William C K Pomerantz

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A Structure-based Design Approach for Generating High Affinity BRD4 D1-Selective Chemical Probes. Journal of Medicinal Chemistry, 2022, 65, 2342-2360.	6.4	19
2	An Innovation 10 Years in the Making: The Stories in the Pages of <i>ACS Medicinal Chemistry Letters</i> . ACS Medicinal Chemistry Letters, 2022, 13, 540-545.	2.8	0
3	Tracking Fluorine during Aqueous Photolysis and Advanced UV Treatment of Fluorinated Phenols and Pharmaceuticals Using a Combined ¹⁹ F-NMR, Chromatography, and Mass Spectrometry Approach. ACS Environmental Au, 2022, 2, 242-252.	7.0	9
4	Development of a Highly Responsive Organofluorine Temperature Sensor for ¹⁹ F Magnetic Resonance Applications. Analytical Chemistry, 2022, 94, 3782-3790.	6.5	4
5	Dihydropyridine Lactam Analogs Targeting BET Bromodomains. ChemMedChem, 2022, 17, e202100407.	3.2	1
6	In This Issue, Volume 13, Issue 4. ACS Medicinal Chemistry Letters, 2022, 13, 515-516.	2.8	0
7	Alternative Mechanisms for DNA Engagement by BET Bromodomain-Containing Proteins. Biochemistry, 2022, 61, 1260-1272.	2.5	4
8	Development of a single culture E. coli expression system for the enzymatic synthesis of fluorinated tyrosine and its incorporation into proteins. Journal of Fluorine Chemistry, 2022, 261-262, 110014.	1.7	0
9	Endothelial p300 Promotes Portal Hypertension and Hepatic Fibrosis Through C Motif Chemokine Ligand 2–Mediated Angiocrine Signaling. Hepatology, 2021, 73, 2468-2483.	7.3	52
10	Selective Nâ€Terminal BET Bromodomain Inhibitors by Targeting Nonâ€Conserved Residues and Structured Water Displacement**. Angewandte Chemie, 2021, 133, 1240-1246.	2.0	0
11	Selective Nâ€Terminal BET Bromodomain Inhibitors by Targeting Non onserved Residues and Structured Water Displacement**. Angewandte Chemie - International Edition, 2021, 60, 1220-1226.	13.8	27
12	¹⁹ F NMR viewed through two different lenses: ligand-observed and protein-observed ¹⁹ F NMR applications for fragment-based drug discovery. RSC Chemical Biology, 2021, 2, 1312-1330.	4.1	35
13	Autophagy-Dependent Sensitization of Triple-Negative Breast Cancer Models to Topoisomerase II Poisons by Inhibition of the Nucleosome Remodeling Factor. Molecular Cancer Research, 2021, 19, 1338-1349.	3.4	9
14	Investigation of the Post-Synthetic Confinement of Fluorous Liquids Inside Mesoporous Silica Nanoparticles. Langmuir, 2021, 37, 5222-5231.	3.5	3
15	Fragment-Based Ligand Discovery Using Protein-Observed ¹⁹ F NMR: A Second Semester Organic Chemistry CURE Project. Journal of Chemical Education, 2021, 98, 1963-1973.	2.3	8
16	Soluble Methane Monooxygenase Component Interactions Monitored by ¹⁹ F NMR. Biochemistry, 2021, 60, 1995-2010.	2.5	8
17	Potent inhibitors of toxic alpha-synuclein identified via cellular time-resolved FRET biosensors. Npj Parkinson's Disease, 2021, 7, 52.	5.3	22
18	Super enhancer regulation of cytokine-induced chemokine production in alcoholic hepatitis. Nature Communications, 2021, 12, 4560.	12.8	37

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19	Controlling Intramolecular Interactions in the Design of Selective, High-Affinity Ligands for the CREBBP Bromodomain. Journal of Medicinal Chemistry, 2021, 64, 10102-10123.	6.4	17
