

Edward D Sturrock

List of Publications by Year in descending order

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86
papers

3,997
citations

147566

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86
docs citations

86
times ranked

3430
citing authors

#	ARTICLE	IF	CITATIONS
1	Probing the Requirements for Dual Angiotensin-Converting Enzyme C-Domain Selective/Neprilysin Inhibition. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 3371-3387.	2.9	3
2	Structural basis for the inhibition of human angiotensinâ€ converting enzyme by fosinoprilat. <i>FEBS Journal</i> , 2022, 289, 6659-6671.	2.2	5
3	<scp>Cryoâ€EM</scp> reveals mechanisms of angiotensin lâ€converting enzyme allostery and dimerization. <i>EMBO Journal</i> , 2022, 41, .	3.5	4
4	Novel ACE mutations mimicking sarcoidosis by increasing blood ACE levels. <i>Translational Research</i> , 2021, 230, 5-20.	2.2	12
5	Angiotensinâ€converting enzyme open for business: structural insights into the subdomain dynamics. <i>FEBS Journal</i> , 2021, 288, 2238-2256.	2.2	21
6	Epitope mapping of novel monoclonal antibodies to human angiotensin lâ€converting enzyme. <i>Protein Science</i> , 2021, 30, 1577-1593.	3.1	7
7	Investigating the antifibrotic potential of Nâ€acetyl serylâ€aspartylâ€lysylâ€proline sequence peptides. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2021, 48, 1558-1565.	0.9	1
8	Selective Inhibition of the C-Domain of ACE (Angiotensin-Converting Enzyme) Combined With Inhibition of NEP (Neprilysin): A Potential New Therapy for Hypertension. <i>Hypertension</i> , 2021, 78, 604-616.	1.3	7
9	Targeting the oncogenic TBX3:nucleolin complex to treat multiple sarcoma subtypes. <i>American Journal of Cancer Research</i> , 2021, 11, 5680-5700.	1.4	0
10	Prospects for SARS-CoV-2 diagnostics, therapeutics and vaccines in Africa. <i>Nature Reviews Microbiology</i> , 2020, 18, 690-704.	13.6	42
11	Molecular Basis for Omapatrilat and Sampatrilat Binding to Neprilysinâ€Implications for Dual Inhibitor Design with Angiotensin-Converting Enzyme. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 5488-5500.	2.9	13
12	ACE-domain selectivity extends beyond direct interacting residues at the active site. <i>Biochemical Journal</i> , 2020, 477, 1241-1259.	1.7	10
13	ACE2 and ACE: structure-based insights into mechanism, regulation and receptor recognition by SARS-CoV. <i>Clinical Science</i> , 2020, 134, 2851-2871.	1.8	47
14	Novel Therapeutic Approaches Targeting the Renin-Angiotensin System and Associated Peptides in Hypertension and Heart Failure. <i>Pharmacological Reviews</i> , 2019, 71, 539-570.	7.1	235
15	Interacting cogs in the machinery of the renin angiotensin system. <i>Biophysical Reviews</i> , 2019, 11, 583-589.	1.5	3
16	Structural basis for the C-domain-selective angiotensin-converting enzyme inhibition by bradykinin-potentiating peptide b (BPPb). <i>Biochemical Journal</i> , 2019, 476, 1553-1570.	1.7	16
17	Crystal structures of sampatrilat and sampatrilatâ€Asp in complex with human ACE â€ a molecular basis for domain selectivity. <i>FEBS Journal</i> , 2018, 285, 1477-1490.	2.2	23
18	Investigation into the Mechanism of Homo- and Heterodimerization of Angiotensin-Converting Enzyme. <i>Molecular Pharmacology</i> , 2018, 93, 344-354.	1.0	4

