

Clemente Capasso

List of Publications by Year in descending order

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1,384
papers

87,075
citations

460

134
h-index

1764

218
g-index

1407
all docs

1407
docs citations

1407
times ranked

31995
citing authors

#	ARTICLE	IF	CITATIONS
1	Carbonic anhydrases: novel therapeutic applications for inhibitors and activators. <i>Nature Reviews Drug Discovery</i> , 2008, 7, 168-181.	21.5	2,702
2	Natural products in drug discovery: advances and opportunities. <i>Nature Reviews Drug Discovery</i> , 2021, 20, 200-216.	21.5	1,990
3	Interfering with pH regulation in tumours as a therapeutic strategy. <i>Nature Reviews Drug Discovery</i> , 2011, 10, 767-777.	21.5	1,340
4	Carbonic anhydrase inhibitors. <i>Medicinal Research Reviews</i> , 2003, 23, 146-189.	5.0	1,126
5	Multiple Binding Modes of Inhibitors to Carbonic Anhydrases: How to Design Specific Drugs Targeting 15 Different Isoforms?. <i>Chemical Reviews</i> , 2012, 112, 4421-4468.	23.0	1,056
6	Carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 1999, 7, 2397-2406.	1.4	808
7	Structure and function of carbonic anhydrases. <i>Biochemical Journal</i> , 2016, 473, 2023-2032.	1.7	688
8	Targeting Tumor Hypoxia: Suppression of Breast Tumor Growth and Metastasis by Novel Carbonic Anhydrase IX Inhibitors. <i>Cancer Research</i> , 2011, 71, 3364-3376.	0.4	662
9	Anticancer and Antiviral Sulfonamides. <i>Current Medicinal Chemistry</i> , 2003, 10, 925-953.	1.2	646
10	Hypoxia activates the capacity of tumor-associated carbonic anhydrase IX to acidify extracellular pH. <i>FEBS Letters</i> , 2004, 577, 439-445.	1.3	620
11	Review Article. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2004, 19, 199-229.	2.5	595
12	How many carbonic anhydrase inhibition mechanisms exist?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 345-360.	2.5	588
13	Carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 3467-3474.	1.0	579
14	Structure-based drug discovery of carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012, 27, 759-772.	2.5	554
15	Carbonic anhydrases as targets for medicinal chemistry. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 4336-4350.	1.4	521
16	Carbonic Anhydrases An Overview. <i>Current Pharmaceutical Design</i> , 2008, 14, 603-614.	0.9	476
17	Non-Zinc Mediated Inhibition of Carbonic Anhydrases: Coumarins Are a New Class of Suicide Inhibitors. <i>Journal of the American Chemical Society</i> , 2009, 131, 3057-3062.	6.6	457
18	Crystal structure of the catalytic domain of the tumor-associated human carbonic anhydrase IX. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 16233-16238.	3.3	451

