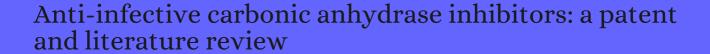
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DOI: 10.1517/13543776.2013.778245 Expert Opinion on Therapeutic Patents, 2013, 23, 693-704.

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199	5-Substituted-(1,2,3-triazol-4-yl)thiophene-2-sulfonamides strongly inhibit human carbonic anhydrases I, II, IX and XII: solution and X-ray crystallographic studies. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 5130-8	3.4	26
198	A class of sulfonamides with strong inhibitory action against the Etarbonic anhydrase from Trypanosoma cruzi. <i>Journal of Medicinal Chemistry</i> , <b>2013</b> , 56, 5773-81	8.3	51
197	Synthesis of novel acridine and bis acridine sulfonamides with effective inhibitory activity against the cytosolic carbonic anhydrase isoforms II and VII. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 5799-	-80 <del>5</del>	29
196	Anion inhibition studies of a Etarbonic anhydrase from Clostridium perfringens. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2013</b> , 23, 6706-10	2.9	42
195	Antimalarial activity of compounds comprising a primary benzene sulfonamide fragment. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2013</b> , 23, 6114-7	2.9	25
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193	Biochemical characterization of the Earbonic anhydrase from the marine diatom Thalassiosira weissflogii, TweCA. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2014</b> , 29, 906-11	5.6	58
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177	Synthesis and carbonic anhydrase I, II, IX and XII inhibition studies of 4-N,N-disubstituted sulfanilamides incorporating 4,4,4-trifluoro-3-oxo-but-1-enyl, phenacylthiourea and imidazol-2(3H)-one/thione moieties. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2014</b> , 24, 1776-9	2.9	21
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167	The zinc coordination pattern in the Etarbonic anhydrase from Plasmodium falciparum is different from all other carbonic anhydrase genetic families. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2015</b> , 25, 1385-9	2.9	95
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161	Inhibition of Earbonic anhydrases from Brucella suis with C-cinnamoyl glycosides incorporating the phenol moiety. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , <b>2015</b> , 30, 1017-20	5.6	13
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142	Dendrimers incorporating benzenesulfonamide moieties strongly inhibit carbonic anhydrase isoforms I-XIV. <b>2015</b> , 13, 6453-7		12
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