

Clemente Capasso

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

1,363
papers

71,124
citations

126
h-index

197
g-index

1,403
ext. papers

77,848
ext. citations

5.2
avg, IF

8.87
L-index

#	Paper	IF	Citations
1363	Discovery of 2,4-thiazolidinedione-tethered coumarins as novel selective inhibitors for carbonic anhydrase IX and XII isoforms.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 531-541	5.6	8
1362	Coumarins effectively inhibit bacterial β -carbonic anhydrases.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 333-338	5.6	6
1361	2-(2-Hydroxyethyl)piperazine derivatives as potent human carbonic anhydrase inhibitors: Synthesis, enzyme inhibition, computational studies and antiglaucoma activity.. <i>European Journal of Medicinal Chemistry</i> , 2022 , 228, 114026	6.8	
1360	The Role of Selenium in Pathologies: An Updated Review.. <i>Antioxidants</i> , 2022 , 11,	7.1	14
1359	Chagas Disease: Drug Development and Parasite Targets. <i>Topics in Medicinal Chemistry</i> , 2022 , 1	0.4	3
1358	Discovery of new carbonic anhydrase IX inhibitors as anticancer agents by tuning the hydrophobic and hydrophilic rims of the active site to encounter the dual-tail approach.. <i>European Journal of Medicinal Chemistry</i> , 2022 , 232, 114190	6.8	7
1357	Inhibition studies of bacterial β -carbonic anhydrases with phenols.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 666-671	5.6	3
1356	Coumarins inhibit β -class carbonic anhydrase from .. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 680-685	5.6	1
1355	Repurposing FDA-approved sulphonamide carbonic anhydrase inhibitors for treatment of .. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 51-61	5.6	6
1354	2-Aminobenzoxazole-appended coumarins as potent and selective inhibitors of tumour-associated carbonic anhydrases.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 168-177	5.6	3
1353	Antiproliferative effects of sulphonamide carbonic anhydrase inhibitors C18, SLC-0111 and acetazolamide on bladder, glioblastoma and pancreatic cancer cell lines.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 280-286	5.6	9
1352	Synthesis, molecular modelling and QSAR study of new phenylacetamide-2-oxoindole benzensulfonamide conjugates as carbonic anhydrase inhibitors with antiproliferative activity.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 701-717	5.6	2
1351	Isocoumarins: a new class of selective carbonic anhydrase IX and XII inhibitors.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 743-748	5.6	5
1350	Inhibition of <i>Schistosoma mansoni</i> carbonic anhydrase by the antiparasitic drug clorsulon: X-ray crystallographic and in vitro studies.. <i>Acta Crystallographica Section D: Structural Biology</i> , 2022 , 78, 321-327	5.5	3
1349	Diversely substituted sulfamides for fragment-based drug discovery of carbonic anhydrase inhibitors: synthesis and inhibitory profile.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 857-865	5.6	
1348	Biological investigation of α -methyl thiosemicarbazones as antimicrobial agents and bacterial carbonic anhydrases inhibitors.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 986-993	5.6	1
1347	5-(Sulfamoyl)thien-2-yl 1,3-oxazole inhibitors of carbonic anhydrase II with hydrophilic periphery.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1005-1011	5.6	1

1346	Perspectives on the design and discovery of β -ketoamide inhibitors for the treatment of novel coronavirus: where do we stand and where do we go?. <i>Expert Opinion on Drug Discovery</i> , 2022 , 1-11	6.2	1
1345	The three-tails approach as a new strategy to improve selectivity of action of sulphonamide inhibitors against tumour-associated carbonic anhydrase IX and XII.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 930-939	5.6	6
1344	Perfusion-Based Bioreactor Culture and Isothermal Microcalorimetry for Preclinical Drug Testing with the Carbonic Anhydrase Inhibitor SLC-0111 in Patient-Derived Neuroblastoma.. <i>International Journal of Molecular Sciences</i> , 2022 , 23,	6.3	3
1343	4-Anilinoquinazoline-based benzenesulfonamides as nanomolar inhibitors of carbonic anhydrase isoforms I, II, IX, and XII: design, synthesis, , and biological studies.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 994-1004	5.6	5
1342	Exploration of 2-phenylquinoline-4-carboxamide linked benzene sulfonamide derivatives as isoform selective inhibitors of transmembrane human carbonic anhydrases.. <i>European Journal of Medicinal Chemistry</i> , 2022 , 234, 114247	6.8	1
1341	Benzoselenoates: A novel class of carbonic anhydrase inhibitors.. <i>Bioorganic Chemistry</i> , 2022 , 122, 105754	5.1	0
1340	Tail-approach based design and synthesis of Arylthiazolylhydrazono-1,2,3-triazoles incorporating sulfanilamide and metanilamide as human carbonic anhydrase I, II, IV and IX inhibitors.. <i>Bioorganic Chemistry</i> , 2022 , 123, 105764	5.1	1
1339	Heterobimetallic complexes containing organometallic acylhydrazone ligands as potential inhibitors of human carbonic anhydrases.. <i>Journal of Inorganic Biochemistry</i> , 2022 , 232, 111814	4.2	0
1338	Dithiocarbamates effectively inhibit the β -carbonic anhydrase from .. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1-8	5.6	2
1337	Heterologous expression and biochemical characterisation of the recombinant β -carbonic anhydrase (MpaCA) from the warm-blooded vertebrate pathogen .. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 62-68	5.6	2
1336	Novel 3-(6-methylpyridin-2-yl)coumarin-based chalcones as selective inhibitors of cancer-related carbonic anhydrases IX and XII endowed with anti-proliferative activity.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1043-1052	5.6	5
1335	New 1H-indole-2,3-dione 3-thiosemicarbazones with 3-sulfamoylphenyl moiety as selective carbonic anhydrase inhibitors.. <i>Archiv Der Pharmazie</i> , 2022 , e2200023	4.3	0
1334	The inhibitory effect of boric acid on hypoxia-regulated tumour-associated carbonic anhydrase IX.. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1340-1345	5.6	2
1333	4-(3-Alkyl/benzyl-guanidino)benzenesulfonamides as selective carbonic anhydrase VII inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1568-1576	5.6	7
1332	Cloning, purification, kinetic and anion inhibition studies of a recombinant β -carbonic anhydrase from the Atlantic salmon parasite platyhelminth <i>Gyrodactylus salaris</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1577-1586	5.6	4
1331	Selective inhibition of carbonic anhydrase IX by sulphonylated 1,2,3-triazole incorporated benzenesulphonamides capable of inducing apoptosis. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1454-1463	5.6	2
1330	Development of Praziquantel sulphonamide derivatives as antischistosomal drugs. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2022 , 37, 1479-1494	5.6	1
1329	A decade of tail-approach based design of selective as well as potent tumor associated carbonic anhydrase inhibitors. <i>Bioorganic Chemistry</i> , 2022 , 126, 105920	5.1	5

1328	Insights into the effect of elaborating coumarin-based aryl enaminones with sulfonamide or carboxylic acid functionality on carbonic anhydrase inhibitory potency and selectivity. <i>Bioorganic Chemistry</i> , 2022 , 126, 105888	5.1	3
1327	Beta-Carbonic Anhydrase 1 from Trichomonas Vaginalis as New Antiprotozoan Drug Target. <i>Topics in Medicinal Chemistry</i> , 2021 , 1	0.4	
1326	Class Carbonic Anhydrases as Antiplasmodial Drug Targets: Current State of the Art and Hurdles to Develop New Antimalarials. <i>Topics in Medicinal Chemistry</i> , 2021 , 1	0.4	
1325	Targeting Carbonic Anhydrases from Trypanosoma cruzi and Leishmania spp. as a Therapeutic Strategy to Obtain New Antiprotozoal Drugs. <i>Topics in Medicinal Chemistry</i> , 2021 , 1	0.4	1
1324	Post-translational modifications in tumor-associated carbonic anhydrases. <i>Amino Acids</i> , 2021 ,	3.5	5
1323	Amine- and Amino Acid-Based Compounds as Carbonic Anhydrase Activators. <i>Molecules</i> , 2021 , 26,	4.8	5
1322	Carbonic anhydrase inhibitors: an update on experimental agents for the treatment and imaging of hypoxic tumors. <i>Expert Opinion on Investigational Drugs</i> , 2021 ,	5.9	21
1321	Natural inspired ligustrazine-based SLC-0111 analogues as novel carbonic anhydrase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021 , 228, 114008	6.8	4
1320	Application of the dual-tail approach for the design and synthesis of novel Thiopyrimidine-Benzenesulfonamide hybrids as selective carbonic anhydrase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021 , 228, 114004	6.8	3
1319	Carbonic Anhydrase Inhibition with Sulfonamides Incorporating Pyrazole- and Pyridazinecarboxamide Moieties Provides Examples of Isoform-Selective Inhibitors. <i>Molecules</i> , 2021 , 26,	4.8	1
1318	Glyco-Coated CdSe/ZnS Quantum Dots as Nanoprobes for Carbonic Anhydrase IX Imaging in Cancer Cells.. <i>ACS Applied Nano Materials</i> , 2021 , 4, 14153-14160	5.6	2
1317	Synthesis and Applications of Organic Selenols. <i>Advanced Synthesis and Catalysis</i> , 2021 , 363, 5360	5.6	6
1316	Synthesis, biological evaluation, and in silico studies of potential activators of apoptosis and carbonic anhydrase inhibitors on isatin-5-sulfonamide scaffold.. <i>European Journal of Medicinal Chemistry</i> , 2021 , 228, 113997	6.8	3
1315	Microbiota, Bacterial Carbonic Anhydrases, and Modulators of Their Activity: Links to Human Diseases?. <i>Mediators of Inflammation</i> , 2021 , 2021, 6926082	4.3	5
1314	Carbonic Anhydrase Inhibitors: Designing Isozyme-Specific Inhibitors as Therapeutic Agents. <i>Progress in Drug Research Fortschritte Der Arzneimittelforschung Progres Des Recherches Pharmaceutiques</i> , 2021 , 221-235		
1313	Vanillin enones as selective inhibitors of the cancer associated carbonic anhydrase isoforms IX and XII. The out of the active site pocket for the design of selective inhibitors?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 2118-2127	5.6	
1312	Biochemical and structural characterization of beta-carbonic anhydrase from the parasite Trichomonas vaginalis. <i>Journal of Molecular Medicine</i> , 2021 , 1	5.5	1
1311	Design and development of novel series of indole-3-sulfonamide ureido derivatives as selective carbonic anhydrase II inhibitors. <i>Archiv Der Pharmazie</i> , 2021 , e2100333	4.3	1

1310	Novel benzenesulfonamide-bearing pyrazoles and 1,2,4-thiadiazoles as selective carbonic anhydrase inhibitors. <i>Archiv Der Pharmazie</i> , 2021 , e2100241	4.3	1
1309	Development of Novel Quinoline-Based Sulfonamides as Selective Cancer-Associated Carbonic Anhydrase Isoform IX Inhibitors. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	10
1308	Selective Inhibition of Carbonic Anhydrases by Carvacrol and Thymol Could Impair Biofilm Production and the Release of Outer Membrane Vesicles. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	10
1307	Inhibition of the Carbonic anhydrase from the protozoan pathogen with sulphonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 329-334	5.6	3
1306	Discovery of a novel series of indolylchalcone-benzenesulfonamide hybrids acting as selective carbonic anhydrase II inhibitors. <i>Bioorganic Chemistry</i> , 2021 , 108, 104647	5.1	8
1305	Multitargeting application of proline-derived peptidomimetics addressing cancer-related human matrix metalloproteinase 9 and carbonic anhydrase II. <i>European Journal of Medicinal Chemistry</i> , 2021 , 214, 113260	6.8	2
1304	Structure-Activity Relationship Studies of Acetazolamide-Based Carbonic Anhydrase Inhibitors with Activity against. <i>ACS Infectious Diseases</i> , 2021 , 7, 1969-1984	5.5	23
1303	Discovery of Potent Carbonic Anhydrase Inhibitors as Effective Anticonvulsant Agents: Drug Design, Synthesis, and In Vitro and In Vivo Investigations. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 3100-3114	8.3	10
1302	Coronaviruses. <i>Expert Opinion on Therapeutic Patents</i> , 2021 , 31, 291-294	6.8	4
1301	Development of novel benzofuran-based SLC-0111 analogs as selective cancer-associated carbonic anhydrase isoform IX inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021 , 216, 113283	6.8	20
1300	Benzyl alcohol inhibits carbonic anhydrases by anchoring to the zinc coordinated water molecule. <i>Biochemical and Biophysical Research Communications</i> , 2021 , 548, 217-221	3.4	4
1299	Synthesis and Biological Evaluation of Coumarin-Linked 4-Anilinomethyl-1,2,3-Triazoles as Potent Inhibitors of Carbonic Anhydrases IX and XIII Involved in Tumorigenesis. <i>Metabolites</i> , 2021 , 11,	5.6	2
1298	A Highlight on the Inhibition of Fungal Carbonic Anhydrases as Drug Targets for the Antifungal Armamentarium. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	5
1297	Role of Carbonic Anhydrase in Cerebral Ischemia and Carbonic Anhydrase Inhibitors as Putative Protective Agents. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3
1296	Advances in the discovery of novel agents for the treatment of glaucoma. <i>Expert Opinion on Drug Discovery</i> , 2021 , 16, 1209-1225	6.2	8
1295	Emerging role of carbonic anhydrase inhibitors. <i>Clinical Science</i> , 2021 , 135, 1233-1249	6.5	41
1294	Chromene-Containing Aromatic Sulfonamides with Carbonic Anhydrase Inhibitory Properties. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	1
1293	Synthesis of Azasugar Sulfonamide conjugates and their Evaluation as Inhibitors of Carbonic Anhydrases: the Azasugar Approach to Selectivity. <i>European Journal of Organic Chemistry</i> , 2021 , 2021, 2604-2614	3.2	1

1292	Small-molecule CD73 inhibitors for the immunotherapy of cancer: a patent and literature review (2017-present). <i>Expert Opinion on Therapeutic Patents</i> , 2021 , 31, 867-876	6.8	4
1291	Comprehensive study on potent and selective carbonic anhydrase inhibitors: Synthesis, bioactivities and molecular modelling studies of 4-(3-(2-arylidenehydrazine-1-carbonyl)-5-(thiophen-2-yl)-1H-pyrazole-1-yl) benzenesulfonamides. <i>European Journal of Medicinal Chemistry</i> , 2021 , 217, 113351	6.8	13
1290	Quinoline-sulfamoyl carbamates/sulfamide derivatives: Synthesis, cytotoxicity, carbonic anhydrase activity, and molecular modelling studies. <i>Bioorganic Chemistry</i> , 2021 , 110, 104778	5.1	6
1289	Inhibition of Carbonic Anhydrase IX Promotes Apoptosis through Intracellular pH Level Alterations in Cervical Cancer Cells. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	6
1288	Identification of N-phenyl-2-(phenylsulfonyl)acetamides/propanamides as new SLC-0111 analogues: Synthesis and evaluation of the carbonic anhydrase inhibitory activities. <i>European Journal of Medicinal Chemistry</i> , 2021 , 218, 113360	6.8	13
1287	Insertion of metal carbenes into the anilinic N-H bond of unprotected aminobenzenesulfonamides delivers low nanomolar inhibitors of human carbonic anhydrase IX and XII isoforms. <i>European Journal of Medicinal Chemistry</i> , 2021 , 218, 113352	6.8	3
1286	Taurultams incorporating arylsulfonamide: First in vitro inhibition studies of H and E class Carbonic Anhydrases from <i>Vibrio cholerae</i> and <i>Burkholderia pseudomallei</i> . <i>European Journal of Medicinal Chemistry</i> , 2021 , 219, 113444	6.8	2
1285	Synthesis and Human Carbonic Anhydrase I, II, IX, and XII Inhibition Studies of Sulphonamides Incorporating Mono-, Bi- and Tricyclic Imide Moieties. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	1
1284	Structural Insights into Carbonic Anhydrase (SmCA) Inhibition by Selenoureido-Substituted Benzenesulfonamides. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 10418-10428	8.3	3
1283	Novel 3-substituted coumarins as selective human carbonic anhydrase IX and XII inhibitors: Synthesis, biological and molecular dynamics analysis. <i>European Journal of Medicinal Chemistry</i> , 2021 , 209, 112897	6.8	19
1282	Ninhydrins inhibit carbonic anhydrases directly binding to the metal ion. <i>European Journal of Medicinal Chemistry</i> , 2021 , 209, 112875	6.8	11
1281	Radiotracers for positron emission tomography (PET) targeting tumour-associated carbonic anhydrase isoforms. <i>European Journal of Medicinal Chemistry</i> , 2021 , 213, 113046	6.8	6
1280	Protease inhibitors targeting the main protease and papain-like protease of coronaviruses. <i>Expert Opinion on Therapeutic Patents</i> , 2021 , 31, 309-324	6.8	13
1279	Activation of carbonic anhydrases from human brain by amino alcohol oxime ethers: towards human carbonic anhydrase VII selective activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 48-57	5.6	5
1278	Activation of the carbonic anhydrase from the protozoan pathogen with amines and amino acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 758-763	5.6	3
1277	Anion inhibition studies of the carbonic anhydrases from. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1061-1066	5.6	6
1276	PEG Linker Length Strongly Affects Tumor Cell Killing by PEGylated Carbonic Anhydrase Inhibitors in Hypoxic Carcinomas Expressing Carbonic Anhydrase IX. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3
1275	Anti-breast cancer action of carbonic anhydrase IX inhibitor 4-[4-(4-Benzo[1,3]dioxol-5-ylmethyl-piperazin-1-yl)-benzylidene-hydrazinocarbonyl]-benzenesulfonamide (BSM-0004): and studies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 954-963	5.6	4

1274	Zeta-carbonic anhydrases show CS hydrolase activity: A new metabolic carbon acquisition pathway in diatoms?. <i>Computational and Structural Biotechnology Journal</i> , 2021 , 19, 3427-3436	6.8	4
1273	Biochemical profiling of anti-HIV prodrug Elsulfavirine (Elpida) and its active form VM1500A against a panel of twelve human carbonic anhydrase isoforms. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1056-1060	5.6	1
1272	Anion inhibition studies of the Zn(II)-bound β -carbonic anhydrase from the Gram-negative bacterium. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 372-376	5.6	13
1271	A Story on Carbon Dioxide and Its Hydration 2021 , 115-131		
1270	Effect of Sulfonamides and Their Structurally Related Derivatives on the Activity of β -Carbonic Anhydrase from. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	10
1269	Tetrahydroquinazole-based secondary sulphonamides as carbonic anhydrase inhibitors: synthesis, biological evaluation against isoforms I, II, IV, and IX, and computational studies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1874-1883	5.6	2
1268	Multitargeting approaches involving carbonic anhydrase inhibitors: hybrid drugs against a variety of disorders. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1702-1714	5.6	14
1267	Effect of amino acids and amines on the activity of the recombinant β -carbonic anhydrase from the Gram-negative bacterium. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1000-1006	5.6	3
1266	An anion and small molecule inhibition study of the β -carbonic anhydrase from. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1088-1092	5.6	5
1265	Is carbonic anhydrase inhibition useful as a complementary therapy of Covid-19 infection?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1230-1235	5.6	9
1264	Handling drug-target selectivity: A study on ureido containing Carbonic Anhydrase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021 , 212, 113035	6.8	8
1263	Phenols from L. and L. and their activity against carbonic anhydrase. <i>Natural Product Research</i> , 2021 , 1-7	2.3	7
1262	Design and synthesis of benzenesulfonamide-linked imidazo[2,1-b][1,3,4]thiadiazole derivatives as carbonic anhydrase I and II inhibitors. <i>Archiv Der Pharmazie</i> , 2021 , 354, e2100028	4.3	2
1261	Carbonic Anhydrases: New Perspectives on Protein Functional Role and Inhibition in. <i>Frontiers in Microbiology</i> , 2021 , 12, 629163	5.7	18
1260	Synthesis of new 7-amino-3,4-dihydroquinolin-2(1H)-one-peptide derivatives and their carbonic anhydrase enzyme inhibition, antioxidant, and cytotoxic activities. <i>Archiv Der Pharmazie</i> , 2021 , 354, e2100022	4.3	2
1259	Design, synthesis and biochemical evaluation of novel carbonic anhydrase inhibitors triggered by structural knowledge on hCA VII. <i>Bioorganic and Medicinal Chemistry</i> , 2021 , 44, 116279	3.4	0
1258	Modulating the Efficacy of Carbonic Anhydrase Inhibitors through Fluorine Substitution. <i>Angewandte Chemie - International Edition</i> , 2021 , 60, 23068-23082	16.4	4
1257	New Sulfanilamide Derivatives Incorporating Heterocyclic Carboxamide Moieties as Carbonic Anhydrase Inhibitors. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	3

1256	Genome-wide synthetic lethal screen unveils novel CAIX-NFS1/xCT axis as a targetable vulnerability in hypoxic solid tumors. <i>Science Advances</i> , 2021 , 7,	14.3	29
1255	Quantum mechanical study on the activation mechanism of human carbonic anhydrase VII cluster model with bis-histamine schiff bases and bis-spinaceamine derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2021 , 44, 116276	3.4	3
1254	Binding site comparison for coumarin inhibitors and amine/amino acid activators of human carbonic anhydrases. <i>European Journal of Medicinal Chemistry</i> , 2021 , 226, 113875	6.8	7
1253	Determination of intracellular protein-ligand binding affinity by competition binding in-cell NMR. <i>Acta Crystallographica Section D: Structural Biology</i> , 2021 , 77, 1270-1281	5.5	4
1252	Synthesis, Crystal Structure, Inhibitory Activity and Molecular Docking of Coumarins/Sulfonamides Containing Triazolyl Pyridine Moiety as Potent Selective Carbonic Anhydrase IX and XII Inhibitors. <i>Crystals</i> , 2021 , 11, 1076	2.3	4
1251	Novel triazole-sulfonamide bearing pyrimidine moieties with carbonic anhydrase inhibitory action: Design, synthesis, computational and enzyme inhibition studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021 , 48, 128249	2.9	7
1250	Carbonic Anhydrase IV Selective Inhibitors Counteract the Development of Colitis-Associated Visceral Pain in Rats. <i>Cells</i> , 2021 , 10,	7.9	1
1249	QM and QM/MM study on inhibition mechanism of polyphenolic compounds as non-classical inhibitors of human carbonic anhydrase (II). <i>Theoretical Chemistry Accounts</i> , 2021 , 140, 1	1.9	0
1248	Deciphering the key heterocyclic scaffolds in targeting microtubules, kinases and carbonic anhydrases for cancer drug development. <i>Pharmacology & Therapeutics</i> , 2021 , 225, 107860	13.9	9
1247	Novel benzenesulfonamides aryl and arylsulfone conjugates adopting tail/dual tail approaches: Synthesis, carbonic anhydrase inhibitory activity and molecular modeling studies. <i>European Journal of Medicinal Chemistry</i> , 2021 , 221, 113486	6.8	11
1246	Bacterial carbonic anhydrases: underexploited antibacterial therapeutic targets. <i>Future Medicinal Chemistry</i> , 2021 , 13, 1619-1622	4.1	5
1245	Investigation of 3-sulfamoyl coumarins against cancer-related IX and XII isoforms of human carbonic anhydrase as well as cancer cells leads to the discovery of 2-oxo-2H-benzo[h]chromene-3-sulfonamide - A new caspase-activating proapoptotic agent. <i>European Journal of Medicinal Chemistry</i> , 2021 , 225, 113800	6.8	10
1244	Exploring of tumor-associated carbonic anhydrase isoenzyme IX and XII inhibitory effects and cytotoxicities of the novel N-aryl-1-(4-sulfamoylphenyl)-5-(thiophen-2-yl)-1H-pyrazole-3-carboxamides. <i>Bioorganic Chemistry</i> , 2021 , 115, 105194	5.1	6
1243	Natural inspired piperine-based sulfonamides and carboxylic acids as carbonic anhydrase inhibitors: Design, synthesis and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2021 , 225, 113800	6.8	6
1242	Evaluating the efficiency of enzyme accelerated CO capture: chemical kinetics modelling for interpreting measurement results. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 394-401	5.6	2
1241	Selective inhibition of carbonic anhydrase IX and XII by coumarin and psoralen derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 685-692	5.6	8
1240	Inhibition of human carbonic anhydrases from the pathogenic bacterium with aromatic sulphonamides and clinically licenced drugs - a joint docking/molecular dynamics study. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 469-479	5.6	7
1239	Natural products in drug discovery: advances and opportunities. <i>Nature Reviews Drug Discovery</i> , 2021 , 20, 200-216	64.1	522

1238	Activation of carbonic anhydrase isoforms involved in modulation of emotional memory and cognitive disorders with histamine agonists, antagonists and derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 719-726	5.6	8
1237	Protective effects of carbonic anhydrase inhibition in brain ischaemia and models. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 964-976	5.6	4
1236	An overview on the recently discovered iota-carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 1988-1995	5.6	17
1235	Reconsidering anion inhibitors in the general context of drug design studies of modulators of activity of the classical enzyme carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 561-580	5.6	51
1234	Challenges and Promises for Obtaining New Antiprotozoal Drugs: What's Going Wrong?. <i>Topics in Medicinal Chemistry</i> , 2021 , 1	0.4	2
1233	Aromatic Sulfonamides including a Sulfonic Acid Tail: New Membrane Impermeant Carbonic Anhydrase Inhibitors for Targeting Selectively the Cancer-Associated Isoforms.. <i>International Journal of Molecular Sciences</i> , 2021 , 23,	6.3	4
1232	Novel 1,3,5-Triazinyl Aminobenzenesulfonamides Incorporating Aminoalcohol, Aminochalcone and Aminostilbene Structural Motifs as Potent Anti-VRE Agents, and Carbonic Anhydrases I, II, VII, IX, and XII Inhibitors.. <i>International Journal of Molecular Sciences</i> , 2021 , 23,	6.3	3
1231	One-Pot Procedure for the Synthesis of Asymmetric Substituted Ureido Benzene Sulfonamides as Effective Inhibitors of Carbonic Anhydrase Enzymes.. <i>Journal of Medicinal Chemistry</i> , 2021 ,	8.3	1
1230	Ureidosulfocoumarin Derivatives As Selective and Potent Carbonic Anhydrase IX and XII Inhibitors.. <i>ChemMedChem</i> , 2021 , e202100725	3.7	1
1229	Carbonic Anhydrases: Versatile and Useful Biocatalysts in Chemistry and Biochemistry. <i>Catalysts</i> , 2020 , 10, 1008	4	13
1228	Bacterial β -carbonic anhydrase: a new active class of carbonic anhydrase identified in the genome of the Gram-negative bacterium. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1060-1068	5.6	54
1227	Use of an immobilised thermostable -CA (SspCA) for enhancing the metabolic efficiency of the freshwater green microalga. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 913-920	5.6	8
1226	The Carbonic Anhydrase IX inhibitor SLC-0111 as emerging agent against the mesenchymal stem cell-derived pro-survival effects on melanoma cells. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1185-1193	5.6	15
1225	Effect of Carbonic Anhydrase IX inhibitors on human endothelial cell survival. <i>Pharmacological Research</i> , 2020 , 159, 104964	10.2	4
1224	Synthesis of calix[4]azacrown substituted sulphonamides with antioxidant, acetylcholinesterase, butyrylcholinesterase, tyrosinase and carbonic anhydrase inhibitory action. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1215-1223	5.6	13
1223	Development of oxathiino[6,5-b]pyridine 2,2-dioxide derivatives as selective inhibitors of tumor-related carbonic anhydrases IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2020 , 200, 112300	6.8	11
1222	Iodoquinazolinones bearing benzenesulfonamide as human carbonic anhydrase I, II, IX and XII inhibitors: Synthesis, biological evaluation and radiosensitizing activity. <i>European Journal of Medicinal Chemistry</i> , 2020 , 200, 112449	6.8	8
1221	Synthesis and carbonic anhydrase activating properties of a series of 2-amino-imidazolines structurally related to clonidine. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1003-1010	5.6	3

1220	Carbonic anhydrase modulation of emotional memory. Implications for the treatment of cognitive disorders. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1206-1214	5.6	31
1219	Crystal Structure of a Tetrameric Type II β -Carbonic Anhydrase from the Pathogenic Bacterium. <i>Molecules</i> , 2020 , 25,	4.8	5
1218	Sulfonamide Inhibitors of Human Carbonic Anhydrases Designed through a Three-Tails Approach: Improving Ligand/Isoform Matching and Selectivity of Action. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 7422-7444	8.3	35
1217	The Effect of Substituted Benzene-Sulfonamides and Clinically Licensed Drugs on the Catalytic Activity of CynT2, a Carbonic Anhydrase Crucial for Life Cycle. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	12
1216	The role of carbonic anhydrases in extinction of contextual fear memory. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 16000-16008	11.5	18
1215	New coumarin/sulfocoumarin linked phenylacrylamides as selective transmembrane carbonic anhydrase inhibitors: Synthesis and in-vitro biological evaluation. <i>Bioorganic and Medicinal Chemistry</i> , 2020 , 28, 115586	3.4	8
1214	Synthetic Strategies and Computational Inhibition Activity Study for Triazinyl-Substituted Benzenesulfonamide Conjugates with Polar and Hydrophobic Amino Acids as Inhibitors of Carbonic Anhydrases. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	5
1213	Carbonic anhydrase from extremophiles and their potential use in biotechnological applications 2020 , 295-306		1
1212	Biochemical and structural characterisation of a protozoan beta-carbonic anhydrase from. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1292-1299	5.6	18
1211	An update on drug interaction considerations in the therapeutic use of carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , 2020 , 16, 297-307	5.5	22
1210	Appliance of the piperidinyl-hydrazidoureido linker to benzenesulfonamide compounds: Synthesis, in vitro and in silico evaluation of potent carbonic anhydrase II, IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 98, 103728	5.1	10
1209	Activation Effects of Carnosine- and Histidine-Containing Dipeptides on Human Carbonic Anhydrases: A Comprehensive Study. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	10
1208	Anion Inhibition Studies of the β -Class Carbonic Anhydrase CAS3 from the Filamentous Ascomycete. <i>Metabolites</i> , 2020 , 10,	5.6	3
1207	Tail approach synthesis of novel benzenesulfonamides incorporating 1,3,4-oxadiazole hybrids as potent inhibitor of carbonic anhydrase I, II, IX, and XII isoenzymes. <i>European Journal of Medicinal Chemistry</i> , 2020 , 193, 112219	6.8	13
1206	Design, synthesis and biological evaluation of coumarin linked 1,2,4-oxadiazoles as selective carbonic anhydrase IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 98, 103739	5.1	11
1205	Sulfonamide Inhibition Studies of an β -Carbonic Anhydrase from , a Platyhelminth Parasite Responsible for Schistosomiasis. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	13
1204	Exploring the multiple binding modes of inhibitors to carbonic anhydrases for novel drug discovery. <i>Expert Opinion on Drug Discovery</i> , 2020 , 15, 671-686	6.2	61
1203	Tellurides Bearing Sulfonamides as Novel Inhibitors of Leishmanial Carbonic Anhydrase with Potent Antileishmanial Activity. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 4306-4314	8.3	17

