## Ryan Choi

## List of Publications by Year in descending order

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Ργλη Cho

#	Article	IF	CITATIONS
1	Repurposing the Kinase Inhibitor Mavelertinib for Giardiasis Therapy. Antimicrobial Agents and Chemotherapy, 2022, 66, .	3.2	3
2	One health therapeutics: Target-Based drug development for cryptosporidiosis and other apicomplexa diseases. Veterinary Parasitology, 2021, 289, 109336.	1.8	16
3	Pyrrolopyrimidine Bumped Kinase Inhibitors for the Treatment of Cryptosporidiosis. ACS Infectious Diseases, 2021, 7, 1200-1207.	3.8	3
4	Repurposing Infectious Disease Hits as Anti- <i>Cryptosporidium</i> Leads. ACS Infectious Diseases, 2021, 7, 1275-1282.	3.8	8
5	High-throughput screening of the ReFRAME, Pandemic Box, and COVID Box drug repurposing libraries against SARS-CoV-2 nsp15 endoribonuclease to identify small-molecule inhibitors of viral activity. PLoS ONE, 2021, 16, e0250019.	2.5	27
6	In vitro activity, safety and in vivo efficacy of the novel bumped kinase inhibitor BKI-1748 in non-pregnant and pregnant mice experimentally infected with Neospora caninum tachyzoites and Toxoplasma gondii oocysts. International Journal for Parasitology: Drugs and Drug Resistance, 2021, 16, 90-101.	3.4	17
7	Endochin-like quinolones (ELQs) and bumped kinase inhibitors (BKIs): Synergistic and additive effects of combined treatments against Neospora caninum infection in vitro and in vivo. International Journal for Parasitology: Drugs and Drug Resistance, 2021, 17, 92-106.	3.4	7
8	A short-term treatment with BKI-1294 does not protect foetuses from sheep experimentally infected with Neospora caninum tachyzoites during pregnancy. International Journal for Parasitology: Drugs and Drug Resistance, 2021, 17, 176-185.	3.4	5
9	Reduced treatment frequencies with bumped kinase inhibitor 1369 are effective against porcine cystoisosporosis. International Journal for Parasitology: Drugs and Drug Resistance, 2020, 14, 37-45.	3.4	3
10	Comparative assessment of the effects of bumped kinase inhibitors on early zebrafish embryo development and pregnancy in mice. International Journal of Antimicrobial Agents, 2020, 56, 106099.	2.5	12
11	Taming the Boys for Global Good: Contraceptive Strategy to Stop Malaria Transmission. Molecules, 2020, 25, 2773.	3.8	6
12	Structures of glyceraldehyde 3â€phosphate dehydrogenase in <scp><i>Neisseria gonorrhoeae</i></scp> and <i>Chlamydia trachomatis</i> . Protein Science, 2020, 29, 768-778.	7.6	10
13	Bumped Kinase Inhibitors as therapy for apicomplexan parasitic diseases: lessons learned. International Journal for Parasitology, 2020, 50, 413-422.	3.1	37
14	Evaluation of in vitro and in vivo antibiotic efficacy against a novel bioluminescent Shigella flexneri. Scientific Reports, 2019, 9, 13567.	3.3	8
15	P-Glycoprotein–Mediated Efflux Reduces the In Vivo Efficacy of a Therapeutic Targeting the Gastrointestinal Parasite Cryptosporidium. Journal of Infectious Diseases, 2019, 220, 1188-1198.	4.0	7
16	Treatment with Bumped Kinase Inhibitor 1294 Is Safe and Leads to Significant Protection against Abortion and Vertical Transmission in Sheep Experimentally Infected with Toxoplasma gondii during Pregnancy. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	23
17	Development of 5-Aminopyrazole-4-carboxamide-based Bumped-Kinase Inhibitors for Cryptosporidiosis Therapy. Journal of Medicinal Chemistry, 2019, 62, 3135-3146.	6.4	27
18	Bumped kinase inhibitor 1369 is effective against Cystoisospora suis in vivo and in vitro. International Journal for Parasitology: Drugs and Drug Resistance, 2019, 10, 9-19.	3.4	12