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19	Ureido-Substituted Benzenesulfonamides Potently Inhibit Carbonic Anhydrase IX and Show Antimetastatic Activity in a Model of Breast Cancer Metastasis. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 1896-1902.	2.9	443
20	Unexpected Nanomolar Inhibition of Carbonic Anhydrase by COX-2-Selective Celecoxib: A New Pharmacological Opportunities Due to Related Binding Site Recognition. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 550-557.	2.9	426
21	Protease inhibitors of the sulfonamide type: Anticancer, antiinflammatory, and antiviral agents. <i>Medicinal Research Reviews</i> , 2003, 23, 535-558.	5.0	385
22	Recent Developments in Targeting Carbonic Anhydrase IX for Cancer Therapeutics. <i>Oncotarget</i> , 2012, 3, 84-97.	0.8	365
23	Deciphering the Mechanism of Carbonic Anhydrase Inhibition with Coumarins and Thiocoumarins. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 335-344.	2.9	363
24	Targeting tumor-associated carbonic anhydrase IX in cancer therapy. <i>Trends in Pharmacological Sciences</i> , 2006, 27, 566-573.	4.0	362
25	Advances in structure-based drug discovery of carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Discovery</i> , 2017, 12, 61-88.	2.5	356
26	An overview of the alpha-, beta- and gamma-carbonic anhydrases from <i>Bacteria</i> : can bacterial carbonic anhydrases shed new light on evolution of bacteria?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015, 30, 325-332.	2.5	328
27	Discovery of a new family of carbonic anhydrases in the malaria pathogen <i>Plasmodium falciparum</i> – The Γ -carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 4389-4396.	1.0	297
28	Carbonic Anhydrase Inhibitors. Synthesis of Water-Soluble, Topically Effective, Intraocular Pressure-Lowering Aromatic/Heterocyclic Sulfonamides Containing Cationic or Anionic Moieties: Is the Tail More Important than the Ring?. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 2641-2650.	2.9	278
29	Carbonic anhydrases: from biomedical applications of the inhibitors and activators to biotechnological use for CO ₂ capture. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013, 28, 229-230.	2.5	278
30	Antiglaucoma carbonic anhydrase inhibitors: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2013, 23, 705-716.	2.4	273
31	Carbonic Anhydrase Activators: X-ray Crystallographic and Spectroscopic Investigations for the Interaction of Isozymes I and II with Histamine,. <i>Biochemistry</i> , 1997, 36, 10384-10392.	1.2	269
32	Carbonic anhydrase inhibitors and activators for novel therapeutic applications. <i>Future Medicinal Chemistry</i> , 2011, 3, 1165-1180.	1.1	260
33	Biochemical Characterization of CA IX, One of the Most Active Carbonic Anhydrase Isozymes. <i>Journal of Biological Chemistry</i> , 2008, 283, 27799-27809.	1.6	258
34	The Warburg Effect and the Hallmarks of Cancer. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2017, 17, 164-170.	0.9	258
35	Sulfa and trimethoprim-like drugs – antimetabolites acting as carbonic anhydrase, dihydropteroate synthase and dihydrofolate reductase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014, 29, 379-387.	2.5	255
36	Diuretics with carbonic anhydrase inhibitory action: a patent and literature review (2005 – 2013). <i>Expert Opinion on Therapeutic Patents</i> , 2013, 23, 681-691.	2.4	252

