

Santiago Castanys

List of Publications by Year in descending order

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103
papers

4,394
citations

94433

37
h-index

118850

62
g-index

104
all docs

104
docs citations

104
times ranked

3462
citing authors

#	ARTICLE	IF	CITATIONS
1	Genomic and transcriptomic alterations in <i>Leishmania donovani</i> lines experimentally resistant to antileishmanial drugs. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2018, 8, 246-264.	3.4	49
2	Functional role of highly conserved residues of the N-terminal tail and first transmembrane segment of a P4-ATPase. <i>Biochemical Journal</i> , 2018, 475, 887-899.	3.7	1
3	<i>Leishmania</i> LABC2 transporter is involved in ATP-dependent transport of thiols. <i>Biochemical Journal</i> , 2018, 475, 87-97.	3.7	4
4	<i>Leishmania</i> LABC1 and LABC2 transporters are involved in virulence and oxidative stress: functional linkage with autophagy. <i>Parasites and Vectors</i> , 2017, 10, 267.	2.5	16
5	Symmetrical Pyridinium-Phanes and "Diazacyclophanes" Promising Heterocyclic Scaffolds for the Development of Anti-Leishmanial Agents. , 2016, , .		0
6	The LABC2 Transporter from the Protozoan Parasite <i>Leishmania</i> Is Involved in Antimony Resistance. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 3489-3496.	3.2	34
7	Decreased antimony uptake and overexpression of genes of thiol metabolism are associated with drug resistance in a canine isolate of <i>Leishmania infantum</i> . <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2016, 6, 133-139.	3.4	24
8	Optimization by Molecular Fine Tuning of Dihydro- $\hat{1}^2$ -agarofuran Sesquiterpenoids as Reversers of P-Glycoprotein-Mediated Multidrug Resistance. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1880-1890.	6.4	21
9	Genomic and Molecular Characterization of Miltefosine Resistance in <i>Leishmania infantum</i> Strains with Either Natural or Acquired Resistance through Experimental Selection of Intracellular Amastigotes. <i>PLoS ONE</i> , 2016, 11, e0154101.	2.5	80
10	Restoration of Chemosensitivity in P-Glycoprotein-Dependent Multidrug-Resistant Cells by Dihydro- $\hat{1}^2$ -agarofuran Sesquiterpenes from <i>Celastrus vulcanicola</i> . <i>Journal of Natural Products</i> , 2015, 78, 736-745.	3.0	20
11	Antileishmanial activity of sp ² -iminosugar derivatives. <i>RSC Advances</i> , 2015, 5, 21812-21822.	3.6	27
12	Experimental Resistance to Drug Combinations in <i>Leishmania donovani</i> : Metabolic and Phenotypic Adaptations. <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 2242-2255.	3.2	47
13	Fitness of <i>Leishmania donovani</i> Parasites Resistant to Drug Combinations. <i>PLoS Neglected Tropical Diseases</i> , 2015, 9, e0003704.	3.0	28
14	Mechanisms of Action of Substituted $\hat{1}^2$ -Amino Alkanols on <i>Leishmania donovani</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 1211-1218.	3.2	4
15	High-Throughput Screening Platform for Natural Product-Based Drug Discovery Against 3 Neglected Tropical Diseases: Human African Trypanosomiasis, Leishmaniasis, and Chagas Disease. <i>Journal of Biomolecular Screening</i> , 2015, 20, 82-91.	2.6	70
16	Design, synthesis and anti-leishmanial activity of novel symmetrical bispyridinium cyclophanes. <i>European Journal of Medicinal Chemistry</i> , 2015, 89, 362-369.	5.5	9
17	Functional role of evolutionarily highly conserved residues, N-glycosylation level and domains of the <i>Leishmania</i> miltefosine transporter-Cdc50 subunit. <i>Biochemical Journal</i> , 2014, 459, 83-94.	3.7	17
18	Identification of specific reversal agents for <i>Leishmania</i> ABC14-mediated antimony resistance by flavonoid and trolox derivative screening. <i>Journal of Antimicrobial Chemotherapy</i> , 2014, 69, 664-672.	3.0	13