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19	Optimization of Methionyl tRNA-Synthetase Inhibitors for Treatment of <i>Cryptosporidium</i> Infection. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	37
20	Pharmacokinetics and In Vivo Efficacy of Pyrazolopyrimidine, Pyrrolopyrimidine, and 5-Aminopyrazole-4-Carboxamide Bumped Kinase Inhibitors against Toxoplasmosis. Journal of Infectious Diseases, 2019, 219, 1464-1473.	4.0	13
21	Safety and efficacy of the bumped kinase inhibitor BKI-1553 in pregnant sheep experimentally infected with Neospora caninum tachyzoites. International Journal for Parasitology: Drugs and Drug Resistance, 2018, 8, 112-124.	3.4	28
22	Therapeutic Efficacy of Bumped Kinase Inhibitor 1369 in a Pig Model of Acute Diarrhea Caused by Cryptosporidium hominis. Antimicrobial Agents and Chemotherapy, 2018, 62, .	3.2	31
23	Toxoplasma Calcium-Dependent Protein Kinase 1 Inhibitors: Probing Activity and Resistance Using Cellular Thermal Shift Assays. Antimicrobial Agents and Chemotherapy, 2018, 62, .	3.2	12
24	7H-Pyrrolo[2,3-d]pyrimidin-4-amine-Based Inhibitors of Calcium-Dependent Protein Kinase 1 Have Distinct Inhibitory and Oral Pharmacokinetic Characteristics Compared with 1H-Pyrazolo[3,4-d]pyrimidin-4-amine-Based Inhibitors. ACS Infectious Diseases, 2018, 4, 516-522.	3.8	5
25	Screening of the Pathogen Box for inhibitors with dual efficacy against Giardia lamblia and Cryptosporidium parvum. PLoS Neglected Tropical Diseases, 2018, 12, e0006673.	3.0	37
26	Extended-spectrum antiprotozoal bumped kinase inhibitors: A review. Experimental Parasitology, 2017, 180, 71-83.	1.2	71
27	Development of a murine vertical transmission model for Toxoplasma gondii oocyst infection and studies on the efficacy of bumped kinase inhibitor (BKI)-1294 and the naphthoquinone buparvaquone against congenital toxoplasmosis. Journal of Antimicrobial Chemotherapy, 2017, 72, 2334-2341.	3.0	52
28	5-Aminopyrazole-4-Carboxamide-Based Compounds Prevent the Growth of Cryptosporidium parvum. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	17
29	Bumped-Kinase Inhibitors for Cryptosporidiosis Therapy. Journal of Infectious Diseases, 2017, 215, 1275-1284.	4.0	52
30	Advances in bumped kinase inhibitors for human and animal therapy for cryptosporidiosis. International Journal for Parasitology, 2017, 47, 753-763.	3.1	30
31	Necessity of Bumped Kinase Inhibitor Gastrointestinal Exposure in Treating Cryptosporidium Infection. Journal of Infectious Diseases, 2017, 216, 55-63.	4.0	44
32	Biochemical and Structural Characterization of Selective Allosteric Inhibitors of the <i>Plasmodium falciparum</i> Drug Target, Prolyl-tRNA-synthetase. ACS Infectious Diseases, 2017, 3, 34-44.	3.8	45
33	Selective inhibition of Sarcocystis neurona calcium-dependent protein kinase 1 for equine protozoal myeloencephalitis therapy. International Journal for Parasitology, 2016, 46, 871-880.	3.1	22
34	5-Aminopyrazole-4-carboxamide analogues are selective inhibitors of Plasmodium falciparum microgametocyte exflagellation and potential malaria transmission blocking agents. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 5487-5491.	2.2	15
35	Development of an Orally Available and Central Nervous System (CNS) Penetrant <i>Toxoplasma gondii</i> Calcium-Dependent Protein Kinase 1 ( <i>Tg</i> CDPK1) Inhibitor with Minimal Human Ether-a-go-go-Related Gene (hERC) Activity for the Treatment of <i>Toxoplasmosis</i> . Journal of Medicinal Chemistry, 2016, 59, 6531-6546.	6.4	81
36	Bumped kinase inhibitor prohibits egression in Babesia bovis. Veterinary Parasitology, 2016, 215, 22-28.	1.8	19

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37	Brucella melitensis Methionyl-tRNA-Synthetase (MetRS), a Potential Drug Target for Brucellosis. PLoS ONE, 2016, 11, e0160350.	2.5	21
38	Increasing the structural coverage of tuberculosis drug targets. Tuberculosis, 2015, 95, 142-148.	1.9	103
39	SAR Studies of 5-Aminopyrazole-4-carboxamide Analogues as Potent and Selective Inhibitors of <i>Toxoplasma gondii</i> CDPK1. ACS Medicinal Chemistry Letters, 2015, 6, 1184-1189.	2.8	32
40	The gatekeeper residue and beyond: homologous calcium-dependent protein kinases as drug development targets for veterinarian Apicomplexa parasites. Parasitology, 2014, 141, 1499-1509.	1.5	47
41	Potent and Selective Inhibitors of CDPK1 from <i>T. gondii</i> and <i>C. parvum</i> Based on a 5-Aminopyrazole-4-carboxamide Scaffold. ACS Medicinal Chemistry Letters, 2014, 5, 40-44.	2.8	49
42	A Specific Inhibitor of PfCDPK4 Blocks Malaria Transmission: Chemical-genetic Validation. Journal of Infectious Diseases, 2014, 209, 275-284.	4.0	83
43	Development of potent and selective Plasmodium falciparum calcium-dependent protein kinase 4 (PfCDPK4) inhibitors that block the transmission of malaria to mosquitoes. European Journal of Medicinal Chemistry, 2014, 74, 562-573.	5.5	54
44	Immobilized metal-affinity chromatography protein-recovery screening is predictive of crystallographic structure success. Acta Crystallographica Section F: Structural Biology Communications, 2011, 67, 998-1005.	0.7	79