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37	Carbonic anhydrase inhibitors: E7070, a sulfonamide anticancer agent, potently inhibits cytosolic isozymes I and II, and transmembrane, tumor-associated isozyme IX. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 217-223.	1.0	251
38	The Role of Carbonic Anhydrase 9 in Regulating Extracellular and Intracellular pH in Three-dimensional Tumor Cell Growths. <i>Journal of Biological Chemistry</i> , 2009, 284, 20299-20310.	1.6	249
39	Antiobesity carbonic anhydrase inhibitors: a literature and patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2013, 23, 725-735.	2.4	246
40	Applications of carbonic anhydrase inhibitors and activators in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2002, 12, 217-242.	2.4	243
41	A Smallâ€Molecule Drug Conjugate for the Treatment of Carbonic Anhydrase IX Expressing Tumors. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 4231-4235.	7.2	242
42	Bacterial Carbonic Anhydrases as Drug Targets: Toward Novel Antibiotics?. <i>Frontiers in Pharmacology</i> , 2011, 2, 34.	1.6	229
43	Exploiting the hydrophobic and hydrophilic binding sites for designing carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Discovery</i> , 2013, 8, 793-810.	2.5	229
44	Glycosyl Coumarin Carbonic Anhydrase IX and XII Inhibitors Strongly Attenuate the Growth of Primary Breast Tumors. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 8271-8277.	2.9	228
45	Anticancer carbonic anhydrase inhibitors: a patent review (2008 â€“ 2013). <i>Expert Opinion on Therapeutic Patents</i> , 2013, 23, 737-749.	2.4	226
46	Highly Active Antiretroviral Therapy: Current State of the Art, New Agents and Their Pharmacological Interactions Useful for Improving Therapeutic Outcome. <i>Current Pharmaceutical Design</i> , 2005, 11, 1805-1843.	0.9	222
47	Carbonic anhydrase inhibitors: SAR and X-ray crystallographic study for the interaction of sugar sulfamates/sulfamides with isozymes I, II and IV. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 841-845.	1.0	221
48	Carbonic Anhydrase in the Scleractinian Coral <i>Stylophora pistillata</i> . <i>Journal of Biological Chemistry</i> , 2008, 283, 25475-25484.	1.6	221
49	Diuretics: From Classical Carbonic Anhydrase Inhibitors to Novel Applications of the Sulfonamides. <i>Current Pharmaceutical Design</i> , 2008, 14, 641-648.	0.9	219
50	Carbonic anhydrase inhibitors. Inhibition of the transmembrane isozyme XII with sulfonamidesâ€”a new target for the design of antitumor and antiglaucoma drugs?. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 963-969.	1.0	212
51	Dithiocarbamates Strongly Inhibit Carbonic Anhydrases and Show Antiglaucoma Action in Vivo. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 1721-1730.	2.9	211
52	Characterization of CA XIII, a Novel Member of the Carbonic Anhydrase Isozyme Family. <i>Journal of Biological Chemistry</i> , 2004, 279, 2719-2727.	1.6	210
53	Carbonic Anhydrase Inhibitors as Anticonvulsant Agents. <i>Current Topics in Medicinal Chemistry</i> , 2007, 7, 855-864.	1.0	209
54	Carbonic anhydrase inhibitors. Inhibition of human erythrocyte isozymes I and II with a series of antioxidant phenols. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 3207-3211.	1.4	207

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55	Inhibition of carbonic anhydrase IX targets primary tumors, metastases, and cancer stem cells: Three for the price of one. <i>Medicinal Research Reviews</i> , 2018, 38, 1799-1836.	5.0	207
56	Carbonic Anhydrase Inhibitors. Design of Fluorescent Sulfonamides as Probes of Tumor-Associated Carbonic Anhydrase IX That Inhibit Isozyme IX-Mediated Acidification of Hypoxic Tumors. <i>Journal of Medicinal Chemistry</i> , 2005, 48, 4834-4841.	2.9	205
57	Polyamines Inhibit Carbonic Anhydrases by Anchoring to the Zinc-Coordinated Water Molecule. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 5511-5522.	2.9	205
58	Carbonic anhydrase inhibitors. Inhibition of mammalian isoforms Iâ€“XIV with a series of natural product polyphenols and phenolic acids. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 2159-2164.	1.4	204
59	Anti-infective carbonic anhydrase inhibitors: a patent and literature review. <i>Expert Opinion on Therapeutic Patents</i> , 2013, 23, 693-704.	2.4	203
60	Sulfonamides: a patent review (2008 â€“ 2012). <i>Expert Opinion on Therapeutic Patents</i> , 2012, 22, 747-758.	2.4	201
61	Carbonic Anhydrase Inhibitors: X-ray and Molecular Modeling Study for the Interaction of a Fluorescent Antitumor Sulfonamide with Isozyme II and IX. <i>Journal of the American Chemical Society</i> , 2006, 128, 8329-8335.	6.6	200
62	Selective hydrophobic pocket binding observed within the carbonic anhydrase II active site accommodate different 4-substituted-ureido-benzenesulfonamides and correlate to inhibitor potency. <i>Chemical Communications</i> , 2010, 46, 8371.	2.2	200
63	Sulfocoumarins (1,2-Benzoxathiine-2,2-dioxides): A Class of Potent and Isoform-Selective Inhibitors of Tumor-Associated Carbonic Anhydrases. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 293-300.	2.9	199
64	Carbonic Anhydrase Inhibition and the Management of Hypoxic Tumors. <i>Metabolites</i> , 2017, 7, 48.	1.3	197
65	Carbonic Anhydrases as Drug Targets - An Overview. <i>Current Topics in Medicinal Chemistry</i> , 2007, 7, 825-833.	1.0	195
66	Carbonic anhydrase inhibitors as emerging agents for the treatment and imaging of hypoxic tumors. <i>Expert Opinion on Investigational Drugs</i> , 2018, 27, 963-970.	1.9	195
67	Structure and Inhibition of the CO ₂ -Sensing Carbonic Anhydrase Can2 from the Pathogenic Fungus <i>Cryptococcus neoformans</i> . <i>Journal of Molecular Biology</i> , 2009, 385, 1207-1220.	2.0	193
68	Sulfamates and their therapeutic potential. <i>Medicinal Research Reviews</i> , 2005, 25, 186-228.	5.0	191
69	The #945; and #946; Classes Carbonic Anhydrases from <i>Helicobacter pylori</i> as Novel Drug Targets. <i>Current Pharmaceutical Design</i> , 2008, 14, 622-630.	0.9	188
70	Synthesis and Carbonic Anhydrase Isoenzymes I, II, IX, and XII Inhibitory Effects of Dimethoxybromophenol Derivatives Incorporating Cyclopropane Moieties. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 640-650.	2.9	187
71	Carbonic anhydrase inhibitors: Interactions of phenols with the 12 catalytically active mammalian isoforms (CA Iâ€“XIV). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 1583-1587.	1.0	186
72	(In)organic anions as carbonic anhydrase inhibitors. <i>Journal of Inorganic Biochemistry</i> , 2012, 111, 117-129.	1.5	186

