

Enrico Di Cera

List of Publications by Year in descending order

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107
papers

4,914
citations

81900

39
h-index

98798

67
g-index

112
all docs

112
docs citations

112
times ranked

4149
citing authors

#	ARTICLE	IF	CITATIONS
1	The active site region plays a critical role in Na ⁺ binding to thrombin. <i>Journal of Biological Chemistry</i> , 2022, 298, 101458.	3.4	4
2	Cryo-EM structure of the prothrombin-prothrombinase complex. <i>Blood</i> , 2022, 139, 3463-3473.	1.4	19
3	Cryo-EM structures of human coagulation factors V and Va. <i>Blood</i> , 2021, 137, 3137-3144.	1.4	29
4	Role of sequence and position of the cleavage sites in prothrombin activation. <i>Journal of Biological Chemistry</i> , 2021, 297, 100955.	3.4	8
5	Zymogen and activated protein C have similar structural architecture. <i>Journal of Biological Chemistry</i> , 2020, 295, 15236-15244.	3.4	8
6	19F NMR reveals the conformational properties of free thrombin and its zymogen precursor prethrombin-2. <i>Journal of Biological Chemistry</i> , 2020, 295, 8227-8235.	3.4	7
7	Mechanisms of ligand binding. <i>Biophysics Reviews</i> , 2020, 1, 011303.	2.7	33
8	VE-1902 is a direct thrombin inhibitor with reversible covalent mechanism of action shows efficacy with reduced bleeding in rodent models of thrombosis. <i>Thrombosis Research</i> , 2020, 190, 112-121.	1.7	8
9	Role of the activation peptide in the mechanism of protein C activation. <i>Scientific Reports</i> , 2020, 10, 11079.	3.3	10
10	Residues W215, E217 and E192 control the allosteric E* ⁺ -E equilibrium of thrombin. <i>Scientific Reports</i> , 2019, 9, 12304.	3.3	7
11	Probing prothrombin structure by limited proteolysis. <i>Scientific Reports</i> , 2019, 9, 6125.	3.3	7
12	Role of the I16-D194 ionic interaction in the trypsin fold. <i>Scientific Reports</i> , 2019, 9, 18035.	3.3	9
13	Structure of prothrombin in the closed form reveals new details on the mechanism of activation. <i>Scientific Reports</i> , 2018, 8, 2945.	3.3	28
14	Interplay between conformational selection and zymogen activation. <i>Scientific Reports</i> , 2018, 8, 4080.	3.3	22
15	Enhancing the anticoagulant profile of meizothrombin. <i>Biomolecular Concepts</i> , 2018, 9, 169-175.	2.2	10
16	Reversible covalent direct thrombin inhibitors. <i>PLoS ONE</i> , 2018, 13, e0201377.	2.5	13
17	Intrinsic thermodynamics of high affinity inhibitor binding to recombinant human carbonic anhydrase IV. <i>European Biophysics Journal</i> , 2018, 47, 271-290.	2.2	14
18	Induced Fit Is a Special Case of Conformational Selection. <i>Biochemistry</i> , 2017, 56, 2853-2859.	2.5	39

