medicinal Chemistry</i> , 2022, 236, 114324.	5.5	9
2	Identifying the major lactate transporter of <i>Toxoplasma gondii</i> tachyzoites. <i>Scientific Reports</i> , 2021, 11, 6787.	3.3	10
3	An Open Drug Discovery Competition: Experimental Validation of Predictive Models in a Series of Novel Antimalarials. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 16450-16463.	6.4	8
4	A 4-cyano-3-methylisoquinoline inhibitor of <i>Plasmodium falciparum</i> growth targets the sodium efflux pump PfATP4. <i>Scientific Reports</i> , 2019, 9, 10292.	3.3	20
5	Characterization of the ATP4 ion pump in <i>Toxoplasma gondii</i> . <i>Journal of Biological Chemistry</i> , 2019, 294, 5720-5734.	3.4	18
6	Cell Swelling Induced by the Antimalarial KAE609 (Cipargamin) and Other PfATP4-Associated Antimalarials. <i>Antimicrobial Agents and Chemotherapy</i> , 2018, 62, .	3.2	33
7	Protein kinase A negatively regulates Ca <sup>2+</sup> signalling in <i>Toxoplasma gondii</i> . <i>PLoS Biology</i> , 2018, 16, e2005642.	5.6	65
8	Biochemical characterization and chemical inhibition of PfATP4-associated Na <sup>+</sup> -ATPase activity in <i>Plasmodium falciparum</i> membranes. <i>Journal of Biological Chemistry</i> , 2018, 293, 13327-13337.	3.4	32
9	Defense Peptides Engineered from Human Platelet Factor 4 Kill <i>Plasmodium</i> by Selective Membrane Disruption. <i>Cell Chemical Biology</i> , 2018, 25, 1140-1150.e5.	5.2	13
10	Diverse antimalarials from whole-cell phenotypic screens disrupt malaria parasite ion and volume homeostasis. <i>Scientific Reports</i> , 2018, 8, 8795.	3.3	36
11	A forward genetic screen identifies a negative regulator of rapid Ca <sup>2+</sup> -dependent cell egress (MS1) in the intracellular parasite <i>Toxoplasma gondii</i> . <i>Journal of Biological Chemistry</i> , 2017, 292, 7662-7674.	3.4	27
12	Biochemical and Structural Characterization of Selective Allosteric Inhibitors of the <i>Plasmodium falciparum</i> Drug Target, Prolyl-tRNA-synthetase. <i>ACS Infectious Diseases</i> , 2017, 3, 34-44.	3.8	45
13	The Malaria Parasite's Lactate Transporter PfFNT Is the Target of Antiplasmodial Compounds Identified in Whole Cell Phenotypic Screens. <i>PLoS Pathogens</i> , 2017, 13, e1006180.	4.7	37
14	Open Source Drug Discovery with the Malaria Box Compound Collection for Neglected Diseases and Beyond. <i>PLoS Pathogens</i> , 2016, 12, e1005763.	4.7	244
15	Globally prevalent PfMDR1 mutations modulate <i>Plasmodium falciparum</i> susceptibility to artemisinin-based combination therapies. <i>Nature Communications</i> , 2016, 7, 11553.	12.8	208
16	Verapamil-Sensitive Transport of Quinacrine and Methylene Blue via the <i>Plasmodium falciparum</i> Chloroquine Resistance Transporter Reduces the Parasite's Susceptibility to these Tricyclic Drugs. <i>Journal of Infectious Diseases</i> , 2016, 213, 800-810.	4.0	22
17	Molecular Mechanisms for Drug Hypersensitivity Induced by the Malaria Parasite's Chloroquine Resistance Transporter. <i>PLoS Pathogens</i> , 2016, 12, e1005725.	4.7	29
18	Balancing drug resistance and growth rates via compensatory mutations in the <i>Plasmodium falciparum</i> chloroquine resistance transporter. <i>Molecular Microbiology</i> , 2015, 97, 381-395.	2.5	47

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19	A lactate and formate transporter in the intraerythrocytic malaria parasite, <i>Plasmodium falciparum</i> . <i>Nature Communications</i> , 2015, 6, 6721.	12.8	56