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73	Tumor-associated Carbonic Anhydrase 9 Spatially Coordinates Intracellular pH in Three-dimensional Multicellular Growths. <i>Journal of Biological Chemistry</i> , 2008, 283, 20473-20483.	1.6	185
74	Imaging of CA IX with fluorescent labelled sulfonamides distinguishes hypoxic and (re)-oxygenated cells in a xenograft tumour model. <i>Radiotherapy and Oncology</i> , 2009, 92, 423-428.	0.3	185
75	Carbonic Anhydrase Inhibitors. The Mitochondrial Isozyme VB as a New Target for Sulfonamide and Sulfamate Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2005, 48, 7860-7866.	2.9	179
76	Carbonic Anhydrase Inhibitors: Inhibition of Isozymes I, II, and IX with Triazole-Linked O-Glycosides of Benzene Sulfonamides. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 1651-1657.	2.9	179
77	Carbonic anhydrase inhibitors. Zonisamide is an effective inhibitor of the cytosolic isozyme II and mitochondrial isozyme V: solution and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 2315-2320.	1.0	176
78	Carbonic anhydrases in anthozoan corals: A review. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 1437-1450.	1.4	174
79	Bacterial, fungal and protozoan carbonic anhydrases as drug targets. <i>Expert Opinion on Therapeutic Targets</i> , 2015, 19, 1689-1704.	1.5	174
80	Therapeutic potential of sulfamides as enzyme inhibitors. <i>Medicinal Research Reviews</i> , 2006, 26, 767-792.	5.0	173
81	Rosmarinic acid inhibits some metabolic enzymes including glutathione S-transferase, lactoperoxidase, acetylcholinesterase, butyrylcholinesterase and carbonic anhydrase isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 1698-1702.	2.5	173
82	Sulfonamides and their isosters as carbonic anhydrase inhibitors. <i>Future Medicinal Chemistry</i> , 2014, 6, 1149-1165.	1.1	172
83	In Vitro Inhibition of Human Carbonic Anhydrase I and II Isozymes with Natural Phenolic Compounds. <i>Chemical Biology and Drug Design</i> , 2011, 77, 494-499.	1.5	170
84	In vitro inhibition of \pm -carbonic anhydrase isozymes by some phenolic compounds. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 4259-4262.	1.0	170
85	A Novel Class of Carbonic Anhydrase Inhibitors: Glycoconjugate Benzene Sulfonamides Prepared by Click-Tailing. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 6539-6548.	2.9	168
86	Saccharin Inhibits Carbonic Anhydrases: Possible Explanation for its Unpleasant Metallic Aftertaste. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 7697-7699.	7.2	168
87	Taking advantage of tumor cell adaptations to hypoxia for developing new tumor markers and treatment strategies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2009, 24, 1-39.	2.5	167
88	New strategies for targeting the hypoxic tumour microenvironment in breast cancer. <i>Cancer Treatment Reviews</i> , 2013, 39, 171-179.	3.4	167
89	Indisulam: an anticancer sulfonamide in clinical development. <i>Expert Opinion on Investigational Drugs</i> , 2003, 12, 283-287.	1.9	166
90	Carbonic anhydrase IX: Biochemical and crystallographic characterization of a novel antitumor target. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2010, 1804, 404-409.	1.1	166