1202	Looking toward the Rim of the Active Site Cavity of Druggable Human Carbonic Anhydrase Isoforms. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 1000-1005	4.3	1
1201	Anion Inhibition Studies of the Beta-Carbonic Anhydrase from. <i>Molecules</i> , 2020 , 25,	4.8	10
1200	Synthesis, computational studies and assessment of inhibitory activity of umbelliferon-based compounds against tumour-associated carbonic anhydrase isoforms IX and XII. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1442-1449	5.6	3
1199	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	2.7	82
1198	Structural and biochemical characterization of novel carbonic anhydrases from <i>Phaeodactylum tricornutum</i> . <i>Acta Crystallographica Section D: Structural Biology</i> , 2020 , 76, 676-686	5.5	6
1197	Perfluoroalkyl Substances of Significant Environmental Concern Can Strongly Inhibit Human Carbonic Anhydrase Isozymes. <i>Analytical Chemistry</i> , 2020 , 92, 4614-4622	7.8	16
1196	7-Acylamino-3H-1,2-benzoxathiepine 2,2-dioxides as new isoform-selective carbonic anhydrase IX and XII inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 650-656	5.6	9
1195	New Dihydrothiazole Benzenesulfonamides: Looking for Selectivity toward Carbonic Anhydrase Isoforms I, II, IX, and XII. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 852-856	4.3	2
1194	Sulfonamide Inhibition Studies of the β -Class Carbonic Anhydrase CAS3 from the Filamentous Ascomycete. <i>Molecules</i> , 2020 , 25,	4.8	3
1193	Bioorganometallic derivatives of 4-hydrazino-benzenesulphonamide as carbonic anhydrase inhibitors: synthesis, characterisation and biological evaluation. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 622-628	5.6	6
1192	1,3-Dipolar Cycloaddition, HPLC Enantioseparation, and Docking Studies of Saccharin/Isoxazole and Saccharin/Isoxazoline Derivatives as Selective Carbonic Anhydrase IX and XII Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 2470-2488	8.3	24
1191	A class of carbonic anhydrase IX/XII - selective carboxylate inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 549-554	5.6	2
1190	Evaluation of Thio- and Seleno-Acetamides Bearing Benzenesulfonamide as Inhibitor of Carbonic Anhydrases from Different Pathogenic Bacteria. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	9
1189	Coumarins from as inhibitors of the tumour-associated carbonic anhydrases IX and XII: isolation, biological studies and in silico evaluation. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 539-548	5.6	14
1188	Synthesis of some N-aroyle-2-oxindole benzenesulfonamide conjugates with carbonic anhydrase inhibitory activity. <i>Bioorganic Chemistry</i> , 2020 , 96, 103635	5.1	10
1187	New thiopyrimidine-benzenesulfonamide conjugates as selective carbonic anhydrase II inhibitors: synthesis, in vitro biological evaluation, and molecular docking studies. <i>Bioorganic and Medicinal Chemistry</i> , 2020 , 28, 115329	3.4	8
1186	Sulfonamide Inhibition Profile of the β -Carbonic Anhydrase from , An Opportunistic Pathogen Triggering Scalp Conditions. <i>Metabolites</i> , 2020 , 10,	5.6	9
1185	Pharmacological Inhibition of CA-IX Impairs Tumor Cell Proliferation, Migration and Invasiveness. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	14

1184	Benzylaminoethureido-Tailed Benzenesulfonamides: Design, Synthesis, Kinetic and X-ray Investigations on Human Carbonic Anhydrases. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	10
1183	Synthesis and carbonic anhydrase inhibition studies of sulfonamide based indole-1,2,3-triazole chalcone hybrids. <i>Bioorganic Chemistry</i> , 2020 , 99, 103839	5.1	14
1182	1,2,4-Triazole-based anticonvulsant agents with additional ROS scavenging activity are effective in a model of pharmacoresistant epilepsy. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 993-1002	5.6	11
1181	Aryl-4,5-dihydro-1-pyrazole-1-carboxamide Derivatives Bearing a Sulfonamide Moiety Show Single-digit Nanomolar-to-Subnanomolar Inhibition Constants against the Tumor-associated Human Carbonic Anhydrases IX and XII. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	3
1180	Dual P-Glycoprotein and CA XII Inhibitors: A New Strategy to Reverse the P-gp Mediated Multidrug Resistance (MDR) in Cancer Cells. <i>Molecules</i> , 2020 , 25,	4.8	17
1179	Development of Thiazolidinones as Fungal Carbonic Anhydrase Inhibitors. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	11
1178	Experimental Carbonic Anhydrase Inhibitors for the Treatment of Hypoxic Tumors. <i>Journal of Experimental Pharmacology</i> , 2020 , 12, 603-617	3	41
1177	Carbonic Anhydrase 2020 , 77-90		
1176	Expanding the anticancer potential of 1,2,3-triazoles via simultaneously targeting Cyclooxygenase-2, 15-lipoxygenase and tumor-associated carbonic anhydrases. <i>European Journal of Medicinal Chemistry</i> , 2020 , 200, 112439	6.8	15
1175	Design, synthesis, inhibition and toxicological evaluation of human carbonic anhydrases I, II and IX inhibitors in 5-nitroimidazole series. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 109-117	5.6	12
1174	Phosphonamidates are the first phosphorus-based zinc binding motif to show inhibition of Eclass carbonic anhydrases from bacteria, fungi, and protozoa. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 59-64	5.6	7
1173	Direct and straightforward access to substituted alkyl selenols as novel carbonic anhydrase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020 , 185, 111811	6.8	19
1172	Synthesis, biological and molecular dynamics investigations with a series of triazolopyrimidine/triazole-based benzenesulfonamides as novel carbonic anhydrase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020 , 185, 111843	6.8	21
1171	Bioisosteric Development of Multitarget Nonsteroidal Anti-Inflammatory Drug-Carbonic Anhydrases Inhibitor Hybrids for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 2325-2342	8.3	14
1170	Why hasn't there been more progress in new Chagas disease drug discovery?. <i>Expert Opinion on Drug Discovery</i> , 2020 , 15, 145-158	6.2	29
1169	Inhibition survey with phenolic compounds against the Eand Eclass carbonic anhydrases from the marine diatom and protozoan. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 377-382	5.6	6
1168	Synthesis and human carbonic anhydrase I, II, VA, and XII inhibition with novel amino acid-sulphonamide conjugates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 489-497	5.6	6
1167	Synthesis and selective inhibitory effects of some 2-oxindole benzenesulfonamide conjugates on human carbonic anhydrase isoforms CA I, CA II, CA IX and CAXII. <i>Bioorganic Chemistry</i> , 2020 , 95, 103514	5.1	8

1166	Benzimidazole derivatives as potent and isoform selective tumor-associated carbonic anhydrase IX/XII inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 95, 103544	5.1	11
1165	Pyrrolo and pyrrolopyrimidine sulfonamides act as cytotoxic agents in hypoxia via inhibition of transmembrane carbonic anhydrases. <i>European Journal of Medicinal Chemistry</i> , 2020 , 188, 112021	6.8	9
1164	inhibition of -carbonic anhydrase 3 with Mono- and dithiocarbamates and evaluation of their toxicity using zebrafish developing embryos. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 65-71	5.6	11
1163	Carbonic anhydrase IX as a novel candidate in liquid biopsy. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 255-260	5.6	11
1162	Novel benzofuran-based sulphonamides as selective carbonic anhydrases IX and XII inhibitors: synthesis and biological evaluation. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 298-305	5.6	13
1161	Design, synthesis and molecular modelling studies of some pyrazole derivatives as carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 289-297	5.6	23
1160	Novel sulphonamides incorporating triazene moieties show powerful carbonic anhydrase I and II inhibitory properties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 325-329	5.6	18
1159	Aryl derivatives of 3H-1,2-benzoxathiepine 2,2-dioxide as carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 245-254	5.6	11
1158	Plasmatic exosomes from prostate cancer patients show increased carbonic anhydrase IX expression and activity and low pH. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 280-288	5.6	30
1157	Benzothiazole derivatives as anticancer agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 265-279	5.6	60
1156	Sulfonamide/sulfamate switch with a series of piperazinyllureido derivatives: Synthesis, kinetic and in silico evaluation as carbonic anhydrase isoforms I, II, IV, and IX inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020 , 186, 111896	6.8	11
1155	Effects of New NSAID-CAI Hybrid Compounds in Inflammation and Lung Fibrosis. <i>Biomolecules</i> , 2020 , 10,	5.9	6
1154	Synthesis and biological evaluation of some coumarin hybrids as selective carbonic anhydrase IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 104, 104272	5.1	12
1153	3-Methylthiazolo[3,2-a]benzimidazole-benzenesulfonamide conjugates as novel carbonic anhydrase inhibitors endowed with anticancer activity: Design, synthesis, biological and molecular modeling studies. <i>European Journal of Medicinal Chemistry</i> , 2020 , 207, 112745	6.8	24
1152	Structural Basis of Nanomolar Inhibition of Tumor-Associated Carbonic Anhydrase IX: X-Ray Crystallographic and Inhibition Study of Lipophilic Inhibitors with Acetazolamide Backbone. <i>Journal of Medicinal Chemistry</i> , 2020 , 63, 13064-13075	8.3	11
1151	Inhibition of the newly discovered β -carbonic anhydrase from the protozoan pathogen <i>Trichomonas vaginalis</i> with inorganic anions and small molecules. <i>Journal of Inorganic Biochemistry</i> , 2020 , 213, 111274	4.2	8
1150	Sulphonamide inhibition profile of β -carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1834-1839	5.6	8
1149	Novel insights on saccharin- and acesulfame-based carbonic anhydrase inhibitors: design, synthesis, modelling investigations and biological activity evaluation. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1891-1905	5.6	7

1148	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	14.4	73
1147	Activation studies of the β -carbonic anhydrases from with amino acids and amines. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1379-1386	5.6	8
1146	Chagas Disease: Perspectives on the Past and Present and Challenges in Drug Discovery. <i>Molecules</i> , 2020 , 25,	4.8	13
1145	Nontargeted Identification of Plasma Proteins O-, N-, and S-Transmethylated by O-Methyl Organophosphates. <i>Analytical Chemistry</i> , 2020 , 92, 15420-15428	7.8	2
1144	Inhibitory activity against carbonic anhydrase IX and XII as a candidate selection criterion in the development of new anticancer agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1555-1561	5.6	16
1143	Synthesis, Computational Studies and Assessment of in Vitro Activity of Squalene Derivatives as Carbonic Anhydrase Inhibitors. <i>ChemMedChem</i> , 2020 , 15, 2052-2057	3.7	2
1142	Coumarin carbonic anhydrase inhibitors from natural sources. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1462-1470	5.6	30
1141	A measurement system for the evaluation of efficiency of enzyme accelerated CO ₂ capture systems based on modeling 2020 ,		1
1140	Carbonic Anhydrase Inhibitors Targeting Metabolism and Tumor Microenvironment. <i>Metabolites</i> , 2020 , 10,	5.6	70
1139	In Silico-Guided Identification of New Potent Inhibitors of Carbonic Anhydrases Expressed in. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 2294-2299	4.3	6
1138	Catechols: a new class of carbonic anhydrase inhibitors. <i>Chemical Communications</i> , 2020 , 56, 13033-13036	5.8	13
1137	Benzylaminoethylureido-Tailed Benzenesulfonamides Show Potent Inhibitory Activity against Bacterial Carbonic Anhydrases. <i>ChemMedChem</i> , 2020 , 15, 2444-2447	3.7	4
1136	Toxicity evaluation of sulfamides and coumarins that efficiently inhibit human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1765-1772	5.6	4
1135	-carbonic anhydrase: characterisation and effects of simple aromatic/heterocyclic sulphonamide inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1545-1554	5.6	18
1134	Benzoxaboroles: New Potent Inhibitors of the Carbonic Anhydrases of the Pathogenic Bacterium. <i>ACS Medicinal Chemistry Letters</i> , 2020 , 11, 2277-2284	4.3	9
1133	Antibacterial carbonic anhydrase inhibitors: an update on the recent literature. <i>Expert Opinion on Therapeutic Patents</i> , 2020 , 30, 963-982	6.8	36
1132	A structure-based approach towards the identification of novel antichagasic compounds: carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 21-30	5.6	8
1131	Activation studies of the β -carbonic anhydrases from with amines and amino acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 824-830	5.6	4

1130	Benzoxepinones: A new isoform-selective class of tumor associated carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2020 , 28, 115496	3.4	14
1129	Synthesis, characterisation, biological evaluation and studies of sulphonamide Schiff bases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 950-962	5.6	41
1128	Preparation, carbonic anhydrase enzyme inhibition and antioxidant activity of novel 7-amino-3,4-dihydroquinolin-2(1H)-one derivatives incorporating mono or dipeptide moiety. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1021-1026	5.6	5
1127	The antibiotic furagin and its derivatives are isoform-selective human carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020 , 35, 1011-1020	5.6	13
1126	Antibacterial activity of ethoxzolamide against strains SS1 and 26695. <i>Gut Pathogens</i> , 2020 , 12, 20	5.4	15
1125	Synthesis and Biological Evaluation of Imidazo[2,1-b]Thiazole based Sulfonyl Piperazines as Novel Carbonic Anhydrase II Inhibitors. <i>Metabolites</i> , 2020 , 10,	5.6	9
1124	Drug Screening in Human Cells by NMR Spectroscopy Allows the Early Assessment of Drug Potency. <i>Angewandte Chemie</i> , 2020 , 132, 6597-6601	3.6	4
1123	Drug Screening in Human Cells by NMR Spectroscopy Allows the Early Assessment of Drug Potency. <i>Angewandte Chemie - International Edition</i> , 2020 , 59, 6535-6539	16.4	25
1122	Dihydropteroate Synthase (Sulfonamides) and Dihydrofolate Reductase Inhibitors 2019 , 163-172		1
1121	Seeking new approach for therapeutic treatment of cholera disease via inhibition of bacterial carbonic anhydrases: experimental and theoretical studies for sixteen benzenesulfonamide derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1186-1192	5.6	7
1120	Exploration of the residues modulating the catalytic features of human carbonic anhydrase XIII by a site-specific mutagenesis approach. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1506-1510	5.6	6
1119	Novel method of treating macular degeneration: a patent evaluation (WO2018/107005). <i>Expert Opinion on Therapeutic Patents</i> , 2019 , 29, 749-752	6.8	3
1118	Crystal structure and chemical inhibition of essential schistosome host-interactive virulence factor carbonic anhydrase SmCA. <i>Communications Biology</i> , 2019 , 2, 333	6.7	19
1117	Anti- activity of ethoxzolamide. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1660-1663	3.6	23
1116	Power architectures for the integration of photovoltaic generation systems in DC-microgrids. <i>Energy Procedia</i> , 2019 , 159, 34-41	2.3	4
1115	Design of a Hybrid Propulsion Architecture for Midsized Boats. <i>Energy Procedia</i> , 2019 , 158, 2954-2959	2.3	7
1114	Continued exploration and tail approach synthesis of benzenesulfonamides containing triazole and dual triazole moieties as carbonic anhydrase I, II, IV and IX inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 183, 111698	6.8	21
1113	Exploring new structural features of the 4-[(3-methyl-4-aryl-2,3-dihydro-1,3-thiazol-2-ylidene)amino]benzenesulphonamide scaffold for the inhibition of human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1524-1533	5.6	5

1112	Synthesis and exploration of 2-morpholino-4-phenylthiazol-5-yl acrylamide derivatives for their effects against carbonic anhydrase I, II, IX and XII isoforms as a non-sulfonamide class of inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 115090	3.4	6
1111	Synthesis of coumarin-sulfonamide derivatives and determination of their cytotoxicity, carbonic anhydrase inhibitory and molecular docking studies. <i>European Journal of Medicinal Chemistry</i> , 2019 , 183, 111702	6.8	32
1110	The management of glaucoma and macular degeneration. <i>Expert Opinion on Therapeutic Patents</i> , 2019 , 29, 745-747	6.8	23
1109	A New Kid on the Block? Carbonic Anhydrases as Possible New Targets in Alzheimer's Disease. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	34
1108	Activation of α and β class of carbonic anhydrases with amines and amino acids: a review. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1652-1659	5.6	18
1107	β -Carbonic anhydrases are strongly activated by spinaceamine derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 800-804	3.4	16
1106	Inhibition of acetylcholinesterase and butyrylcholinesterase with uracil derivatives: kinetic and computational studies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 429-437	5.6	41
1105	Phaeodactylum tricornutum as a model organism for testing the membrane penetrability of sulphonamide carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 510-518	5.6	13
1104	Click-tailed benzenesulfonamides as potent bacterial carbonic anhydrase inhibitors for targeting Mycobacterium tuberculosis and Vibrio cholerae. <i>Bioorganic Chemistry</i> , 2019 , 86, 183-186	5.1	9
1103	Novel approaches for designing drugs that interfere with pH regulation. <i>Expert Opinion on Drug Discovery</i> , 2019 , 14, 231-248	6.2	23
1102	Discovery of new organoselenium compounds as antileishmanial agents. <i>Bioorganic Chemistry</i> , 2019 , 86, 339-345	5.1	23
1101	Design, synthesis and biological evaluation of coumarin-3-carboxamides as selective carbonic anhydrase IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 86, 386-392	5.1	20
1100	Benzenesulfonamides incorporating nitrogenous bases show effective inhibition of β -carbonic anhydrases from the pathogenic fungi Cryptococcus neoformans, Candida glabrata and Malassezia globosa. <i>Bioorganic Chemistry</i> , 2019 , 86, 39-43	5.1	4
1099	Selenols: a new class of carbonic anhydrase inhibitors. <i>Chemical Communications</i> , 2019 , 55, 648-651	5.8	42
1098	Novel Diamide-Based Benzenesulfonamides as Selective Carbonic Anhydrase IX Inhibitors Endowed with Antitumor Activity: Synthesis, Biological Evaluation and In Silico Insights. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	16
1097	Appraisal of anti-protozoan activity of nitroaromatic benzenesulfonamides inhibiting carbonic anhydrases from and. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1164-1171	5.6	14
1096	Synthesis and biological evaluation of novel 8-substituted quinoline-2-carboxamides as carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1172-1177	5.6	11
1095	Synthesis of a new series of 3-functionalised-1-phenyl-1,2,3-triazole sulfamoylbenzamides as carbonic anhydrase I, II, IV and IX inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1199-1209	5.6	7

1094	Ring opening reactions of heterocycles with selenium and tellurium nucleophiles. <i>New Journal of Chemistry</i> , 2019 , 43, 11451-11468	3.6	29
1093	Activation of human α -carbonic anhydrase isoforms I, II, IV and VII with bis-histamine schiff bases and bis-spinaceamine substituted derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1193-1198	5.6	13
1092	New phenolic Mannich bases with piperazines and their bioactivities. <i>Bioorganic Chemistry</i> , 2019 , 90, 103057	5.1	34
1091	arginase: biochemical characterization and inhibition by naturally occurring phenolic substances. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1100-1109	5.6	15
1090	Cloning, Purification, and Characterization of a α -Carbonic Anhydrase from , an Opportunistic Pathogen Involved in Dandruff and Seborrheic Dermatitis. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	17
1089	Syntesis of thio- and seleno-acetamides bearing benzenesulfonamide as potent inhibitors of human carbonic anhydrase II and XII. <i>Bioorganic Chemistry</i> , 2019 , 89, 102984	5.1	11
1088	Indole-Based Hydrazones Containing A Sulfonamide Moiety as Selective Inhibitors of Tumor-Associated Human Carbonic Anhydrase Isoforms IX and XII. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	9
1087	A non-catalytic function of carbonic anhydrase IX contributes to the glycolytic phenotype and pH regulation in human breast cancer cells. <i>Biochemical Journal</i> , 2019 , 476, 1497-1513	3.8	17
1086	Fibrate-based N-acylsulphonamides targeting carbonic anhydrases: synthesis, biochemical evaluation, and docking studies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1051-1061	5.6	12
1085	Pain Relieving Effect of-NSAIDs-CAIs Hybrid Molecules: Systemic and Intra-Articular Treatments against Rheumatoid Arthritis. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	17
1084	Extrinsic acidosis suppresses glycolysis and migration while increasing network formation in pulmonary microvascular endothelial cells. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2019 , 317, L188-L201	5.8	12
1083	Thermostability enhancement of the α -carbonic anhydrase from Sulfurihydrogenibium yellowstonense by using the anchoring-and-self-labelling-protein-tag system (ASL). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 946-954	5.6	8
1082	Direct biocatalysed synthesis of first sulfur-, selenium- and tellurium- containing l-ascorbyl hybrid derivatives with radical trapping and GPx-like properties. <i>Chemical Communications</i> , 2019 , 55, 5705-5708	5.8	35
1081	Activation Studies of the α -Carbonic Anhydrases from the Antarctic Marine Bacteria and with Amino Acids and Amines. <i>Marine Drugs</i> , 2019 , 17,	6	7
1080	Carbonic Anhydrase Inhibitor-NO Donor Hybrids and Their Pharmacological Applications 2019 , 229-242		6
1079	Synthesis of benzenesulfonamides linked to quinazoline scaffolds as novel carbonic anhydrase inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 87, 78-90	5.1	25
1078	Synthesis, biological activity and multiscale molecular modeling studies of bis-coumarins as selective carbonic anhydrase IX and XII inhibitors with effective cytotoxicity against hepatocellular carcinoma. <i>Bioorganic Chemistry</i> , 2019 , 87, 838-850	5.1	32
1077	Comparison of the Sulfonamide Inhibition Profiles of the α -Carbonic Anhydrase Isoforms (SpiCA1, SpiCA2 and SpiCA3) Encoded by the Genome of the Scleractinian Coral. <i>Marine Drugs</i> , 2019 , 17,	6	3

1076	Novel 2-indolinones containing a sulfonamide moiety as selective inhibitors of candida β -carbonic anhydrase enzyme. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 528-531	5.6	9
1075	Novel 8-Substituted Coumarins That Selectively Inhibit Human Carbonic Anhydrase IX and XII. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	14
1074	Seleno-Michael Reaction of Stable Functionalised Alkyl Selenols: A Versatile Tool for the Synthesis of Acyclic and Cyclic Unsymmetrical Alkyl and Vinyl Selenides. <i>Advanced Synthesis and Catalysis</i> , 2019 , 361, 2337-2346	5.6	26
1073	Sulfur, selenium and tellurium containing amines act as effective carbonic anhydrase activators. <i>Bioorganic Chemistry</i> , 2019 , 87, 516-522	5.1	29
1072	Novel synthesized SLC-0111 thiazole and thiadiazole analogues: Determination of their carbonic anhydrase inhibitory activity and molecular modeling studies. <i>Bioorganic Chemistry</i> , 2019 , 87, 794-802	5.1	28
1071	Synthesis and biological evaluation of coumarin-1,3,4-oxadiazole hybrids as selective carbonic anhydrase IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 87, 765-772	5.1	26
1070	The Possible Role of in Gastric Cancer and Its Management. <i>Frontiers in Oncology</i> , 2019 , 9, 75	5.3	39
1069	Activation Studies of the β -Carbonic Anhydrase from the Pathogenic Protozoan with Amino Acids and Amines. <i>Metabolites</i> , 2019 , 9,	5.6	7
1068	An AGT-based protein-tag system for the labelling and surface immobilization of enzymes on E. coli outer membrane. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 490-499	5.6	11
1067	Pyridazinone-substituted benzenesulfonamides display potent inhibition of membrane-bound human carbonic anhydrase IX and promising antiproliferative activity against cancer cell lines. <i>European Journal of Medicinal Chemistry</i> , 2019 , 168, 301-314	6.8	13
1066	Design, synthesis and biological activity of selective hCAs inhibitors based on 2-(benzylsulfinyl)benzoic acid scaffold. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1400-1413	5.6	14
1065	Synthesis and biological evaluation of novel 3-(quinolin-4-ylamino)benzenesulfonamides AQ3 as carbonic anhydrase isoforms I and II inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1457-1464	5.6	20
1064	Advances in the structural annotation of human carbonic anhydrases and impact on future drug discovery. <i>Expert Opinion on Drug Discovery</i> , 2019 , 14, 1175-1197	6.2	96
1063	Carbonic anhydrase inhibitors as diuretics 2019 , 287-309		
1062	Carbonic anhydrases from pathogens 2019 , 387-417		
1061	Carbonic anhydrase activators and their potential in the pharmaceutical field 2019 , 477-492		
1060	Mechanism of action of carbonic anhydrase inhibitors 2019 , 245-255		1
1059	Biotechnologic applications of carbonic anhydrases from extremophiles 2019 , 495-514		

1058	New anthranilic acid-incorporating N-benzenesulfonamidophthalimides as potent inhibitors of carbonic anhydrases I, II, IX, and XII: Synthesis, in vitro testing, and in silico assessment. <i>European Journal of Medicinal Chemistry</i> , 2019 , 181, 111573	6.8	11
1057	Structure-activity relationship with pyrazoline-based aromatic sulfamates as carbonic anhydrase isoforms I, II, IX and XII inhibitors: Synthesis and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2019 , 182, 111638	6.8	15
1056	Sulfonamides incorporating piperazine bioisosteres as potent human carbonic anhydrase I, II, IV and IX inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 91, 103130	5.1	6
1055	Bis-benzoxaboroles: Design, Synthesis, and Biological Evaluation as Carbonic Anhydrase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2019 , 10, 1205-1210	4.3	9
1054	Synthesis and Evaluation of Carbonic Anhydrase Inhibitors with Carbon Monoxide Releasing Properties for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2019 , 62, 7233-7249	8.3	26
1053	Anion Inhibition Profile of the β -Carbonic Anhydrase from the Opportunist Pathogenic Fungus Involved in Dandruff and Seborrheic Dermatitis. <i>Metabolites</i> , 2019 , 9,	5.6	10
1052	The first activation study of the β -carbonic anhydrases from the pathogenic bacteria and with amines and amino acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1178-1185	5.6	6
1051	A computer-assisted discovery of novel potential anti-obesity compounds as selective carbonic anhydrase VA inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 181, 111565	6.8	17
1050	Carbonic anhydrase inhibitors for the treatment of neuropathic pain and arthritis 2019 , 367-386		0
1049	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	13.3	91
1048	β -Carbonic anhydrases 2019 , 107-129		1
1047	Human carbonic anhydrases 2019 , 151-185		12
1046	Synthesis, biological evaluation and in silico studies with 4-benzylidene-2-phenyl-5(4H)-imidazolone-based benzenesulfonamides as novel selective carbonic anhydrase IX inhibitors endowed with anticancer activity. <i>Bioorganic Chemistry</i> , 2019 , 90, 103102	5.1	17
1045	-Carbonic Anhydrases: Novel Targets for Developing Antituberculosis Drugs. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	16
1044	<i>Pseudomonas aeruginosa</i> β -carbonic anhydrase, psCA1, is required for calcium deposition and contributes to virulence. <i>Cell Calcium</i> , 2019 , 84, 102080	4	16
1043	Extending the Inhibition Profiles of Coumarin-Based Compounds Against Human Carbonic Anhydrases: Synthesis, Biological, and In Silico Evaluation. <i>Molecules</i> , 2019 , 24,	4.8	3
1042	Carbonic anhydrase inhibition and the management of glaucoma: a literature and patent review 2013-2019. <i>Expert Opinion on Therapeutic Patents</i> , 2019 , 29, 781-792	6.8	44
1041	Extending the β -class carbonic anhydrases inhibition profiles with phenolic compounds. <i>Bioorganic Chemistry</i> , 2019 , 93, 103336	5.1	7