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19	4-Amino Bis-Pyridinium Derivatives as Novel Antileishmanial Agents. <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 4103-4112.	3.2	27
20	Mechanisms of Miltefosine Resistance in <i>Leishmania</i> . , 2013, , 351-379.		3
21	A New ABC Half-Transporter in <i>Leishmania major</i> Is Involved in Resistance to Antimony. <i>Antimicrobial Agents and Chemotherapy</i> , 2013, 57, 3719-3730.	3.2	56
22	LABCG2, a New ABC Transporter Implicated in Phosphatidylserine Exposure, Is Involved in the Infectivity and Pathogenicity of <i>Leishmania</i> . <i>PLoS Neglected Tropical Diseases</i> , 2013, 7, e2179.	3.0	23
23	<i>Leishmania donovani</i> Develops Resistance to Drug Combinations. <i>PLoS Neglected Tropical Diseases</i> , 2012, 6, e1974.	3.0	93
24	Oxazolo[3,2-a]pyridine. A new structural scaffold for the reversal of multi-drug resistance in <i>Leishmania</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 6272-6275.	2.2	18
25	Leishmanicidal and Reversal Multidrug Resistance Constituents from <i>Aeonium lindleyi</i> . <i>Planta Medica</i> , 2011, 77, 77-80.	1.3	18
26	Non-reducing trisaccharide fatty acid monoesters: Novel detergents in membrane biochemistry. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2011, 1808, 717-726.	2.6	19
27	A new ATP-binding cassette protein is involved in intracellular haem trafficking in <i>Leishmania</i> . <i>Molecular Microbiology</i> , 2011, 79, 1430-1444.	2.5	36
28	Overcoming human P-glycoprotein-dependent multidrug resistance with novel dihydro- β -agarofuran sesquiterpenes. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 4915-4923.	5.5	19
29	Terpenoids from <i>Maytenus</i> Species and Assessment of Their Reversal Activity against a Multidrug-Resistant <i>Leishmania tropica</i> Line. <i>Chemistry and Biodiversity</i> , 2011, 8, 2291-2298.	2.1	12
30	The 8-Aminoquinoline Analogue Sitamaquine Causes Oxidative Stress in <i>Leishmania donovani</i> Promastigotes by Targeting Succinate Dehydrogenase. <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 4204-4210.	3.2	65
31	Sitamaquine Overcomes ABC-Mediated Resistance to Miltefosine and Antimony in <i>Leishmania</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 3838-3844.	3.2	41
32	Uptake of the antileishmania drug tafenoquine follows a sterol-dependent diffusion process in <i>Leishmania</i> . <i>Journal of Antimicrobial Chemotherapy</i> , 2011, 66, 2562-2565.	3.0	9
33	Increased Glycolytic ATP Synthesis Is Associated with Tafenoquine Resistance in <i>Leishmania major</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2011, 55, 1045-1052.	3.2	28
34	CDC50A plays a key role in the uptake of the anticancer drug perifosine in human carcinoma cells. <i>Biochemical Pharmacology</i> , 2010, 80, 793-800.	4.4	24
35	Disruption of the Lipid-Transporting LdMT-LdRos3 Complex in <i>Leishmania donovani</i> Affects Membrane Lipid Asymmetry but Not Host Cell Invasion. <i>PLoS ONE</i> , 2010, 5, e12443.	2.5	32
36	Tafenoquine, an Antiplasmodial 8-Aminoquinoline, Targets <i>Leishmania</i> Respiratory Complex III and Induces Apoptosis. <i>Antimicrobial Agents and Chemotherapy</i> , 2010, 54, 5344-5351.	3.2	82