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19	Rational Design of Protein C Activators. <i>Scientific Reports</i> , 2017, 7, 44596.	3.3	6
20	Molecular Mechanisms of Enzyme Activation by Monovalent Cations. <i>Journal of Biological Chemistry</i> , 2016, 291, 20840-20848.	3.4	80
21	Thrombin Cleavage of Plasmodium falciparum Erythrocyte Membrane Protein 1 Inhibits Cytoadherence. <i>MBio</i> , 2016, 7, .	4.1	9
22	Structural Architecture of Prothrombin in Solution Revealed by Single Molecule Spectroscopy. <i>Journal of Biological Chemistry</i> , 2016, 291, 18107-18116.	3.4	26
23	Data publication with the structural biology data grid supports live analysis. <i>Nature Communications</i> , 2016, 7, 10882.	12.8	113
24	Loop Electrostatics Asymmetry Modulates the Preexisting Conformational Equilibrium in Thrombin. <i>Biochemistry</i> , 2016, 55, 3984-3994.	2.5	17
25	How the Linker Connecting the Two Kringles Influences Activation and Conformational Plasticity of Prothrombin. <i>Journal of Biological Chemistry</i> , 2016, 291, 6071-6082.	3.4	28
26	Potassium and the K ⁺ /H ⁺ Exchanger Kha1p Promote Binding of Copper to ApoFet3p Multi-copper Ferroxidase. <i>Journal of Biological Chemistry</i> , 2016, 291, 9796-9806.	3.4	20
27	Why Ser and Not Thr Brokers Catalysis in the Trypsin Fold. <i>Biochemistry</i> , 2015, 54, 1457-1464.	2.5	12
28	John A. Schellman, 1924â€“2014. <i>Biophysical Chemistry</i> , 2015, 199, 51.	2.8	0
29	Kinetic Dissection of the Pre-existing Conformational Equilibrium in the Trypsin Fold. <i>Journal of Biological Chemistry</i> , 2015, 290, 22435-22445.	3.4	31
30	Prothrombin structure: unanticipated features and opportunities. <i>Expert Review of Proteomics</i> , 2014, 11, 653-655.	3.0	11
31	The linker connecting the two kringles plays a key role in prothrombin activation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 7630-7635.	7.1	37
32	Essential role of conformational selection in ligand binding. <i>Biophysical Chemistry</i> , 2014, 186, 13-21.	2.8	92
33	Special issue on conformational selection. <i>Biophysical Chemistry</i> , 2014, 186, 1-2.	2.8	3
34	Conformational Selection Is a Dominant Mechanism of Ligand Binding. <i>Biochemistry</i> , 2013, 52, 5723-5729.	2.5	110
35	Crystal Structure of Prothrombin Reveals Conformational Flexibility and Mechanism of Activation. <i>Journal of Biological Chemistry</i> , 2013, 288, 22734-22744.	3.4	42
36	Rapid Interruption Of Occlusive Thrombus Formation By The Protein C Activator Enzyme EWE Thrombin (ProCase) In Primates. <i>Blood</i> , 2013, 122, 202-202.	1.4	0

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37	In vitro veritas: 90 years of biochemistry at Saint Louis University. <i>Missouri Medicine</i> , 2013, 110, 297-301.	0.3	0
38	Conformational Selection or Induced Fit? A Critical Appraisal of the Kinetic Mechanism. <i>Biochemistry</i> , 2012, 51, 5894-5902.	2.5	269
39	Conformational selection in trypsin-like proteases. <i>Current Opinion in Structural Biology</i> , 2012, 22, 421-431.	5.7	79
40	Exposure of R169 controls protein C activation and autoactivation. <i>Blood</i> , 2012, 120, 664-670.	1.4	23
41	Crystal Structures of Prethrombin-2 Reveal Alternative Conformations under Identical Solution Conditions and the Mechanism of Zymogen Activation. <i>Biochemistry</i> , 2011, 50, 10195-10202.	2.5	40
42	Crystallographic and Kinetic Evidence of Allostery in a Trypsin-like Protease. <i>Biochemistry</i> , 2011, 50, 6301-6307.	2.5	44
43	Allostery in trypsin-like proteases suggests new therapeutic strategies. <i>Trends in Biotechnology</i> , 2011, 29, 577-585.	9.3	80
44	Rigidification of the autolysis loop enhances Na ⁺ binding to thrombin. <i>Biophysical Chemistry</i> , 2011, 159, 6-13.	2.8	21
45	Structural basis of thrombin- α 1-protease-activated receptor interactions. <i>IUBMB Life</i> , 2011, 63, 375-382.	3.4	25
46	Introduction to mini-theme issue on α 1-protease-activated receptor signaling. <i>IUBMB Life</i> , 2011, 63, 373-374.	3.4	0
47	Thrombin as an Anticoagulant. <i>Progress in Molecular Biology and Translational Science</i> , 2011, 99, 145-184.	1.7	17
48	Crystal structure of prethrombin-1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 19278-19283.	7.1	37
49	Crystal Structure of Thrombin Bound to the Uncleaved Extracellular Fragment of PAR1. <i>Journal of Biological Chemistry</i> , 2010, 285, 15393-15398.	3.4	56
50	Engineering Thrombin for Selective Specificity toward Protein C and PAR1. <i>Journal of Biological Chemistry</i> , 2010, 285, 19145-19152.	3.4	38
51	Evidence of the E [*] E Equilibrium from Rapid Kinetics of Na ⁺ Binding to Activated Protein C and Factor Xa [*] . <i>Journal of Physical Chemistry B</i> , 2010, 114, 16125-16130.	2.6	20
52	Combinatorial Enzyme Design Probes Allostery and Cooperativity in the Trypsin Fold. <i>Journal of Molecular Biology</i> , 2010, 399, 306-319.	4.2	9
53	Mutant N143P Reveals How Na ⁺ Activates Thrombin. <i>Journal of Biological Chemistry</i> , 2009, 284, 36175-36185.	3.4	31
54	Mechanism of the Anticoagulant Activity of Thrombin Mutant W215A/E217A. <i>Journal of Biological Chemistry</i> , 2009, 284, 24098-24105.	3.4	23