1040	Carbonic anhydrases 2019 , 3-16		11
1039	Multivalent Carbonic Anhydrases Inhibitors. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	11
1038	State of the Art on Carbonic Anhydrase Modulators for Biomedical Purposes. <i>Current Medicinal Chemistry</i> , 2019 , 26, 2558-2573	4.3	9
1037	Investigation of inhibitory properties of some hydrazone compounds on hCA I, hCA II and AChE enzymes. <i>Bioorganic Chemistry</i> , 2019 , 86, 316-321	5.1	80
1036	Exploring benzoxaborole derivatives as carbonic anhydrase inhibitors: a structural and computational analysis reveals their conformational variability as a tool to increase enzyme selectivity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 1498-1505	5.6	5
1035	Discovery of new ureido benzenesulfonamides incorporating 1,3,5-triazine moieties as carbonic anhydrase I, II, IX and XII inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 1588-1594	3.4	32
1034	Inhibition of β and γ class carbonic anhydrases from bacteria, fungi, algae, diatoms and protozoans with famotidine. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 644-650	5.6	30
1033	Continued exploration of 1,2,4-oxadiazole periphery for carbonic anhydrase-targeting primary arene sulfonamides: Discovery of subnanomolar inhibitors of membrane-bound hCA IX isoform that selectively kill cancer cells in hypoxic environment. <i>European Journal of Medicinal Chemistry</i> , 2019 , 184, 92-105	6.8	41
1032	Inhibition of bacterial β and γ class carbonic anhydrases with selenazoles incorporating benzenesulfonamide moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 244-249	5.6	12
1031	Comparison of blood carbonic anhydrase activity of athletes performing interval and continuous running exercise at high altitude. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 218-224	5.6	6
1030	Synthesis carbonic anhydrase enzyme inhibition and antioxidant activity of novel benzothiazole derivatives incorporating glycine, methionine, alanine, and phenylalanine moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 343-349	5.6	13
1029	SLC-0111 enamionone analogs, 3/4-(3-aryl-3-oxopropenyl) aminobenzenesulfonamides, as novel selective subnanomolar inhibitors of the tumor-associated carbonic anhydrase isoform IX. <i>Bioorganic Chemistry</i> , 2019 , 83, 549-558	5.1	40
1028	Synthesis and anti-inflammatory activity of sulfonamides and carboxylates incorporating trimellitimides: Dual cyclooxygenase/carbonic anhydrase inhibitory actions. <i>Bioorganic Chemistry</i> , 2019 , 84, 260-268	5.1	30
1027	Identification and characterization of the hCA in the outer membrane vesicles produced by <i>Helicobacter pylori</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 189-195	5.6	24
1026	Synthesis of novel benzenesulfonamide bearing 1,2,3-triazole linked hydroxy-trifluoromethylpyrazolines and hydrazones as selective carbonic anhydrase isoforms IX and XII inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 85, 198-208	5.1	16
1025	Prostate cancer cells and exosomes in acidic condition show increased carbonic anhydrase IX expression and activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 272-278	5.6	40
1024	(Hetero)aryl substituted thiazol-2,4-yl scaffold as human carbonic anhydrase I, II, VII and XIV activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 224-229	5.6	7
1023	Nanoscale Ion Emitters in Native Mass Spectrometry for Measuring Ligand-Protein Binding Affinities. <i>ACS Central Science</i> , 2019 , 5, 308-318	16.8	59

1022	New sulfonamides containing organometallic-acylhydrazones: synthesis, characterisation and biological evaluation as inhibitors of human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 451-458	5.6	8
1021	4-Substituted benzenesulfonamides featuring cyclic imides moieties exhibit potent and isoform-selective carbonic anhydrase II/IX inhibition. <i>Bioorganic Chemistry</i> , 2019 , 83, 198-204	5.1	19
1020	The carbonic anhydrase IX inhibitor SLC-0111 sensitises cancer cells to conventional chemotherapy. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019 , 34, 117-123	5.6	52
1019	Synthesis and carbonic anhydrase inhibitory properties of novel 4-(2-aminoethyl)benzenesulfonamide-dipeptide conjugates. <i>Bioorganic Chemistry</i> , 2019 , 83, 414-423	5.1	14
1018	Enhancement of the tail hydrophobic interactions within the carbonic anhydrase IX active site via structural extension: Design and synthesis of novel N-substituted isatins-SLC-0111 hybrids as carbonic anhydrase inhibitors and antitumor agents. <i>European Journal of Medicinal Chemistry</i> , 2019 , 162, 147-160	6.8	63
1017	Exploring structural properties of potent human carbonic anhydrase inhibitors bearing a 4-(cycloalkylamino-1-carbonyl)benzenesulfonamide moiety. <i>European Journal of Medicinal Chemistry</i> , 2019 , 163, 443-452	6.8	16
1016	-Nitrosulfonamides as Carbonic Anhydrase Inhibitors: A Promising Chemotype for Targeting Chagas Disease and Leishmaniasis. <i>ACS Medicinal Chemistry Letters</i> , 2019 , 10, 413-418	4.3	14
1015	3-Aminobenzenesulfonamides incorporating acylthiourea moieties selectively inhibit the tumor-associated carbonic anhydrase isoform IX over the off-target isoforms I, II and IV. <i>Bioorganic Chemistry</i> , 2019 , 82, 123-128	5.1	4
1014	1,2,4-Trisubstituted imidazolinones with dual carbonic anhydrase and p38 mitogen-activated protein kinase inhibitory activity. <i>Bioorganic Chemistry</i> , 2019 , 82, 109-116	5.1	9
1013	Design, synthesis and biological evaluation of novel ureido benzenesulfonamides incorporating 1,3,5-triazine moieties as potent carbonic anhydrase IX inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 82, 117-122	5.1	32
1012	Phosphorus versus Sulfur: Discovery of Benzenephosphonamidates as Versatile Sulfonamide-Mimic Chemotypes Acting as Carbonic Anhydrase Inhibitors. <i>Chemistry - A European Journal</i> , 2019 , 25, 1188-1192	4.8	31
1011	Carbon- versus sulphur-based zinc binding groups for carbonic anhydrase inhibitors?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 485-495	5.6	86
1010	Carbonic anhydrase activators. <i>Future Medicinal Chemistry</i> , 2018 , 10, 561-573	4.1	86
1009	Synthesis of N'-phenyl-N-hydroxyureas and investigation of their inhibitory activities on human carbonic anhydrases. <i>Bioorganic Chemistry</i> , 2018 , 78, 1-6	5.1	9
1008	Synthesis and biological evaluation of novel pyrazoline-based aromatic sulfamates with potent carbonic anhydrase isoforms II, IV and IX inhibitory efficacy. <i>Bioorganic Chemistry</i> , 2018 , 77, 633-639	5.1	16
1007	Activation studies of the β and ϵ carbonic anhydrases from the pathogenic bacterium <i>Vibrio cholerae</i> with amines and amino acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 227-233	5.6	16
1006	pH regulators to target the tumor immune microenvironment in human hepatocellular carcinoma. <i>Oncotmunology</i> , 2018 , 7, e1445452	7.2	37
1005	Synthesis and Biological Evaluation of 4-Sulfamoylphenyl/Sulfocoumarin Carboxamides as Selective Inhibitors of Carbonic Anhydrase Isoforms hCA II, IX, and XII. <i>ChemMedChem</i> , 2018 , 13, 1165-1171	3.7	8

1004	Evaluation of Tc-sulfonamide and sulfocoumarin derivatives for imaging carbonic anhydrase IX expression. <i>Journal of Inorganic Biochemistry</i> , 2018 , 185, 63-70	4.2	18
1003	Activation studies with amines and amino acids of the β -carbonic anhydrase from the pathogenic protozoan <i>Leishmania donovani</i> chagasi. <i>Bioorganic Chemistry</i> , 2018 , 78, 406-410	5.1	17
1002	Deciphering the Mechanism of Human Carbonic Anhydrases Inhibition with Sulfocoumarins: Computational and Experimental Studies. <i>Chemistry - A European Journal</i> , 2018 , 24, 7840-7844	4.8	51
1001	Natural Polyphenols Selectively Inhibit β -Carbonic Anhydrase from the Dandruff-Producing Fungus <i>Malassezia globosa</i> : Activity and Modeling Studies. <i>ChemMedChem</i> , 2018 , 13, 816-823	3.7	24
1000	2-Benzylpiperazine: A new scaffold for potent human carbonic anhydrase inhibitors. Synthesis, enzyme inhibition, enantioselectivity, computational and crystallographic studies and in vivo activity for a new class of intraocular pressure lowering agents. <i>European Journal of Medicinal Chemistry</i> , 2018 , 151, 363-375	6.8	27
999	Design, Synthesis, and X-ray of Selenides as New Class of Agents for Prevention of Diabetic Cerebrovascular Pathology. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 462-467	4.3	17
998	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	14.4	159
997	Dual-tail arylsulfone-based benzenesulfonamides differently match the hydrophobic and hydrophilic halves of human carbonic anhydrases active sites: Selective inhibitors for the tumor-associated hCA IX isoform. <i>European Journal of Medicinal Chemistry</i> , 2018 , 152, 1-9	6.8	49
996	Design and synthesis of novel 1,3-diaryltriazene-substituted sulfonamides as potent and selective carbonic anhydrase II inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 77, 542-547	5.1	38
995	Anion inhibition studies of a beta carbonic anhydrase from the malaria mosquito <i>Anopheles gambiae</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 359-363	5.6	5
994	Discovery of thiazolin-4-one-based aromatic sulfamates as a new class of carbonic anhydrase isoforms I, II, IV, and IX inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 77, 293-299	5.1	20
993	Synthesis, biological evaluation and computational studies of novel iminothiazolidinone benzenesulfonamides as potent carbonic anhydrase II and IX inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 77, 381-386	5.1	21
992	A Remarkable Influence of a Trifluoromethyl Group on the Reactions of β -Mercaptoalcohols with Fluorinated β -Bromo enones. <i>European Journal of Organic Chemistry</i> , 2018 , 2018, 3716-3723	3.2	22
991	Sulfonamide inhibition studies of two β -carbonic anhydrases from the ascomycete fungus <i>Sordaria macrospora</i> , CAS1 and CAS2. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 390-396	5.6	9
990	New anticancer drug candidates sulfonamides as selective hCA IX or hCA XII inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 77, 411-419	5.1	67
989	Synthesis and biological evaluation of novel N,N'-diaryl cyanoguanidines acting as potent and selective carbonic anhydrase II inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 77, 245-251	5.1	29
988	Crystal Structure of Carbonic Anhydrase II in Complex with an Activating Ligand: Implications in Neuronal Function. <i>Molecular Neurobiology</i> , 2018 , 55, 7431-7437	6.2	19
987	Protonography and anion inhibition profile of the β -carbonic anhydrase (CruCA4) identified in the Mediterranean red coral <i>Corallium rubrum</i> . <i>Bioorganic Chemistry</i> , 2018 , 76, 281-287	5.1	10

986	Investigation of piperazines as human carbonic anhydrase I, II, IV and VII activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 303-308	5.6	5
985	Carbonic anhydrases from <i>Trypanosoma cruzi</i> and <i>Leishmania donovani</i> chagasi are inhibited by benzoxaboroles. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 286-289	5.6	42
984	Supercharging protein ions in native mass spectrometry using theta capillary nanoelectrospray ionization mass spectrometry and cyclic alkylcarbonates. <i>Analytica Chimica Acta</i> , 2018 , 1003, 1-9	6.6	16
983	Plasmatic carbonic anhydrase IX as a diagnostic marker for clear cell renal cell carcinoma. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 234-240	5.6	12
982	Inhibition studies of <i>Brucella suis</i> β -carbonic anhydrases with a series of 4-substituted pyridine-3-sulphonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 255-259	5.6	7
981	Structural investigations on coumarins leading to chromeno[4,3-c]pyrazol-4-ones and pyrano[4,3-c]pyrazol-4-ones: New scaffolds for the design of the tumor-associated carbonic anhydrase isoforms IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2018 , 146, 47-59	6.8	36
980	Activation studies with amines and amino acids of the β -carbonic anhydrase encoded by the Rv3273 gene from the pathogenic bacterium <i>Mycobacterium tuberculosis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 364-369	5.6	16
979	The β -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> is potently activated by amines and amino acids. <i>Bioorganic Chemistry</i> , 2018 , 77, 1-5	5.1	16
978	The first activation study of a β -carbonic anhydrase: TweCA from the diatom <i>Thalassiosira weissflogii</i> is effectively activated by amines and amino acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 680-685	5.6	16
977	Fluoroenesulphonamides: N-sulphonylurea isosteres showing nanomolar selective cancer-related transmembrane human carbonic anhydrase inhibition. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 804-808	5.6	8
976	Benzoxaborole compounds for therapeutic uses: a patent review (2010- 2018). <i>Expert Opinion on Therapeutic Patents</i> , 2018 , 28, 493-504	6.8	70
975	Antileishmanial activity of sulphonamide nanoemulsions targeting the β -carbonic anhydrase from <i>Leishmania</i> species. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 850-857	5.6	30
974	Anticancer effects of new dibenzenesulfonamides by inducing apoptosis and autophagy pathways and their carbonic anhydrase inhibitory effects on hCA I, hCA II, hCA IX, hCA XII isoenzymes. <i>Bioorganic Chemistry</i> , 2018 , 78, 290-297	5.1	33
973	Inhibition studies on a panel of human carbonic anhydrases with N1-substituted secondary sulfonamides incorporating thiazolinone or imidazolone-indole tails. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 629-638	5.6	27
972	Mono- and di-thiocarbamate inhibition studies of the β -carbonic anhydrase TweCA from the marine diatom <i>Thalassiosira weissflogii</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 707-713	5.6	15
971	Discovery of Benzenesulfonamide Derivatives as Carbonic Anhydrase Inhibitors with Effective Anticonvulsant Action: Design, Synthesis, and Pharmacological Evaluation. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 3151-3165	8.3	18
970	Synthesis of novel 4-functionalized 1,5-diaryl-1,2,3-triazoles containing benzenesulfonamide moiety as carbonic anhydrase I, II, IV and IX inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018 , 150, 678-686	6.8	30
969	Sulfonamide carbonic anhydrase inhibitors: Zinc coordination and tail effects influence inhibitory efficacy and selectivity for different isoforms. <i>Inorganica Chimica Acta</i> , 2018 , 470, 128-132	2.7	6

968	Heterocyclic periphery in the design of carbonic anhydrase inhibitors: 1,2,4-Oxadiazol-5-yl benzenesulfonamides as potent and selective inhibitors of cytosolic hCA II and membrane-bound hCA IX isoforms. <i>Bioorganic Chemistry</i> , 2018 , 76, 88-97	5.1	37
967	Novel carbonic anhydrase IX-targeted therapy enhances the anti-tumour effects of cisplatin in small cell lung cancer. <i>International Journal of Cancer</i> , 2018 , 142, 191-201	7.5	19
966	Comparison of the Anion Inhibition Profiles of the hCA Isoforms (SpiCA1, SpiCA2 and SpiCA3) from the Scleractinian Coral <i>Stylophora pistillata</i> . <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	7
965	Carbonic anhydrase inhibitors as antitumor/antimetastatic agents: a patent review (2008-2018). <i>Expert Opinion on Therapeutic Patents</i> , 2018 , 28, 729-740	6.8	121
964	Inhibition of α -glucosidase and α -carbonic anhydrases from bacteria and diatoms with N'-aryl-N-hydroxy-ureas. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1194-1198	5.6	16
963	Novel hydrazido benzenesulfonamides-isatin conjugates: Synthesis, carbonic anhydrase inhibitory activity and molecular modeling studies. <i>European Journal of Medicinal Chemistry</i> , 2018 , 157, 28-36	6.8	39
962	Synthesis, X-ray structure, in silico calculation, and carbonic anhydrase inhibitory properties of benzylimidazole metal complexes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1150-1159	5.6	5
961	Novel 6- and 7-Substituted Coumarins with Inhibitory Action against Lipoxygenase and Tumor-Associated Carbonic Anhydrase IX. <i>Molecules</i> , 2018 , 23,	4.8	20
960	Synthesis of novel isoindoline-1,3-dione-based oximes and benzenesulfonamide hydrazones as selective inhibitors of the tumor-associated carbonic anhydrase IX. <i>Bioorganic Chemistry</i> , 2018 , 80, 706-713	5.1	25
959	The Crystal Structure of a hCA VII Variant Provides Insights into the Molecular Determinants Responsible for Its Catalytic Behavior. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	19
958	Biomedical applications of prokaryotic carbonic anhydrases. <i>Expert Opinion on Therapeutic Patents</i> , 2018 , 28, 745-754	6.8	67
957	Development of a Fingerprint-Based Scoring Function for the Prediction of the Binding Mode of Carbonic Anhydrase II Inhibitors. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	10
956	Amino Acids as Building Blocks for Carbonic Anhydrase Inhibitors. <i>Metabolites</i> , 2018 , 8,	5.6	14
955	Benzamide-4-Sulfonamides Are Effective Human Carbonic Anhydrase I, II, VII, and IX Inhibitors. <i>Metabolites</i> , 2018 , 8,	5.6	10
954	Activation studies with amines and amino acids of the α -carbonic anhydrase from the pathogenic protozoan <i>Trypanosoma cruzi</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2018 , 26, 4187-4190	3.4	12
953	Discovery of potent anti-convulsant carbonic anhydrase inhibitors: Design, synthesis, in vitro and in vivo appraisal. <i>European Journal of Medicinal Chemistry</i> , 2018 , 156, 430-443	6.8	13
952	Synthesis of novel benzenesulfamide derivatives with inhibitory activity against human cytosolic carbonic anhydrase I and II and <i>Vibrio cholerae</i> α - and β -class enzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1125-1136	5.6	14
951	Discovery of Novel Nonsteroidal Anti-Inflammatory Drugs and Carbonic Anhydrase Inhibitors Hybrids (NSAIDs-CAIs) for the Management of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 4961-4977	8.3	37

950	Synthesis of novel dipeptide sulfonamide conjugates with effective carbonic anhydrase I, II, IX, and XII inhibitory properties. <i>Bioorganic Chemistry</i> , 2018 , 81, 311-318	5.1	13
949	Inhibition of carbonic anhydrases by a substrate analog: benzyl carbamate directly coordinates the catalytic zinc ion mimicking bicarbonate binding. <i>Chemical Communications</i> , 2018 , 54, 10312-10315	5.8	14
948	Structural Mapping of Anion Inhibitors to α -Carbonic Anhydrase psCA3 from <i>Pseudomonas aeruginosa</i> . <i>ChemMedChem</i> , 2018 , 13, 2024-2029	3.7	19
947	Design and synthesis of novel benzenesulfonamide containing 1,2,3-triazoles as potent human carbonic anhydrase isoforms I, II, IV and IX inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018 , 155, 545-551	6.8	26
946	Targeting Tumor Associated Carbonic Anhydrases IX and XII: Highly Isozyme Selective Coumarin and Psoralen Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 725-729	4.3	27
945	Nitroimidazole-based inhibitors DTP338 and DTP348 are safe for zebrafish embryos and efficiently inhibit the activity of human CA IX in <i>Xenopus</i> oocytes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1064-1073	5.6	13
944	Are Carbonic Anhydrases Suitable Targets to Fight Protozoan Parasitic Diseases?. <i>Current Medicinal Chemistry</i> , 2018 , 25, 5266-5278	4.3	18
943	Unprotected primary sulfonamide group facilitates ring-forming cascade en route to polycyclic [1,4]oxazepine-based carbonic anhydrase inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 76, 140-146	5.1	11
942	Inhibitory effects and structural insights for a novel series of coumarin-based compounds that selectively target human CA IX and CA XII carbonic anhydrases. <i>European Journal of Medicinal Chemistry</i> , 2018 , 143, 276-282	6.8	45
941	Nanoemulsions of sulfonamide carbonic anhydrase inhibitors strongly inhibit the growth of <i>Trypanosoma cruzi</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 139-146	5.6	41
940	Crystal structure of the human carbonic anhydrase II adduct with 1-(4-sulfamoylphenyl-ethyl)-2,4,6-triphenylpyridinium perchlorate, a membrane-impermeant, isoform selective inhibitor. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 151-157	5.6	24
939	First evaluation of organotellurium derivatives as carbonic anhydrase I, II, IV, VII and IX inhibitors. <i>Bioorganic Chemistry</i> , 2018 , 76, 268-272	5.1	29
938	Sulphonamide inhibition studies of the α -carbonic anhydrase from the bacterial pathogen <i>Clostridium perfringens</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 31-36	5.6	12
937	Exponential Activation of Carbonic Anhydrase by Encapsulation in Dynameric Host Matrices with Chiral Discrimination. <i>Chemistry - A European Journal</i> , 2018 , 24, 715-720	4.8	10
936	Comparison of the amine/amino acid activation profiles of the β and α -carbonic anhydrases from the pathogenic bacterium <i>Burkholderia pseudomallei</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 25-30	5.6	15
935	Characterization of technical grade carbonic anhydrase as biocatalyst for CO ₂ capture in potassium carbonate solutions 2018 , 8, 279-291		10
934	Improving the carbonic anhydrase inhibition profile of the sulfamoylphenyl pharmacophore by attachment of carbohydrate moieties. <i>Bioorganic Chemistry</i> , 2018 , 76, 61-66	5.1	9
933	Bioactive isoflavones from <i>Pueraria lobata</i> root and starch: Different extraction techniques and carbonic anhydrase inhibition. <i>Food and Chemical Toxicology</i> , 2018 , 112, 441-447	4.7	27

932	Sulfonamide Inhibition Studies of a New α -Carbonic Anhydrase from the Pathogenic Protozoan. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	7
931	Carbonic Anhydrase Inhibitors as Novel Drugs against Mycobacterial α -Carbonic Anhydrases: An Update on and Studies. <i>Molecules</i> , 2018 , 23,	4.8	14
930	Selective inhibition of carbonic anhydrase IX over carbonic anhydrase XII in breast cancer cells using benzene sulfonamides: Disconnect between activity and growth inhibition. <i>PLoS ONE</i> , 2018 , 13, e0207417	3.7	26
929	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	5.9	139
928	4-Hydroxy-3-nitro-5-ureido-benzenesulfonamides Selectively Target the Tumor-Associated Carbonic Anhydrase Isoforms IX and XII Showing Hypoxia-Enhanced Antiproliferative Profiles. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 10860-10874	8.3	41
927	Cloning, Characterization and Anion Inhibition Studies of a α -Carbonic Anhydrase from the Pathogenic Protozoan. <i>Molecules</i> , 2018 , 23,	4.8	7
926	Carbonic anhydrase inhibition with a series of novel benzenesulfonamide-triazole conjugates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1565-1574	5.6	22
925	Discovering a new class of antifungal agents that selectively inhibits microbial carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1537-1544	5.6	13
924	Novel thiazolidinone-containing compounds, without the well-known sulphonamide zinc-binding group acting as human carbonic anhydrase IX inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1299-1308	5.6	16
923	Synthesis of different thio-scaffolds bearing sulfonamide with subnanomolar carbonic anhydrase II and IX inhibitory properties and X-ray investigations for their inhibitory mechanism. <i>Bioorganic Chemistry</i> , 2018 , 81, 642-648	5.1	30
922	Steroids interfere with human carbonic anhydrase activity by using alternative binding mechanisms. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1453-1459	5.6	56
921	Immobilization of carbonic anhydrase for enhancement of CO ₂ reactive absorption. <i>New Biotechnology</i> , 2018 , 44, S44	6.4	
920	Discovery of novel 1,3-diaryltriazene sulfonamides as carbonic anhydrase I, II, VII, and IX inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 1575-1580	5.6	31
919	Performance evaluation of an all-electric waterbus supplied by hybrid energy storage systems 2018 ,		3
918	Blocking HIF signaling via novel inhibitors of CA9 and APE1/Ref-1 dramatically affects pancreatic cancer cell survival. <i>Scientific Reports</i> , 2018 , 8, 13759	4.9	29
917	Famotidine, an Antiulcer Agent, Strongly Inhibits and Human Carbonic Anhydrases. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 1035-1038	4.3	27
916	Tumor-associated carbonic anhydrase isoform IX and XII inhibitory properties of certain isatin-bearing sulfonamides endowed with in vitro antitumor activity towards colon cancer. <i>Bioorganic Chemistry</i> , 2018 , 81, 425-432	5.1	42
915	New azafluorenones with cytotoxic and carbonic anhydrase inhibitory properties: 2-Aryl-4-(4-hydroxyphenyl)-5H-indeno[1,2-b]pyridin-5-ones. <i>Bioorganic Chemistry</i> , 2018 , 81, 433-439	5.1	51

914	Applications of carbonic anhydrases inhibitors in renal and central nervous system diseases. <i>Expert Opinion on Therapeutic Patents</i> , 2018 , 28, 713-721	6.8	81
913	Tuning the Dual Inhibition of Carbonic Anhydrase and Cyclooxygenase by Dihydrothiazole Benzenesulfonamides. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 1045-1050	4.3	14
912	Discovery of new 2, 5-disubstituted 1,3-selenazoles as selective human carbonic anhydrase IX inhibitors with potent anti-tumor activity. <i>European Journal of Medicinal Chemistry</i> , 2018 , 157, 1214-1222	6.8	22
911	A case study of a DC-microgrid for the smart integration of renewable sources with the urban electric mobility 2018 ,		5
910	Heterocoumarins Are Selective Carbonic Anhydrase IX and XII Inhibitors with Cytotoxic Effects against Cancer Cells Lines. <i>ACS Medicinal Chemistry Letters</i> , 2018 , 9, 947-951	4.3	26
909	Dioxygen, an unexpected carbonic anhydrase ligand. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 999-1005	5.6	12
908	Evaluation of sulphonamide derivatives acting as inhibitors of human carbonic anhydrase isoforms I, II and Mycobacterium tuberculosis β -class enzyme Rv3273. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 962-971	5.6	22
907	Selenides bearing benzenesulfonamide show potent inhibition activity against carbonic anhydrases from pathogenic bacteria <i>Vibrio cholerae</i> and <i>Burkholderia pseudomallei</i> . <i>Bioorganic Chemistry</i> , 2018 , 79, 319-322	5.1	12
906	Design, synthesis and X-ray crystallography of selenides bearing benzenesulfonamide moiety with neuropathic pain modulating effects. <i>European Journal of Medicinal Chemistry</i> , 2018 , 154, 210-219	6.8	31
905	Discovery of β -Adrenergic Receptors Blocker-Carbonic Anhydrase Inhibitor Hybrids for Multitargeted Antiglaucoma Therapy. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 5380-5394	8.3	45
904	Activation of β - and β -carbonic anhydrases from pathogenic bacteria with tripeptides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018 , 33, 945-950	5.6	28
903	The zinc - but not cadmium - containing β -carbonic from the diatom <i>Thalassiosira weissflogii</i> is potently activated by amines and amino acids. <i>Bioorganic Chemistry</i> , 2018 , 80, 261-265	5.1	17
902	A Straightforward Access to Stable β -Functionalized Alkyl Selenols. <i>Advanced Synthesis and Catalysis</i> , 2018 , 360, 3367-3375	5.6	30
901	The first activation studies of the β -carbonic anhydrase from the malaria parasite <i>Plasmodium falciparum</i> with amines and amino acids. <i>Bioorganic Chemistry</i> , 2018 , 80, 94-98	5.1	20
900	Kinetic characterization of carbonic anhydrase immobilized on magnetic nanoparticles as biocatalyst for CO ₂ capture. <i>Biochemical Engineering Journal</i> , 2018 , 138, 1-11	4.2	18
899	Discovery of 4-Hydroxy-3-(3-(phenylureido)benzenesulfonamides as SLC-0111 Analogues for the Treatment of Hypoxic Tumors Overexpressing Carbonic Anhydrase IX. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 6328-6338	8.3	42
898	Carbonic anhydrase inhibitory properties of some uracil derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 74-77	5.6	26
897	Supported ionic liquid membranes immobilized with carbonic anhydrases for CO ₂ transport at high temperatures. <i>Journal of Membrane Science</i> , 2017 , 528, 225-230	9.6	45

896	Synthesis and biological evaluation of cyclic imides incorporating benzenesulfonamide moieties as carbonic anhydrase I, II, IV and IX inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1666-1671	3-4	27
895	Carbonic anhydrases from Trypanosoma and Leishmania as anti-protozoan drug targets. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1543-1555	3-4	46
894	Synthesis of isoxazole-containing sulfonamides with potent carbonic anhydrase II and VII inhibitory properties. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1456-1464	3-4	20
893	Novel indolin-2-one-based sulfonamides as carbonic anhydrase inhibitors: Synthesis, in vitro biological evaluation against carbonic anhydrases isoforms I, II, IV and VII and molecular docking studies. <i>European Journal of Medicinal Chemistry</i> , 2017 , 127, 521-530	6.8	44
892	Design and Synthesis of Novel Nonsteroidal Anti-Inflammatory Drugs and Carbonic Anhydrase Inhibitors Hybrids (NSAIDs-CAIs) for the Treatment of Rheumatoid Arthritis. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 1159-1170	8.3	94
891	Investigating the antiplasmodial activity of primary sulfonamide compounds identified in open source malaria data. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2017 , 7, 61-70	4	9
890	Carbonic anhydrases activation with 3-amino-1H-1,2,4-triazole-1-carboxamides: Discovery of subnanomolar isoform II activators. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1681-1686	3-4	23
889	Synthesis of bulky-tailed sulfonamides incorporating pyrido[2,3-d][1,2,4]triazolo[4,3-a]pyrimidin-1(5H)-yl moieties and evaluation of their carbonic anhydrases I, II, IV and IX inhibitory effects. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2210-2217	3-4	30
888	Evaluation of selenide, diselenide and selenoheterocycle derivatives as carbonic anhydrase I, II, IV, VII and IX inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2518-2523	3-4	38
887	Synthesis and human/bacterial carbonic anhydrase inhibition with a series of sulfonamides incorporating phthalimido moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2524-2529	3-4	23
886	Lead Development of Thiazolylsulfonamides with Carbonic Anhydrase Inhibitory Action. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 3154-3164	8.3	10
885	Carbonic anhydrase activation enhances object recognition memory in mice through phosphorylation of the extracellular signal-regulated kinase in the cortex and the hippocampus. <i>Neuropharmacology</i> , 2017 , 118, 148-156	5.5	57
884	Biochemical characterization of the native carbonic anhydrase purified from the mantle of the Mediterranean mussel, <i>Mytilus galloprovincialis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 632-639	5.6	24
883	Comparison of the anion inhibition profiles of the human and carbonic anhydrases from the pathogenic bacterium <i>Burkholderia pseudomallei</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2010-2015	3-4	6
882	Discovery of Benzenesulfonamides with Potent Human Carbonic Anhydrase Inhibitory and Effective Anticonvulsant Action: Design, Synthesis, and Pharmacological Assessment. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 2456-2469	8.3	38
881	Design, synthesis and evaluation of F-labeled cationic carbonic anhydrase IX inhibitors for PET imaging. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 722-730	5.6	39
880	N-Substituted and ring opened saccharin derivatives selectively inhibit transmembrane, tumor-associated carbonic anhydrases IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3583-3589	3-4	29
879	Novel sulfonamide-containing 2-indolinones that selectively inhibit tumor-associated alpha carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3714-3718	3-4	18