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37	Low Plasma Membrane Expression of the Miltefosine Transport Complex Renders <i>Leishmania braziliensis</i> Refractory to the Drug. <i>Antimicrobial Agents and Chemotherapy</i> , 2009, 53, 1305-1313.	3.2	73
38	Novel dihydro- δ^2 -agarofuran sesquiterpenes as potent modulators of human P-glycoprotein dependent multidrug resistance. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 5166.	2.8	30
39	New terpenoids from <i>Maytenus apurimacensis</i> as MDR reversal agents in the parasite <i>Leishmania</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2008, 16, 1425-1430.	3.0	28
40	The anti-tumor alkylphospholipid perifosine is internalized by an ATP-dependent translocase activity across the plasma membrane of human KB carcinoma cells. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2008, 1778, 530-540.	2.6	25
41	Bis-pyranobenzoquinones as a New Family of Reversal Agents of the Multidrug Resistance Phenotype Mediated by P-Glycoprotein in Mammalian Cells and the Protozoan Parasite <i>Leishmania</i> . <i>Journal of Medicinal Chemistry</i> , 2008, 51, 7132-7143.	6.4	33
42	Characterization of an ABCG-Like Transporter from the Protozoan Parasite <i>Leishmania</i> with a Role in Drug Resistance and Transbilayer Lipid Movement. <i>Antimicrobial Agents and Chemotherapy</i> , 2008, 52, 3573-3579.	3.2	96
43	Sitamaquine Sensitivity in <i>Leishmania</i> Species Is Not Mediated by Drug Accumulation in Acidocalcisomes. <i>Antimicrobial Agents and Chemotherapy</i> , 2008, 52, 4030-4036.	3.2	33
44	Biological Evaluation, Structure-Activity Relationships, and Three-Dimensional Quantitative Structure-Activity Relationship Studies of Dihydro- δ^2 -agarofuran Sesquiterpenes as Modulators of P-Glycoprotein-Dependent Multidrug Resistance. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 4808-4817.	6.4	39
45	Inactivation of the miltefosine transporter, LdMT, causes miltefosine resistance that is conferred to the amastigote stage of <i>Leishmania donovani</i> and persists in vivo. <i>International Journal of Antimicrobial Agents</i> , 2007, 30, 229-235.	2.5	118
46	A novel ATP-binding cassette transporter from <i>Leishmania</i> is involved in transport of phosphatidylcholine analogues and resistance to alkyl-phospholipids. <i>Molecular Microbiology</i> , 2007, 64, 1141-1153.	2.5	78
47	Insights into the molecular mechanism of action of Celastraceae sesquiterpenes as specific, non-transported inhibitors of human P-glycoprotein. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2006, 1758, 98-110.	2.6	12
48	Mechanisms of experimental resistance of <i>Leishmania</i> to miltefosine: Implications for clinical use. <i>Drug Resistance Updates</i> , 2006, 9, 26-39.	14.4	172
49	Combination of Suboptimal Doses of Inhibitors Targeting Different Domains of LtrMDR1 Efficiently Overcomes Resistance of <i>Leishmania</i> spp. to Miltefosine by Inhibiting Drug Efflux. <i>Antimicrobial Agents and Chemotherapy</i> , 2006, 50, 3102-3110.	3.2	45
50	Phospholipid Translocation and Miltefosine Potency Require Both <i>L. donovani</i> Miltefosine Transporter and the New Protein LdRos3 in <i>Leishmania</i> Parasites. <i>Journal of Biological Chemistry</i> , 2006, 281, 23766-23775.	3.4	149
51	Dihydro- δ^2 -Agarofuran Sesquiterpenes: A New Class of Reversal Agents of the Multidrug Resistance Phenotype Mediated by P-Glycoprotein in the Protozoan Parasite <i>Leishmania</i> . <i>Current Pharmaceutical Design</i> , 2005, 11, 3125-3139.	1.9	46
52	Flavonoid Structure-Activity Studies Identify 6-Prenylchrysin and Tectochrysin as Potent and Specific Inhibitors of Breast Cancer Resistance Protein ABCG2. <i>Cancer Research</i> , 2005, 65, 4852-4860.	0.9	181
53	Reversion of Human Pgp-Dependent Multidrug Resistance by New Sesquiterpenes from <i>Zinowewiacostaricensis</i> . <i>Journal of Medicinal Chemistry</i> , 2005, 48, 4266-4275.	6.4	31
54	Functional expression of a multidrug P-glycoprotein transporter of <i>Leishmania</i> . <i>Biochemical and Biophysical Research Communications</i> , 2005, 329, 502-507.	2.1	20