20	<sup>1</sup> H-NMR metabolite profiles of different strains of <i>Plasmodium falciparum</i> . <i>Bioscience Reports</i> , 2014, 34, e00150.	2.4	22
21	(+)-SJ733, a clinical candidate for malaria that acts through ATP4 to induce rapid host-mediated clearance of <i>Plasmodium</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, E5455-62.	7.1	199
22	Diverse chemotypes disrupt ion homeostasis in the malaria parasite. <i>Molecular Microbiology</i> , 2014, 94, 327-339.	2.5	79
23	Membrane transport in the malaria parasite and its host erythrocyte. <i>Biochemical Journal</i> , 2014, 457, 1-18.	3.7	70
24	Quinine Dimers Are Potent Inhibitors of the <i>Plasmodium falciparum</i> Chloroquine Resistance Transporter and Are Active against Quinoline-Resistant <i>P. falciparum</i> . <i>ACS Chemical Biology</i> , 2014, 9, 722-730.	3.4	34
25	Chlorpheniramine Analogues Reverse Chloroquine Resistance in <i>Plasmodium falciparum</i> by Inhibiting PfCRT. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 576-581.	2.8	18
26	Degrees of chloroquine resistance in <i>Plasmodium</i> – Is the redox system involved?. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2012, 2, 47-57.	3.4	37
27	PfCRT and its role in antimalarial drug resistance. <i>Trends in Parasitology</i> , 2012, 28, 504-514.	3.3	223
28	Differential Drug Efflux or Accumulation Does Not Explain Variation in the Chloroquine Response of <i>Plasmodium falciparum</i> Strains Expressing the Same Isoform of Mutant PfCRT. <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 2310-2318.	3.2	14
29	Molecular Markers of <i>Plasmodium</i> Resistance to Antimalarials. , 2011, , 249-280.		5
30	Efflux of a range of antimalarial drugs and $\text{Ca}^{2+}$ -chloroquine resistance reversers <sup>TM</sup> from the digestive vacuole in malaria parasites with mutant PfCRT. <i>Molecular Microbiology</i> , 2010, 77, 1039-1051.	2.5	39
31	An Acid-loading Chloride Transport Pathway in the Intraerythrocytic Malaria Parasite, <i>Plasmodium falciparum</i> . <i>Journal of Biological Chemistry</i> , 2010, 285, 18615-18626.	3.4	8
32	A polymorphic drug pump in the malaria parasite. <i>Molecular Microbiology</i> , 2008, 70, 775-779.	2.5	6
33	Common dietary flavonoids inhibit the growth of the intraerythrocytic malaria parasite. <i>BMC Research Notes</i> , 2008, 1, 26.	1.4	122
34	Chloroquine Resistance-Confering Mutations in <i>pfprt</i> Give Rise to a Chloroquine-Associated H <sup>+</sup> Leak from the Malaria Parasite's Digestive Vacuole. <i>Antimicrobial Agents and Chemotherapy</i> , 2008, 52, 4374-4380.	3.2	46
35	A verapamil-sensitive chloroquine-associated H <sup>+</sup> leak from the digestive vacuole in chloroquine-resistant malaria parasites. <i>Journal of Cell Science</i> , 2008, 121, 1624-1632.	2.0	51
36	Feedback Inhibition of Pantothenate Kinase Regulates Pantothenol Uptake by the Malaria Parasite. <i>Journal of Biological Chemistry</i> , 2007, 282, 25395-25405.	3.4	19

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37	Bacteriophage-encoded glucosyltransferase GtrII of <i>Shigella flexneri</i> : membrane topology and identification of critical residues. <i>Biochemical Journal</i> , 2005, 389, 137-143.	3.7	27
38	Choline uptake into the malaria parasite is energized by the membrane potential. <i>Biochemical and Biophysical Research Communications</i> , 2004, 320, 311-317.	2.1	50