

# CITATION REPORT

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## Anti-infective carbonic anhydrase inhibitors: a patent and literature review

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#	Paper	IF	Citations
199	5-Substituted-(1,2,3-triazol-4-yl)thiophene-2-sulfonamides strongly inhibit human carbonic anhydrases I, II, IX and XII: solution and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 5130-8	3.4	26
198	A class of sulfonamides with strong inhibitory action against the $\beta$ carbonic anhydrase from <i>Trypanosoma cruzi</i> . <i>Journal of Medicinal Chemistry</i> , <b>2013</b> , 56, 5773-81	8.3	51
197	Synthesis of novel acridine and bis acridine sulfonamides with effective inhibitory activity against the cytosolic carbonic anhydrase isoforms II and VII. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 5799-804	3.4	29
196	Anion inhibition studies of a $\beta$ carbonic anhydrase from <i>Clostridium perfringens</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2013</b> , 23, 6706-10	2.9	42
195	Antimalarial activity of compounds comprising a primary benzene sulfonamide fragment. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2013</b> , 23, 6114-7	2.9	25
194	Anion inhibition studies of the $\beta$ carbonic anhydrase from the protozoan pathogen <i>Trypanosoma cruzi</i> , the causative agent of Chagas disease. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 4472-6	3.4	45
193	Biochemical characterization of the $\beta$ carbonic anhydrase from the marine diatom <i>Thalassiosira weissflogii</i> , TweCA. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2014</b> , 29, 906-11	5.6	58
192	Sulfonamides and their isosters as carbonic anhydrase inhibitors. <i>Future Medicinal Chemistry</i> , <b>2014</b> , 6, 1149-65	4.1	133
191	Biochemical characterization of the $\beta$ carbonic anhydrase from the oral pathogen <i>Porphyromonas gingivalis</i> , PgiCA. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2014</b> , 29, 532-7	5.6	62
190	A class of 4-sulfamoylphenyl- $\beta$ -aminoalkyl ethers with effective carbonic anhydrase inhibitory action and antiglaucoma effects. <i>Journal of Medicinal Chemistry</i> , <b>2014</b> , 57, 9673-86	8.3	44
189	Structural insights on carbonic anhydrase inhibitory action, isoform selectivity, and potency of sulfonamides and coumarins incorporating arylsulfonylureido groups. <i>Journal of Medicinal Chemistry</i> , <b>2014</b> , 57, 9152-67	8.3	46
188	Anion inhibition study of the $\beta$ carbonic anhydrase (CahB1) from the cyanobacterium <i>Coleofasciculus chthonoplastes</i> (ex- <i>Microcoleus chthonoplastes</i> ). <i>Bioorganic and Medicinal Chemistry</i> , <b>2014</b> , 22, 1667-71	3.4	22
187	6-Triazolyl-substituted sulfocoumarins are potent, selective inhibitors of the tumor-associated carbonic anhydrases IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 1256-60	2.9	52
186	Sulfonamide inhibition studies of two $\beta$ carbonic anhydrases from the bacterial pathogen <i>Legionella pneumophila</i> . <i>Bioorganic and Medicinal Chemistry</i> , <b>2014</b> , 22, 2939-46	3.4	41
185	Inhibition of carbonic anhydrases from the extremophilic bacteria <i>Sulfurihydrogenibium yellowstonense</i> (SspCA) and <i>S. azorensis</i> (SazCA) with a new series of sulfonamides incorporating aroylhydrazone-, [1,2,4]triazolo[3,4-b][1,3,4]thiadiazinyl- or 2-(arylsulfonylureido)thiazole-, 1,2,4-triazolo[3,4-b][1,3,4]thiadiazinyl- and 1,2,4-triazolo[3,4-b][1,3,4]thiadiazinyl- groups. <i>Bioorganic and Medicinal Chemistry</i> , <b>2014</b> , 22, 1667-71	3.4	42
184	Sulfonamide inhibition studies of the $\beta$ carbonic anhydrase from the diatom <i>Thalassiosira weissflogii</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 275-9	2.9	47
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182	Discovery of a new family of carbonic anhydrases in the malaria pathogen Plasmodium falciparum—the $\beta$ -carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 4389-4396	2.9	258
181	Synthesis of sulfonamides with effective inhibitory action against Porphyromonas gingivalis $\beta$ -carbonic anhydrase. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 4006-10	2.9	20
180	Anion inhibition study of the $\beta$ -class carbonic anhydrase (PgiCAB) from the oral pathogen Porphyromonas gingivalis. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 4402-4406	2.9	27
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178	Anion inhibition studies of two new $\beta$ -carbonic anhydrases from the bacterial pathogen Legionella pneumophila. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 1127-32	2.9	44
177	Synthesis and carbonic anhydrase I, II, IX and XII inhibition studies of 4-N,N-disubstituted sulfanilamides incorporating 4,4,4-trifluoro-3-oxo-but-1-enyl, phenacylthiourea and imidazol-2(3H)-one/thione moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 1776-9	2.9	21
176	Anion inhibition studies of two $\beta$ -carbonic anhydrases from Lotus japonicus, LjCAA1 and LjCAA2. <b>2014</b> , 136, 67-72		15
175	Sulfonamide inhibition studies of the $\beta$ -carbonic anhydrase from the oral pathogen Porphyromonas gingivalis. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 240-4	2.9	46
174	Overview of carbonic anhydrase families/isoforms. <b>2014</b> , 6-16		0
173	Developing Novel Bacterial Targets: Carbonic Anhydrases as Antibacterial Drug Targets. <b>2014</b> , 31-46		1
172	Crystal structure and kinetic studies of a tetrameric type II $\beta$ -carbonic anhydrase from the pathogenic bacterium Vibrio cholerae. <b>2015</b> , 71, 2449-56		83
171	Editorial: New Antimicrobial Therapeutics. <i>Current Medicinal Chemistry</i> , <b>2015</b> , 22, 2112-5	4.3	3
170	Design and synthesis of benzothiazole-6-sulfonamides acting as highly potent inhibitors of carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 4989-4999	3.4	23
169	Protonography, a powerful tool for analyzing the activity and the oligomeric state of the $\beta$ -carbonic anhydrase identified in the genome of Porphyromonas gingivalis. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 3747-50	3.4	39
168	Cloning, characterization and anion inhibition study of a $\beta$ -class carbonic anhydrase from the caries producing pathogen Streptococcus mutans. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 2995-3001	3.4	24
167	The zinc coordination pattern in the $\beta$ -carbonic anhydrase from Plasmodium falciparum is different from all other carbonic anhydrase genetic families. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2015</b> , 25, 1385-9	2.9	95
166	Cloning, characterization and anion inhibition studies of a $\beta$ -carbonic anhydrase from the Antarctic cyanobacterium Nostoc commune. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2015</b> , 25, 4970-4975	2.9	12
165	Nanoparticles for controlled release of anti-biofilm agents WO2014130994 (A1): a patent evaluation. <i>Expert Opinion on Therapeutic Patents</i> , <b>2015</b> , 25, 945-8	6.8	