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55	The overexpression of an intracellular ABCA-like transporter alters phospholipid trafficking in Leishmania. <i>Biochemical and Biophysical Research Communications</i> , 2005, 330, 349-355.	2.1	36
56	Characterization of an ABCA-like transporter involved in vesicular trafficking in the protozoan parasite <i>Trypanosoma cruzi</i> . <i>Molecular Microbiology</i> , 2004, 54, 632-646.	2.5	32
57	SAR Studies of Dihydro- $\hat{1}^2$ -agarofuran Sesquiterpenes as Inhibitors of the Multidrug-Resistance Phenotype in a <i>Leishmania tropicalis</i> Line Overexpressing a P-Glycoprotein-Like Transporter. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 576-587.	6.4	43
58	Celastraceae Sesquiterpenes as a New Class of Modulators That Bind Specifically to Human P-Glycoprotein and Reverse Cellular Multidrug Resistance. <i>Cancer Research</i> , 2004, 64, 7130-7138.	0.9	72
59	RU49953: a non-hormonal steroid derivative that potently inhibits P-glycoprotein and reverts cellular multidrug resistance. <i>Cellular and Molecular Life Sciences</i> , 2003, 60, 526-535.	5.4	18
60	Characterisation of <i>Leishmania donovani</i> promastigotes resistant to hexadecylphosphocholine (miltefosine). <i>International Journal of Antimicrobial Agents</i> , 2003, 22, 380-387.	2.5	157
61	The overexpression of a new ABC transporter in <i>Leishmania</i> is related to phospholipid trafficking and reduced infectivity. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2003, 1612, 195-207.	2.6	49
62	Rapid transport of phospholipids across the plasma membrane of <i>Leishmania infantum</i> . <i>Biochemical and Biophysical Research Communications</i> , 2003, 306, 250-255.	2.1	21
63	<i>Leishmania donovani</i> Resistance to Miltefosine Involves a Defective Inward Translocation of the Drug. <i>Antimicrobial Agents and Chemotherapy</i> , 2003, 47, 2397-2403.	3.2	165
64	Functional Cloning of the Miltefosine Transporter. <i>Journal of Biological Chemistry</i> , 2003, 278, 49965-49971.	3.4	189
65	Multidrug Resistance Phenotype Mediated by the P-Glycoprotein-Like Transporter in <i>Leishmania</i> : A Search for Reversal Agents. <i>Current Drug Targets</i> , 2002, 3, 311-333.	2.1	66
66	Two casein kinase 1 isoforms are differentially expressed in <i>Trypanosoma cruzi</i> . <i>Molecular and Biochemical Parasitology</i> , 2002, 124, 23-36.	1.1	16
67	Modulation by flavonoids of cell multidrug resistance mediated by P-glycoprotein and related ABC transporters. <i>Cellular and Molecular Life Sciences</i> , 2002, 59, 307-322.	5.4	230
68	ABC transporters in the protozoan parasite <i>Leishmania</i> . <i>International Microbiology</i> , 2001, 4, 159-166.	2.4	39
69	Chemosensitization of a Multidrug-Resistant <i>Leishmania tropicalis</i> Line by New Sesquiterpenes from <i>Maytenus magellanica</i> and <i>Maytenus chubutensis</i> . <i>Journal of Medicinal Chemistry</i> , 2001, 44, 4668-4676.	6.4	79
70	Alkyl-Lysophospholipid Resistance in Multidrug-Resistant <i>Leishmania tropicalis</i> and Chemosensitization by a Novel P-Glycoprotein-Like Transporter Modulator. <i>Antimicrobial Agents and Chemotherapy</i> , 2001, 45, 2468-2474.	3.2	120
71	High-Affinity Binding of Silybin Derivatives to the Nucleotide-Binding Domain of a <i>Leishmania tropicalis</i> P-Glycoprotein-Like Transporter and Chemosensitization of a Multidrug-Resistant Parasite to Daunomycin. <i>Antimicrobial Agents and Chemotherapy</i> , 2001, 45, 439-446.	3.2	46
72	<i>Trypanosoma cruzi</i> : Molecular Cloning of a Gene Coding for a Putative Vacuolar Protein. <i>Experimental Parasitology</i> , 2000, 94, 129-131.	1.2	1

