Rex Pratt

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.

87 2,165 28 42 g-index

87 2,233 5.1 4.83 ext. papers ext. citations avg, IF L-index

#	Paper	IF	Citations
87	Detection of an enzyme isomechanism by means of the kinetics of covalent inhibition. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2021 , 1869, 140681	4	O
86	Elusive structural changes of New Delhi metallo-flactamase revealed by ultraviolet photodissociation mass spectrometry. <i>Chemical Science</i> , 2020 , 11, 8999-9010	9.4	4
85	A Lysine-Targeted Affinity Label for Serine-Lactamase Also Covalently Modifies New Delhi Metallo-Lactamase-1 (NDM-1). <i>Biochemistry</i> , 2019 , 58, 2834-2843	3.2	16
84	Specificity of extended O-aryloxycarbonyl hydroxamates as inhibitors of a class C llactamase. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 1430-1436	3.4	2
83	Kinetic Evidence for a Second Ligand Binding Site on Streptococcus pneumoniae Penicillin-Binding Protein 2x. <i>Biochemistry</i> , 2018 , 57, 1758-1766	3.2	2
82	Specificity and mechanism of mandelamide hydrolase catalysis. <i>Archives of Biochemistry and Biophysics</i> , 2017 , 618, 23-31	4.1	
81	Penicillin acylase and O-aryloxycarbonyl hydroxamates: Two acyl-enzymes, one leading to hydrolysis, the other to inactivation. <i>Archives of Biochemistry and Biophysics</i> , 2017 , 614, 65-71	4.1	4
80	Synthesis and Kinetic Analysis of Two Conformationally Restricted Peptide Substrates of Escherichia coli Penicillin-Binding Protein 5. <i>Biochemistry</i> , 2016 , 55, 4065-76	3.2	1
79	ELactamases: Why and How. Journal of Medicinal Chemistry, 2016, 59, 8207-20	8.3	27
78	A New Covalent Inhibitor of Class C Lactamases Reveals Extended Active Site Specificity. <i>Biochemistry</i> , 2015 , 54, 7375-84	3.2	7
77	Neutral £Lactams Inactivate High Molecular Mass Penicillin-Binding Proteins of Class B1, Including PBP2a of MRSA. <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 154-7	4.3	4
76	Inhibition of DD-peptidases by a specific trifluoroketone: crystal structure of a complex with the Actinomadura R39 DD-peptidase. <i>Biochemistry</i> , 2013 , 52, 2128-38	3.2	5
75	Kinetics of action of a two-stage pro-inhibitor of serine 🛭 actamases. <i>Biochemistry</i> , 2013 , 52, 7060-70	3.2	10
74	Dual substrate specificity of Bacillus subtilis PBP4a. <i>Biochemistry</i> , 2013 , 52, 2627-37	3.2	5
73	Covalent inhibition of serine 🛘 actamases by novel hydroxamic acid derivatives. <i>Biochemistry</i> , 2013 , 52, 3712-20	3.2	8
72	4-quinolones as noncovalent inhibitors of high molecular mass penicillin-binding proteins. <i>ACS Medicinal Chemistry Letters</i> , 2012 , 3, 592-5	4.3	15
71	Kinetics and stereochemistry of hydrolysis of an N-(phenylacetyl)-Ehydroxyglycine ester catalyzed by serine Elactamases and DD-peptidases. <i>Organic and Biomolecular Chemistry</i> , 2012 , 10, 7356-62	3.9	

(2007-2012)