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91	Carbonic anhydrase inhibitors as emerging drugs for the treatment of obesity. <i>Expert Opinion on Emerging Drugs</i> , 2008, 13, 383-392.	1.0	165
92	An Overview of the Bacterial Carbonic Anhydrases. <i>Metabolites</i> , 2017, 7, 56.	1.3	165
93	Carbonic Anhydrases and Metabolism. <i>Metabolites</i> , 2018, 8, 25.	1.3	164
94	Review on plug-in electric vehicle charging architectures integrated with distributed energy sources for sustainable mobility. <i>Applied Energy</i> , 2017, 207, 438-464.	5.1	162
95	In vitro inhibition of salicylic acid derivatives on human cytosolic carbonic anhydrase isozymes I and II. <i>Bioorganic and Medicinal Chemistry</i> , 2008, 16, 9101-9105.	1.4	160
96	Carbonic anhydrase inhibitors as antitumor/antimetastatic agents: a patent review (2008-2018). <i>Expert Opinion on Therapeutic Patents</i> , 2018, 28, 729-740.	2.4	160
97	Modulation of carbonic anhydrase activity and its applications in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2004, 14, 667-702.	2.4	159
98	Carbonic anhydrase inhibitors and activators and their use in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2006, 16, 1627-1664.	2.4	158
99	Carbonic Anhydrase Inhibitors: Stacking with Phe131 Determines Active Site Binding Region of Inhibitors As Exemplified by the X-ray Crystal Structure of a Membrane-Impermeant Antitumor Sulfonamide Complexed with Isozyme II. <i>Journal of Medicinal Chemistry</i> , 2005, 48, 5721-5727.	2.9	157
100	Imaging the hypoxia surrogate marker CA IX requires expression and catalytic activity for binding fluorescent sulfonamide inhibitors. <i>Radiotherapy and Oncology</i> , 2007, 83, 367-373.	0.3	157
101	Carbonic anhydrase inhibitors: The \hat{I}^2 -carbonic anhydrase from <i>Helicobacter pylori</i> is a new target for sulfonamide and sulfamate inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 3585-3594.	1.0	157
102	Dithiocarbamates: a new class of carbonic anhydrase inhibitors. <i>Crystallographic and kinetic investigations</i> . <i>Chemical Communications</i> , 2012, 48, 1868.	2.2	157
103	Specific inhibition of carbonic anhydrase IX activity enhances the in vivo therapeutic effect of tumor irradiation. <i>Radiotherapy and Oncology</i> , 2011, 99, 424-431.	0.3	156
104	Nonaromatic Sulfonamide Group as an Ideal Anchor for Potent Human Carbonic Anhydrase Inhibitors: Role of Hydrogen-Bonding Networks in Ligand Binding and Drug Design. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 3583-3587.	2.9	154
105	Carbonic Anhydrase Inhibitors: DNA Cloning and Inhibition Studies of the \hat{I}^{\pm} -Carbonic Anhydrase from <i>Helicobacter pylori</i> , A New Target for Developing Sulfonamide and Sulfamate Gastric Drugs. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 2117-2126.	2.9	154
106	Metal-Based Antibacterial and Antifungal Agents: Synthesis, Characterization, and In Vitro Biological Evaluation of Co(II), Cu(II), Ni(II), and Zn(II) Complexes with Amino Acid-Derived Compounds. <i>Bioinorganic Chemistry and Applications</i> , 2006, 2006, 1-13.	1.8	154
107	Progress in the development of human carbonic anhydrase inhibitors and their pharmacological applications: Where are we today?. <i>Medicinal Research Reviews</i> , 2020, 40, 2485-2565.	5.0	154
108	Carbonic Anhydrase Inhibitors: Synthesis of Water-Soluble, Aminoacyl/Dipeptidyl Sulfonamides Possessing Long-Lasting Intraocular Pressure-Lowering Properties via the Topical Route. <i>Journal of Medicinal Chemistry</i> , 1999, 42, 3690-3700.	2.9	153