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73	Evolutionary relationships in <i>Trypanosoma cruzi</i> : molecular phylogenetics supports the existence of a new major lineage of strains. <i>Gene</i> , 2000, 246, 331-338.	2.2	37
74	Characterization of a new ATP-binding cassette transporter in <i>Trypanosoma cruzi</i> associated to a L1Tc retrotransposon. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1999, 1489, 428-432.	2.4	21
75	Different catalytic activities of hexokinase and phosphofructokinase in wild type and glucantime-resistant <i>Leishmania promastigotes</i> appears not causatively related to resistance. <i>European Journal of Protistology</i> , 1999, 35, 338-341.	1.5	1
76	New Natural Sesquiterpenes as Modulators of Daunomycin Resistance in a Multidrug-Resistant <i>Leishmania tropica</i> Line. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 4388-4393.	6.4	63
77	Correlation between the Affinity of Flavonoids Binding to the Cytosolic Site of <i>Leishmania tropica</i> Multidrug Transporter and Their Efficiency To Revert Parasite Resistance to Daunomycin. <i>Biochemistry</i> , 1999, 38, 1736-1743.	2.5	77
78	Involvement of thiol metabolism in resistance to glucantime in <i>leishmania tropica</i> . <i>Biochemical Pharmacology</i> , 1998, 56, 1201-1208.	4.4	36
79	A new member of YER057c family in <i>Trypanosoma cruzi</i> is adjacent to an ABC-transporter. <i>Gene</i> , 1998, 220, 1-12.	2.2	12
80	Altered Drug Membrane Permeability in a Multidrug-Resistant <i>Leishmania tropica</i> Line. <i>Biochemical Pharmacology</i> , 1998, 55, 131-139.	4.4	74
81	A pteridine reductase gene <i>ptr1</i> contiguous to a P-glycoprotein confers resistance to antifolates in <i>Trypanosoma cruzi</i> Note: nucleotide sequence data reported in this paper have been submitted to the GenBank™ data base with the accession number U61221.1. <i>Molecular and Biochemical Parasitology</i> , 1997, 90, 525-535.	1.1	55
82	Molecular characterization of a P-glycoprotein-related <i>tcpgp2</i> gene in <i>Trypanosoma cruzi</i> . <i>Molecular and Biochemical Parasitology</i> , 1996, 75, 145-157.	1.1	42
83	A protein secreted by <i>Trypanosoma cruzi</i> capable of inducing the entry of inert particles into HeLa cells. <i>International Journal for Parasitology</i> , 1995, 25, 1213-1225.	3.1	6
84	Increased P-Type ATPase Activity in <i>Leishmania tropica</i> Resistant to Methotrexate. <i>Biochemical and Biophysical Research Communications</i> , 1994, 199, 855-861.	2.1	11
85	<i>Trypanosoma cruzi</i> : Sequence of the ATP-Binding Site of a P-Glycoprotein Gene. <i>Experimental Parasitology</i> , 1994, 79, 63-67.	1.2	16
86	P-Glycoprotein overexpression in methotrexate-resistant <i>Leishmania tropica</i> . <i>Biochemical Pharmacology</i> , 1994, 47, 1939-1947.	4.4	57
87	Amplification of the H locus in <i>Leishmania infantum</i> . <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 1994, 1227, 188-194.	3.8	13
88	Possible coexistence of two independent mechanisms contributing to anthracycline resistance in leukaemia P388 cells. <i>European Journal of Cancer</i> , 1993, 29, 2144-2150.	2.8	15
89	Pyridinium azolate betaines and their derivatives: a new class of antiprotozoal agents. <i>European Journal of Medicinal Chemistry</i> , 1990, 25, 309-319.	5.5	20
90	<i>Trypanosoma cruzi</i> : Calcium ion movement during internalization in host HeLa cells. <i>International Journal for Parasitology</i> , 1990, 20, 673-676.	3.1	21

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91	Purification of a glycoprotein excreted by Trypanosoma cruzi to increase the permeability of the host-cell membrane. Biochemical and Biophysical Research Communications, 1990, 166, 736-742.	2.1	5
92	Inhibitory Effect of New Pyrimidine Bases on Trypanosoma Cruzi. Archiv Der Pharmazie, 1989, 322, 843-846.	4.1	3
93	New Antiparasitic Agents. Chemotherapy, 1988, 34, 127-133.	1.6	9
94	Antitrypanosomal Action of Cis-Diamminedichloroplatinum (II) Analogs. Journal of Parasitology, 1987, 73, 272.	0.7	10
95	Antiamebic Activity of New Acridinic Derivatives against Naegleria and Acanthamoeba Species in vitro. Chemotherapy, 1987, 33, 18-21.	1.6	4
96	Activity against Trypanosoma cruzi of New Analogues of Nifurtimox. Archiv Der Pharmazie, 1987, 320, 115-120.	4.1	27
97	Inhibition of lysosomal fusion by Trypanosoma cruzi in peritoneal macrophages. International Journal for Parasitology, 1986, 16, 629-632.	3.1	21
98	Effect of poly-L-lysine and neuraminidase on the infectivity of Trypanosoma cruzi in cultured HeLa cells. Zeitschrift für Parasitenkunde (Berlin, Germany), 1985, 71, 429-433.	0.8	2
99	Isolation and purification of amastigotes of Trypanosoma cruzi from cultured Vero cells. Zeitschrift für Parasitenkunde (Berlin, Germany), 1985, 71, 15-17.	0.8	8
100	Effect of interferon on the infectivity of Trypanosoma cruzi in cultured HeLa cells. International Journal for Parasitology, 1985, 15, 167-170.	3.1	4
101	Purification of metacyclic forms of Trypanosoma cruzi by Percoll discontinuous gradient centrifugation. Zeitschrift für Parasitenkunde (Berlin, Germany), 1984, 70, 443-449.	0.8	30
102	Some factors affecting the in vitro invasion of HeLa cells by Trypanosoma cruzi. International Journal for Parasitology, 1984, 14, 253-257.	3.1	30
103	Resistance to Reinfection of HeLa Cells Parasitized by Trypanosoma cruzi. Journal of Parasitology, 1984, 70, 825.	0.7	4