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19	The Design and Development of a Potent and Selective Novel Diprolyl Derivative That Binds to the N-Domain of Angiotensin-I Converting Enzyme. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 344-359.	2.9	20
20	Molecular Basis for Multiple Omapatrilat Binding Sites within the ACE C-Domain: Implications for Drug Design. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 10141-10154.	2.9	22
21	Established and novel pathophysiological mechanisms of pericardial injury and constrictive pericarditis. <i>World Journal of Cardiology</i> , 2018, 10, 87-96.	0.5	25
22	The Dynamic Nonprime Binding of Sampatrilat to the C-Domain of Angiotensin-Converting Enzyme. <i>Journal of Chemical Information and Modeling</i> , 2016, 56, 2486-2494.	2.5	12
23	Kinetic and structural characterization of amyloid β peptide hydrolysis by human angiotensin α -converting enzyme. <i>FEBS Journal</i> , 2016, 283, 1060-1076.	2.2	19
24	The influence of angiotensin converting enzyme mutations on the kinetics and dynamics of N α -domain selective inhibition. <i>FEBS Journal</i> , 2016, 283, 3941-3961.	2.2	18
25	The effect of structural motifs on the ectodomain shedding of human angiotensin-converting enzyme. <i>Biochemical and Biophysical Research Communications</i> , 2016, 481, 111-116.	1.0	6
26	Structural basis of Ac-SDKP hydrolysis by Angiotensin-I converting enzyme. <i>Scientific Reports</i> , 2015, 5, 13742.	1.6	18
27	Characterisation of the flavin adenine dinucleotide binding region of <i>Myxococcus xanthus</i> protoporphyrinogen oxidase. <i>Biochemistry and Biophysics Reports</i> , 2015, 4, 306-311.	0.7	3
28	Pharmacodynamic effects of C-domain-specific ACE inhibitors on the renin-angiotensin system in myocardial infarcted rats. <i>JRAAS - Journal of the Renin-Angiotensin-Aldosterone System</i> , 2015, 16, 1149-1158.	1.0	24
29	A Novel Angiotensin I-Converting Enzyme Mutation (S333W) Impairs N-Domain Enzymatic Cleavage of the Anti-Fibrotic Peptide, AcSDKP. <i>PLoS ONE</i> , 2014, 9, e88001.	1.1	19
30	Effects of a domain-selective ACE inhibitor in a mouse model of chronic angiotensin II-dependent hypertension. <i>Clinical Science</i> , 2014, 127, 57-63.	1.8	27
31	Fragment-based design for the development of N-domain-selective angiotensin-1-converting enzyme inhibitors. <i>Clinical Science</i> , 2014, 126, 305-313.	1.8	36
32	Angiotensin-I converting enzyme (ACE): structure, biological roles, and molecular basis for chloride ion dependence. <i>Biological Chemistry</i> , 2014, 395, 1135-1149.	1.2	43
33	Molecular and Thermodynamic Mechanisms of the Chloride-dependent Human Angiotensin-I-converting Enzyme (ACE). <i>Journal of Biological Chemistry</i> , 2014, 289, 1798-1814.	1.6	29
34	Pharmacokinetic evaluation of lisinopril-tryptophan, a novel C-domain ACE inhibitor. <i>European Journal of Pharmaceutical Sciences</i> , 2014, 56, 113-119.	1.9	12
35	Crystal structures of highly specific phosphinic tripeptide enantiomers in complex with the angiotensin α -converting enzyme. <i>FEBS Journal</i> , 2014, 281, 943-956.	2.2	27
36	Interkingdom Pharmacology of Angiotensin-I Converting Enzyme Inhibitor Phosphonates Produced by Actinomycetes. <i>ACS Medicinal Chemistry Letters</i> , 2014, 5, 346-351.	1.3	26