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55	Serine proteases. IUBMB Life, 2009, 61, 510-515.	3.4	306
56	Serine proteases. IUBMB Life, 2009, 61, spcone-spcone.	3.4	0
57	Kinetics of Allosteric Activation. Methods in Enzymology, 2009, 466, 259-271.	1.0	6
58	How I became a biochemist. IUBMB Life, 2008, 60, 859-861.	3.4	0
59	Thrombin. Molecular Aspects of Medicine, 2008, 29, 203-254.	6.4	282
60	Thrombin Mutant W215A/E217A Acts as a Platelet GPIb Antagonist. Arteriosclerosis, Thrombosis, and Vascular Biology, 2008, 28, 329-334.	2.4	41
61	Important Role of the Cys-191â€“Cys-220 Disulfide Bond in Thrombin Function and Allostery. Journal of Biological Chemistry, 2007, 282, 27165-27170.	3.4	29
62	Relative antithrombotic and antihemostatic effects of protein C activator versus low-molecular-weight heparin in primates. Blood, 2007, 109, 3733-3740.	1.4	49
63	Thrombin allostery. Physical Chemistry Chemical Physics, 2007, 9, 1291.	2.8	46
64	Mechanism of Na ⁺ binding to thrombin resolved by ultra-rapid kinetics. Biophysical Chemistry, 2007, 131, 111-114.	2.8	26
65	Mutation of W215 Compromises Thrombin Cleavage of Fibrinogen, but Not of PAR1 or Protein C. Annals of the New York Academy of Sciences, 2006, 936, 456-458.	3.8	1
66	Thrombin: A paradigm for enzymes allosterically activated by monovalent cations. Rendiconti Lincei, 2006, 17, 97-113.	2.2	1
67	A Structural Perspective on Enzymes Activated by Monovalent Cations. Journal of Biological Chemistry, 2006, 281, 1305-1308.	3.4	115
68	Crystal Structure of Thrombin in a Self-inhibited Conformation. Journal of Biological Chemistry, 2006, 281, 32922-32928.	3.4	45
69	Rapid Kinetics of Na ⁺ Binding to Thrombin. Journal of Biological Chemistry, 2006, 281, 40049-40056.	3.4	62
70	Safe Interruption of Ongoing Thrombosis by the Protein C Activator Thrombin Analog, W215A/E217A; a Comparison to Enoxaparin in Primates.. Blood, 2006, 108, 909-909.	1.4	2
71	Thrombomodulin-Dependent Protein C Activator Treatment Improves the Short-Term Outcome of Experimental Ischemic Stroke in Mice.. Blood, 2006, 108, 895-895.	1.4	0
72	Thrombomodulin Changes the Molecular Surface of Interaction and the Rate of Complex Formation between Thrombin and Protein C. Journal of Biological Chemistry, 2005, 280, 7956-7961.	3.4	46