70	Crossover inhibition as an indicator of convergent evolution of enzyme mechanisms: a Elactamase and a N-terminal nucleophile hydrolase. <i>FEBS Letters</i> , 2012 , 586, 4186-9	3.8	8
69	Inhibition of bacterial DD-peptidases (penicillin-binding proteins) in membranes and in vivo by peptidoglycan-mimetic boronic acids. <i>Biochemistry</i> , 2012 , 51, 2804-11	3.2	21
68	Unexpected tricovalent binding mode of boronic acids within the active site of a penicillin-binding protein. <i>Journal of the American Chemical Society</i> , 2011 , 133, 10839-48	16.4	34
67	Kinetics of reactions of the Actinomadura R39 DD-peptidase with specific substrates. <i>Biochemistry</i> , 2011 , 50, 376-87	3.2	9
66	Substrate specificity of low-molecular mass bacterial DD-peptidases. <i>Biochemistry</i> , 2011 , 50, 10091-101	3.2	17
65	Crystal structure of a complex between the Actinomadura R39 DD-peptidase and a peptidoglycan-mimetic boronate inhibitor: interpretation of a transition state analogue in terms of catalytic mechanism. <i>Biochemistry</i> , 2010 , 49, 6411-9	3.2	28
64	Serendipitous discovery of Ehydroxyalkyl esters as Elactamase substrates. <i>Biochemistry</i> , 2010 , 49, 10496-	5,026	1
63	Structural relationship between the active sites of Elactam-recognizing and amidase signature enzymes: convergent evolution?. <i>Biochemistry</i> , 2010 , 49, 9688-97	3.2	32
62	Crystal structures of covalent complexes of Elactam antibiotics with Escherichia coli penicillin-binding protein 5: toward an understanding of antibiotic specificity. <i>Biochemistry</i> , 2010 , 49, 8094-104	3.2	39
61	Substituted aryl malonamates as new serine beta-lactamase substrates: structure-activity studies. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 282-91	3.4	7
60	Approaches to the simultaneous inactivation of metallo- and serine-beta-lactamases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 1618-22	2.9	28
59	Intramolecular cooperativity in the reaction of diacyl phosphates with serine beta-lactamases. <i>Biochemistry</i> , 2009 , 48, 8293-8	3.2	4
58	Inhibition of class A and C beta-lactamases by diaroyl phosphates. <i>Biochemistry</i> , 2009 , 48, 8285-92	3.2	10
57	Crystal structures of complexes of bacterial DD-peptidases with peptidoglycan-mimetic ligands: the substrate specificity puzzle. <i>Journal of Molecular Biology</i> , 2008 , 381, 383-93	6.5	40
56	Kinetics and mechanism of inhibition of a serine beta-lactamase by O-aryloxycarbonyl hydroxamates. <i>Biochemistry</i> , 2008 , 47, 12037-46	3.2	20
55	Substrate specificity of bacterial DD-peptidases (penicillin-binding proteins). <i>Cellular and Molecular Life Sciences</i> , 2008 , 65, 2138-55	10.3	71
54	Beta-ketophosphonates as beta-lactamase inhibitors: Intramolecular cooperativity between the hydrophobic subsites of a class D beta-lactamase. <i>Bioorganic and Medicinal Chemistry</i> , 2008 , 16, 6987-94	.3.4	35
53	Reactions of peptidoglycan-mimetic beta-lactams with penicillin-binding proteins in vivo and in membranes. <i>ACS Chemical Biology</i> , 2007 , 2, 620-4	4.9	12

52	Crystal structure of the Bacillus subtilis penicillin-binding protein 4a, and its complex with a peptidoglycan mimetic peptide. <i>Journal of Molecular Biology</i> , 2007 , 371, 528-39	6.5	48
51	O-aryloxycarbonyl hydroxamates: new beta-lactamase inhibitors that cross-link the active site. <i>Journal of the American Chemical Society</i> , 2007 , 129, 9548-9	16.4	27
50	Reactivity of penicillin-binding proteins with peptidoglycan-mimetic beta-lactams: what's wrong with these enzymes?. <i>Biochemistry</i> , 2006 , 45, 15873-83	3.2	36
49	Deacylation transition states of a bacterial DD-peptidase. <i>Biochemistry</i> , 2006 , 45, 13074-82	3.2	7
48	Synthesis and beta-lactamase reactivity of alpha-substituted phenaceturates. <i>Bioorganic and Medicinal Chemistry</i> , 2006 , 14, 7023-33	3.4	15
47	Synthesis and reactivity with beta-lactamases of a monobactam bearing a retro-amide side chain. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 869-71	2.9	5
46	Synthesis and evaluation of ketophosph(on)ates as beta-lactamase inhibitors. <i>Journal of Organic Chemistry</i> , 2006 , 71, 4778-85	4.2	19
45	Transpeptidation reactions of a specific substrate catalyzed by the streptomyces R61 DD-peptidase: characterization of a chromogenic substrate and acyl acceptor design. <i>Biochemistry</i> , 2005 , 44, 9971-9	3.2	24
44	The D-methyl group in beta-lactamase evolution: evidence from the Y221G and GC1 mutants of the class C beta-lactamase of Enterobacter cloacae P99. <i>Biochemistry</i> , 2005 , 44, 7543-52	3.2	11
43	Inhibition of class D beta-lactamases by diaroyl phosphates. <i>Biochemistry</i> , 2005 , 44, 16121-9	3.2	16
42	Transpeptidation reactions of a specific substrate catalyzed by the Streptomyces R61 DD-peptidase: the structural basis of acyl acceptor specificity. <i>Biochemistry</i> , 2005 , 44, 9961-70	3.2	17
41	Crystal structures of complexes between the R61 DD-peptidase and peptidoglycan-mimetic beta-lactams: a non-covalent complex with a "perfect penicillin". <i>Journal of Molecular Biology</i> , 2005 , 345, 521-33	6.5	53
40	Inhibition of class D beta-lactamases by acyl phosphates and phosphonates. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 4410-2	5.9	18
39	Kinetic and structural consequences of the leaving group in substrates of a class C beta-lactamase. <i>Bioorganic and Medicinal Chemistry</i> , 2004 , 12, 1537-42	3.4	10
38	Benzopyranones with retro-amide side chains as (inhibitory) beta-lactamase substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004 , 14, 5117-20	2.9	4
37	Synthesis and evaluation of new substrate analogues of streptomyces R61 DD-peptidase: dissection of a specific ligand. <i>Journal of Organic Chemistry</i> , 2004 , 69, 7472-8	4.2	12
36	Kinetics of turnover of cefotaxime by the Enterobacter cloacae P99 and GCl beta-lactamases: two free enzyme forms of the P99 beta-lactamase detected by a combination of pre- and post-steady state kinetics. <i>Biochemistry</i> , 2004 , 43, 2664-72	3.2	13
35	The perfect penicillin? Inhibition of a bacterial DD-peptidase by peptidoglycan-mimetic beta-lactams. <i>Journal of the American Chemical Society</i> , 2004 , 126, 8122-3	16.4	32