164	Inhibition of carbonic anhydrase isoforms I, II, IX and XII with Schiff's bases incorporating iminoureido moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 901-7	5.6	13
163	N-glycosyl-N-hydroxysulfamides as potent inhibitors of <i>Brucella suis</i> carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 1010-2	5.6	6
162	A new class of quinazoline-sulfonamides acting as efficient inhibitors against the $\beta$ -carbonic anhydrase from <i>Trypanosoma cruzi</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 581-5	5.6	19
161	Inhibition of $\beta$ -carbonic anhydrases from <i>Brucella suis</i> with C-cinnamoyl glycosides incorporating the phenol moiety. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 1017-20	5.6	13
160	The $\beta$ -class carbonic anhydrases as drug targets for antimalarial agents. <b>2015</b> , 19, 551-63		135
159	Protonography, a technique applicable for the analysis of $\beta$ -carbonic anhydrase activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 920-4	5.6	44
158	Exploring QSARs of some benzenesulfonamides incorporating cyanoacrylamide moieties as a carbonic anhydrase inhibitors (specifically against tumor-associated isoforms IX and XII). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 519-23	5.6	7
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154	Synthesis and carbonic anhydrase I, II, IX and XII inhibitory activity of sulfamates incorporating piperazinyl-ureido moieties. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 5619-25	3.4	14
153	Bacterial, fungal and protozoan carbonic anhydrases as drug targets. <b>2015</b> , 19, 1689-704		153
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148	Fluorinated pyrrolidines and piperidines incorporating tertiary benzenesulfonamide moieties are selective carbonic anhydrase II inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 737-45	5.6	30
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146	A Key Opinion Leader interview: insight into the research and career of Prof. Claudiu T Supuran. <i>Expert Opinion on Therapeutic Patents</i> , <b>2015</b> , 25, 501-5	6.8	3
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144	7-Amino-3,4-dihydro-1H-quinolin-2-one, a compound similar to the substituted coumarins, inhibits $\alpha$ -carbonic anhydrases without hydrolysis of the lactam ring. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 773-7	5.6	21
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142	Dendrimers incorporating benzenesulfonamide moieties strongly inhibit carbonic anhydrase isoforms I-XIV. <b>2015</b> , 13, 6453-7		12
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138	Inhibition of mammalian carbonic anhydrase isoforms I-XIV with a series of phenolic acid esters. <i>Bioorganic and Medicinal Chemistry</i> , <b>2015</b> , 23, 7181-8	3.4	24
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127	New light on bacterial carbonic anhydrases phylogeny based on the analysis of signal peptide sequences. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2016</b> , 31, 1254-60	5.6	59
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116	Synthesis and carbonic anhydrase inhibitory activities of new thienyl-substituted pyrazoline benzenesulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2016</b> , 31, 1-5	5.6	39
115	Bacterial Carbonic Anhydrases. <i>Topics in Medicinal Chemistry</i> , <b>2016</b> , 135-152	0.4	2
114	Monothiocarbamates Strongly Inhibit Carbonic Anhydrases in Vitro and Possess Intraocular Pressure Lowering Activity in an Animal Model of Glaucoma. <i>Journal of Medicinal Chemistry</i> , <b>2016</b> , 59, 5857-67	8.3	47
113	Synthesis of 4,5-disubstituted-2-thioxo-1,2,3,4-tetrahydropyrimidines and investigation of their acetylcholinesterase, butyrylcholinesterase, carbonic anhydrase I/II inhibitory and antioxidant activities. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2016</b> , 31, 1-9	5.6	92
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111	Dithiocarbamates with potent inhibitory activity against the <i>Saccharomyces cerevisiae</i> $\beta$ carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2016</b> , 31, 132-6	5.6	15



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93	Metal complexes of benzimidazole derived sulfonamide: Synthesis, molecular structures and antimicrobial activity. <i>Inorganica Chimica Acta</i> , <b>2016</b> , 443, 179-185	2.7	33

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