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73	Efficient Barrier Protective Signaling by Activated Protein C Is Mechanistically Linked to Protein C Activation on Endothelial Cells.. Blood, 2005, 106, 28-28.	1.4	1
74	The Conformation of the Activation Peptide of Protein C Is Influenced by Ca ²⁺ and Na ⁺ Binding. Journal of Biological Chemistry, 2004, 279, 38519-38524.	3.4	32
75	Molecular Dissection of Na ⁺ Binding to Thrombin. Journal of Biological Chemistry, 2004, 279, 31842-31853.	3.4	161
76	The Anticoagulant Thrombin Mutant W215A/E217A Has a Collapsed Primary Specificity Pocket. Journal of Biological Chemistry, 2004, 279, 39824-39828.	3.4	51
77	Thrombin: a paradigm for enzymes allosterically activated by monovalent cations. Comptes Rendus - Biologies, 2004, 327, 1065-1076.	0.2	29
78	Sustained Pharmacological Activation of Protein C (PC) in Baboons.. Blood, 2004, 104, 3499-3499.	1.4	0
79	Thrombin Interactions. Chest, 2003, 124, 11S-17S.	0.8	115
80	The Thrombin Mutant W215A/E217A Shows Safe and Potent Anticoagulant and Antithrombotic Effects in Vivo. Journal of Biological Chemistry, 2002, 277, 27581-27584.	3.4	71
81	Evolution of enzyme cascades from embryonic development to blood coagulation. Trends in Biochemical Sciences, 2002, 27, 67-74.	7.5	301
82	Dissecting substrate recognition by thrombin using the inactive mutant S195A. Biophysical Chemistry, 2002, 100, 315-323.	2.8	29
83	Replacement of thrombin residue G184 with Lys or Arg fails to mimic Na ⁺ binding. Proteins: Structure, Function and Bioinformatics, 2001, 43, 315-318.	2.6	7
84	Molecular mapping of thrombin-receptor interactions. Proteins: Structure, Function and Bioinformatics, 2001, 45, 107-116.	2.6	99
85	Determinants of Thrombin Specificity. Annals of the New York Academy of Sciences, 2001, 936, 133-146.	3.8	26
86	A simple method for the determination of individual rate constants for substrate hydrolysis by serine proteases. Protein Science, 2000, 9, 1589-1593.	7.6	47
87	Rational Design of a Potent Anticoagulant Thrombin. Journal of Biological Chemistry, 2000, 275, 39827-39830.	3.4	78
88	Mutation of W215 Compromises Thrombin Cleavage of Fibrinogen, but Not of PAR-1 or Protein C α . Biochemistry, 2000, 39, 8095-8101.	2.5	51
89	Role of residue Y99 in tissue plasminogen activator. Protein Science, 2000, 9, 619-622.	7.6	6
90	Defining epitopes: It's not as easy as it seems. Nature Biotechnology, 1999, 17, 936-937.	17.5	64

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91	Conserved water molecules in the specificity pocket of serine proteases and the molecular mechanism of Na ⁺ binding. , 1998, 30, 34-42.		42
92	Role of P225 and the C136â€C201 disulfide bond in tissue plasminogen activator. Protein Science, 1998, 7, 1728-1737.	7.6	20
93	Histoplasma acquisition of calcium and expression of CBP1 during intracellular parasitism. Molecular Microbiology, 1998, 27, 531-539.	2.5	65
94	Synthesis and Characterization of More Potent Analogues of Hirudin Fragment 1â€47 Containing Non-Natural Amino Acids,. Biochemistry, 1998, 37, 13507-13515.	2.5	35
95	Site-Specific Thermodynamics:Â Understanding Cooperativity in Molecular Recognition. Chemical Reviews, 1998, 98, 1563-1592.	47.7	82
96	Site-Specific Analysis of Mutational Effects in Proteins. Advances in Protein Chemistry, 1998, 51, 59-119.	4.4	44
97	Kinetic Pathway for the Slow to Fast Transition of Thrombin. Journal of Biological Chemistry, 1997, 272, 30275-30282.	3.4	49
98	Energetics of Thrombinâ€Thrombomodulin Interaction. Biochemistry, 1997, 36, 6674-6681.	2.5	66
99	Site-specific dissection of substrate recognition by thrombin. Nature Biotechnology, 1997, 15, 891-895.	17.5	60
100	Rational engineering of activity and specificity in a serine protease. Nature Biotechnology, 1997, 15, 146-149.	17.5	101
101	Release of Fibrinopeptides by the Slow and Fast Forms of Thrombinâ€. Biochemistry, 1996, 35, 4417-4426.	2.5	84
102	Site-Specific thermodynamics of ising networks: A theorem for linearly connected subsystems. Biopolymers, 1994, 34, 673-678.	2.4	5
103	Thermodynamic basis of site-specific cooperativity. Biopolymers, 1994, 34, 1001-1005.	2.4	2
104	Looking at Fokker-Planck dynamics with a noisy instrument. Journal of Statistical Physics, 1993, 71, 1179-1190.	1.2	2
105	Meanâ€field treatment of local binding processes. Journal of Chemical Physics, 1992, 96, 6515-6522.	3.0	9
106	Thrombin is a sodium ion activated enzyme. Biochemistry, 1992, 31, 11721-11730.	2.5	247
107	Stochastic linkage: Effect of random fluctuations on a twoâ€state process. Journal of Chemical Physics, 1991, 95, 5082-5086.	3.0	47