34	The crystal structure of phosphonate-inhibited D-Ala-D-Ala peptidase reveals an analogue of a tetrahedral transition state. <i>Biochemistry</i> , 2003 , 42, 1199-208	3.2	55
33	New substrates for beta-lactam-recognizing enzymes: aryl malonamates. <i>Biochemistry</i> , 2003 , 42, 6719-2.	5 3.2	7
32	Functional evolution of the serine Elactamase active site. Perkin Transactions II RSC, 2002, 851-861		41
31	Structures of two kinetic intermediates reveal species specificity of penicillin-binding proteins. Journal of Molecular Biology, 2002 , 322, 111-22	6.5	78
30	The synthesis and evaluation of benzofuranones as beta-lactamase substrates. <i>Bioorganic and Medicinal Chemistry</i> , 2001 , 9, 1175-83	3.4	28
29	Inverse acyl phosph(on)ates: substrates or inhibitors of beta-lactam-recognizing enzymes?. <i>Bioorganic Chemistry</i> , 2001 , 29, 271-81	5.1	8
28	Potential Substrates/Inhibitors of Lactam-Recognizing Enzymes. European Journal of Organic	3.2	13
27	Chemistry, 2001 , 2001, 141-149 Mechanism of reaction of acyl phosph(on)ates with the beta-lactamase of Enterobacter cloacae P99. <i>Biochemistry</i> , 2001 , 40, 4610-21	3.2	21
26	Dipeptide binding to the extended active site of the Streptomyces R61 D-alanyl-D-alanine-peptidase: the path to a specific substrate. <i>Biochemistry</i> , 2000 , 39, 12200-9	3.2	34
25	A "cephalosporin-like" cyclic depsipeptide: synthesis and reaction with beta-lactam-recognizing enzymes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999 , 9, 341-6	2.9	4
24	On the importance of a methyl group in beta-lactamase evolution: free energy profiles and molecular modeling. <i>Biochemistry</i> , 1999 , 38, 10499-510	3.2	32
23	Beta-secondary and solvent deuterium kinetic isotope effects on catalysis by the Streptomyces R61 DD-peptidase: comparisons with a structurally similar class C beta-lactamase. <i>Biochemistry</i> , 1999 , 38, 1469-77	3.2	24
22	Synthesis and Reactivity with beta-Lactamases of "Penicillin-like" Cyclic Depsipeptides. <i>Journal of Organic Chemistry</i> , 1999 , 64, 713-720	4.2	35
21	Salicyloyl Cyclic Phosphate, a B enicillin-LikeInhibitor of ELactamases. <i>Journal of the American Chemical Society</i> , 1998 , 120, 3004-3006	16.4	12
20	Inhibition of Serine ELactamases by Acyl Phosph(on)ates: [A New Source of Inert Acyl [and Phosphyl] Enzymes. <i>Journal of the American Chemical Society</i> , 1998 , 120, 4264-4268	16.4	18
19	Reaction of soluble penicillin-binding protein 2a of methicillin-resistant Staphylococcus aureus with beta-lactams and acyclic substrates: kinetics in homogeneous solution. <i>Biochemical Journal</i> , 1998 , 332 (Pt 3), 755-61	3.8	48
18	Effectiveness of Tetrahedral Adducts as Transition-State Analogs and Inhibitors of the Class C Lactamase of Enterobacter cloacae P99. <i>Journal of the American Chemical Society</i> , 1997 , 119, 1529-1538	3 ^{16.4}	56
17	Structure-activity studies of the inhibition of serine beta-lactamases by phosphonate monoesters. Bioorganic and Medicinal Chemistry, 1997 , 5, 1783-8	3.4	24