878	Synthesis of novel acyl selenoureido benzenesulfonamides as carbonic anhydrase I, II, VII and IX inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3567-3573	3.4	34
877	Kinetic properties and affinities for sulfonamide inhibitors of an α -carbonic anhydrase (CruCA4) involved in coral biomineralization in the Mediterranean red coral <i>Corallium rubrum</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3525-3530	3.4	11
876	Probing Molecular Interactions between Human Carbonic Anhydrases (hCAs) and a Novel Class of Benzenesulfonamides. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 4316-4326	8.3	30
875	Coumarins and other fused bicyclic heterocycles with selective tumor-associated carbonic anhydrase isoforms inhibitory activity. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 677-683	3.4	29
874	Production and covalent immobilisation of the recombinant bacterial carbonic anhydrase (SspCA) onto magnetic nanoparticles. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 759-766	5.6	19
873	ECA-specific inhibitor dithiocarbamate Fc14-584B: a novel antimycobacterial agent with potential to treat drug-resistant tuberculosis. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 832-840	5.6	29
872	3H-1,2-benzoxathiepine 2,2-dioxides: a new class of isoform-selective carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 767-775	5.6	32
871	Inhibition of the α -carbonic anhydrase from <i>Vibrio cholerae</i> with amides and sulfonamides incorporating imidazole moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 798-804	5.6	25
870	Sulfonamide inhibition profiles of the α -carbonic anhydrase from the pathogenic bacterium <i>Francisella tularensis</i> responsible of the febrile illness tularemia. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3555-3561	3.4	14
869	Synthesis and biological evaluation of novel aromatic and heterocyclic bis-sulfonamide Schiff bases as carbonic anhydrase I, II, VII and IX inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 3093-3097	3.4	42
868	Microwave-assisted synthesis and bioevaluation of new sulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 369-374	5.6	37
867	Synthesis of an acridine orange sulfonamide derivative with potent carbonic anhydrase IX inhibitory action. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 701-706	5.6	10
866	Inhibition of <i>Malassezia globosa</i> carbonic anhydrase with phenols. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2577-2582	3.4	34
865	Synthesis and carbonic anhydrase inhibition of a series of SLC-0111 analogs. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2569-2576	3.4	62
864	Structure activity study of carbonic anhydrase IX: Selective inhibition with ureido-substituted benzenesulfonamides. <i>European Journal of Medicinal Chemistry</i> , 2017 , 132, 184-191	6.8	43
863	3D QSAR studies, pharmacophore modeling, and virtual screening of diarylpyrazole-benzenesulfonamide derivatives as a template to obtain new inhibitors, using human carbonic anhydrase II as a model protein. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 688-700	5.6	11
862	Sulfonamide inhibition profile of the α -carbonic anhydrase identified in the genome of the pathogenic bacterium <i>Burkholderia pseudomallei</i> the etiological agent responsible of melioidosis. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017 , 27, 490-495	2.9	21
861	Benzenesulfonamide bearing imidazothiadiazole and thiazolotriazole scaffolds as potent tumor associated human carbonic anhydrase IX and XII inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 1286-1293	3.4	24

- 860 Dithiocarbamates effectively inhibit the α -carbonic anhydrase from the dandruff-producing fungus *Malassezia globosa*. *Bioorganic and Medicinal Chemistry*, **2017**, 25, 1260-1265 3.4 33
- 859 Insights into the role of reactive sulfhydryl groups of Carbonic Anhydrase III and VII during oxidative damage. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 5-12 5.6 26
- 858 Discovery of curcumin inspired sulfonamide derivatives as a new class of carbonic anhydrase isoforms I, II, IX, and XII inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1274-1281 5.6 24
- 857 Dual targeting of cancer-related human matrix metalloproteinases and carbonic anhydrases by chiral N-(biarylsulfonyl)-phosphonic acids. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1260-1264 5.6 4
- 856 Psychoactive substances belonging to the amphetamine class potentially activate brain carbonic anhydrase isoforms VA, VB, VII, and XII. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1253-1259 5.6 24
- 855 Carbonic anhydrase IX inhibition affects viability of cancer cells adapted to extracellular acidosis. *Journal of Molecular Medicine*, **2017**, 95, 1341-1353 5.5 58
- 854 Acyl selenoureido benzensulfonamides show potent inhibitory activity against carbonic anhydrases from the pathogenic bacterium *Vibrio cholerae*. *Bioorganic Chemistry*, **2017**, 75, 170-172 5.1 15
- 853 Benzoxaboroles as Efficient Inhibitors of the α -Carbonic Anhydrases from Pathogenic Fungi: Activity and Modeling Study. *ACS Medicinal Chemistry Letters*, **2017**, 8, 1194-1198 4.3 37
- 852 Carbonic Anhydrase from *Porphyromonas Gingivalis* as a Drug Target. *Pathogens*, **2017**, 6, 4.5 33
- 851 Bioactive Natural Product and Superacid Chemistry for Lead Compound Identification: A Case Study of Selective hCA III and L-Type Ca Current Inhibitors for Hypotensive Agent Discovery. *Molecules*, **2017**, 22, 4.8 2
- 850 Sequence Analysis, Kinetic Constants, and Anion Inhibition Profile of the Nacrein-Like Protein (CgINAP2X1) from the Pacific Oyster *Magallana gigas* (*Ex-Crassostrea gigas*). *Marine Drugs*, **2017**, 15, 6 2
- 849 Activation Profile Analysis of CruCA4, an α -Carbonic Anhydrase Involved in Skeleton Formation of the Mediterranean Red Coral, *Corallium rubrum*. *Molecules*, **2017**, 23, 4.8 3
- 848 Synthesis, biological activity and multiscale molecular modeling studies for coumaryl-carboxamide derivatives as selective carbonic anhydrase IX inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1042-1052 5.6 23
- 847 The synthesis of novel sulfamides derived from β -benzylphenethylamines as acetylcholinesterase, butyrylcholinesterase and carbonic anhydrase enzymes inhibitors. *Bioorganic Chemistry*, **2017**, 74, 238-250 5.1 55
- 846 Synthesis and biological evaluation of aminomethyl and alkoxymethyl derivatives as carbonic anhydrase, acetylcholinesterase and butyrylcholinesterase inhibitors. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1174-1182 5.6 67
- 845 Synthesis and biological evaluation of benzenesulphonamide-bearing 1,4,5-trisubstituted-1,2,3-triazoles possessing human carbonic anhydrase I, II, IV, and IX inhibitory activity. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2017**, 32, 1187-1194 5.6 30
- 844 Development of sulfonamides incorporating phenylacrylamido functionalities as carbonic anhydrase isoforms I, II, IX and XII inhibitors. *Bioorganic and Medicinal Chemistry*, **2017**, 25, 5726-5732 3.4 8
- 843 Sulfocoumarin-, Coumarin-, 4-Sulfamoylphenyl-Bearing Indazole-3-carboxamide Hybrids: Synthesis and Selective Inhibition of Tumor-Associated Carbonic Anhydrase Isozymes IX and XII. *ChemMedChem*, **2017**, 12, 1578-1584 3.7 26

842	Primary mono- and bis-sulfonamides obtained via regiospecific sulfochlorination of N-arylpyrazoles: inhibition profile against a panel of human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 920-934	5.6	14
841	Natural extracellular nanovesicles and photodynamic molecules: is there a future for drug delivery?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 908-916	5.6	35
840	Synthesis and carbonic anhydrase inhibition of polycyclic imides incorporating N-benzenesulfonamide moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 5373-5379	3.4	21
839	Cloning, expression and purification of the β -carbonic anhydrase from the mantle of the Mediterranean mussel, <i>Mytilus galloprovincialis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1029-1035	5.6	11
838	Novel 4/3-((4-oxo-5-(2-oxoindolin-3-ylidene)thiazolidin-2-ylidene)amino) benzenesulfonamides: Synthesis, carbonic anhydrase inhibitory activity, anticancer activity and molecular modelling studies. <i>European Journal of Medicinal Chemistry</i> , 2017 , 139, 250-262	6.8	89
837	Review on plug-in electric vehicle charging architectures integrated with distributed energy sources for sustainable mobility. <i>Applied Energy</i> , 2017 , 207, 438-464	10.7	110
836	Discovery of New Selenoureido Analogues of 4-(4-Fluorophenylureido)benzenesulfonamide as Carbonic Anhydrase Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 963-968	4.3	51
835	Inhibition of the β -carbonic anhydrase from the dandruff-producing fungus <i>Malassezia globosa</i> with monothiocarbamates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1064-1070	5.6	31
834	A one-step procedure for immobilising the thermostable carbonic anhydrase (SspCA) on the surface membrane of <i>Escherichia coli</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1120-1128	5.6	16
833	Anion inhibitors of the β -carbonic anhydrase from the pathogenic bacterium responsible of tularemia, <i>Francisella tularensis</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 4800-4804	3.4	10
832	Synthesis and carbonic anhydrase I, II, VII, and IX inhibition studies with a series of benzo[d]thiazole-5- and 6-sulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1071-1078	5.6	39
831	Dialkyl Dicyanofumarates as Oxidizing Reagents for the Conversion of Thiols into Disulfides and Selenols into Diselenides. <i>European Journal of Organic Chemistry</i> , 2017 , 2017, 6831-6839	3.2	20
830	Discovery of New Sulfonamide Carbonic Anhydrase IX Inhibitors Incorporating Nitrogenous Bases. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 1314-1319	4.3	46
829	Synthesis and biological evaluation of histamine Schiff bases as carbonic anhydrase I, II, IV, VII, and IX activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1305-1312	5.6	41
828	Synthesis of Novel Selenides Bearing Benzenesulfonamide Moieties as Carbonic Anhydrase I, II, IV, VII, and IX Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 1213-1217	4.3	32
827	Carbonic anhydrase I, II, IV and IX inhibition with a series of 7-amino-3,4-dihydroquinolin-2(1H)-one derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 885-892	5.6	8
826	-Acylbenzenesulfonamide Dihydro-1,3,4-oxadiazole Hybrids: Seeking Selectivity toward Carbonic Anhydrase Isoforms. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 792-796	4.3	22
825	Integration between Super-capacitors and ZEBRA Batteries as High Performance Hybrid Storage System for Electric Vehicles. <i>Energy Procedia</i> , 2017 , 105, 2539-2544	2.3	13

824	3-Hydroxy-1H-quinazoline-2,4-dione as a New Scaffold To Develop Potent and Selective Inhibitors of the Tumor-Associated Carbonic Anhydrases IX and XII. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 6428-6439	8.3	22
823	Advances in structure-based drug discovery of carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Discovery</i> , 2017 , 12, 61-88	6.2	298
822	A class of carbonic anhydrase I - selective activators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 37-46	5.6	31
821	Anion inhibition profiles of the β -carbonic anhydrase from the pathogenic bacterium <i>Burkholderia pseudomallei</i> responsible of melioidosis and highly drug resistant to common antibiotics. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 575-580	3.4	11
820	<i>Burkholderia pseudomallei</i> β -carbonic anhydrase is strongly activated by amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017 , 27, 77-80	2.9	23
819	Isatin: a privileged scaffold for the design of carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 68-73	5.6	42
818	Bortezomib inhibits mammalian carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 5064-5067	3.1	7
817	Open saccharin-based secondary sulfonamides as potent and selective inhibitors of cancer-related carbonic anhydrase IX and XII isoforms. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 51-59	5.6	39
816	Designing, synthesis and bioactivities of 4-[3-(4-hydroxyphenyl)-5-aryl-4,5-dihydro-pyrazol-1-yl]benzenesulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 169-175	5.6	34
815	5-Substituted-benzylsulfanyl-thiophene-2-sulfonamides with effective carbonic anhydrase inhibitory activity: Solution and crystallographic investigations. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 857-863	3.4	10
814	Structure-Activity Relationships of Benzenesulfonamide-Based Inhibitors towards Carbonic Anhydrase Isoform Specificity. <i>ChemBioChem</i> , 2017 , 18, 213-222	3.8	35
813	Novel Sulfamide-Containing Compounds as Selective Carbonic Anhydrase I Inhibitors. <i>Molecules</i> , 2017 , 22,	4.8	18
812	Targeting <i>Malassezia</i> species for Novel Synthetic and Natural Antidandruff Agents. <i>Current Medicinal Chemistry</i> , 2017 , 24, 2392-2412	4.3	23
811	Carbonic Anhydrase Inhibition and the Management of Hypoxic Tumors. <i>Metabolites</i> , 2017 , 7,	5.6	151
810	An Overview of the Bacterial Carbonic Anhydrases. <i>Metabolites</i> , 2017 , 7,	5.6	117
809	Comparison of the Sulfonamide Inhibition Profiles of the β and γ -Carbonic Anhydrases from the Pathogenic Bacterium <i>Burkholderia pseudomallei</i> . <i>Molecules</i> , 2017 , 22,	4.8	21
808	Five- and Six-Membered Nitrogen-Containing Compounds as Selective Carbonic Anhydrase Activators. <i>Molecules</i> , 2017 , 22,	4.8	12
807	Inhibition of Bacterial Carbonic Anhydrases as a Novel Approach to Escape Drug Resistance. <i>Current Topics in Medicinal Chemistry</i> , 2017 , 17, 1237-1248	3	40

806	The Warburg Effect and the Hallmarks of Cancer. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2017 , 17, 164-170	2.2	179
805	Insights into the binding mode of sulphamates and sulphamides to hCA II: crystallographic studies and binding free energy calculations. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2017 , 32, 1002-1011	5.6	25
804	Rethinking the Combination of Proton Exchanger Inhibitors in Cancer Therapy. <i>Metabolites</i> , 2017 , 8,	5.6	41
803	Hydroxamic acid derivatives: a promising scaffold for rational compound optimization in Chagas disease. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 964-73	5.6	21
802	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, 1095-101	5.6	101
801	Expression and characterization of a recombinant psychrophilic β -carbonic anhydrase (NcoCA) identified in the genome of the Antarctic cyanobacteria belonging to the genus <i>Nostoc</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 810-7	5.6	6
800	The history and rationale of using carbonic anhydrase inhibitors in the treatment of peptic ulcers. In memoriam Ioan Pădău (1932-2015). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 527-33	5.6	50
799	The effects of some bromophenols on human carbonic anhydrase isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 603-7	5.6	69
798	The effects of some avermectins on bovine carbonic anhydrase enzyme. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 773-8	5.6	41
797	A magnificent enzyme superfamily: carbonic anhydrases, their purification and characterization. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 689-94	5.6	105
796	Synthesis and inhibition potency of novel ureido benzenesulfonamides incorporating GABA as tumor-associated carbonic anhydrase IX and XII inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 205-11	5.6	14
795	Spirobisnaphthalenes effectively inhibit carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 503-7	5.6	24
794	Identification and inhibition of carbonic anhydrases from nematodes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 176-184	5.6	15
793	CA IX stratification based on cancer treatment: a patent evaluation of US2016/0002350. <i>Expert Opinion on Therapeutic Patents</i> , 2016 , 26, 1105-1109	6.8	3
792	Microwave-assisted extraction, HPLC analysis, and inhibitory effects on carbonic anhydrase I, II, VA, and VII isoforms of 14 blueberry Italian cultivars. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1-6	5.6	43
791	Mycobacterial carbonic anhydrase inhibition with phenolic acids and esters: kinetic and computational investigations. <i>Organic and Biomolecular Chemistry</i> , 2016 , 14, 8322-30	3.9	23
790	Anion inhibition profiles of the complete domain of the β -carbonic anhydrase from <i>Plasmodium falciparum</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 4410-4414	3.4	30
789	Cloning, expression and purification of the complete domain of the β -carbonic anhydrase from <i>Plasmodium falciparum</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 54-59	5.6	50

788	Overexpression of the transmembrane carbonic anhydrase isoforms IX and XII in the inflamed synovium. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 60-63	5.6	72
787	Regulation of HIF1 α under Hypoxia by APE1/Ref-1 Impacts CA9 Expression: Dual Targeting in Patient-Derived 3D Pancreatic Cancer Models. <i>Molecular Cancer Therapeutics</i> , 2016 , 15, 2722-2732	6.1	70
786	Structure-Activity Relationship for Sulfonamide Inhibition of Helicobacter pylori β -Carbonic Anhydrase. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 11098-11109	8.3	40
785	Protozoan Carbonic Anhydrases. <i>Topics in Medicinal Chemistry</i> , 2016 , 111-133	0.4	1
784	Synthesis and carbonic anhydrase inhibitory activities of new thienyl-substituted pyrazoline benzenesulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1-5	5.6	39
783	Synthesis and bioactivity studies of 1-aryl-3-(2-hydroxyethylthio)-1-propanones. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 105-109	5.6	12
782	Structure and function of carbonic anhydrases. <i>Biochemical Journal</i> , 2016 , 473, 2023-32	3.8	524
781	Bacterial Carbonic Anhydrases. <i>Topics in Medicinal Chemistry</i> , 2016 , 135-152	0.4	2
780	Synthesis and carbonic anhydrase inhibitory properties of novel chalcone substituted benzenesulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 5867-5870	2.9	32
779	Benzenesulfonamides Incorporating Flexible Triazole Moieties Are Highly Effective Carbonic Anhydrase Inhibitors: Synthesis and Kinetic, Crystallographic, Computational, and Intraocular Pressure Lowering Investigations. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 10692-10704	8.3	73
778	Multicomponent chemistry in the synthesis of carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 185-199	5.6	13
777	A new hexapeptide from the leader peptide of rMnSOD enters cells through the oestrogen receptor to deliver therapeutic molecules. <i>Scientific Reports</i> , 2016 , 6, 18691	4.9	5
776	Systems engineering approach for eco-comparison among power-train configurations of hybrid bus 2016 ,		2
775	Active Components of Essential Oils as Anti-Obesity Potential Drugs Investigated by in Silico Techniques. <i>Journal of Agricultural and Food Chemistry</i> , 2016 , 64, 5295-300	5.7	26
774	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> , 2016 , 59, 5857-67	8.3	47
773	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> , 2016 , 31, 1-9	5.6	92
772	1,2-Benzisothiazole Derivatives Bearing 4-, 5-, or 6-Alkyl/arylcarboxamide Moieties Inhibit Carbonic Anhydrase Isoform IX (CAIX) and Cell Proliferation under Hypoxic Conditions. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 6547-52	8.3	15
771	Design, synthesis and biological evaluation of N-(5-methyl-isoxazol-3-yl)-1,3,4-thiadiazol-2-yl)-4-(3-substitutedphenylureido) benzenesulfonamides as human carbonic anhydrase isoenzymes I, II, VII and XII inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 174-179	5.6	21

770	Microwave assisted synthesis of novel acridine-acetazolamide conjugates and investigation of their inhibition effects on human carbonic anhydrase isoforms hCA I, II, IV and VII. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 3548-55	3.4	9
769	Synthesis of diaryl ethers with acetylcholinesterase, butyrylcholinesterase and carbonic anhydrase inhibitory actions. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 79-85	5.6	101
768	Synthesis 4-[2-(2-mercapto-4-oxo-4H-quinazolin-3-yl)-ethyl]-benzenesulfonamides with subnanomolar carbonic anhydrase II and XII inhibitory properties. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 4100-4107	3.4	9
767	Synthesis of 4-[2-(3,4-dimethoxybenzyl)cyclopentyl]-1,2-dimethoxybenzene Derivatives and Evaluations of Their Carbonic Anhydrase Isoenzymes Inhibitory Effects. <i>Chemical Biology and Drug Design</i> , 2016 , 87, 594-607	2.9	41
766	Dithiocarbamates with potent inhibitory activity against the <i>Saccharomyces cerevisiae</i> β -carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 132-6	5.6	15
765	The human carbonic anhydrase isoenzymes I and II (hCA I and II) inhibition effects of trimethoxyindane derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 152-7	5.6	83
764	Experimental evaluation of DC charging architecture for fully-electrified low-power two-wheeler. <i>Applied Energy</i> , 2016 , 162, 1428-1438	10.7	28
763	A novel library of saccharin and acesulfame derivatives as potent and selective inhibitors of carbonic anhydrase IX and XII isoforms. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 1095-105	3.4	44
762	Amido/ureidosubstituted benzenesulfonamides-isatin conjugates as low nanomolar/subnanomolar inhibitors of the tumor-associated carbonic anhydrase isoform XII. <i>European Journal of Medicinal Chemistry</i> , 2016 , 110, 259-66	6.8	62
761	Carbonic anhydrase activators: Activation of the β -carbonic anhydrase from <i>Malassezia globosa</i> with amines and amino acids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1381-5	2.9	26
760	Inhibition of carbonic anhydrase isoforms I, II, IX and XII with secondary sulfonamides incorporating benzothiazole scaffolds. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1306-11	5.6	15
759	Sulfonamide inhibition studies of the β -carbonic anhydrase from the gammaproteobacterium <i>Thiomicrospira crunogena</i> XCL-2, TcrCA. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 401-405	2.9	1
758	Cloning, characterization and anion inhibition studies of a β -carbonic anhydrase from the Antarctic bacterium <i>Colwellia psychrerythraea</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 835-40	3.4	38
757	Synthesis of 4-sulfamoylphenyl-benzylamine derivatives with inhibitory activity against human carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 982-8	3.4	26
756	4-Arylbenzenesulfonamides as Human Carbonic Anhydrase Inhibitors (hCAIs): Synthesis by Pd Nanocatalyst-Mediated Suzuki-Miyaura Reaction, Enzyme Inhibition, and X-ray Crystallographic Studies. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 721-32	8.3	32
755	The inhibitory effects of phenolic Mannich bases on carbonic anhydrase I and II isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1678-81	5.6	34
754	Synthesis, characterization and carbonic anhydrase inhibitory activity of novel benzothiazole derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1221-5	5.6	12
753	A new procedure for the cloning, expression and purification of the β -carbonic anhydrase from the pathogenic yeast <i>Malassezia globosa</i> , an anti-dandruff drug target. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1156-61	5.6	22

752	How many carbonic anhydrase inhibition mechanisms exist?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 345-60	5.6	485
751	Synthesis and inhibitory properties of some carbamates on carbonic anhydrase and acetylcholine esterase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1484-91	5.6	28
750	Inhibitory effects of benzimidazole containing new phenolic Mannich bases on human carbonic anhydrase isoforms hCA I and II. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1540-4	5.6	12
749	Drug interaction considerations in the therapeutic use of carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , 2016 , 12, 423-31	5.5	68
748	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, 1698-702	5.6	134
747	Development of 3-(4-aminosulphonyl)-phenyl-2-mercapto-3H-quinazolin-4-ones as inhibitors of carbonic anhydrase isoforms involved in tumorigenesis and glaucoma. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 1402-7	3.4	9
746	Dynamic encapsulation and activation of carbonic anhydrase in multivalent dynameric host matrices. <i>Chemical Communications</i> , 2016 , 52, 4053-5	5.8	23
745	Carbonic anhydrase inhibition and cytotoxicity studies of Mannich base derivatives of thymol. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1375-80	5.6	31
744	Sulfonamide inhibition studies of the β -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 1115-20	3.4	51
743	Sulfonamide inhibition studies of the β -carbonic anhydrase from the Antarctic bacterium <i>Colwellia psychrerythraea</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1253-9	2.9	11
742	Anion inhibition studies of the β -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1406-10	2.9	18
741	Synthesis of N-alkyl (aryl)-tetra pyrimidine thiones and investigation of their human carbonic anhydrase I and II inhibitory effects. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1192-5 ⁶	5.6	18
740	Inhibition of carbonic anhydrase from <i>Trypanosoma cruzi</i> for the management of Chagas disease: an underexplored therapeutic opportunity. <i>Future Medicinal Chemistry</i> , 2016 , 8, 311-24	4.1	27
739	Synthesis and carbonic anhydrase I, II, IV and XII inhibitory properties of N-protected amino acid - sulfonamide conjugates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1476-83	5.6	13
738	Isatin analogs as novel inhibitors of <i>Candida</i> spp. β -carbonic anhydrase enzymes. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 1648-52	3.4	18
737	Sulfonamide inhibition studies of the β -carbonic anhydrase from the newly discovered bacterium <i>Enterobacter</i> sp. B13. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1821-6	2.9	5
736	Inhibitory effects of isatin Mannich bases on carbonic anhydrases, acetylcholinesterase, and butyrylcholinesterase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1498-501	5.6	98
735	Comparison of the sulfonamide inhibition profiles of the β and γ -carbonic anhydrases from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 1941-6	2.9	42

734	Recombinant thermoactive phosphoenolpyruvate carboxylase (PEPC) from <i>Thermosynechococcus elongatus</i> and its coupling with mesophilic/thermophilic bacterial carbonic anhydrases (CAs) for the conversion of CO ₂ to oxaloacetate. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 220-5	3.4	13
733	Sulfonamides incorporating heteropolycyclic scaffolds show potent inhibitory action against carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 921-7	3.4	17
732	Kinetic and X-ray crystallographic investigations on carbonic anhydrase isoforms I, II, IX and XII of a thioureido analog of SLC-0111. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 976-81	3.4	59
731	Development of certain new 2-substituted-quinazolin-4-yl-aminobenzenesulfonamide as potential antitumor agents. <i>European Journal of Medicinal Chemistry</i> , 2016 , 109, 247-53	6.8	32
730	The human carbonic anhydrase isoenzymes I and II inhibitory effects of some hydroperoxides, alcohols, and acetates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1248-53	5.6	13
729	Thioxocoumarins Show an Alternative Carbonic Anhydrase Inhibition Mechanism Compared to Coumarins. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 462-73	8.3	62
728	Fluorescent sulfonamide carbonic anhydrase inhibitors incorporating 1,2,3-triazole moieties: Kinetic and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 104-12	3.4	15
727	Sulfamide derivatives with selective carbonic anhydrase VII inhibitory action. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 894-901	3.4	18
726	Carbonic anhydrase inhibition for the management of cerebral ischemia: in vivo evaluation of sulfonamide and coumarin inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 894-9	5.6	78
725	Inhibition of carbonic anhydrase isoforms I, II, IV, VII and XII with carboxylates and sulfonamides incorporating phthalimide/phthalic anhydride scaffolds. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 20-5	3.4	31
724	Synthesis and carbonic anhydrase inhibitory properties of amino acid - coumarin/quinolinone conjugates incorporating glycine, alanine and phenylalanine moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1198-202	5.6	22
723	Cloning, expression and biochemical characterization of a β -carbonic anhydrase from the soil bacterium <i>Enterobacter</i> sp. B13. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1111-8	5.6	5
722	Interaction of anions with a newly characterized alpha carbonic anhydrase from <i>Halomonas</i> sp. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1119-23	5.6	10
721	The synthesis of (Z)-4-oxo-4-(arylamino)but-2-enoic acids derivatives and determination of their inhibition properties against human carbonic anhydrase I and II isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 939-45	5.6	15
720	In silico modeling of β -carbonic anhydrase inhibitors from the fungus <i>Malassezia globosa</i> as antidandruff agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 417-24	5.6	8
719	An Update on Natural Products with Carbonic Anhydrase Inhibitory Activity. <i>Current Pharmaceutical Design</i> , 2016 , 22, 1570-91	3.3	15
718	An Overview of the Carbonic Anhydrases from Two Pathogens of the Oral Cavity: <i>Streptococcus mutans</i> and <i>Porphyromonas gingivalis</i> . <i>Current Topics in Medicinal Chemistry</i> , 2016 , 16, 2359-68	3	54
717	Non-Classical Inhibition of Carbonic Anhydrase. <i>International Journal of Molecular Sciences</i> , 2016 , 17,	6.3	74