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37	Peptidyl-Dipeptidase A/Angiotensin I-Converting Enzyme. , 2013, , 480-494.		13
38	Antifibrotic peptide <i>N</i> -acetyl-L-serine-L-aspartyl-L-proline (Ac-SDKP): Opportunities for angiotensin-converting enzyme inhibitor design. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2013, 40, 535-541.	0.9	7
39	C domain-selective inhibition of angiotensin-converting enzyme. <i>JRAAS - Journal of the Renin-Angiotensin-Aldosterone System</i> , 2013, 14, 189-192.	1.0	9
40	Scientific letter: Ac-SDKP (N-acetyl-seryl-aspartyl-lysyl-proline) and Galectin-3 levels in tuberculous pericardial effusion: implications for pathogenesis and prevention of pericardial constriction. <i>Heart</i> , 2012, 98, 1326.1-1328.	1.2	16
41	Structural basis of peptide recognition by the angiotensin-converting enzyme homologue An _{CE} from <i>Drosophila melanogaster</i> . <i>FEBS Journal</i> , 2012, 279, 4525-4534.	2.2	21
42	Molecular recognition and regulation of human angiotensin-I converting enzyme (ACE) activity by natural inhibitory peptides. <i>Scientific Reports</i> , 2012, 2, 717.	1.6	127
43	New ketomethylene inhibitor analogues: synthesis and assessment of structural determinants for N-domain selective inhibition of angiotensin-converting enzyme. <i>Biological Chemistry</i> , 2012, 393, 485-493.	1.2	11
44	Shedding the load of hypertension: The proteolytic processing of angiotensin-converting enzyme. <i>South African Medical Journal</i> , 2012, 102, 461.	0.2	14
45	An Angiotensin I-Converting Enzyme Mutation (Y465D) Causes a Dramatic Increase in Blood ACE via Accelerated ACE Shedding. <i>PLoS ONE</i> , 2011, 6, e25952.	1.1	37
46	Structural characterization of angiotensin-converting enzyme in complex with a selenium analogue of captopril. <i>FEBS Journal</i> , 2011, 278, 3644-3650.	2.2	33
47	The significance of the $\text{C}\hat{\pm}$ substituent in the selective inhibition of matrix metalloproteinases 1 and 9. <i>Biological Chemistry</i> , 2011, 392, 1003-10.	1.2	0
48	Characterization of domain-selective inhibitor binding in angiotensin-converting enzyme using a novel derivative of lisinopril. <i>Biochemical Journal</i> , 2010, 428, 67-74.	1.7	38
49	The N Domain of Human Angiotensin-I-converting Enzyme. <i>Journal of Biological Chemistry</i> , 2010, 285, 35685-35693.	1.6	76
50	High-Resolution Crystal Structures of <i>Drosophila melanogaster</i> Angiotensin-Converting Enzyme in Complex with Novel Inhibitors and Antihypertensive Drugs. <i>Journal of Molecular Biology</i> , 2010, 400, 502-517.	2.0	65
51	Investigating the Domain Specificity of Phosphinic Inhibitors RXP380 and RXP407 in Angiotensin-Converting Enzyme. <i>Biochemistry</i> , 2009, 48, 8405-8412.	1.2	42
52	Angiotensin-Converting Enzyme - New Insights into Structure, Biological Significance and Prospects for Domain-Selective Inhibitors. <i>Current Enzyme Inhibition</i> , 2009, 5, 134-147.	0.3	13
53	Probing the Basis of Domain-Dependent Inhibition Using Novel Ketone Inhibitors of Angiotensin-Converting Enzyme. <i>Biochemistry</i> , 2008, 47, 5942-5950.	1.2	53
54	Mapping of Conformational mAb Epitopes to the C Domain of Human Angiotensin I-Converting Enzyme. <i>Journal of Proteome Research</i> , 2008, 7, 3396-3411.	1.8	26

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55	The role of glycosylation and domain interactions in the thermal stability of human angiotensin-converting enzyme. <i>Biological Chemistry</i> , 2008, 389, 1153-1161.	1.2	17
56	Simulated Interactions between Angiotensin-Converting Enzyme and Substrate Gonadotropin-Releasing Hormone: Novel Insights into Domain Selectivity. <i>Biochemistry</i> , 2007, 46, 8753-8765.	1.2	24
57	Fine Epitope Mapping of Monoclonal Antibody 5F1 Reveals Anticatalytic Activity toward the N Domain of Human Angiotensin-Converting Enzyme. <i>Biochemistry</i> , 2007, 46, 9019-9031.	1.2	24
58	The Structure of Testis Angiotensin-Converting Enzyme in Complex with the C Domain-Specific Inhibitor RXPA380. <i>Biochemistry</i> , 2007, 46, 5473-5478.	1.2	88
59	Structure of Testis ACE Glycosylation Mutants and Evidence for Conserved Domain Movement. <i>Biochemistry</i> , 2006, 45, 12654-12663.	1.2	53
60	Homologous substitution of ACE C-domain regions with N-domain sequences: effect on processing, shedding, and catalytic properties. <i>Biological Chemistry</i> , 2006, 387, 1043-51.	1.2	17
61	Crystal Structure of the N Domain of Human Somatic Angiotensin I-converting Enzyme Provides a Structural Basis for Domain-specific Inhibitor Design. <i>Journal of Molecular Biology</i> , 2006, 357, 964-974.	2.0	150
62	A high-throughput fluorimetric assay for angiotensin I-converting enzyme. <i>Nature Protocols</i> , 2006, 1, 1961-1964.	5.5	49
63	A continuous fluorescence resonance energy transfer angiotensin I-converting enzyme assay. <i>Nature Protocols</i> , 2006, 1, 1971-1976.	5.5	84
64	Synthesis of novel keto-ACE analogues as domain-selective angiotensin I-converting enzyme inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 4612-4615.	1.0	35
65	Synthesis and molecular modeling of a lisinopril-tryptophan analogue inhibitor of angiotensin I-converting enzyme. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 4616-4619.	1.0	34
66	Novel ketomethylene inhibitors of angiotensin I-converting enzyme (ACE): inhibition and molecular modelling. <i>Biological Chemistry</i> , 2006, 387, 461-6.	1.2	12
67	The N domain of somatic angiotensin-converting enzyme negatively regulates ectodomain shedding and catalytic activity. <i>Biochemical Journal</i> , 2005, 389, 739-744.	1.7	43
68	Development of Domain-Selective Angiotensin I-Converting Enzyme Inhibitors. <i>Annals of the New York Academy of Sciences</i> , 2005, 1056, 160-175.	1.8	22
69	Monoclonal Antibodies 1B3 and 5C8 as Probes for Monitoring the Integrity of the C-Terminal End of Soluble Angiotensin-Converting Enzyme. <i>Hybridoma</i> , 2005, 24, 14-26.	0.5	25
70	Localization of an N-Domain Region of Angiotensin-Converting Enzyme Involved in the Regulation of Ectodomain Shedding Using Monoclonal Antibodies. <i>Journal of Proteome Research</i> , 2005, 4, 258-267.	1.8	30
71	Positional-Scanning Combinatorial Libraries of Fluorescence Resonance Energy Transfer Peptides for Defining Substrate Specificity of the Angiotensin I-Converting Enzyme and Development of Selective C-Domain Substrates. <i>Biochemistry</i> , 2004, 43, 15729-15736.	1.2	32
72	Structural Details on the Binding of Antihypertensive Drugs Captopril and Enalaprilat to Human Testicular Angiotensin I-Converting Enzyme. <i>Biochemistry</i> , 2004, 43, 8718-8724.	1.2	240