20	4-Methyl-1,2,3-Triazoles as <i>N</i> -Acetyl-Lysine Mimics Afford Potent BET Bromodomain Inhibitors with Improved Selectivity. Journal of Medicinal Chemistry, 2021, 64, 10497-10511.	6.4	22
21	Opportunity knocks for uncovering the new function of an understudied nucleosome remodeling complex member, the bromodomain PHD finger transcription factor, BPTF. Current Opinion in Chemical Biology, 2021, 63, 57-67.	6.1	11
22	In This Issue, Volume 12, Issue 9. ACS Medicinal Chemistry Letters, 2021, 12, 1357-1358.	2.8	0
23	New Design Rules for Developing Potent Cell-Active Inhibitors of the Nucleosome Remodeling Factor (NURF) via BPTF Bromodomain Inhibition. Journal of Medicinal Chemistry, 2021, 64, 13902-13917.	6.4	14
24	Siteâ€Specific 5â€Formyl Cytosine Mediated DNAâ€Histone Crossâ€Links: Synthesis and Polymerase Bypass by Human DNA Polymerase Î∙. Angewandte Chemie - International Edition, 2021, 60, 26489-26494.	13.8	7
25	Multidimensional Nanoparticle Characterization through Ion Mobility-Mass Spectrometry. Analytical Chemistry, 2020, 92, 2503-2510.	6.5	16
26	2-Fluorotyrosine is a valuable but understudied amino acid for protein-observed 19F NMR. Journal of Biomolecular NMR, 2020, 74, 61-69.	2.8	14
27	In This Issue, Volume 11, Issue 10 ("Medicinal Chemistry: From Targets to Therapies―Special Issue). ACS Medicinal Chemistry Letters, 2020, 11, 1783-1784.	2.8	0
28	Quantifying the Selectivity of Protein–Protein and Small Molecule Interactions with Fluorinated Tandem Bromodomain Reader Proteins. ACS Chemical Biology, 2020, 15, 3038-3049.	3.4	4
29	In this Issue, Volume 11, Issue 7. ACS Medicinal Chemistry Letters, 2020, 11, 1492-1493.	2.8	0
30	Combined Protein- and Ligand-Observed NMR Workflow to Screen Fragment Cocktails against Multiple Proteins: A Case Study Using Bromodomains. Molecules, 2020, 25, 3949.	3.8	10
31	NMR Analyses of Acetylated H2A.Z Isoforms Identify Differential Binding Interactions with the Bromodomain of the NURF Nucleosome Remodeling Complex. Biochemistry, 2020, 59, 1871-1880.	2.5	11
32	New inhibitors for the BPTF bromodomain enabled by structural biology and biophysical assay development. Organic and Biomolecular Chemistry, 2020, 18, 5174-5182.	2.8	14
33	Efficient Synthesis of 1,4-Thiazepanones and 1,4-Thiazepanes as 3D Fragments for Screening Libraries. Organic Letters, 2020, 22, 3946-3950.	4.6	12
34	Systematically Mitigating the p38α Activity of Triazole-based BET Inhibitors. ACS Medicinal Chemistry Letters, 2019, 10, 1296-1301.	2.8	22
35	SAR by (Protein-Observed) ¹⁹ F NMR. Accounts of Chemical Research, 2019, 52, 3407-3418.	15.6	42
36	Selectivity, ligand deconstruction, and cellular activity analysis of a BPTF bromodomain inhibitor. Organic and Biomolecular Chemistry, 2019, 17, 2020-2027.	2.8	18

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37	In This Issue, Volume 10, Issue 1. ACS Medicinal Chemistry Letters, 2019, 10, 1-1.	2.8	Ο
38	In This Issue, Volume 10, Issue 10. ACS Medicinal Chemistry Letters, 2019, 10, 1359-1360.	2.8	0
39	Evaluating the Advantages of Using 3D-Enriched Fragments for Targeting BET Bromodomains. ACS Medicinal Chemistry Letters, 2019, 10, 1648-1654.	2.8	26
40	Applied Biophysics for Bromodomain Drug Discovery. Topics in Medicinal Chemistry, 2019, , 287-337.	0.8	3