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109	Regulation of pH by Carbonic Anhydrase 9 Mediates Survival of Pancreatic Cancer Cells With Activated KRAS in Response to Hypoxia. <i>Gastroenterology</i> , 2019, 157, 823-837.	0.6	153
110	Carbonic anhydrase inhibitors - Part 49: Synthesis of substituted ureido and thioureido derivatives of aromatic/heterocyclic sulfonamides with increased affinities for isozyme I. <i>European Journal of Medicinal Chemistry</i> , 1998, 33, 83-93.	2.6	152
111	7,8-Disubstituted- but not 6,7-disubstituted coumarins selectively inhibit the transmembrane, tumor-associated carbonic anhydrase isoforms IX and XII over the cytosolic ones I and II in the low nanomolar/subnanomolar range. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 7255-7258.	1.0	152
112	Direct Extracellular Interaction between Carbonic Anhydrase IV and the Human NBC1 Sodium/Bicarbonate Co-Transporter. <i>Biochemistry</i> , 2003, 42, 12321-12329.	1.2	151
113	Carbonic anhydrase inhibitors. Antioxidant polyphenols effectively inhibit mammalian isoforms I-XV. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 5050-5053.	1.0	151
114	Carbonic anhydrase inhibitors: The first selective, membrane-impermeant inhibitors targeting the tumor-associated isozyme IX. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 869-873.	1.0	150
115	Are Carbonic Anhydrase Inhibitors Suitable for Obtaining Antiobesity Drugs ?. <i>Current Pharmaceutical Design</i> , 2008, 14, 655-660.	0.9	150
116	A new approach to antiglaucoma drugs: carbonic anhydrase inhibitors with or without NO donating moieties. Mechanism of action and preliminary pharmacology. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012, 27, 138-147.	2.5	150
117	Carbonic Anhydrase Inhibitors. Design of Selective, Membrane-Impermeant Inhibitors Targeting the Human Tumor-Associated Isozyme IX. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 2337-2347.	2.9	149
118	Antimetastatic Effect of Sulfamate Carbonic Anhydrase IX Inhibitors in Breast Carcinoma Xenografts. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 5591-5600.	2.9	149
119	Bacterial protease inhibitors. <i>Medicinal Research Reviews</i> , 2002, 22, 329-372.	5.0	147
120	Carbonic anhydrase inhibitors: X-ray crystallographic structure of the adduct of human isozyme II with EMATE, a dual inhibitor of carbonic anhydrases and steroid sulfatase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 231-234.	1.0	147
121	Metal binding and antibacterial activity of ciprofloxacin complexes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2005, 20, 303-307.	2.5	147
122	Carbonic Anhydrase Inhibitors: Clash with Ala65 as a Means for Designing Inhibitors with Low Affinity for the Ubiquitous Isozyme II, Exemplified by the Crystal Structure of the Topiramate Sulfamide Analogue. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 7024-7031.	2.9	147
123	Zinc Complexes of Benzothiazole-derived Schiff Bases with Antibacterial Activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2003, 18, 259-263.	2.5	146
124	The β -class carbonic anhydrases as drug targets for antimalarial agents. <i>Expert Opinion on Therapeutic Targets</i> , 2015, 19, 551-563.	1.5	146
125	Carbonic anhydrase inhibitors. Inhibition of the human cytosolic isozyme VII with aromatic and heterocyclic sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 971-976.	1.0	145
126	Carbonic Anhydrase Inhibitors: A Water-Soluble 4-Sulfamoylphenylthioureas as Topical Intraocular Pressure-Lowering Agents with Long-Lasting Effects. <i>Journal of Medicinal Chemistry</i> , 2000, 43, 4884-4892.	2.9	143