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73	Crystal structure of the human angiotensin-converting enzyme-lisinopril complex. <i>Nature</i> , 2003, 421, 551-554.	13.7	738
74	Ace revisited: A new target for structure-based drug design. <i>Nature Reviews Drug Discovery</i> , 2003, 2, 891-902.	21.5	285
75	Angiotensin-Converting Enzyme-2 (ACE2): A Comparative Modeling of the Active Site, Specificity Requirements, and Chloride Dependence. <i>Biochemistry</i> , 2003, 42, 13185-13192.	1.2	164
76	Deglycosylation, processing and crystallization of human testis angiotensin-converting enzyme. <i>Biochemical Journal</i> , 2003, 371, 437-442.	1.7	64
77	Defining the boundaries of the testis angiotensin I-converting enzyme ectodomain. <i>Biochemical and Biophysical Research Communications</i> , 2002, 297, 1225-1230.	1.0	20
78	Roles of the juxtamembrane and extracellular domains of angiotensin-converting enzyme in ectodomain shedding. <i>Biochemical Journal</i> , 2001, 358, 185-192.	1.7	35
79	A study of crystal matrix extract and urinary prothrombin fragment 1 from a stone-prone and stone-free population. <i>Urological Research</i> , 2001, 29, 83-88.	1.5	22
80	Shedding of somatic angiotensin-converting enzyme (ACE) is inefficient compared with testis ACE despite cleavage at identical stalk sites. <i>Biochemical Journal</i> , 2000, 347, 711.	1.7	18
81	Shedding of somatic angiotensin-converting enzyme (ACE) is inefficient compared with testis ACE despite cleavage at identical stalk sites. <i>Biochemical Journal</i> , 2000, 347, 711-718.	1.7	74
82	Modulation of Juxtamembrane Cleavage (Shedding) of Angiotensin-Converting Enzyme by Stalk Glycosylation: Evidence for an Alternative Shedding Protease. <i>Biochemistry</i> , 1999, 38, 10388-10397.	1.2	36
83	Phorbol Ester-Induced Juxtamembrane Cleavage of Angiotensin-Converting Enzyme is not Inhibited by a Disulfide-Bridged Stalk. <i>Biochemical Society Transactions</i> , 1999, 27, A56-A56.	1.6	0
84	Identification of N-Linked Glycosylation Sites in Human Testis Angiotensin-converting Enzyme and Expression of an Active Deglycosylated Form. <i>Journal of Biological Chemistry</i> , 1997, 272, 3511-3519.	1.6	66
85	Limited Proteolysis of Human Kidney Angiotensin-Converting Enzyme and Generation of Catalytically Active N- and C-Terminal Domains. <i>Biochemical and Biophysical Research Communications</i> , 1997, 236, 16-19.	1.0	30
86	Assignment of Free and Disulfide-Bonded Cysteine Residues in Testis Angiotensin-Converting Enzyme: Functional Implications. <i>Biochemistry</i> , 1996, 35, 9560-9566.	1.2	26