716	Phenols and Polyphenols as Carbonic Anhydrase Inhibitors. <i>Molecules</i> , 2016 , 21,	4.8	49
715	Legionella pneumophila Carbonic Anhydrases: Underexplored Antibacterial Drug Targets. <i>Pathogens</i> , 2016 , 5,	4.5	37
714	Coral Carbonic Anhydrases: Regulation by Ocean Acidification. <i>Marine Drugs</i> , 2016 , 14,	6	32
713	Synthesis and carbonic anhydrase inhibitory effects of new N-glycosylsulfonamides incorporating the phenol moiety. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 3892-5	2.9	7
712	Discovery of New Potential Anti-Infective Compounds Based on Carbonic Anhydrase Inhibitors by Rational Target-Focused Repurposing Approaches. <i>ChemMedChem</i> , 2016 , 11, 1904-14	3.7	41
711	New light on bacterial carbonic anhydrases phylogeny based on the analysis of signal peptide sequences. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1254-60	5.6	59
710	In Vivo Evaluation of Selective Carbonic Anhydrase Inhibitors as Potential Anticonvulsant Agents. <i>ChemMedChem</i> , 2016 , 11, 1812-8	3.7	23
709	9,10-Dibromo-N-aryl-9,10-dihydro-9,10-[3,4]epipyrroloanthracene-12,14-diones: Synthesis and Investigation of Their Effects on Carbonic Anhydrase Isozymes I, II, IX, and XII. <i>Archiv Der Pharmazie</i> , 2016 , 349, 466-74	4.3	28
708	A Combined Crystallographic and Theoretical Study Explains the Capability of Carboxylic Acids to Adopt Multiple Binding Modes in the Active Site of Carbonic Anhydrases. <i>Chemistry - A European Journal</i> , 2016 , 22, 97-100	4.8	34
707	Synthesis of some tetrahydropyrimidine-5-carboxylates, determination of their metal chelating effects and inhibition profiles against acetylcholinesterase, butyrylcholinesterase and carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1531-9	5.6	78
706	N-Nitrosulfonamides: A new chemotype for carbonic anhydrase inhibition. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 3612-7	3.4	29
705	Carbonic anhydrase inhibition and the management of neuropathic pain. <i>Expert Review of Neurotherapeutics</i> , 2016 , 16, 961-8	4.3	104
704	Synthesis of two phloroglucinol derivatives with cinnamyl moieties as inhibitors of the carbonic anhydrase isozymes I and II: an in vitro study. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 208-212	5.6	9
703	A substituted sulfonamide and its Co (II), Cu (II), and Zn (II) complexes as potential antifungal agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 51-62	5.6	40
702	Anion inhibition profiles of β and γ carbonic anhydrases from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 3413-7	3.4	45
701	Synthesis and bioactivity studies on new 4-(3-(4-Substitutedphenyl)-3a,4-dihydro-3H-indeno[1,2-c]pyrazol-2-yl) benzenesulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 1619-24	5.6	100
700	The synthesis of some β -lactams and investigation of their metal-chelating activity, carbonic anhydrase and acetylcholinesterase inhibition profiles. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 79-88	5.6	80
699	Lansoprazole and carbonic anhydrase IX inhibitors synergize against human melanoma cells. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 119-125	5.6	43

698	The anticonvulsant sulfamide JNJ-26990990 and its S,S-dioxide analog strongly inhibit carbonic anhydrases: solution and X-ray crystallographic studies. <i>Organic and Biomolecular Chemistry</i> , 2016 , 14, 4853-8	3.9	21
697	Novel sulfonamide bearing coumarin scaffolds as selective inhibitors of tumor associated carbonic anhydrase isoforms IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 2882-2886	3.4	34
696	Pyrazolylbenzo[d]imidazoles as new potent and selective inhibitors of carbonic anhydrase isoforms hCA IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 2907-2913	3.4	25
695	PEGylated Bis-Sulfonamide Carbonic Anhydrase Inhibitors Can Efficiently Control the Growth of Several Carbonic Anhydrase IX-Expressing Carcinomas. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 5077-88	8.3	45
694	Synthesis of 4-(thiazol-2-ylamino)-benzenesulfonamides with carbonic anhydrase I, II and IX inhibitory activity and cytotoxic effects against breast cancer cell lines. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 3043-3051	3.4	43
693	An Unusual Natural Product Primary Sulfonamide: Synthesis, Carbonic Anhydrase Inhibition, and Protein X-ray Structures of Psammaphin C. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 5462-70	8.3	29
692	Synthesis, cytotoxicity and carbonic anhydrase inhibitory activities of new pyrazolines. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 20-24	5.6	40
691	Synthesis and bioactivities of halogen bearing phenolic chalcones and their corresponding bis Mannich bases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 125-131	5.6	40
690	Benzoxaborole as a new chemotype for carbonic anhydrase inhibition. <i>Chemical Communications</i> , 2016 , 52, 11983-11986	5.8	60
689	Experimental set-up of DC PEV charging station supported by open and interoperable communication technologies 2016 ,		4
688	Cloning, expression, purification and sulfonamide inhibition profile of the complete domain of the β -carbonic anhydrase from Plasmodium falciparum. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 4184-90	2.9	34
687	Bortezomib inhibits bacterial and fungal β -carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry</i> , 2016 , 24, 4406-4409	3.4	24
686	Protonography, a technique applicable for the analysis of β -carbonic anhydrase activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 920-4	5.6	44
685	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> , 2015 , 30, 519-23	5.6	7
684	A failed tentative to design a super carbonic anhydrase having the biochemical properties of the most thermostable CA (SspCA) and the fastest (SazCA) enzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 989-94	5.6	11
683	Plasmonic Particles that Hit Hypoxic Cells. <i>Advanced Functional Materials</i> , 2015 , 25, 316-323	15.6	34
682	Dipotassium-trioxohydroxytetrafluorotriborate, $K_2[B_3O_6(OH)]$, is a potent inhibitor of human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 341-4	5.6	23
681	Investigation of arenesulfonyl-2-imidazolidinones as potent carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 81-4	5.6	36

- 680 Synthesis of pro-apoptotic indapamide derivatives as anticancer agents. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2015**, 30, 967-80 5.6 6
- 679 Synthesis of 6-aryl-substituted sulfocoumarins and investigation of their carbonic anhydrase inhibitory action. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 1430-6 3.4 33
- 678 Biochemical characterization of recombinant β -carbonic anhydrase (PgiCAB) identified in the genome of the oral pathogenic bacterium *Porphyromonas gingivalis*. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2015**, 30, 366-70 5.6 64
- 677 A class of sulfonamide carbonic anhydrase inhibitors with neuropathic pain modulating effects. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 1828-40 3.4 103
- 676 Synthesis of a novel affinity gel for the purification of carbonic anhydrases. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2015**, 30, 240-4 5.6 20
- 675 Synthesis of 3,4-dihydropyrrrolidine-2,5-dione and 3,5-dihydroxybenzoic acid derivatives and evaluation of the carbonic anhydrase I and II inhibition. *Journal of Enzyme Inhibition and Medicinal Chemistry*, **2015**, 30, 896-900 5.6 17
- 674 Sulfonamide inhibition studies of the β -carbonic anhydrase from the Antarctic bacterium *Pseudoalteromonas haloplanktis*. *Bioorganic and Medicinal Chemistry Letters*, **2015**, 25, 3550-5 2.9 28
- 673 Acatalytic Carbonic Anhydrases (CAs VIII, X, XI) **2015**, 239-245 2
- 672 Bacterial Carbonic Anhydrases as Drug Targets **2015**, 275-288 2
- 671 Synthesis and carbonic anhydrase I, II, IX and XII inhibitory activity of sulfamates incorporating piperazinyl-ureido moieties. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 5619-25 3.4 14
- 670 Synthesis of sulfonamides incorporating piperazinyl-ureido moieties and their carbonic anhydrase I, II, IX and XII inhibitory activity. *Bioorganic and Medicinal Chemistry Letters*, **2015**, 25, 3850-3 2.9 23
- 669 Bacterial, fungal and protozoan carbonic anhydrases as drug targets. *Expert Opinion on Therapeutic Targets*, **2015**, 19, 1689-704 6.4 153
- 668 Carbonic anhydrase inhibitors: Design, synthesis and structural characterization of new heteroaryl-N-carbonylbenzenesulfonamides targeting druggable human carbonic anhydrase isoforms. *European Journal of Medicinal Chemistry*, **2015**, 102, 223-32 6.8 18
- 667 Mapping Selective Inhibition of the Cancer-Related Carbonic Anhydrase IX Using Structure-Activity Relationships of Glucosyl-Based Sulfamates. *Journal of Medicinal Chemistry*, **2015**, 58, 6630-8 8.3 22
- 666 Hit Recycling: Discovery of a Potent Carbonic Anhydrase Inhibitor by in Silico Target Fishing. *ACS Chemical Biology*, **2015**, 10, 1964-9 4.9 16
- 665 Sulfonamide inhibition study of the β -class carbonic anhydrase from the caries producing pathogen *Streptococcus mutans*. *Bioorganic and Medicinal Chemistry Letters*, **2015**, 25, 2291-7 2.9 27
- 664 Cloning, characterization and anion inhibition studies of a new β -carbonic anhydrase from the Antarctic bacterium *Pseudoalteromonas haloplanktis*. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 4403-4409²⁵ 3.4
- 663 Inhibition studies of bacterial, fungal and protozoan β -class carbonic anhydrases with Schiff bases incorporating sulfonamide moieties. *Bioorganic and Medicinal Chemistry*, **2015**, 23, 4181-4187 3.4 24

662	Acetazolamide for the treatment of idiopathic intracranial hypertension. <i>Expert Review of Neurotherapeutics</i> , 2015 , 15, 851-6	4.3	108
661	Structure and inhibition studies of a type II beta-carbonic anhydrase psCA3 from <i>Pseudomonas aeruginosa</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 4831-4838	3.4	49
660	Exploring carbonic anhydrase inhibition with multimeric coumarins displayed on a fullerene scaffold. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 7445-51	3.9	32
659	New 4-[(3-cyclohexyl-4-aryl-2,3-dihydro-1,3-thiazol-2-ylidene)amino]benzene-1-sulfonamides, synthesis and inhibitory activity toward carbonic anhydrase I, II, IX, XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 3281-4	2.9	16
658	Sulfonamide bearing pyrazolylpyrazolines as potent inhibitors of carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 3208-12	2.9	33
657	Carbonic Anhydrases From Extremophiles and Their Biotechnological Applications 2015 , 311-324		
656	Probing the 'bipolar' nature of the carbonic anhydrase active site: aromatic sulfonamides containing 1,3-oxazol-5-yl moiety as picomolar inhibitors of cytosolic CA I and CA II isoforms. <i>European Journal of Medicinal Chemistry</i> , 2015 , 101, 334-47	6.8	44
655	Fluorinated pyrrolidines and piperidines incorporating tertiary benzenesulfonamide moieties are selective carbonic anhydrase II inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 737-45	5.6	30
654	Discovery of potent carbonic anhydrase and acetylcholine esterase inhibitors: novel sulfamoylcarbamates and sulfamides derived from acetophenones. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3592-602	3.4	119
653	The carbonic anhydrase from the malaria mosquito <i>Anopheles gambiae</i> is highly inhibited by sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2303-9	3.4	20
652	Synthesis of a new series of dithiocarbamates with effective human carbonic anhydrase inhibitory activity and antiglaucoma action. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2368-76	3.4	34
651	Designing carbonic anhydrase inhibitors for the treatment of breast cancer. <i>Expert Opinion on Drug Discovery</i> , 2015 , 10, 591-7	6.2	34
650	6-Substituted sulfocoumarins are selective carbonic anhydrase IX and XII inhibitors with significant cytotoxicity against colorectal cancer cells. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 3975-83	8.3	75
649	Discovery of novel isatin-based sulfonamides with potent and selective inhibition of the tumor-associated carbonic anhydrase isoforms IX and XII. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 6493-9	3.9	46
648	New amide derivatives of Probenecid as selective inhibitors of carbonic anhydrase IX and XII: biological evaluation and molecular modelling studies. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2975-81	3.4	30
647	7-Amino-3,4-dihydro-1H-quinolin-2-one, a compound similar to the substituted coumarins, inhibits carbonic anhydrases without hydrolysis of the lactam ring. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 773-7	5.6	21
646	Sulfonamide inhibition studies of the carbonic anhydrase from the Antarctic cyanobacterium <i>Nostoc commune</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 1728-34	3.4	32
645	Dendrimers incorporating benzenesulfonamide moieties strongly inhibit carbonic anhydrase isoforms I-XIV. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 6453-7	3.9	12

644	C-glycosides incorporating the 6-methoxy-2-naphthyl moiety are selective inhibitors of fungal and bacterial carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 857-61	5.6	23
643	Synthesis of Schiff base derivatives of 4-(2-aminoethyl)-benzenesulfonamide with inhibitory activity against carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 2377-81	2.9	30
642	N-Acylsulfonamides strongly inhibit human carbonic anhydrase isoenzymes I and II. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2598-605	3.4	128
641	Poly(amidoamine) Dendrimers with Carbonic Anhydrase Inhibitory Activity and Antiglaucoma Action. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 4039-45	8.3	28
640	Synthesis of novel acridine bis-sulfonamides with effective inhibitory activity against the carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 6573-80	3.4	23
639	Carbonic anhydrase IX inhibition is an effective strategy for osteosarcoma treatment. <i>Expert Opinion on Therapeutic Targets</i> , 2015 , 19, 1593-605	6.4	23
638	Anion inhibition studies of the dandruff-producing fungus <i>Malassezia globosa</i> α -carbonic anhydrase MgCA. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 5194-8	2.9	25
637	Inhibition of mammalian carbonic anhydrase isoforms I-XIV with a series of phenolic acid esters. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 7181-8	3.4	24
636	Discovery of 1,1'-Biphenyl-4-sulfonamides as a New Class of Potent and Selective Carbonic Anhydrase XIV Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 8564-72	8.3	34
635	New natural product carbonic anhydrase inhibitors incorporating phenol moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 7219-25	3.4	31
634	Exploring new Probenecid-based carbonic anhydrase inhibitors: Synthesis, biological evaluation and docking studies. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 5311-8	3.4	38
633	Click-tailed coumarins with potent and selective inhibitory action against the tumor-associated carbonic anhydrases IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 6955-66	3.4	61
632	<i>Ascaris lumbricoides</i> α -carbonic anhydrase: a potential target enzyme for treatment of ascariasis. <i>Parasites and Vectors</i> , 2015 , 8, 479	4	20
631	Carbonic anhydrase IX inhibitors in cancer therapy: an update. <i>Future Medicinal Chemistry</i> , 2015 , 7, 1407-14	4.4	106
630	Phosphate Chemical Probes Designed for Location Specific Inhibition of Intracellular Carbonic Anhydrases. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 7580-90	8.3	9
629	Isatin-pyrazole benzenesulfonamide hybrids potently inhibit tumor-associated carbonic anhydrase isoforms IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2015 , 103, 583-93	6.8	77
628	Acetazolamide protects steatotic liver grafts against cold ischemia reperfusion injury. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2015 , 355, 191-8	4.7	15
627	Anion and sulfonamide inhibition studies of an α -carbonic anhydrase from the Antarctic hemoglobinless fish <i>Chionodraco hamatus</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 5485-9	2.9	1

626	Poly(amidoamine) dendrimers show carbonic anhydrase inhibitory activity against β and γ class enzymes. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 6794-8	3.4	26
625	Efficient Expression and Crystallization System of Cancer-Associated Carbonic Anhydrase Isoform IX. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 9004-9	8.3	94
624	Carbonic anhydrase and acetylcholinesterase inhibitory effects of carbamates and sulfamoylcarbamates. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 316-20	5.6	105
623	Sulfonamide inhibition studies of the β class carbonic anhydrase from the malaria pathogen Plasmodium falciparum. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 526-31	3.4	48
622	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, 640-50	8.3	164
621	Carbonic anhydrase inhibitors: guaiacol and catechol derivatives effectively inhibit certain human carbonic anhydrase isoenzymes (hCA I, II, IX and XII). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 586-91	5.6	105
620	Inhibition studies of quinazoline-sulfonamide derivatives against the β CA (PgiCA) from the pathogenic bacterium, Porphyromonas gingivalis. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 592-6	5.6	43
619	Targeting tumour hypoxia to prevent cancer metastasis. From biology, biosensing and technology to drug development: the METOXIA consortium. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 689-721	5.6	79
618	Inhibition of human carbonic anhydrase isozymes I, II, IX and XII with a new series of sulfonamides incorporating aroylhydrazone-, [1,2,4]triazolo[3,4-b][1,3,4]thiadiazinyl- or 2-(cyanophenylmethylene)-1,3,4-thiadiazol-3(2H)-yl moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 52-6	5.6	38
617	Out of the active site binding pocket for carbonic anhydrase inhibitors. <i>Chemical Communications</i> , 2015 , 51, 302-5	5.8	96
616	6-Substituted 1,2-benzoxathiine-2,2-dioxides are isoform-selective inhibitors of human carbonic anhydrases IX, XII and VA. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 77-80	3.9	30
615	Drosophila melanogaster: a model organism for controlling Dipteran vectors and pests. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 505-13	5.6	33
614	New series of sulfonamides containing amino acid moiety act as effective and selective inhibitors of tumor-associated carbonic anhydrase XII. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 430-4	5.6	30
613	A new affinity gel for the purification of β carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 224-8	5.6	16
612	Protonography, a new technique for the analysis of carbonic anhydrase activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 277-82	5.6	75
611	Synthesis and biological activity of novel thiourea derivatives as carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 75-80	5.6	57
610	An overview of the alpha-, beta- and gamma-carbonic anhydrases from Bacteria: can bacterial carbonic anhydrases shed new light on evolution of bacteria?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 325-32	5.6	279
609	Recent advances in the discovery of zinc-binding motifs for the development of carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 321-4	5.6	68

608	Crystal structure and kinetic studies of a tetrameric type II β -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2015 , 71, 2449-56		83
607	Exploration of anionic inhibition of the β -carbonic anhydrase from <i>Thiomicrospira crunogena</i> XCL-2 gammaproteobacterium: A potential bio-catalytic agent for industrial CO ₂ removal. <i>Chemical Engineering Science</i> , 2015 , 138, 575-580	4.4	10
606	Carbonic Anhydrase II as Target for Drug Design 2015 , 51-90		2
605	Human Carbonic Anhydrases: Catalytic Properties, Structural Features, and Tissue Distribution 2015 , 17-30		5
604	Nitric oxide donors and selective carbonic anhydrase inhibitors: a dual pharmacological approach for the treatment of glaucoma, cancer and osteoporosis. <i>Molecules</i> , 2015 , 20, 5667-79	4.8	29
603	Carbonic Anhydrase Protects Fatty Liver Grafts against Ischemic Reperfusion Damage. <i>PLoS ONE</i> , 2015 , 10, e0134499	3.7	6
602	Hydroxylamine-O-sulfonamide is a versatile lead compound for the development of carbonic anhydrase inhibitors. <i>Chemical Communications</i> , 2015 , 51, 11519-22	5.8	9
601	Experimental study of a DC charging station for full electric and plug in hybrid vehicles. <i>Applied Energy</i> , 2015 , 152, 131-142	10.7	66
600	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> , 2015 , 23, 4989-4999	3.4	23
599	Protonography, a powerful tool for analyzing the activity and the oligomeric state of the β -carbonic anhydrase identified in the genome of <i>Porphyromonas gingivalis</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3747-50	3.4	39
598	New pyrazolo[4,3-e][1,2,4]triazine sulfonamides as carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 3674-80	3.4	33
597	Cloning, characterization and anion inhibition study of a β -class carbonic anhydrase from the caries producing pathogen <i>Streptococcus mutans</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 2995-3001	3.4	24
596	The zinc coordination pattern in the β -carbonic anhydrase from <i>Plasmodium falciparum</i> is different from all other carbonic anhydrase genetic families. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 1385-9	2.9	95
595	Crystal structure of the most catalytically effective carbonic anhydrase enzyme known, SazCA from the thermophilic bacterium <i>Sulfurihydrogenibium azorense</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 2002-6	2.9	60
594	Cloning, characterization and anion inhibition studies of a β -carbonic anhydrase from the Antarctic cyanobacterium <i>Nostoc commune</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 4970-4975	2.9	12
593	Nanoparticles for controlled release of anti-biofilm agents WO2014130994 (A1): a patent evaluation. <i>Expert Opinion on Therapeutic Patents</i> , 2015 , 25, 945-8	6.8	
592	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> , 2015 , 30, 901-7	5.6	13
591	N-glycosyl-N-hydroxysulfamides as potent inhibitors of <i>Brucella suis</i> carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 1010-2	5.6	6

590	Laboratory Bench to Test ZEBRA Battery Plus Super-Capacitor Based Propulsion Systems for Urban Electric Transportation. <i>Energy Procedia</i> , 2015 , 75, 1956-1961	2.3	13
589	Benzenesulfonamides incorporating bulky aromatic/heterocyclic tails with potent carbonic anhydrase inhibitory activity. <i>Bioorganic and Medicinal Chemistry</i> , 2015 , 23, 7751-64	3.4	16
588	A new class of quinazoline-sulfonamides acting as efficient inhibitors against the β -carbonic anhydrase from <i>Trypanosoma cruzi</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 581-5	5.6	19
587	Inhibition of β -carbonic anhydrases from <i>Brucella suis</i> with C-cinnamoyl glycosides incorporating the phenol moiety. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 1017-20	5.6	13
586	β -Carbonic Anhydrases Possess Thioesterase Activity. <i>ACS Medicinal Chemistry Letters</i> , 2015 , 6, 292-5	4.3	28
585	Computational investigation of the selectivity of salen and tetrahydrosalen compounds towards the tumor-associated hCA XII isozyme. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 114-8	5.6	36
584	Carbonic anhydrase inhibitors with dual-tail moieties to match the hydrophobic and hydrophilic halves of the carbonic anhydrase active site. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 1494-501	8.3	69
583	The β -class carbonic anhydrases as drug targets for antimalarial agents. <i>Expert Opinion on Therapeutic Targets</i> , 2015 , 19, 551-63	6.4	135
582	The impact of hydroquinone on acetylcholine esterase and certain human carbonic anhydrase isoenzymes (hCA I, II, IX, and XII). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2015 , 30, 941-6	5.6	74
581	Structural Basis for the Inhibition of <i>Helicobacter pylori</i> β -Carbonic Anhydrase by Sulfonamides. <i>PLoS ONE</i> , 2015 , 10, e0127149	3.7	37
580	An Overview of the Selectivity and Efficiency of the Bacterial Carbonic Anhydrase Inhibitors. <i>Current Medicinal Chemistry</i> , 2015 , 22, 2130-9	4.3	87
579	Dual Cyclooxygenase and Carbonic Anhydrase Inhibition by Nonsteroidal Anti-Inflammatory Drugs for the Treatment of Cancer. <i>Current Medicinal Chemistry</i> , 2015 , 22, 2812-8	4.3	33
578	Carbonic Anhydrases: An Overview 2015 , 3-13		11
577	Benzenesulfonamide bearing 1,2,4-triazole scaffolds as potent inhibitors of tumor associated carbonic anhydrase isoforms hCA IX and hCA XII. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 1873-82	3.4	37
576	Anion inhibition study of the β -carbonic anhydrase (CahB1) from the cyanobacterium <i>Coleofasciculus chthonoplastes</i> (ex- <i>Microcoleus chthonoplastes</i>). <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 1667-71	3.4	22
575	6-Triazolyl-substituted sulfocoumarins are potent, selective inhibitors of the tumor-associated carbonic anhydrases IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 1256-60	2.9	52
574	A small-molecule drug conjugate for the treatment of carbonic anhydrase IX expressing tumors. <i>Angewandte Chemie - International Edition</i> , 2014 , 53, 4231-5	16.4	210
573	Carbonic anhydrase inhibitors. Inhibition of human cytosolic isoforms I and II with (reduced) Schiff's bases incorporating sulfonamide, carboxylate and carboxymethyl moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2867-74	3.4	22

572	Experimental analysis on the performance of lithium based batteries for road full electric and hybrid vehicles. <i>Applied Energy</i> , 2014 , 136, 921-930	10.7	98
571	Structure-based screening for the discovery of new carbonic anhydrase VII inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2014 , 71, 105-11	6.8	41
570	Combining the tail and the ring approaches for obtaining potent and isoform-selective carbonic anhydrase inhibitors: solution and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 334-40	3.4	86
569	Sulfonamide inhibition studies of two β -carbonic anhydrases from the bacterial pathogen <i>Legionella pneumophila</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 2939-46	3.4	41
568	Dual carbonic anhydrase/matrix metalloproteinase inhibitors incorporating bisphosphonic acid moieties targeting bone tumors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 2617-20	2.9	20
567	4-Functionalized 1,3-diarylpyrazoles bearing benzenesulfonamide moiety as selective potent inhibitors of the tumor associated carbonic anhydrase isoforms IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2014 , 76, 284-90	6.8	40
566	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-(cyanophenylmethylene)-1,3,4-thiadiazol-3(2H)-yl moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 5883-90	3.4	42
565	Sulfonamide inhibition studies of the β -carbonic anhydrase from the diatom <i>Thalassiosira weissflogii</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 275-9	2.9	47
564	Cloning, characterization and anion inhibition study of the β -class carbonic anhydrase (TweCA) from the marine diatom <i>Thalassiosira weissflogii</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 531-7	3.4	56
563	Structural insights into carbonic anhydrase IX isoform specificity of carbohydrate-based sulfamates. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 8635-45	8.3	47
562	Inhibition of carbonic anhydrase isoforms I, II, IX and XII with novel Schiff bases: identification of selective inhibitors for the tumor-associated isoforms over the cytosolic ones. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 5883-90	3.4	11
561	Carbonic anhydrase inhibitory activity of sulfonamides and carboxylic acids incorporating cyclic imide scaffolds. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 5185-9	2.9	41
560	Shading the TRF2 recruiting function: a new horizon in drug development. <i>Journal of the American Chemical Society</i> , 2014 , 136, 16708-11	16.4	16
559	Quinazoline-sulfonamides with potent inhibitory activity against the β -carbonic anhydrase from <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 5133-40	3.4	35
558	Mono- and di-halogenated histamine, histidine and carnosine derivatives are potent carbonic anhydrase I, II, VII, XII and XIV activators. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 4752-8	3.4	16
557	Discovery of a new family of carbonic anhydrases in the malaria pathogen <i>Plasmodium falciparum</i> —the β -carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 4389-4396	2.9	258
556	Immobilization of carbonic anhydrase for biomimetic CO ₂ capture in slurry absorber. <i>New Biotechnology</i> , 2014 , 31, S20-S21	6.4	2
555	Cyclic tertiary sulfamates: selective inhibition of the tumor-associated carbonic anhydrases IX and XII by N- and O-substituted acesulfame derivatives. <i>European Journal of Medicinal Chemistry</i> , 2014 , 84, 240-6	6.8	37

554	Synthesis of a new series of N-substituted 4-(2-aminoethyl)benzenesulfonamides and their inhibitory effect on human carbonic anhydrase cytosolic isozymes I and II and transmembrane tumor-associated isozymes IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2014 , 84, 59-67	6.8	15
553	Ethylene bis-imidazoles are highly potent and selective activators for isozymes VA and VII of carbonic anhydrase, with a potential nootropic effect. <i>Chemical Communications</i> , 2014 , 50, 5980-3	5.8	44
552	Crystal structures of two tetrameric α -carbonic anhydrases from the filamentous ascomycete <i>Sordaria macrospora</i> . <i>FEBS Journal</i> , 2014 , 281, 1759-72	5.7	35
551	Furazan and furoxan sulfonamides are strong α -carbonic anhydrase inhibitors and potential antiglaucoma agents. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3913-21	3.4	25
550	Synthesis of sulfonamides with effective inhibitory action against <i>Porphyromonas gingivalis</i> α -carbonic anhydrase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 4006-10	2.9	20
549	Attachment of carbohydrates to methoxyaryl moieties leads to highly selective inhibitors of the cancer associated carbonic anhydrase isoforms IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 5308-14	3.4	28
548	Anion inhibition study of the β -class carbonic anhydrase (PgiCAB) from the oral pathogen <i>Porphyromonas gingivalis</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 4402-4406	2.9	27
547	Sulfonamides with Potent Inhibitory Action and Selectivity against the α -Carbonic Anhydrase from <i>Vibrio cholerae</i> . <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 826-30	4.3	23
546	Synthesis of 6-tetrazolyl-substituted sulfocoumarins acting as highly potent and selective inhibitors of the tumor-associated carbonic anhydrase isoforms IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 1522-8	3.4	45
545	Chemometric modeling of breast cancer associated carbonic anhydrase IX inhibitors belonging to the ureido-substituted benzene sulfonamide class. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 877-83	5.6	8
544	Sulfonamides incorporating fluorine and 1,3,5-triazine moieties are effective inhibitors of three β -class carbonic anhydrases from <i>Mycobacterium tuberculosis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 686-9	5.6	38
543	Selective inhibition of human carbonic anhydrases by novel amide derivatives of probenecid: synthesis, biological evaluation and molecular modelling studies. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3982-8	3.4	34
542	Sulfonamide inhibition study of the carbonic anhydrases from the bacterial pathogen <i>Porphyromonas gingivalis</i> : the β -class (PgiCAB) versus the β -class (PgiCA) enzymes. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 4537-43	3.4	32
541	Substituted benzene sulfonamides incorporating 1,3,5-triazinyl moieties potently inhibit human carbonic anhydrases II, IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 1310-4	2.9	16
540	Anion inhibition studies of two new α -carbonic anhydrases from the bacterial pathogen <i>Legionella pneumophila</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 1127-32	2.9	44
539	Arylamino bisphosphonates: potent and selective inhibitors of the tumor-associated carbonic anhydrase XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 1941-3	2.9	10
538	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> , 2014 , 24, 1776-9	2.9	21
537	Carbonic anhydrase inhibitors. Synthesis of a novel series of 5-substituted 2,4-dichlorobenzenesulfonamides and their inhibition of human cytosolic isozymes I and II and the transmembrane tumor-associated isozymes IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2014 , 82, 17-55	6.8	15