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127	N-Acylsulfonamides strongly inhibit human carbonic anhydrase isoenzymes I and II. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 2598-2605.	1.4	142
128	The effect of caffeic acid phenethyl ester (CAPE) on metabolic enzymes including acetylcholinesterase, butyrylcholinesterase, glutathione S-transferase, lactoperoxidase, and carbonic anhydrase isoenzymes I, II, IX, and XII. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 1095-1101.	2.5	142
129	Carbonic Anhydrase Inhibitors: Anticonvulsant Sulfonamides Incorporating Valproyl and Other Lipophilic Moieties. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 312-320.	2.9	141
130	Carbonic Anhydrase Inhibitors. Inhibition of Tumor-Associated Isozyme IX by Halogenosulfanilamide and Halogenophenylaminobenzolamide Derivatives. <i>Journal of Medicinal Chemistry</i> , 2003, 46, 2187-2196.	2.9	141
131	Efficient Expression and Crystallization System of Cancer-Associated Carbonic Anhydrase Isoform IX. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 9004-9009.	2.9	141
132	A Phase 1 Study of SLC-0111, a Novel Inhibitor of Carbonic Anhydrase IX, in Patients With Advanced Solid Tumors. <i>American Journal of Clinical Oncology: Cancer Clinical Trials</i> , 2020, 43, 484-490.	0.6	141
133	Benzothiazole derivatives as anticancer agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 265-279.	2.5	140
134	Carbonic Anhydrase Inhibitors: Perfluoroalkyl/Aryl-Substituted Derivatives of Aromatic/Heterocyclic Sulfonamides as Topical Intraocular Pressure-Lowering Agents with Prolonged Duration of Action. <i>Journal of Medicinal Chemistry</i> , 2000, 43, 4542-4551.	2.9	139
135	Carbonic anhydrase inhibitors in the treatment and prophylaxis of obesity. <i>Expert Opinion on Therapeutic Patents</i> , 2003, 13, 1545-1550.	2.4	139
136	Carbonic Anhydrase Inhibitors. A General Approach for the Preparation of Water-Soluble Sulfonamides Incorporating Polyamino Polycarboxylate Tails and of Their Metal Complexes Possessing Long-Lasting, Topical Intraocular Pressure-Lowering Properties. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 1466-1476.	2.9	138
137	Carbonic anhydrase inhibitors: Inhibition of the transmembrane isozyme XIV with sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 3828-3833.	1.0	138
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1237	Benzenesulfonamides incorporating nitrogenous bases show effective inhibition of $\hat{1}^2$ -carbonic anhydrases from the pathogenic fungi <i>Cryptococcus neoformans</i> , <i>Candida glabrata</i> and <i>Malassezia globosa</i> . <i>Bioorganic Chemistry</i> , 2019, 86, 39-43.	2.0	8
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