41	Dual Labeling of the CBP/p300 KIX Domain for ¹⁹ Fâ€NMR Leads to Identification of a New Smallâ€Molecule Binding Site. ChemBioChem, 2018, 19, 963-969.	2.6	9
42	Molecular Basis for the N-Terminal Bromodomain-and-Extra-Terminal-Family Selectivity of a Dual Kinase–Bromodomain Inhibitor. Journal of Medicinal Chemistry, 2018, 61, 9316-9334.	6.4	56
43	Design, Synthesis, and Characterization of a Fluorescence Polarization Pan-BET Bromodomain Probe. ACS Medicinal Chemistry Letters, 2018, 9, 1223-1229.	2.8	8
44	Oxygen Sensing with Perfluorocarbon-Loaded Ultraporous Mesostructured Silica Nanoparticles. ACS Nano, 2017, 11, 5623-5632.	14.6	40
45	Quantifying Protein Concentrations Using Smartphone Colorimetry: A New Method for an Established Test. Journal of Chemical Education, 2017, 94, 941-945.	2.3	43
46	Synthesis of Intrinsically Disordered Fluorinated Peptides for Modular Design of Highâ€5ignal ¹⁹ F MRI Agents. Angewandte Chemie - International Edition, 2017, 56, 6440-6444.	13.8	37
47	Synthesis of Intrinsically Disordered Fluorinated Peptides for Modular Design of High‣ignal ¹⁹ F MRI Agents. Angewandte Chemie, 2017, 129, 6540-6544.	2.0	5
48	Tuning Sulfur Oxidation States on Thioetherâ€Bridged Peptide Macrocycles for Modulation of Protein Interactions. ChemBioChem, 2017, 18, 1836-1844.	2.6	18
49	BET Bromodomain Inhibitors with One-Step Synthesis Discovered from Virtual Screen. Journal of Medicinal Chemistry, 2017, 60, 4805-4817.	6.4	39
50	BPTF Maintains Chromatin Accessibility and the Self-Renewal Capacity of Mammary Gland Stem Cells. Stem Cell Reports, 2017, 9, 23-31.	4.8	43
51	Meeting Proceedings ICBS2016—Translating the Power of Chemical Biology to Clinical Advances. ACS Chemical Biology, 2017, 12, 869-877.	3.4	2
52	Specific Acetylation Patterns of H2A.Z Form Transient Interactions with the BPTF Bromodomain. Biochemistry, 2017, 56, 4607-4615.	2.5	16
53	Throwing Away the Cookbook: Implementing Course-Based Undergraduate Research Experiences (CUREs) in Chemistry. ACS Symposium Series, 2017, , 33-63.	0.5	37
54	Oxidation increases the strength of the methionine-aromatic interaction. Nature Chemical Biology, 2016, 12, 860-866.	8.0	53

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55	Protein-Observed Fluorine NMR Is a Complementary Ligand Discovery Method to ¹ H CPMG Ligand-Observed NMR. ACS Chemical Biology, 2016, 11, 3154-3164.	3.4	40
56	Protein-observed 19F-NMR for fragment screening, affinity quantification and druggability assessment. Nature Protocols, 2016, 11, 1414-1427.	12.0	82
57	Paramagnetic relaxation enhancement for protein-observed ¹⁹ F NMR as an enabling approach for efficient fragment screening. RSC Advances, 2016, 6, 95715-95721.	3.6	13
58	Prediction of ¹⁹ F NMR Chemical Shifts in Labeled Proteins: Computational Protocol and Case Study. Molecular Pharmaceutics, 2016, 13, 2376-2386.	4.6	23
59	Protein-Observed Fluorine NMR: A Bioorthogonal Approach for Small Molecule Discovery. Journal of Medicinal Chemistry, 2016, 59, 5158-5171.	6.4	144
60	Fragment Screening and Druggability Assessment for the CBP/p300 KIX Domain through Proteinâ€Observed ¹⁹ Fâ€NMR Spectroscopy. Angewandte Chemie - International Edition, 2015, 54, 3735-3739.	13.8	52
61	Dual Screening of BPTF and Brd4 Using Protein-Observed Fluorine NMR Uncovers New Bromodomain Probe Molecules. ACS Chemical Biology, 2015, 10, 2246-2256.	3.4	64