536	Carbonic anhydrase inhibitors. Synthesis, and molecular structure of novel series N-substituted N'-(2-arylmethylthio-4-chloro-5-methylbenzenesulfonyl)guanidines and their inhibition of human cytosolic isozymes I and II and the transmembrane tumor-associated isozymes IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2014 , 71, 135-47	6.8	58
535	Anion inhibition studies of two β -carbonic anhydrases from <i>Lotus japonicus</i> , LjCAA1 and LjCAA2. <i>Journal of Inorganic Biochemistry</i> , 2014 , 136, 67-72	4.2	15
534	Oxidation of cyanobenzocycloheptatrienes: Synthesis, photooxygenation reaction and carbonic anhydrase isoenzymes inhibition properties of some new benzotropone derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3537-43	3.4	92
533	Sulfonamide inhibition studies of the β -carbonic anhydrase from the oral pathogen <i>Porphyromonas gingivalis</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 240-4	2.9	46
532	Design, synthesis, and evaluation of hydroxamic acid derivatives as promising agents for the management of Chagas disease. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 298-308	8.3	64
531	Dominant behaviours in the expression of human carbonic anhydrase hCA I activity. <i>Chemical Communications</i> , 2014 , 50, 8043-6	5.8	15
530	Carbonic Anhydrase Inhibition with Benzenesulfonamides and Tetrafluorobenzenesulfonamides Obtained via Click Chemistry. <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 927-30	4.3	41
529	Biochemical characterization of the chloroplastic β -carbonic anhydrase from <i>Flaveria bidentis</i> (L.) "Kuntze". <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 500-4	5.6	16
528	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, 379-87	5.6	212
527	Biochemical properties of a new β -carbonic anhydrase from the human pathogenic bacterium, <i>Vibrio cholerae</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 23-7	5.6	85
526	Novel coumarins and benzocoumarins acting as isoform-selective inhibitors against the tumor-associated carbonic anhydrase IX. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 292-6	5.6	78
525	Overview of carbonic anhydrase families/isoforms 2014 , 6-16		0
524	Next-generation secondary/tertiary sulfonamide carbonic anhydrase inhibitor 2014 , 52-67		
523	Developing Novel Bacterial Targets: Carbonic Anhydrases as Antibacterial Drug Targets 2014 , 31-46		1
522	Targeting Carbonic Anhydrases 2014 ,		7
521	Experimental Analysis of a Zebra Battery Based Propulsion System for Urban Bus under Dynamic Conditions. <i>Energy Procedia</i> , 2014 , 61, 1138-1141	2.3	5
520	Effect of a recombinant manganese superoxide dismutase on prevention of contrast-induced acute kidney injury. <i>Clinical and Experimental Nephrology</i> , 2014 , 18, 424-31	2.5	39
519	Targeting carbonic anhydrases in biotechnology 2014 , 158-169		1

518	Biochemical characterization of the α -carbonic anhydrase from the marine diatom <i>Thalassiosira weissflogii</i> , TweCA. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 906-11	5.6	58
517	Sulfonamides and their isosters as carbonic anhydrase inhibitors. <i>Future Medicinal Chemistry</i> , 2014 , 6, 1149-65	4.1	133
516	Synthesis and evaluation of ¹⁸ F-labeled carbonic anhydrase IX inhibitors for imaging with positron emission tomography. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 249-55	5.6	60
515	Biomimetic CO ₂ capture using a highly thermostable bacterial α -carbonic anhydrase immobilized on a polyurethane foam. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 146-50	5.6	116
514	Carbonic anhydrase inhibitors. Phenols incorporating 2- or 3-pyridyl-ethenylcarbonyl and tertiary amine moieties strongly inhibit <i>Saccharomyces cerevisiae</i> α -carbonic anhydrase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 495-9	5.6	46
513	Biochemical characterization of the β -carbonic anhydrase from the oral pathogen <i>Porphyromonas gingivalis</i> , PgiCA. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 532-7	5.6	62
512	Monoclonal antibodies raised against 167-180 aa sequence of human carbonic anhydrase XII inhibit its enzymatic activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 804-10	5.6	13
511	Natural product polyamines that inhibit human carbonic anhydrases. <i>BioMed Research International</i> , 2014 , 2014, 374079	3	18
510	Hydrophobic substituents of the phenylmethanesulfonamide moiety can be used for the development of new selective carbonic anhydrase inhibitors. <i>BioMed Research International</i> , 2014 , 2014, 523210	3	12
509	A class of 4-sulfamoylphenyl- β -aminoalkyl ethers with effective carbonic anhydrase inhibitory action and antiglaucoma effects. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 9673-86	8.3	44
508	Structural insights on carbonic anhydrase inhibitory action, isoform selectivity, and potency of sulfonamides and coumarins incorporating arylsulfonylureido groups. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 9152-67	8.3	46
507	Inhibition studies of new ureido-substituted sulfonamides incorporating a GABA moiety against human carbonic anhydrase isoforms I-XIV. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 6768-75	3.4	21
506	4-Functionalized 1,3-diarylpyrazoles bearing 6-aminosulfonylbenzothiazole moiety as potent inhibitors of carbonic anhydrase isoforms hCA I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 6945-52	3.4	18
505	The structural comparison between membrane-associated human carbonic anhydrases provides insights into drug design of selective inhibitors. <i>Biopolymers</i> , 2014 , 101, 769-78	2.2	39
504	Carbonic anhydrase inhibitory properties of novel sulfonamide derivatives of aminoindanes and aminotetralins. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 35-42	5.6	92
503	Protozoan, fungal and bacterial carbonic anhydrases targeting for obtaining anti-infectives 2014 , 132-141		2
502	Carbonic anhydrase inhibitors drug design. <i>Sub-Cellular Biochemistry</i> , 2014 , 75, 291-323	5.5	82
501	Carbonic anhydrase inhibitors: synthesis and inhibition of the human carbonic anhydrase isoforms I, II, IX and XII with benzene sulfonamides incorporating 4- and 3-nitrophthalimide moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 1586-95	3.4	31

500	Carbonic anhydrase inhibitors: Synthesis, molecular docking, cytotoxic and inhibition of the human carbonic anhydrase isoforms I, II, IX, XII with novel benzenesulfonamides incorporating pyrrole, pyrrolopyrimidine and fused pyrrolopyrimidine moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2014 , 22, 3684-95	3.4	44
499	Sulfonamide inhibition studies of the β -carbonic anhydrase from <i>Drosophila melanogaster</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014 , 24, 2797-801	2.9	10
498	3D-QSAR CoMFA studies on sulfonamide inhibitors of the Rv3588c β -carbonic anhydrase from <i>Mycobacterium tuberculosis</i> and design of not yet synthesized new molecules. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 449-55	5.6	38
497	Inhibition of mammalian carbonic anhydrases I-XIV with grayanotoxin III: solution and in silico studies. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2014 , 29, 469-75	5.6	31
496	Glaucoma and the applications of carbonic anhydrase inhibitors. <i>Sub-Cellular Biochemistry</i> , 2014 , 75, 349-59	5.9	86
495	Next-generation dithiocarbamate carbonic anhydrase inhibitors 2014 , 114-130		
494	The role of carbonic anhydrase IX in hypoxia control in OSCC. <i>Journal of Oral Pathology and Medicine</i> , 2013 , 42, 1-8	3.3	17
493	Secondary/tertiary benzenesulfonamides with inhibitory action against the cytosolic human carbonic anhydrase isoforms I and II. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 294-8	5.6	71
492	Targeting carbonic anhydrase IX by nitroimidazole based sulfamides enhances the therapeutic effect of tumor irradiation: a new concept of dual targeting drugs. <i>Radiotherapy and Oncology</i> , 2013 , 108, 523-8	5.3	74
491	Hypoxia induced CA9 inhibitory targeting by two different sulfonamide derivatives including acetazolamide in human glioblastoma. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 3949-57	3.4	45
490	Carbonic anhydrase inhibitors. Benzenesulfonamides incorporating cyanoacrylamide moieties strongly inhibit <i>Saccharomyces cerevisiae</i> β -carbonic anhydrase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 3570-5	2.9	16
489	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> , 2013 , 21, 5130-8	3.4	26
488	A class of sulfonamides with strong inhibitory action against the β -carbonic anhydrase from <i>Trypanosoma cruzi</i> . <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 5773-81	8.3	51
487	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> , 2013 , 21, 5799-805	3.4	29
486	Analysis of saponins and phenolic compounds as inhibitors of β -carbonic anhydrase isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 412-7	5.6	46
485	Carbonic anhydrase inhibitors: synthesis and inhibition of the human carbonic anhydrase isoforms I, II, VII, IX and XII with benzene sulfonamides incorporating 4,5,6,7-tetrabromophthalimide moiety. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 5973-82	3.4	18
484	Inhibition of the β -carbonic anhydrases from <i>Mycobacterium tuberculosis</i> with C-cinnamoyl glycosides: identification of the first inhibitor with anti-mycobacterial activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 740-3	2.9	44
483	Anion inhibition studies of a β -carbonic anhydrase from <i>Clostridium perfringens</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 6706-10	2.9	42

482	QSAR studies of sulfamate and sulfamide inhibitors targeting human carbonic anhydrase isozymes I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1404-9	3.4	9
481	Structural study of interaction between brinzolamide and dorzolamide inhibition of human carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 7210-5	3.4	76
480	Salen and tetrahydrosalen derivatives act as effective inhibitors of the tumor-associated carbonic anhydrase XII--a new scaffold for designing isoform-selective inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 6759-63	2.9	37
479	Carbonic anhydrase inhibitors. Synthesis of heterocyclic 4-substituted pyridine-3-sulfonamide derivatives and their inhibition of the human cytosolic isozymes I and II and transmembrane tumor-associated isozymes IX and XII. <i>European Journal of Medicinal Chemistry</i> , 2013 , 69, 701-10	6.8	31
478	Exploiting the hydrophobic and hydrophilic binding sites for designing carbonic anhydrase inhibitors. <i>Expert Opinion on Drug Discovery</i> , 2013 , 8, 793-810	6.2	215
477	Effect of incorporating a thiophene tail in the scaffold of acetazolamide on the inhibition of human carbonic anhydrase isoforms I, II, IX and XII. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 5646-9	2.9	17
476	Restoring catalytic activity to the human carbonic anhydrase (CA) related proteins VIII, X and XI affords isoforms with high catalytic efficiency and susceptibility to anion inhibition. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 256-60	2.9	37
475	Carbonic anhydrase inhibitors: Synthesis and inhibition of the cytosolic mammalian carbonic anhydrase isoforms I, II and VII with benzene sulfonamides incorporating 4,5,6,7-tetrachlorophthalimide moiety. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 5168-74	3.4	15
474	Carbonic anhydrase regulation and CO(2) sensing in the fungal pathogen <i>Candida glabrata</i> involves a novel Rca1p ortholog. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1549-54	3.4	40
473	Inhibition of the alpha- and beta-carbonic anhydrases from the gastric pathogen <i>Helicobacter pylori</i> with anions. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 388-91	5.6	82
472	Glycosidic carbonic anhydrase IX inhibitors: a sweet approach against cancer. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1419-26	3.4	52
471	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	8.3	174
470	Kinetic and in silico analysis of thiazolidin-based inhibitors of carbonic anhydrase isoenzymes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 370-4	5.6	11
469	Carbonic anhydrases in anthozoan corals-A review. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1437-50	5.4	137
468	Heavy metal ion inhibition studies of human, sheep and fish carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 278-82	5.6	35
467	Effect of sulfonamides as carbonic anhydrase VA and VB inhibitors on mitochondrial metabolic energy conversion. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1544-8	3.4	84
466	Pharmacological inhibition of carbonic anhydrase XII interferes with cell proliferation and induces cell apoptosis in T-cell lymphomas. <i>Cancer Letters</i> , 2013 , 333, 76-88	9.9	44
465	Structural study of the location of the phenyl tail of benzene sulfonamides and the effect on human carbonic anhydrase inhibition. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 6674-80	3.4	12

464	An α -carbonic anhydrase from the thermophilic bacterium <i>Sulphurihydrogenibium azorense</i> is the fastest enzyme known for the CO ₂ hydration reaction. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1465-9	3.4	96
463	The extremophilic carbonic anhydrase (CA) from <i>Sulphurihydrogenibium azorense</i> , the fastest CA known, is highly activated by amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 1087-90	2.9	52
462	o-Benzenedisulfonimido-sulfonamides are potent inhibitors of the tumor-associated carbonic anhydrase isoforms CA IX and CA XII. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1386-91	3.4	19
461	Characterization of carbonic anhydrase IX interactome reveals proteins assisting its nuclear localization in hypoxic cells. <i>Journal of Proteome Research</i> , 2013 , 12, 282-92	5.6	37
460	Antiobesity carbonic anhydrase inhibitors: a literature and patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 725-35	6.8	213
459	Inhibition of human carbonic anhydrase isoforms I-XIV with sulfonamides incorporating fluorine and 1,3,5-triazine moieties. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 6929-36	3.4	18
458	Kinetic and anion inhibition studies of a β -carbonic anhydrase (FbiCA 1) from the C4 plant <i>Flaveria bidentis</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 1626-30	2.9	33
457	Carbonic anhydrase inhibitors: inhibition of the β -class enzyme from the pathogenic yeast <i>Candida glabrata</i> with sulfonamides, sulfamates and sulfamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 2647-52	2.9	38
456	Inhibition of tumor-associated human carbonic anhydrase isozymes IX and XII by a new class of substituted-phenylacetamido aromatic sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 5228-32	3.4	19
455	A highly catalytically active β -carbonic anhydrase from the pathogenic anaerobe <i>Porphyromonas gingivalis</i> and its inhibition profile with anions and small molecules. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 4067-71	2.9	58
454	The extremophilic carbonic anhydrase from the thermophilic bacterium <i>Sulphurihydrogenibium azorense</i> is highly inhibited by sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 4521-5	3.4	60
453	Cloning, characterization, and sulfonamide and thiol inhibition studies of an α -carbonic anhydrase from <i>Trypanosoma cruzi</i> , the causative agent of Chagas disease. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 1761-71	8.3	81
452	Anti-infective carbonic anhydrase inhibitors: a patent and literature review. <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 693-704	6.8	192
451	Diuretics with carbonic anhydrase inhibitory action: a patent and literature review (2005 - 2013). <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 681-91	6.8	218
450	Structural effect of phenyl ring compared to thiadiazole based adamantyl-sulfonamides on carbonic anhydrase inhibition. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 2314-2318	3.4	19
449	Characterization, bioinformatic analysis and dithiocarbamate inhibition studies of two new β -carbonic anhydrases, CAH1 and CAH2, from the fruit fly <i>Drosophila melanogaster</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1516-21	3.4	15
448	Metalloenzyme inhibitors for the treatment of Gram-negative bacterial infections: a patent review (2009-2012). <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 777-88	6.8	23
447	The α -carbonic anhydrase from the thermophilic bacterium <i>Sulphurihydrogenibium yellowstonense</i> YO3AOP1 is highly susceptible to inhibition by sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1534-8	3.4	50

446	Carbonic anhydrase inhibitors: benzenesulfonamides incorporating cyanoacrylamide moieties are low nanomolar/subnanomolar inhibitors of the tumor-associated isoforms IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1396-403	3.4	46
445	Secondary and tertiary sulfonamides: a patent review (2008 - 2012). <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 203-13	6.8	67
444	Natural product coumarins that inhibit human carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1539-43	3.4	82
443	Novel small molecule protein arginine deiminase 4 (PAD4) inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 715-9	2.9	28
442	Nothepsin 2013 , 63-69		
441	Dihalogenated sulfanilamides and benzolamides are effective inhibitors of the three β -class carbonic anhydrases from <i>Mycobacterium tuberculosis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 384-7	5.6	71
440	Dithiocarbamates strongly inhibit the β -class carbonic anhydrases from <i>Mycobacterium tuberculosis</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 407-11	5.6	118
439	Inhibition of the β -class carbonic anhydrases from <i>Mycobacterium tuberculosis</i> with carboxylic acids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 392-6	5.6	68
438	Synthesis and carbonic anhydrase inhibitory properties of sulfamides structurally related to dopamine. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 2925-31	3.4	112
437	Anion inhibition studies of the β -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 1636-8	2.9	50
436	Sperm from sneaker male squids exhibit chemotactic swarming to CO ₂ <i>Current Biology</i> , 2013 , 23, 775-81	6.3	43
435	Antiglaucoma carbonic anhydrase inhibitors: a patent review. <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 705-16	6.8	232
434	Root effect hemoglobin may have evolved to enhance general tissue oxygen delivery. <i>Science</i> , 2013 , 340, 1327-9	33.3	106
433	Natural product hybrid and its superacid synthesized analogues: dodoneine and its derivatives show selective inhibition of carbonic anhydrase isoforms I, III, XIII and XIV. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 3790-4	3.4	14
432	Xanthates and trithiocarbonates strongly inhibit carbonic anhydrases and show antiglaucoma effects in vivo. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 4691-700	8.3	82
431	7-Substituted-sulfocoumarins are isoform-selective, potent carbonic anhydrase II inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 4502-10	3.4	62
430	Anticancer carbonic anhydrase inhibitors: a patent review (2008 - 2013). <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 737-49	6.8	208
429	Mono-/dihydroxybenzoic acid esters and phenol pyridinium derivatives as inhibitors of the mammalian carbonic anhydrase isoforms I, II, VII, IX, XII and XIV. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 1564-9	3.4	43

428	Anion inhibition studies of the α -carbonic anhydrase from the protozoan pathogen <i>Trypanosoma cruzi</i> , the causative agent of Chagas disease. <i>Bioorganic and Medicinal Chemistry</i> , 2013 , 21, 4472-6	3.4	45
427	Kinetic study of a novel thermo-stable β -carbonic anhydrase for biomimetic CO ₂ capture. <i>Enzyme and Microbial Technology</i> , 2013 , 53, 271-7	3.8	32
426	New strategies for targeting the hypoxic tumour microenvironment in breast cancer. <i>Cancer Treatment Reviews</i> , 2013 , 39, 171-9	14.4	142
425	Hypoxia-targeting carbonic anhydrase IX inhibitors by a new series of nitroimidazole-sulfonamides/sulfamides/sulfamates. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 8512-20	8.3	68
424	Carbonic anhydrase inhibitors: in vitro inhibition of β -isoforms (hCA I, hCA II, bCA III, hCA IV) by flavonoids. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 283-8	5.6	94
423	Cloning, characterization, and inhibition studies of a β -carbonic anhydrase from <i>Leishmania donovani</i> chagasi, the protozoan parasite responsible for leishmaniasis. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 7372-81	8.3	79
422	Carbonic anhydrase III: a neglected isozyme is stepping into the limelight. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 231-9	5.6	69
421	A new recombinant MnSOD prevents the cyclosporine A-induced renal impairment. <i>Nephrology Dialysis Transplantation</i> , 2013 , 28, 2066-72	4.3	19
420	Human carbonic anhydrase VII protects cells from oxidative damage. <i>Biological Chemistry</i> , 2013 , 394, 1343-8	4.5	29
419	Metronidazole-coumarin conjugates and 3-cyano-7-hydroxy-coumarin act as isoform-selective carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 397-401	5.6	81
418	X-ray structure of the first α -extremo- β -carbonic anhydrase', a dimeric enzyme from the thermophilic bacterium <i>Sulfolobus solfataricus</i> yellowstonense YO3AOP1. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2013 , 69, 1150-9		89
417	Novel antibody to a carbonic anhydrase: patent evaluation of WO2011138279A1. <i>Expert Opinion on Therapeutic Patents</i> , 2013 , 23, 757-60	6.8	6
416	Synthesis of aminocyanopyrazoles via a multi-component reaction and anti-carbonic anhydrase inhibitory activity of their sulfamide derivatives against cytosolic and transmembrane isoforms. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013 , 28, 343-9	5.6	23
415	Cloning, characterization and sulfonamide inhibition studies of an α -carbonic anhydrase from the living fossil sponge <i>Astrosclera willeyana</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 1403-10	3.4	6
414	Novel coumarins and 2-thioxo-coumarins as inhibitors of the tumor-associated carbonic anhydrases IX and XII. <i>Bioorganic and Medicinal Chemistry</i> , 2012 , 20, 2266-73	3.4	101
413	Synthesis and evaluation of near-infrared fluorescent sulfonamide derivatives for imaging of hypoxia-induced carbonic anhydrase IX expression in tumors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 653-7	2.9	44
412	5- and 6-membered (thio)lactones are prodrug type carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 267-70	2.9	54
411	Dithiocarbamates are strong inhibitors of the beta-class fungal carbonic anhydrases from <i>Cryptococcus neoformans</i> , <i>Candida albicans</i> and <i>Candida glabrata</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 859-62	2.9	89

410	Anion inhibition studies of an α -carbonic anhydrase from the living fossil <i>Astrosclera willeyana</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 1314-6	2.9	5
409	Carbonic anhydrase VII is S-glutathionylated without loss of catalytic activity and affinity for sulfonamide inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 1560-4	2.9	46
408	New chemotypes acting as isozyme-selective carbonic anhydrase inhibitors with low affinity for the offtarget cytosolic isoform II. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 2182-5	2.9	46
407	A molecular carrier to transport and deliver cisplatin into endometrial cancer cells. <i>Chemical Biology and Drug Design</i> , 2012 , 80, 9-16	2.9	2
406	Inhibition of beta-carbonic anhydrases from the bacterial pathogen <i>Brucella suis</i> with inorganic anions. <i>Journal of Inorganic Biochemistry</i> , 2012 , 110, 36-9	4.2	27
405	Tricyclic sulfonamides incorporating benzothiopyrano[4,3-c]pyrazole and pyridothiopyrano[4,3-c]pyrazole effectively inhibit α - and β -carbonic anhydrase: X-ray crystallography and solution investigations on 15 isoforms. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 8618-28	8.3	32
404	Antimetastatic effect of sulfamate carbonic anhydrase IX inhibitors in breast carcinoma xenografts. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 5591-600	8.3	123
403	DNA cloning, characterization, and inhibition studies of an α -carbonic anhydrase from the pathogenic bacterium <i>Vibrio cholerae</i> . <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 10742-8	8.3	91
402	Carbonic anhydrases inhibitory effects of new benzenesulfonamides synthesized by using superacid chemistry. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 886-91	5.6	67
401	Anion inhibition studies of the fastest carbonic anhydrase (CA) known, the extremo-CA from the bacterium <i>Sulfurihydrogenibium azorense</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 7142-5	2.9	60
400	Chromone containing sulfonamides as potent carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 744-7	5.6	36
399	Inhibition of β -class cytosolic human carbonic anhydrases I, II, IX and XII, and β -class fungal enzymes by carboxylic acids and their derivatives: new isoform-I selective nanomolar inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 5801-6	2.9	29
398	The first activation study of a bacterial carbonic anhydrase (CA). The thermostable β -CA from <i>Sulfurihydrogenibium yellowstonense</i> YO3AOP1 is highly activated by amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 6324-7	2.9	69
397	Upcoming conferences of interest. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 166-166		
396	Dithiocarbamates: a new class of carbonic anhydrase inhibitors. Crystallographic and kinetic investigations. <i>Chemical Communications</i> , 2012 , 48, 1868-70	5.8	149
395	Synthesis, structure-activity relationship studies, and X-ray crystallographic analysis of arylsulfonamides as potent carbonic anhydrase inhibitors. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 3891-9	8.3	22
394	β -Carbonic anhydrases are sulfatases with cyclic diol monosulfate esters. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 148-54	5.6	63
393	Polypharmacology of sulfonamides: pazopanib, a multitargeted receptor tyrosine kinase inhibitor in clinical use, potently inhibits several mammalian carbonic anhydrases. <i>Chemical Communications</i> , 2012 , 48, 8177-9	5.8	62

392	Serendipitous fragment-based drug discovery: ketogenic diet metabolites and statins effectively inhibit several carbonic anhydrases. <i>Chemical Communications</i> , 2012 , 48, 3551-3	5.8	22
391	Effects of dopaminergic compounds on carbonic anhydrase isozymes I, II, and VI. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 365-9	5.6	27
390	QSARs on human carbonic anhydrase VA and VB inhibitors of some new not yet synthesized, substituted aromatic/heterocyclic sulphonamides as anti-obesity agent. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 666-72	5.6	20
389	Biochemical properties of a novel and highly thermostable bacterial α -carbonic anhydrase from <i>Sulfolobus solfataricus</i> yellowstonense YO3AOP1. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 892-7	5.6	87
388	Molecular cloning, characterization, and inhibition studies of a β -carbonic anhydrase from <i>Malassezia globosa</i> , a potential antidandruff target. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 3513-20	8.3	44
387	CO ₂ permeability of cell membranes is regulated by membrane cholesterol and protein gas channels. <i>FASEB Journal</i> , 2012 , 26, 5182-91	0.9	72
386	(In)organic anions as carbonic anhydrase inhibitors. <i>Journal of Inorganic Biochemistry</i> , 2012 , 111, 117-29	4.2	173
385	Structural and inhibition insights into carbonic anhydrase CDCA1 from the marine diatom <i>Thalassiosira weissflogii</i> . <i>Biochimie</i> , 2012 , 94, 1232-41	4.6	88
384	Ureido-substituted sulfamates show potent carbonic anhydrase IX inhibitory and antiproliferative activities against breast cancer cell lines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 4681-5	2.9	51
383	Anion inhibition studies of an α -carbonic anhydrase from the thermophilic bacterium <i>Sulfolobus solfataricus</i> yellowstonense YO3AOP1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 5630-4	2.9	71
382	Rational drug repositioning guided by an integrated pharmacological network of protein, disease and drug. <i>BMC Systems Biology</i> , 2012 , 6, 80	3.5	62
381	Development of potent carbonic anhydrase inhibitors incorporating both sulfonamide and sulfamide groups. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 6776-83	8.3	43
380	Dual inhibitors for aspartic proteases HIV-1 PR and renin: advancements in AIDS-hypertension-diabetes linkage via molecular dynamics, inhibition assays, and binding free energy calculations. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 5784-96	8.3	33
379	Dithiocarbamates strongly inhibit carbonic anhydrases and show antiglaucoma action in vivo. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 1721-30	8.3	195
378	Toxicity, accumulation, and removal of heavy metals by three aquatic macrophytes. <i>International Journal of Phytoremediation</i> , 2012 , 14, 374-87	3.9	70
377	Structure-based drug discovery of carbonic anhydrase inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 759-72	5.6	483
376	Simple methanesulfonates are hydrolyzed by the sulfatase carbonic anhydrase activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 880-5	5.6	50
375	Sulfapyridine-like benzenesulfonamide derivatives as inhibitors of carbonic anhydrase isoenzymes I, II and VI. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 818-24	5.6	45

374	Carbonic anhydrase inhibitors: inhibition of human and bovine isoenzymes by benzenesulphonamides, cyclitols and phenolic compounds. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 845-8	5.6	67
373	Inhibition of carbonic anhydrase IX as a novel anticancer mechanism. <i>World Journal of Clinical Oncology</i> , 2012 , 3, 98-103	2.5	73
372	Novel therapies for glaucoma: a patent review 2007 - 2011. <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 79-88	6.8	104
371	Protein-protein interactions: inhibition of mammalian carbonic anhydrases I-XV by the murine inhibitor of carbonic anhydrase and other members of the transferrin family. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 5529-35	8.3	25
370	Multiple binding modes of inhibitors to carbonic anhydrases: how to design specific drugs targeting 15 different isoforms?. <i>Chemical Reviews</i> , 2012 , 112, 4421-68	68.1	889
369	Synthesis, characterization and biological studies of sulfonamide Schiff's bases and some of their metal derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 58-68	5.6	37
368	Hydroxamate represents a versatile zinc binding group for the development of new carbonic anhydrase inhibitors. <i>Chemical Communications</i> , 2012 , 48, 8838-40	5.8	58
367	Inhibition of carbonic anhydrase isozymes I and II with natural products extracted from plants, mushrooms and honey. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2012 , 27, 395-402	5.6	27
366	Sulfonamides: a patent review (2008 - 2012). <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 747-58	6.8	167
365	Amide derivatives of benzene-sulfonanilide, pharmaceutical composition thereof and method for cancer treatment using the same (US20120095092). <i>Expert Opinion on Therapeutic Patents</i> , 2012 , 22, 1251-5	6.8	4
364	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, 138-47	5.6	138
363	Microbial enzyme: applications in industry and in bioremediation. <i>Enzyme Research</i> , 2012 , 2012, 980681	2.4	9
362	Inhibition of V-ATPase and carbonic anhydrases as interference strategy with tumor acidification processes. <i>Current Pharmaceutical Design</i> , 2012 , 18, 1407-13	3.3	14
361	Recent developments in targeting carbonic anhydrase IX for cancer therapeutics. <i>Oncotarget</i> , 2012 , 3, 84-97	3.3	325
360	Interfering with pH regulation in tumours as a therapeutic strategy. <i>Nature Reviews Drug Discovery</i> , 2011 , 10, 767-77	64.1	1146
359	Natural product-based phenols as novel probes for mycobacterial and fungal carbonic anhydrases. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1682-92	8.3	85
358	Anticonvulsant 4-aminobenzenesulfonamide derivatives with branched-alkylamide moieties: X-ray crystallography and inhibition studies of human carbonic anhydrase isoforms I, II, VII, and XIV. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 3977-81	8.3	65
357	Glycosyl coumarin carbonic anhydrase IX and XII inhibitors strongly attenuate the growth of primary breast tumors. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 8271-7	8.3	201