16	Kinetics and mechanism of the hydrolysis of depsipeptides catalyzed by the beta-lactamase of Enterobacter cloacae P99. <i>Biochemistry</i> , 1996 , 35, 3595-603	3.2	34
15	8-Hydroxypenillic Acid from 6-Aminopenicillanic Acid: A New Reaction Catalyzed by a Class C Lactamase. <i>Journal of the American Chemical Society</i> , 1996 , 118, 8207-8212	16.4	8
14	Beta-secondary and solvent deuterium kinetic isotope effects on beta-lactamase catalysis. <i>Biochemistry</i> , 1996 , 35, 3604-13	3.2	32
13	Characterization of covalently bound enzyme inhibitors as transition-state analogs by protein stability measurements: phosphonate monoester inhibitors of a beta-lactamase. <i>Biochemistry</i> , 1994 , 33, 116-25	3.2	40
12	Crystallographic structure of a phosphonate derivative of the Enterobacter cloacae P99 cephalosporinase: mechanistic interpretation of a beta-lactamase transition-state analog. <i>Biochemistry</i> , 1994 , 33, 6762-72	3.2	158
11	Relative specificities of a series of beta-lactam-recognizing enzymes towards the side-chains of penicillins and of acyclic thioldepsipeptides. <i>Biochemical Journal</i> , 1994 , 302 (Pt 3), 851-6	3.8	26
10	Effect of side-chain amide thionation on turnover of beta-lactam substrates by beta-lactamases. Further evidence on the question of side-chain hydrogen-bonding in catalysis. <i>Biochemical Journal</i> , 1992 , 286 (Pt 3), 857-62	3.8	4
9	Mechanism of inhibition of the class C beta-lactamase of Enterobacter cloacae P99 by phosphonate monoesters. <i>Biochemistry</i> , 1992 , 31, 5869-78	3.2	49
8	N-(phenylacetyl)glycyl-D-aziridine-2-carboxylate, an acyclic amide substrate of beta-lactamases: importance of the shape of the substrate in beta-lactamase evolution. <i>Biochemistry</i> , 1991 , 30, 3640-9	3.2	38
7	Inhibition of a class C beta-lactamase by a specific phosphonate monoester. <i>Science</i> , 1989 , 246, 917-9	33.3	119
6	Effect of the 3'-leaving group on turnover of cephem antibiotics by a class C beta-lactamase. <i>Biochemical Journal</i> , 1989 , 259, 255-60	3.8	36
5	Accumulation of acyl-enzyme intermediates during turnover of penicillins by the class A beta-lactamase of Staphylococcus aureus PC1. <i>Biochemical Journal</i> , 1988 , 254, 919-22	3.8	23
4	Nucleophilic re-activation of the PC1 beta-lactamase of Staphylococcus aureus and of the DD-peptidase of Streptomyces R61 after their inactivation by cephalosporins and cephamycins. <i>Biochemical Journal</i> , 1987 , 246, 651-8	3.8	11
3	Kinetics and mechanism of the serine beta-lactamase catalyzed hydrolysis of depsipeptides. <i>Biochemistry</i> , 1987 , 26, 3385-95	3.2	80
2	Interactions of cephalosporins with the Streptomyces R61 DD-transpeptidase/carboxypeptidase. Influence of the 3'-substituent. <i>Biochemical Journal</i> , 1986 , 238, 309-12	3.8	18
1	beta-Lactamase-catalyzed hydrolysis of acyclic depsipeptides and acyl transfer to specific amino acid acceptors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1984 , 81, 1302-6	11.5	58