356	Structural basis for the interaction between carbonic anhydrase and 1,2,3,4-tetrahydroisoquinolin-2-ylsulfonamides. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 2522-6	8.3	30
355	Selection of Carbonic Anhydrase IX Inhibitors from One Million DNA-Encoded Compounds. <i>ACS Chemical Biology</i> , 2011 , 6, 336-44	4.9	117
354	Targeting tumor hypoxia: suppression of breast tumor growth and metastasis by novel carbonic anhydrase IX inhibitors. <i>Cancer Research</i> , 2011 , 71, 3364-76	10.1	563
353	Carbonic anhydrase inhibitors: purification and inhibition studies of pigeon (<i>Columba livia</i> var. domestica) red blood cell carbonic anhydrase with sulfonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2011 , 26, 749-53	5.6	18
352	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, 1896-902	8.3	391
351	Pyridinium derivatives of histamine are potent activators of cytosolic carbonic anhydrase isoforms I, II and VII. <i>Organic and Biomolecular Chemistry</i> , 2011 , 9, 2790-800	3.9	25
350	Carbonic anhydrase activators: gold nanoparticles coated with derivatized histamine, histidine, and carnosine show enhanced activatory effects on several mammalian isoforms. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1170-7	8.3	43
349	Specific inhibition of carbonic anhydrase IX activity enhances the in vivo therapeutic effect of tumor irradiation. <i>Radiotherapy and Oncology</i> , 2011 , 99, 424-31	5.3	144
348	Associations of selenium status with cardiometabolic risk factors: an 8-year follow-up analysis of the Olivetti Heart study. <i>Atherosclerosis</i> , 2011 , 217, 274-8	3.1	67
347	Bacterial carbonic anhydrases as drug targets: toward novel antibiotics?. <i>Frontiers in Pharmacology</i> , 2011 , 2, 34	5.6	201
346	In Vitro inhibition of human carbonic anhydrase I and II isozymes with natural phenolic compounds. <i>Chemical Biology and Drug Design</i> , 2011 , 77, 494-9	2.9	154
345	Synthesis and biological profile of new 1,2,3,4-tetrahydroisoquinolines as selective carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 7003-7	3.4	17
344	Carbonic anhydrase I and II activation with mono- and dihalogenated histamine derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 4884-7	2.9	20
343	Synthesis of rhodamine B-benzenesulfonamide conjugates and their inhibitory activity against human and bacterial/fungal carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 5210-3	2.9	11
342	Promiscuity of carbonic anhydrase II. Unexpected ester hydrolysis of carbohydrate-based sulfamate inhibitors. <i>Journal of the American Chemical Society</i> , 2011 , 133, 18452-62	16.4	37
341	Carbonic anhydrase inhibition with natural products: novel chemotypes and inhibition mechanisms. <i>Molecular Diversity</i> , 2011 , 15, 305-16	3.1	56
340	Gene expression profiling of phytoplasma-infected Madagascar periwinkle leaves using differential display. <i>Molecular Biology Reports</i> , 2011 , 38, 2993-3000	2.8	22
339	A new coral carbonic anhydrase in <i>Stylophora pistillata</i> . <i>Marine Biotechnology</i> , 2011 , 13, 992-1002	3.4	72

338	A new α -carbonic anhydrase from <i>Brucella suis</i> , its cloning, characterization, and inhibition with sulfonamides and sulfamates, leading to impaired pathogen growth. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1172-8	3.4	72
337	Conformational variability of different sulfonamide inhibitors with thienyl-acetamido moieties attributes to differential binding in the active site of cytosolic human carbonic anhydrase isoforms. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 3732-8	3.4	45
336	The leader peptide of a human rec. MnSOD as molecular carrier which delivers high amounts of Cisplatin into tumor cells inducing a fast apoptosis in vitro. <i>International Journal of Cancer</i> , 2011 , 128, 453-9	7.5	10
335	Characterization and inhibition studies of an α -carbonic anhydrase from the endangered sturgeon species <i>Acipenser gueldenstaedti</i> . <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2011 , 26, 895-900 ^{5,6}	5.6	46
334	Carbonic anhydrase inhibitors: X-ray crystallographic studies for the binding of N-substituted benzenesulfonamides to human isoform II. <i>Chemical Communications</i> , 2011 , 47, 11636-8	5.8	42
333	Inhibition of the α -carbonic anhydrase from <i>Streptococcus pneumoniae</i> by inorganic anions and small molecules: Toward innovative drug design of anti-infectives?. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 243-8	3.4	69
332	Characterization and anions inhibition studies of an α -carbonic anhydrase from the teleost fish <i>Dicentrarchus labrax</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 744-8	3.4	55
331	Kinetic and docking studies of phenol-based inhibitors of carbonic anhydrase isoforms I, II, IX and XII evidence a new binding mode within the enzyme active site. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1381-9	3.4	89
330	Purification and inhibition studies with anions and sulfonamides of an α -carbonic anhydrase from the Antarctic seal <i>Leptonychotes weddellii</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 1847-51	3.4	7
329	Inhibition studies of the α -carbonic anhydrases from the bacterial pathogen <i>Salmonella enterica</i> serovar Typhimurium with sulfonamides and sulfamates. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 5023-30	3.4	45
328	An inhibitor-like binding mode of a carbonic anhydrase activator within the active site of isoform II. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 2764-8	2.9	25
327	Inhibition of α -carbonic anhydrases with ureido-substituted benzenesulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 102-5	2.9	26
326	Carbonic anhydrase inhibitors. Inhibition studies with anions and sulfonamides of a new cytosolic enzyme from the scleractinian coral <i>Stylophora pistillata</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 710-4	2.9	25
325	Carbonic anhydrase inhibitors. Inhibition of the α -class enzymes from the fungal pathogens <i>Candida albicans</i> and <i>Cryptococcus neoformans</i> with branched aliphatic/aromatic carboxylates and their derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 2521-6	2.9	28
324	Synthesis and crystallographic analysis of new sulfonamides incorporating NO-donating moieties with potent antiglaucoma action. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 3216-21	2.9	42
323	Inhibition studies with anions and small molecules of two novel α -carbonic anhydrases from the bacterial pathogen <i>Salmonella enterica</i> serovar Typhimurium. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 3591-5	2.9	63
322	In vitro inhibition of α -carbonic anhydrase isozymes by some phenolic compounds. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 4259-62	2.9	158
321	Sulfonamides incorporating 1,3,5-triazine moieties selectively and potently inhibit carbonic anhydrase transmembrane isoforms IX, XII and XIV over cytosolic isoforms I and II: Solution and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry</i> , 2011 , 19, 3105-19	3.4	82

320	Therapeutic compounds: patent evaluation of WO2011011652A1. <i>Expert Opinion on Therapeutic Patents</i> , 2011 , 21, 1491-5	6.8	1
319	Acetaldehyde-derived modifications on cytosolic human carbonic anhydrases. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2011 , 26, 862-70	5.6	18
318	Carbonic anhydrase inhibitors and activators for novel therapeutic applications. <i>Future Medicinal Chemistry</i> , 2011 , 3, 1165-80	4.1	240
317	Carbonic anhydrase inhibitors: Inhibition of human erythrocyte isozymes I and II with a series of phenolic acids. <i>Chemical Biology and Drug Design</i> , 2010 , 75, 515-20	2.9	114
316	Selective inhibition of carbonic anhydrase IX decreases cell proliferation and induces ceramide-mediated apoptosis in human cancer cells. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010 , 334, 710-9	4.7	81
315	Inhibition and binding studies of carbonic anhydrase isozymes I, II and IX with benzimidazo[1,2-c][1,2,3]thiadiazole-7-sulphonamides. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2010 , 25, 863-70	5.6	64
314	Carbonic anhydrase inhibition/activation: trip of a scientist around the world in the search of novel chemotypes and drug targets. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3233-45	3.3	108
313	The coumarin-binding site in carbonic anhydrase accommodates structurally diverse inhibitors: the antiepileptic lacosamide as an example and lead molecule for novel classes of carbonic anhydrase inhibitors. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 850-4	8.3	110
312	Deciphering the mechanism of carbonic anhydrase inhibition with coumarins and thiocoumarins. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 335-44	8.3	311
311	Cloning, characterization, and inhibition studies of a beta-carbonic anhydrase from <i>Brucella suis</i> . <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 2277-85	8.3	97
310	Identification of 3,4-Dihydroisoquinoline-2(1H)-sulfonamides as potent carbonic anhydrase inhibitors: synthesis, biological evaluation, and enzyme-ligand X-ray studies. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 2401-8	8.3	42
309	The molecular characterization of a novel GH38 β -mannosidase from the crenarchaeon <i>Sulfolobus solfataricus</i> revealed its ability in de-mannosylating glycoproteins. <i>Biochimie</i> , 2010 , 92, 1895-907	4.6	21
308	Synthesis and biological evaluation of a ^{99m}Tc -labelled sulfonamide conjugate for in vivo visualization of carbonic anhydrase IX expression in tumor hypoxia. <i>Nuclear Medicine and Biology</i> , 2010 , 37, 557-64	2.1	81
307	The first example of a significant active site conformational rearrangement in a carbonic anhydrase-inhibitor adduct: the carbonic anhydrase I-topiramate complex. <i>Organic and Biomolecular Chemistry</i> , 2010 , 8, 3528-33	3.9	34
306	Dietary sodium intake in a sample of adult male population in southern Italy: results of the Olivetti Heart Study. <i>European Journal of Clinical Nutrition</i> , 2010 , 64, 518-24	5.2	29
305	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, 8371-3	5.8	180
304	Polyamines inhibit carbonic anhydrases by anchoring to the zinc-coordinated water molecule. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 5511-22	8.3	184
303	The β -carbonic anhydrases from <i>Mycobacterium tuberculosis</i> as drug targets. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3300-9	3.3	70

302	3-phenyl-1H-indole-5-sulfonamides: structure-based drug design of a promising class of carbonic anhydrase inhibitors. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3317-26	3.3	15
301	Recent advances in structural studies of the carbonic anhydrase family: the crystal structure of human CA IX and CA XIII. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3246-54	3.3	27
300	Brucella carbonic anhydrases: new targets for designing anti-infective agents. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3310-6	3.3	46
299	Saccharomyces cerevisiae β -carbonic anhydrase: inhibition and activation studies. <i>Current Pharmaceutical Design</i> , 2010 , 16, 3327-36	3.3	16
298	Carbonic anhydrase activators: activation of the beta-carbonic anhydrases from the pathogenic fungi Candida albicans and Cryptococcus neoformans with amines and amino acids. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 1034-7	3.4	18
297	Carbonic anhydrase activators. The first activation study of a coral secretory isoform with amino acids and amines. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 2300-2303	3.4	25
296	Nanoscale enzyme inhibitors: fullerenes inhibit carbonic anhydrase by occluding the active site entrance. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 2822-8	3.4	63
295	Coumarinyl-substituted sulfonamides strongly inhibit several human carbonic anhydrase isoforms: solution and crystallographic investigations. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 4873-8	3.4	57
294	Identification of potent and selective human carbonic anhydrase VII (hCA VII) inhibitors. <i>ChemMedChem</i> , 2010 , 5, 823-6	3.7	21
293	Bidentate Zinc chelators for alpha-carbonic anhydrases that produce a trigonal bipyramidal coordination geometry. <i>ChemMedChem</i> , 2010 , 5, 1609-15	3.7	24
292	Inhibition studies of a beta-carbonic anhydrase from Brucella suis with a series of water soluble glycosyl sulfanilamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 2178-82	2.9	48
291	Carbonic anhydrase inhibitors. The beta-carbonic anhydrases from the fungal pathogens Cryptococcus neoformans and Candida albicans are strongly inhibited by substituted-phenyl-1H-indole-5-sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 2508-11	2.9	26
290	Carbonic anhydrase IX: Biochemical and crystallographic characterization of a novel antitumor target. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2010 , 1804, 404-9	4	151
289	Carbonic anhydrase inhibitors. Identification of selective inhibitors of the human mitochondrial isozymes VA and VB over the cytosolic isozymes I and II from a natural product-based phenolic library. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 14-8	3.4	63
288	Carbonic anhydrase inhibitors. Inhibition of mammalian isoforms I-XIV with a series of natural product polyphenols and phenolic acids. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 2159-2164	3.4	190
287	Carbonic anhydrase inhibitors. X-ray crystal studies of the carbonic anhydrase II-trithiocarbonate adduct--an inhibitor mimicking the sulfonamide and urea binding to the enzyme. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 474-8	2.9	74
286	Carbonic anhydrase activators: Activation of the beta-carbonic anhydrase from the pathogenic yeast Candida glabrata with amines and amino acids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 1701-4	2.9	17
285	Carbonic anhydrase inhibitors. Inhibition of transmembrane isoforms IX, XII, and XIV with less investigated anions including trithiocarbonate and dithiocarbamate. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 1548-50	2.9	45

284	Carbonic anhydrase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 3467-74	2.9	538
283	Coumarins incorporating hydroxy- and chloro-moieties selectively inhibit the transmembrane, tumor-associated carbonic anhydrase isoforms IX and XII over the cytosolic ones I and II. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 4511-4	2.9	118
282	Carbonic anhydrase inhibitors. The X-ray crystal structure of human isoform II in adduct with an adamantyl analogue of acetazolamide resides in a less utilized binding pocket than most hydrophobic inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 4376-81	2.9	66
281	Inhibition of the R1 fragment of the cadmium-containing zeta-class carbonic anhydrase from the diatom <i>Thalassiosira weissflogii</i> with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 4745-8	2.9	35
280	Carbonic anhydrase inhibitors. Antioxidant polyphenols effectively inhibit mammalian isoforms I-XV. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 5050-3	2.9	135
279	Crystal structure of the C183S/C217S mutant of human CA VII in complex with acetazolamide. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 5023-6	2.9	73
278	Paraoxon, 4-nitrophenyl phosphate and acetate are substrates of β but not of α and γ carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 6208-12	2.9	46
277	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, 7255-8	2.9	143
276	Characterization of the first beta-class carbonic anhydrase from an arthropod (<i>Drosophila melanogaster</i>) and phylogenetic analysis of beta-class carbonic anhydrases in invertebrates. <i>BMC Biochemistry</i> , 2010 , 11, 28	4.8	57
275	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, 16233-8	11.5	399
274	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, 20299-310	5.4	218
273	Therapeutic applications of glycosidic carbonic anhydrase inhibitors. <i>Medicinal Research Reviews</i> , 2009 , 29, 419-35	14.4	96
272	Novel organometallic cationic ruthenium(II) pentamethylcyclopentadienyl benzenesulfonamide complexes targeted to inhibit carbonic anhydrase. <i>Journal of Biological Inorganic Chemistry</i> , 2009 , 14, 935-45	3.7	27
271	Crystal structure of human carbonic anhydrase XIII and its complex with the inhibitor acetazolamide. <i>Proteins: Structure, Function and Bioinformatics</i> , 2009 , 74, 164-75	4.2	90
270	Which carbonic anhydrases are targeted by the antiepileptic sulfonamides and sulfamates?. <i>Chemical Biology and Drug Design</i> , 2009 , 74, 317-21	2.9	62
269	Carbonic anhydrase inhibitors: inhibition studies of a coral secretory isoform with inorganic anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 650-3	2.9	27
268	Carbonic anhydrase activators. Activation of the membrane-associated isoform XV with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 3430-3	2.9	11
267	Carbonic anhydrase inhibitors. Characterization and inhibition studies of the most active beta-carbonic anhydrase from <i>Mycobacterium tuberculosis</i> , Rv3588c. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 6649-54	2.9	91

266	Nitric oxide-donating carbonic anhydrase inhibitors for the treatment of open-angle glaucoma. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 6565-70	2.9	57
265	Carbonic anhydrase inhibitors. Comparison of chlorthalidone, indapamide, trichloromethiazide, and furosemide X-ray crystal structures in adducts with isozyme II, when several water molecules make the difference. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 1214-21	3.4	53
264	Carbonic anhydrase inhibitors: inhibition of the beta-class enzyme from the yeast <i>Saccharomyces cerevisiae</i> with sulfonamides and sulfamates. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 1158-63	3.4	78
263	Carbonic anhydrase inhibitors. The nematode alpha-carbonic anhydrase of <i>Caenorhabditis elegans</i> CAH-4b is highly inhibited by 2-(hydrazinocarbonyl)-3-substituted-phenyl-1H-indole-5-sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 3212-5	3.4	15
262	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, 3207-11	3.4	194
261	Carbonic anhydrase inhibitors. Inhibition of the beta-class enzymes from the fungal pathogens <i>Candida albicans</i> and <i>Cryptococcus neoformans</i> with aliphatic and aromatic carboxylates. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 2654-7	3.4	66
260	Carbonic anhydrase inhibitors. Inhibition of the human cytosolic isoforms I and II and transmembrane, tumor-associated isoforms IX and XII with boronic acids. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 3649-52	3.4	27
259	Synthesis and evaluation of pharmacological profile of 1-aryl-6,7-dimethoxy-3,4-dihydroisoquinoline-2(1H)-sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 3659-64	3.4	29
258	Carbonic anhydrase inhibitors. Inhibition and homology modeling studies of the fungal beta-carbonic anhydrase from <i>Candida albicans</i> with sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 4503-9	3.4	51
257	Carbonic anhydrase inhibitors. Inhibition studies of a coral secretory isoform by sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 5054-8	3.4	29
256	Carbonic anhydrase inhibitors. Aromatic/heterocyclic sulfonamides incorporating phenacetyl, pyridylacetyl and thienylacetyl tails act as potent inhibitors of human mitochondrial isoforms VA and VB. <i>Bioorganic and Medicinal Chemistry</i> , 2009 , 17, 4894-9	3.4	35
255	Carbonic anhydrase inhibitors: the membrane-associated isoform XV is highly inhibited by inorganic anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 1155-8	2.9	13
254	A thiabendazole sulfonamide shows potent inhibitory activity against mammalian and nematode alpha-carbonic anhydrases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 1371-5	2.9	18
253	Carbonic anhydrase activators: activation of the beta-carbonic anhydrase Nce103 from the yeast <i>Saccharomyces cerevisiae</i> with amines and amino acids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 1662-5	2.9	16
252	Carbonic anhydrase inhibitors. Inhibition of cytosolic isoforms I, II, III, VII and XIII with less investigated inorganic anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 1855-7	2.9	35
251	Carbonic anhydrase activators: activation of human isozymes I, II and IX with phenylsulfonylhydrazido l-histidine derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 2440-3	2.9	21
250	Carbonic anhydrase inhibitors. Inhibition of the fungal beta-carbonic anhydrases from <i>Candida albicans</i> and <i>Cryptococcus neoformans</i> with boronic acids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 2642-5	2.9	42
249	Carbonic anhydrase inhibitors. Phenacetyl-, pyridylacetyl- and thienylacetyl-substituted aromatic sulfonamides act as potent and selective isoform VII inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 3170-3	2.9	33

248	The protein tyrosine kinase inhibitors imatinib and nilotinib strongly inhibit several mammalian alpha-carbonic anhydrase isoforms. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4102-6	2.9	64
247	Carbonic anhydrase inhibitors. Inhibition of the beta-class enzyme from the pathogenic yeast <i>Candida glabrata</i> with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4802-5	2.9	39
246	Carbonic anhydrase inhibitors. Inhibition of the Rv1284 and Rv3273 beta-carbonic anhydrases from <i>Mycobacterium tuberculosis</i> with diazenylbenzenesulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4929-32	2.9	27
245	The proteoglycan region of the tumor-associated carbonic anhydrase isoform IX acts as an intrinsic buffer optimizing CO ₂ hydration at acidic pH values characteristic of solid tumors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 5825-8	2.9	73
244	Carbonic anhydrase inhibitors. Comparison of chlorthalidone and indapamide X-ray crystal structures in adducts with isozyme II: when three water molecules and the keto-enol tautomerism make the difference. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 322-8	8.3	49
243	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, 1207-20	6.5	176
242	Aspartic proteinases in Antarctic fish. <i>Marine Genomics</i> , 2009 , 2, 1-10	1.9	13
241	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 Suppl 1, 1-39	5.6	153
240	Carbonic anhydrase inhibitors. Cloning, characterization, and inhibition studies of a new beta-carbonic anhydrase from <i>Mycobacterium tuberculosis</i> . <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 3116-20	8.3	98
239	Molecular cloning, characterization, and inhibition studies of the Rv1284 beta-carbonic anhydrase from <i>Mycobacterium tuberculosis</i> with sulfonamides and a sulfamate. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 2226-32	8.3	85
238	Discovery of low nanomolar and subnanomolar inhibitors of the mycobacterial beta-carbonic anhydrases Rv1284 and Rv3273. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 4063-7	8.3	75
237	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, 3057-62	16.4	400
236	Carbonic anhydrase inhibitors. Cloning, characterization and inhibition studies of the cytosolic isozyme III with anions. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2009 , 24, 70-6	5.6	32
235	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, 423-8	5.3	173
234	Carbonic anhydrase inhibitors. Comparison of aliphatic sulfamate/bis-sulfamate adducts with isozymes II and IX as a platform for designing tight-binding, more isoform-selective inhibitors. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 5990-8	8.3	19
233	Carbonic anhydrases: novel therapeutic applications for inhibitors and activators. <i>Nature Reviews Drug Discovery</i> , 2008 , 7, 168-81	64.1	2297
232	Development of small molecule carbonic anhydrase IX inhibitors. <i>BJU International</i> , 2008 , 101 Suppl 4, 39-40	5.6	36
231	External pH influences the transcriptional profile of the carbonic anhydrase, CAH-4b in <i>Caenorhabditis elegans</i> . <i>Molecular and Biochemical Parasitology</i> , 2008 , 161, 140-9	1.9	25

230	Carbonic anhydrase inhibitors. Sulfonamide diuretics revisited--old leads for new applications?. <i>Organic and Biomolecular Chemistry</i> , 2008 , 6, 2499-506	3.9	74
229	Biochemical characterization of CA IX, one of the most active carbonic anhydrase isozymes. <i>Journal of Biological Chemistry</i> , 2008 , 283, 27799-27809	5.4	224
228	Carbonic anhydrase inhibitor coated gold nanoparticles selectively inhibit the tumor-associated isoform IX over the cytosolic isozymes I and II. <i>Journal of the American Chemical Society</i> , 2008 , 130, 16130-16144	16.4	99
227	Tumor-associated carbonic anhydrase 9 spatially coordinates intracellular pH in three-dimensional multicellular growths. <i>Journal of Biological Chemistry</i> , 2008 , 283, 20473-83	5.4	172
226	Carbonic anhydrase in the scleractinian coral <i>Stylophora pistillata</i> : characterization, localization, and role in biomineralization. <i>Journal of Biological Chemistry</i> , 2008 , 283, 25475-25484	5.4	192
225	Carbonic anhydrases--an overview. <i>Current Pharmaceutical Design</i> , 2008 , 14, 603-14	3.3	397
224	The alpha and beta classes carbonic anhydrases from <i>Helicobacter pylori</i> as novel drug targets. <i>Current Pharmaceutical Design</i> , 2008 , 14, 622-30	3.3	175
223	Diuretics: from classical carbonic anhydrase inhibitors to novel applications of the sulfonamides. <i>Current Pharmaceutical Design</i> , 2008 , 14, 641-8	3.3	194
222	Carbonic anhydrase activation and the drug design. <i>Current Pharmaceutical Design</i> , 2008 , 14, 708-15	3.3	73
221	Design of zinc binding functions for carbonic anhydrase inhibitors. <i>Current Pharmaceutical Design</i> , 2008 , 14, 615-21	3.3	69
220	The alpha-carbonic anhydrase from the malaria parasite and its inhibition. <i>Current Pharmaceutical Design</i> , 2008 , 14, 631-40	3.3	66
219	Are carbonic anhydrase inhibitors suitable for obtaining antiobesity drugs?. <i>Current Pharmaceutical Design</i> , 2008 , 14, 655-60	3.3	137
218	Anticonvulsant sulfonamides/sulfamates/sulfamides with carbonic anhydrase inhibitory activity: drug design and mechanism of action. <i>Current Pharmaceutical Design</i> , 2008 , 14, 661-71	3.3	110
217	Cloning, polymorphism, and inhibition of beta-carbonic anhydrase of <i>Helicobacter pylori</i> . <i>Journal of Gastroenterology</i> , 2008 , 43, 849-57	6.9	40
216	Carbonic anhydrase inhibitors: binding of indanesulfonamides to the human isoform II. <i>ChemMedChem</i> , 2008 , 3, 473-7	3.7	10
215	Carbonic anhydrase inhibitors: inhibition of human cytosolic isozymes I and II and tumor-associated isozymes IX and XII with S-substituted 4-chloro-2-mercapto-5-methyl-benzenesulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 3933-40	3.4	27
214	Carbonic anhydrase activators: activation of the human tumor-associated isozymes IX and XII with amino acids and amines. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 3530-6	3.4	44
213	Carbonic anhydrase inhibitors: inhibition of mammalian isoforms I-XIV with a series of substituted phenols including paracetamol and salicylic acid. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 7424-8	3.4	114

212	Carbonic anhydrase activators: kinetic and X-ray crystallographic study for the interaction of D- and L-tryptophan with the mammalian isoforms I-XIV. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 8373-8	3.4	60
211	In vitro inhibition of salicylic acid derivatives on human cytosolic carbonic anhydrase isozymes I and II. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 9101-5	3.4	142
210	Carbonic anhydrase inhibitors. Interaction of 2-(hydrazinocarbonyl)-3-phenyl-1H-indole-5-sulfonamide with 12 mammalian isoforms: kinetic and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 152-8	2.9	43
209	Carbonic anhydrase inhibitors. Interaction of 2-N,N-dimethylamino-1,3,4-thiadiazole-5-methanesulfonamide with 12 mammalian isoforms: kinetic and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 999-1005	2.9	20
208	Carbonic anhydrase inhibitors: interactions of phenols with the 12 catalytically active mammalian isoforms (CA I-XIV). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 1583-7	2.9	170
207	Carbonic anhydrase inhibitors. Interaction of indapamide and related diuretics with 12 mammalian isozymes and X-ray crystallographic studies for the indapamide-isozyme II adduct. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2567-73	2.9	59
206	Carbonic anhydrase activators: Activation of the human cytosolic isozyme III and membrane-associated isoform IV with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 4303-7	2.9	29
205	Carbonic anhydrase inhibitors: inhibition of the beta-class enzymes from the fungal pathogens <i>Candida albicans</i> and <i>Cryptococcus neoformans</i> with simple anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 5066-70	2.9	90
204	Carbonic anhydrase inhibitors: inhibition of <i>Plasmodium falciparum</i> carbonic anhydrase with aromatic/heterocyclic sulfonamides-in vitro and in vivo studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 5466-71	2.9	61
203	Carbonic anhydrase activators: activation of the archaeal beta-class (Cab) and gamma-class (Cam) carbonic anhydrases with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 6194-8	2.9	33
202	Carbonic anhydrase inhibitors. Inhibition of the beta-class enzyme from the yeast <i>Saccharomyces cerevisiae</i> with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 6327-31	2.9	44
201	Investigations of the esterase, phosphatase, and sulfatase activities of the cytosolic mammalian carbonic anhydrase isoforms I, II, and XIII with 4-nitrophenyl esters as substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2267-71	2.9	88
200	Carbonic anhydrase inhibitors: the X-ray crystal structure of ethoxzolamide complexed to human isoform II reveals the importance of thr200 and gln92 for obtaining tight-binding inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2669-74	2.9	30
199	Carbonic anhydrase inhibitors: Inhibition of the new membrane-associated isoform XV with phenols. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 3593-6	2.9	56
198	Carbonic anhydrase inhibitors as emerging drugs for the treatment of obesity. <i>Expert Opinion on Emerging Drugs</i> , 2008 , 13, 383-92	3.7	139
197	Carbonic anhydrase inhibitors. DNA cloning, characterization, and inhibition studies of the human secretory isoform VI, a new target for sulfonamide and sulfamate inhibitors. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 381-8	8.3	80
196	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, 1651-7	8.3	169
195	Saccharin inhibits carbonic anhydrases: possible explanation for its unpleasant metallic aftertaste. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 7697-9	16.4	145

194	Carbonic anhydrase activators: the first activation study of the human secretory isoform VI with amino acids and amines. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 5351-7	3-4	42
193	Molecular modeling study for the binding of zonisamide and topiramate to the human mitochondrial carbonic anhydrase isoform VA. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 4152-8	3-4	34
192	Carbonic anhydrases as targets for medicinal chemistry. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 4336-50	3-4	462
191	Design, synthesis, and docking studies of new 1,3,4-thiadiazole-2-thione derivatives with carbonic anhydrase inhibitory activity. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 6975-84	3-4	65
190	Carbonic anhydrase inhibitors: the inhibition profiles of the human mitochondrial isoforms VA and VB with anions are very different. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 6742-7	3-4	16
189	Carbonic anhydrase inhibitors: cloning, characterization, and inhibition studies of the cytosolic isozyme III with sulfonamides. <i>Bioorganic and Medicinal Chemistry</i> , 2007 , 15, 7229-36	3-4	90
188	Carbonic anhydrase activators: an activation study of the human mitochondrial isoforms VA and VB with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 1336-40	2-9	41
187	Carbonic anhydrase inhibitors: binding of an antiglaucoma glycosyl-sulfanilamide derivative to human isoform II and its consequences for the drug design of enzyme inhibitors incorporating sugar moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 1726-31	2-9	35
186	Carbonic anhydrase inhibitors. Inhibition of transmembrane isozymes XII (cancer-associated) and XIV with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 1532-7	2-9	33
185	Carbonic anhydrase inhibitors: the beta-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, 3585-94	2-9	146
184	Carbonic anhydrase activators: activation of the human isoforms VII (cytosolic) and XIV (transmembrane) with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 4107-12	2-9	45
183	Carbonic anhydrase inhibitors. Interaction of the antiepileptic drug sulthiame with twelve mammalian isoforms: kinetic and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 4866-72	2-9	34
182	Phosph(on)ate as a zinc-binding group in metalloenzyme inhibitors: X-ray crystal structure of the antiviral drug foscarnet complexed to human carbonic anhydrase I. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 2210-5	2-9	45
181	Carbonic anhydrase inhibitors: inhibition of human, bacterial, and archaeal isozymes with benzene-1,3-disulfonamides--solution and crystallographic studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 4201-7	2-9	45
180	Purification and characterization of pepsins A1 and A2 from the Antarctic rock cod <i>Trematomus bernacchii</i> . <i>FEBS Journal</i> , 2007 , 274, 6152-66	5-7	33
179	Differential display analysis of gene expression in Etrog citron leaves infected by Citrus viroid III. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 2007 , 1769, 228-35		32
178	Carbonic anhydrase activators: L-Adrenaline plugs the active site entrance of isozyme II, activating better isoforms I, IV, VA, VII, and XIV. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 628-35	2-9	82
177	Carbonic anhydrase inhibitors. Inhibition studies of the human secretory isoform VI with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007 , 17, 1037-42	2-9	32

176	Malarial parasite carbonic anhydrase and its inhibitors. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 909-17	3	40
175	Inhibitors of HIV-1 protease: current state of the art 10 years after their introduction. From antiretroviral drugs to antifungal, antibacterial and antitumor agents based on aspartic protease inhibitors. <i>Current Medicinal Chemistry</i> , 2007 , 14, 2734-48	4.3	115
174	The development of topically acting carbonic anhydrase inhibitors as anti-glaucoma agents. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 849-54	3	49
173	Carbonic anhydrases as drug targets--an overview. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 825-33	3	171
172	Inhibition of the archaeal beta-class (Cab) and gamma-class (Cam) carbonic anhydrases. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 901-8	3	113
171	Carbonic anhydrase inhibitors as anticonvulsant agents. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 855-64	3	178
170	Antiobesity carbonic anhydrase inhibitors. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 879-84	3	82
169	Therapeutic applications of the carbonic anhydrase inhibitors. <i>Therapy: Open Access in Clinical Medicine</i> , 2007 , 4, 355-378		25
168	Imaging the hypoxia surrogate marker CA IX requires expression and catalytic activity for binding fluorescent sulfonamide inhibitors. <i>Radiotherapy and Oncology</i> , 2007 , 83, 367-73	5.3	138
167	Synthesis and antimalarial activity of novel chiral and achiral benzenesulfonamides bearing 1, 3, 4-oxadiazole moieties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2007 , 22, 301-8	5.6	48
166	Carbonic anhydrase inhibitors: Valdecoxib binds to a different active site region of the human isoform II as compared to the structurally related cyclooxygenase II "selective" inhibitor celecoxib. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 437-42	2.9	89
165	Carbonic anhydrase inhibitors: inhibition of the cytosolic human isozyme VII with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 3139-43	2.9	26
164	Carbonic anhydrase activators: activation of isozyme XIII with amino acids and amines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 3955-9	2.9	45
163	Carbonic anhydrase inhibitors: X-ray crystallographic studies for the binding of 5-amino-1,3,4-thiadiazole-2-sulfonamide and 5-(4-amino-3-chloro-5-fluorophenylsulfonamido)-1,3,4-thiadiazole-2-sulfonamide to human isoform II. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 1684-8	2.9	31
162	Carbonic anhydrase activators. Activation of isozymes I, II, IV, VA, VII, and XIV with L- and D-histidine and crystallographic analysis of their adducts with isoform II: engineering proton-transfer processes within the active site of an enzyme. <i>Chemistry - A European Journal</i> , 2006 , 12, 7057-66	4.8	122
161	Therapeutic potential of sulfamides as enzyme inhibitors. <i>Medicinal Research Reviews</i> , 2006 , 26, 767-92	14.4	153
160	Inhibitors of HIV-1 protease: 10 years after. <i>Expert Opinion on Therapeutic Patents</i> , 2006 , 16, 1067-1091	6.8	18
159	Carbonic anhydrase activators. Activation of isoforms I, II, IV, VA, VII, and XIV with L- and D-phenylalanine and crystallographic analysis of their adducts with isozyme II: stereospecific recognition within the active site of an enzyme and its consequences for the drug design. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 3019-27	8.3	116

158	2-substituted estradiol bis-sulfamates, multitargeted antitumor agents: synthesis, in vitro SAR, protein crystallography, and in vivo activity. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 7683-96	8.3	91
157	A novel class of carbonic anhydrase inhibitors: glycoconjugate benzene sulfonamides prepared by "click-tailing". <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 6539-48	8.3	153
156	Indanesulfonamides as carbonic anhydrase inhibitors. Toward structure-based design of selective inhibitors of the tumor-associated isozyme CA IX. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 2743-9	8.3	54
155	Carbonic anhydrase inhibitors: Hypoxia-activatable sulfonamides incorporating disulfide bonds that target the tumor-associated isoform IX. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 5544-51	8.3	93
154	Oxovanadium(IV) complexes of hydrazides: potential antifungal agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2006 , 21, 37-42	5.6	31
153	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, 7024-31	8.3	142
152	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, 8329-35	16.4	186
151	Carbonic anhydrase inhibitors: DNA cloning and inhibition studies of the alpha-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, 2117-26	8.3	137
150	Carbonic anhydrase inhibitors and activators and their use in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2006 , 16, 1627-1664	6.8	143
149	Targeting tumor-associated carbonic anhydrase IX in cancer therapy. <i>Trends in Pharmacological Sciences</i> , 2006 , 27, 566-73	13.2	321
148	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, 83131	4.2	124
147	Carbonic anhydrase inhibitors: cloning and sulfonamide inhibition studies of a carboxyterminal truncated alpha-carbonic anhydrase from <i>Helicobacter pylori</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 2182-8	2.9	36
146	Carbonic anhydrase activators: the first X-ray crystallographic study of an adduct of isoform I. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 5152-6	2.9	74
145	Metal detoxification and homeostasis in Antarctic Notothenioids. A comparative survey on evolution, expression and functional properties of fish and mammal metallothioneins. <i>Reviews in Environmental Science and Biotechnology</i> , 2006 , 5, 253-267	13.9	4
144	Metal detoxification and homeostasis in Antarctic Notothenioids. A comparative survey on evolution, expression and functional properties of fish and mammal metallothioneins 2006 , 369-383		
143	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, 4834-41	8.3	192
142	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, 5721-7	8.3	150
141	In-vitro antibacterial, antifungal and cytotoxic activities of some coumarins and their metal complexes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2005 , 20, 333-40	5.6	111

140	Effect of cadmium on gene expression in the liverwort <i>Lunularia cruciata</i> . <i>Gene</i> , 2005 , 356, 153-9	3.8	17
139	Carbonic anhydrase inhibitors. The mitochondrial isozyme VB as a new target for sulfonamide and sulfamate inhibitors. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 7860-6	8.3	161
138	Carbonic anhydrase inhibitors: synthesis and inhibition of cytosolic/membrane-associated carbonic anhydrase isozymes I, II, and IX with sulfonamides incorporating hydrazino moieties. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 2121-5	8.3	69
137	Metal binding and antibacterial activity of ciprofloxacin complexes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2005 , 20, 303-7	5.6	130
136	Structural and functional studies of vertebrate metallothioneins: cross-talk between domains in the absence of physical contact. <i>Biochemical Journal</i> , 2005 , 391, 95-103	3.8	14
135	Carbonic anhydrase inhibitors. Inhibition of <i>Plasmodium falciparum</i> carbonic anhydrase with aromatic sulfonamides: towards antimalarials with a novel mechanism of action?. <i>Bioorganic and Medicinal Chemistry</i> , 2005 , 13, 483-9	3.4	75
134	Carbonic anhydrase inhibitors: synthesis and inhibition of cytosolic/tumor-associated carbonic anhydrase isozymes I, II, and IX with bis-sulfamates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 579-84	2.9	41
133	Carbonic anhydrase inhibitors. Inhibition of the human cytosolic isozyme VII with aromatic and heterocyclic sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 971-6	2.9	128
132	Carbonic anhydrase inhibitors: X-ray crystal structure of a benzenesulfonamide strong CA II and CA IX inhibitor bearing a pentafluorophenylaminothioureido tail in complex with isozyme II. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 1937-42	2.9	38
131	Carbonic anhydrase inhibitors: synthesis and inhibition of cytosolic/tumor-associated carbonic anhydrase isozymes I, II, IX, and XII with N-hydroxysulfamides--a new zinc-binding function in the design of inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 2353-8	2.9	43
130	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, 2315-20	2.9	166
129	Carbonic anhydrase inhibitors. Inhibition of cytosolic/tumor-associated carbonic anhydrase isozymes I, II, IX, and XII with Schiff's bases incorporating chromone and aromatic sulfonamide moieties, and their zinc complexes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 3096-101	2.9	106
128	Carbonic anhydrase activators: X-ray crystal structure of the adduct of human isozyme II with L-histidine as a platform for the design of stronger activators. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 5136-41	2.9	91
127	Sulfamates and their therapeutic potential. <i>Medicinal Research Reviews</i> , 2005 , 25, 186-228	14.4	169
126	Carbonic anhydrase inhibitors. Inhibition of the transmembrane isozyme XII with sulfonamides-a new target for the design of antitumor and antiglaucoma drugs?. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 963-9	2.9	199
125	Carbonic anhydrase inhibitors: novel sulfonamides incorporating 1,3,5-triazine moieties as inhibitors of the cytosolic and tumour-associated carbonic anhydrase isozymes I, II and IX. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 3102-8	2.9	129
124	Carbonic anhydrase inhibitors: design of thioureido sulfonamides with potent isozyme II and XII inhibitory properties and intraocular pressure lowering activity in a rabbit model of glaucoma. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 3821-7	2.9	25
123	Carbonic anhydrase inhibitors: inhibition of the transmembrane isozyme XIV with sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 3828-33	2.9	125

122	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, 1805-43	3.3	199
121	Characterization of CA XIII, a novel member of the carbonic anhydrase isozyme family. <i>Journal of Biological Chemistry</i> , 2004 , 279, 2719-27	5.4	187
120	COX-2 selective inhibitors, carbonic anhydrase inhibition and anticancer properties of sulfonamides belonging to this class of pharmacological agents. <i>Mini-Reviews in Medicinal Chemistry</i> , 2004 , 4, 625-32	3.2	116
119	Protein tyrosine kinase inhibitors as anticancer agents. <i>Expert Opinion on Therapeutic Patents</i> , 2004 , 14, 35-53	6.8	29
118	Carbonic anhydrases: current state of the art, therapeutic applications and future prospects. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2004 , 19, 199-229	5.6	532
117	Accumulation, localisation, and toxic effects of cadmium in the liverwort <i>Lunularia cruciata</i> . <i>Protoplasma</i> , 2004 , 223, 53-61	3.4	57
116	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, 217-23	2.9	235
115	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, 231-4	2.9	137
114	Carbonic anhydrase inhibitors. Inhibition of the newly isolated murine isozyme XIII with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 5435-9	2.9	42
113	Carbonic anhydrase inhibitors. Inhibition of the prokariotic beta and gamma-class enzymes from Archaea with sulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 6001-6	2.9	77
112	Carbonic anhydrase inhibitors: aromatic and heterocyclic sulfonamides incorporating adamantyl moieties with strong anticonvulsant activity. <i>Bioorganic and Medicinal Chemistry</i> , 2004 , 12, 2717-26	3.4	82
111	Carbonic anhydrase inhibitors: synthesis and inhibition of cytosolic/tumor-associated carbonic anhydrase isozymes I, II, and IX with sulfonamides incorporating 1,2,4-triazine moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 5427-33	2.9	90
110	Carbonic anhydrase inhibitors: X-ray crystallographic structure of the adduct of human isozyme II with the antipsychotic drug sulpiride. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 337-41	2.9	62
109	Carbonic anhydrase inhibitors: the first selective, membrane-impermeant inhibitors targeting the tumor-associated isozyme IX. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 869-73	2.9	140
108	Carbonic anhydrase inhibitors: X-ray crystallographic structure of the adduct of human isozyme II with a topically acting antiglaucoma sulfonamide. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 2357-61	2.9	46
107	Carbonic anhydrase inhibitors. Inhibition of the zinc and cobalt gamma-class enzyme from the archaeon <i>Methanosarcina thermophila</i> with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 3327-31	2.9	14
106	Carbonic anhydrase inhibitors. Inhibition of the beta-class enzyme from the methanoarchaeon <i>Methanobacterium thermoautotrophicum</i> (Cab) with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 4563-7	2.9	47
105	Carbonic anhydrase inhibitors: inhibition of the membrane-bound human isozyme IV with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 5769-73	2.9	26

104	Benzolamide is not a membrane-impermeant carbonic anhydrase inhibitor. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2004 , 19, 269-73	5.6	31
103	Carbonic anhydrase inhibitors. Design of selective, membrane-impermeant inhibitors targeting the human tumor-associated isozyme IX. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 2337-47	8.3	145
102	Carbonic anhydrase inhibitors. Inhibition of the zinc and cobalt γ -class enzyme from the archaeon <i>Methanosarcina thermophila</i> with anions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 3327-3331	2.9	57
101	Unexpected nanomolar inhibition of carbonic anhydrase by COX-2-selective celecoxib: new pharmacological opportunities due to related binding site recognition. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 550-7	8.3	381
100	Modulation of carbonic anhydrase activity and its applications in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2004 , 14, 667-702	6.8	148
99	Carbonic anhydrase inhibitors: the first on-resin screening of a 4-sulfamoylphenylthiourea library. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 5224-9	8.3	46
98	Adaptive evolution and functional divergence of pepsin gene family. <i>Gene</i> , 2004 , 333, 81-90	3.8	38
97	Identification of genes expressed in response to phytoplasma infection in leaves of <i>Prunus armeniaca</i> by messenger RNA differential display. <i>Gene</i> , 2004 , 332, 29-34	3.8	46
96	Gene amplification and cold adaptation of pepsin in Antarctic fish. A possible strategy for food digestion at low temperature. <i>Gene</i> , 2004 , 336, 195-205	3.8	31
95	Hypoxia activates the capacity of tumor-associated carbonic anhydrase IX to acidify extracellular pH. <i>FEBS Letters</i> , 2004 , 577, 439-45	3.8	556
94	Quantum theoretic QSAR of benzene derivatives: some enzyme inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2004 , 19, 237-48	5.6	17
93	New advances in HIV entry inhibitors development. <i>Current Drug Targets Infectious Disorders</i> , 2004 , 4, 339-55		31
92	Designing of Novel Carbonic Anhydrase Inhibitors and Activators. <i>Current Medicinal Chemistry Cardiovascular and Hematological Agents</i> , 2004 , 2, 51-70		2
91	Designing of novel carbonic anhydrase inhibitors and activators. <i>Current Medicinal Chemistry Cardiovascular and Hematological Agents</i> , 2004 , 2, 49-68		27
90	Anticancer and antiviral sulfonamides. <i>Current Medicinal Chemistry</i> , 2003 , 10, 925-53	4.3	557
89	Phylogenetic divergence of fish and mammalian metallothionein: relationships with structural diversification and organismal temperature. <i>Journal of Molecular Evolution</i> , 2003 , 57 Suppl 1, S250-7	3.1	21
88	Solution structure of MT _{nc} , a novel metallothionein from the Antarctic fish <i>Notothenia coriiceps</i> . <i>Structure</i> , 2003 , 11, 435-43	5.2	47
87	Carbonic anhydrase inhibitors. <i>Medicinal Research Reviews</i> , 2003 , 23, 146-89	14.4	1062

86	Protease inhibitors of the sulfonamide type: anticancer, antiinflammatory, and antiviral agents. <i>Medicinal Research Reviews</i> , 2003 , 23, 535-58	14.4	320
85	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, 841-5	2.9	209
84	Hydroxyurea is a carbonic anhydrase inhibitor. <i>Bioorganic and Medicinal Chemistry</i> , 2003 , 11, 2241-6	3.4	36
83	Carbonic anhydrase inhibitors. Inhibition of tumor-associated isozyme IX by halogenosulfanilamide and halogenophenylaminobenzolamide derivatives. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 2187-96	8.3	133
82	Zinc complexes of benzothiazole-derived Schiff bases with antibacterial activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2003 , 18, 259-63	5.6	130
81	Direct extracellular interaction between carbonic anhydrase IV and the human NBC1 sodium/bicarbonate co-transporter. <i>Biochemistry</i> , 2003 , 42, 12321-9	3.2	142
80	Indisulam: an anticancer sulfonamide in clinical development. <i>Expert Opinion on Investigational Drugs</i> , 2003 , 12, 283-7	5.9	146
79	Carbonic anhydrase inhibitors. inhibition of cytosolic isozymes I and II and transmembrane, cancer-associated isozyme IX with anions. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2003 , 18, 403-6	5.6	53
78	Carbonic anhydrase inhibitors in the treatment and prophylaxis of obesity. <i>Expert Opinion on Therapeutic Patents</i> , 2003 , 13, 1545-1550	6.8	135
77	Bacterial protease inhibitors. <i>Medicinal Research Reviews</i> , 2002 , 22, 329-72	14.4	128
76	Carbonic anhydrase activators: human isozyme II is strongly activated by oligopeptides incorporating the carboxyterminal sequence of the bicarbonate anion exchanger AE1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002 , 12, 1177-80	2.9	60
75	Stability and conformational dynamics of metallothioneins from the antarctic fish <i>Notothenia coriiceps</i> and mouse. <i>Proteins: Structure, Function and Bioinformatics</i> , 2002 , 46, 259-67	4.2	27
74	Carbonic anhydrase inhibitors: anticonvulsant sulfonamides incorporating valproyl and other lipophilic moieties. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 312-20	8.3	129
73	Sulfonamide derivatives with protease inhibitory action as anticancer, anti-inflammatory and antiviral agents. <i>Expert Opinion on Therapeutic Patents</i> , 2002 , 12, 1307-1327	6.8	46
72	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, 3583-7	8.3	144
71	Unsymmetrical 1,1'-disubstituted ferrocenes: synthesis of Co(ii), Cu(ii), Ni(ii) and Zn(ii) chelates of ferrocenyl -1-thiadiazolo-1'-tetrazole, -1-thiadiazolo-1'-triazole and -1-tetrazolo-1'-triazole with antimicrobial properties. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2002 , 17, 261-6	5.6	115
70	Applications of carbonic anhydrase inhibitors and activators in therapy. <i>Expert Opinion on Therapeutic Patents</i> , 2002 , 12, 217-242	6.8	228
69	Carbonic anhydrase activators: design of high affinity isozymes I, II, and IV activators, incorporating tri-/tetrasubstituted-pyridinium-azole moieties. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 504-10	8.3	68

68	Carbonic anhydrase activators: high affinity isozymes I, II, and IV activators, incorporating a beta-alanyl-histidine scaffold. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 284-91	8.3	60
67	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, 1466-76	8.3	127
66	Identification of cadmium-sensitive genes in the Antarctic fish <i>Chionodraco hamatus</i> by messenger RNA differential display. <i>Gene</i> , 2002 , 299, 117-24	3.8	32
65	Structural characterization and thermal stability of <i>Notothenia coriiceps</i> metallothionein. <i>Biochemical Journal</i> , 2001 , 354, 291-9	3.8	18
64	Structural characterization and thermal stability of <i>Notothenia coriiceps</i> metallothionein. <i>Biochemical Journal</i> , 2001 , 354, 291-299	3.8	22
63	Carbonic anhydrase inhibitors: synthesis of sulfonamides incorporating dtpa tails and of their zinc complexes with powerful topical antiglaucoma properties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2001 , 11, 575-82	2.9	106
62	Protease inhibitors: synthesis of a series of bacterial collagenase inhibitors of the sulfonyl amino acyl hydroxamate type. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 2253-8	8.3	34
61	Arylsulfonyl-N,N-dialkyl-dithiocarbamates as tumor cell growth inhibitors: novel agents targeting beta-tubulin?. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2001 , 16, 55-63		9
60	Structural and functional analysis of metal regulatory elements in the promoter region of genes encoding metallothionein isoforms in the Antarctic fish <i>Chionodraco hamatus</i> (icefish). <i>Gene</i> , 2001 , 274, 199-208	3.8	37
59	Bacterial proteases: current therapeutic use and future prospects for the development of new antibiotics. <i>Expert Opinion on Therapeutic Patents</i> , 2001 , 11, 221-259	6.8	99
58	Transition Metal Ion Complexes of Schiff-bases. Synthesis, Characterization and Antibacterial Properties. <i>Metal-Based Drugs</i> , 2001 , 8, 137-43		73
57	Arylsulfonyl-N,N-diethyl-dithiocarbamates: a novel class of antitumor agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000 , 10, 1887-91	2.9	70
56	Carbonic anhydrase inhibitors: sulfonamides incorporating furan-, thiophene- and pyrrole-carboxamido groups possess strong topical intraocular pressure lowering properties as aqueous suspensions. <i>Bioorganic and Medicinal Chemistry</i> , 2000 , 8, 2145-55	3.4	61
55	Mechanism of cyanamide hydration catalyzed by carbonic anhydrase II suggested by cryogenic X-ray diffraction. <i>Biochemistry</i> , 2000 , 39, 12391-7	3.2	42
54	Tissue-specific regulation of metallothionein and metallothionein mRNA accumulation in the Antarctic notothenioid, <i>Notothenia coriiceps</i> . <i>Polar Biology</i> , 2000 , 23, 17-23	2	16
53	Susceptibility to heavy metals and cadmium accumulation in aerobic and anaerobic thermophilic microorganisms isolated from deep-sea hydrothermal vents. <i>Current Microbiology</i> , 2000 , 41, 201-5	2.4	31
52	Antifungal Activity of Ag(I) and Zn(II) Complexes of Sulfacetamide Derivatives. <i>Metal-Based Drugs</i> , 2000 , 7, 49-54		22
51	Metallothionein in Antarctic notothenioids: Genetic polymorphism and differential gene expression. <i>Italian Journal of Zoology</i> , 2000 , 67, 13-20		1

50	Protease inhibitors: synthesis of potent bacterial collagenase and matrix metalloproteinase inhibitors incorporating N-4-nitrobenzylsulfonylglycine hydroxamate moieties. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 1858-65	8.3	56
49	Carbonic anhydrase activators: synthesis of high affinity isozymes I, II and IV activators, derivatives of 4-(4-tosylureido-amino acyl)ethyl-1H-imidazole (histamine derivatives). <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2000 , 15, 139-61		28
48	Aspartic proteinases from Antarctic fish. A biochemical and molecular approach. <i>Italian Journal of Zoology</i> , 2000 , 67, 21-26		
47	Carbonic anhydrase inhibitors: inhibition of isozymes I, II and IV by sulfamide and sulfamic acid derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2000 , 15, 443-53		28
46	Carbonic anhydrase inhibitors: water-soluble 4-sulfamoylphenylthioureas as topical intraocular pressure-lowering agents with long-lasting effects. <i>Journal of Medicinal Chemistry</i> , 2000 , 43, 4884-92	8.3	137
45	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, 4542-51	8.3	128
44	Novel carbonic anhydrase isozymes I, II and IV activators incorporating sulfonyl-histamino moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999 , 9, 2043-8	2.9	31
43	Carbonic anhydrase inhibitors: synthesis of water-soluble, topically effective intraocular pressure lowering aromatic/heterocyclic sulfonamides containing 8-quinoline-sulfonyl moieties: is the tail more important than the ring?. <i>Bioorganic and Medicinal Chemistry</i> , 1999 , 7, 2397-406	3.4	148
42	Carbonic anhydrase activators: amino acyl/dipeptidyl histamine derivatives bind with high affinity to isozymes I, II and IV and act as efficient activators. <i>Bioorganic and Medicinal Chemistry</i> , 1999 , 7, 2915-23	3.4	25
41	Carbonic anhydrase inhibitors [Part 57: Quantum chemical QSAR of a group of 1,3,4-thiadiazole- and 1,3,4-thiadiazoline disulfonamides with carbonic anhydrase inhibitory properties. <i>European Journal of Medicinal Chemistry</i> , 1999 , 34, 41-50	6.8	115
40	Carbonic anhydrase inhibitors. Part 61. Quantum chemical QSAR of a group of benzenedisulfonamides. <i>European Journal of Medicinal Chemistry</i> , 1999 , 34, 463-474	6.8	74
39	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, 2641-50	8.3	250
38	Carbonic anhydrase catalyzes cyanamide hydration to urea: is it mimicking the physiological reaction?. <i>Journal of Biological Inorganic Chemistry</i> , 1999 , 4, 528-36	3.7	52
37	Cathepsin D from the liver of the antarctic icefish <i>Chionodraco hamatus</i> exhibits unusual activity and stability at high temperatures ¹ . <i>BBA - Proteins and Proteomics</i> , 1999 , 1431, 64-73		31
36	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, 3690-700	8.3	146
35	Carbonic anhydrase inhibitors: N-cyanosulfonamides, a new class of high affinity isozyme II and IV inhibitors. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 1999 , 14, 289-306		16
34	Accumulation of untranslated metallothionein mRNA in antarctic hemoglobinless fish (icefish) 1999 , 167-172		2
33	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	6.8	139

32	Molecular cloning and sequence determination of a novel aspartic proteinase from Antarctic fish. <i>BBA - Proteins and Proteomics</i> , 1998 , 1387, 457-61		23
31	Carbonic anhydrase inhibitors [Part 53. Synthesis of substituted-pyridinium derivatives of aromatic sulfonamides: The first non-polymeric membrane-impermeable inhibitors with selectivity for isozyme IV. <i>European Journal of Medicinal Chemistry</i> , 1998 , 33, 577-594	6.8	70
30	Carbonic anhydrase inhibitors [Part 29 1: Interaction of isozymes I, II and IV with benzolamide-like derivatives. <i>European Journal of Medicinal Chemistry</i> , 1998 , 33, 739-751	6.8	130
29	Carbonic anhydrase inhibitors [Part 52. Metal complexes of heterocyclic sulfonamides: A new class of strong topical intraocular pressure-lowering agents in rabbits. <i>European Journal of Medicinal Chemistry</i> , 1998 , 33, 247-254	6.8	120
28	Carbonic anhydrase inhibitors: inhibition of isozymes I, II and IV with N-hydroxysulfonamides--a novel class of intraocular pressure lowering agents. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 1998 , 13, 267-84		26
27	Cadmium-induced differential accumulation of metallothionein isoforms in the Antarctic icefish, which exhibits no basal metallothionein protein but high endogenous mRNA levels. <i>Biochemical Journal</i> , 1998 , 332 (Pt 2), 475-81	3.8	57
26	Metallothionein in Antarctic Fish 1998 , 151-161		1
25	Difference in hepatic metallothionein content in Antarctic red-blooded and haemoglobinless fish: undetectable metallothionein levels in haemoglobinless fish is accompanied by accumulation of untranslated metallothionein mRNA. <i>Biochemical Journal</i> , 1997 , 322 (Pt 1), 207-11	3.8	45
24	Carbonic anhydrase activators: X-ray crystallographic and spectroscopic investigations for the interaction of isozymes I and II with histamine. <i>Biochemistry</i> , 1997 , 36, 10384-92	3.2	246
23	Novel aromatic/heterocyclic sulfonamides and their metal complexes as inhibitors of carbonic anhydrase isozymes I, II and IV. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 1997 , 12, 37-51		62
22	Unique structural features of the monomeric Cu,Zn superoxide dismutase from Escherichia coli, revealed by X-ray crystallography. <i>Journal of Molecular Biology</i> , 1997 , 274, 408-20	6.5	73
21	Carbonic Anhydrase Activators. Part 19 Spectroscopic and Kinetic Investigations for the Interaction of Isozymes I and II With Primary Amines. <i>Metal-Based Drugs</i> , 1997 , 4, 221-7		3
20	PCR amplification and cloning of metallothionein complementary DNAs in temperate and Antarctic sea urchin characterized by a large difference in egg metallothionein content. <i>Cellular and Molecular Life Sciences</i> , 1997 , 53, 472-7	10.3	15
19	Crystal structure of the bovine alpha-chymotrypsin:Kunitz inhibitor complex. An example of multiple protein:protein recognition sites. <i>Journal of Molecular Recognition</i> , 1997 , 10, 26-35	2.6	37
18	Is cyanate a carbonic anhydrase substrate?. <i>Proteins: Structure, Function and Bioinformatics</i> , 1997 , 27, 272-8	4.2	42
17	Identification of a high-molecular-weight cadmium-binding protein in copper-resistant <i>Bacillus acidocaldarius</i> cells. <i>Research in Microbiology</i> , 1996 , 147, 287-96	4	12
16	Complexes With Biologically Active Ligands. Part 2. Preparation of Copper(II) Complexes of Positively-Charged Derivatives of Aminogluthethimide. <i>Metal-Based Drugs</i> , 1996 , 3, 57-62		1
15	Complexes with biologically active ligands. Part 4. Coordination compounds of chlorothiazide with transition metal ions behave as strong carbonic anhydrase inhibitors. <i>Metal-Based Drugs</i> , 1996 , 3, 79-83		2

14	Isolation and characterisation of zinc-binding proteins distinct from metallothionein from the eggs of the sea urchin <i>Strongylocentrotus intermedius</i> . <i>Marine Biology</i> , 1996 , 126, 225-230	2.5	3
13	Crystallization and preliminary X-ray analysis of the monomeric Cu,Zn superoxide dismutase from <i>Escherichia coli</i> . <i>Protein Science</i> , 1996 , 5, 2125-7	6.3	10
12	Metal Complexes of 1,3,4-Thiadiazole-2,5-Disulfonamide are Strong Dual Carbonic Anhydrase Inhibitors, although the Ligand Possesses very Weak such Properties. <i>Metal-Based Drugs</i> , 1995 , 2, 331-6		9
11	Isolation and primary structure determination of a metallothionein from <i>Paracentrotus lividus</i> (Echinodermata, Echinoidea). <i>Comparative Biochemistry and Physiology - B Biochemistry and Molecular Biology</i> , 1995 , 111, 329-36	2.3	33
10	Carbonic anhydrase activators. 3: structure-activity correlations for a series of isozyme II activators. <i>Journal of Pharmaceutical Sciences</i> , 1994 , 83, 768-73	3.9	100
9	Metal-binding proteins in eggs of various sea urchin species. <i>Cell Biology International</i> , 1994 , 18, 47-53	4.5	14
8	Carbonic anhydrase activators. VII. Isozyme II activation by bisazoly-methanes, -ethanes and related azoles. <i>Biological and Pharmaceutical Bulletin</i> , 1993 , 16, 1236-9	2.3	27
7	Botulinus Toxin, Tetanus Toxin, and Anthrax Lethal Factor Inhibitors705-720		4
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