Stuart L Schreiber

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.

45,102 90 309 211 h-index g-index citations papers 16.1 52,081 341 7.59 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
309	Bicyclic azetidines target acute and chronic stages of Toxoplasma gondii by inhibiting parasite phenylalanyl t-RNA synthetase <i>Nature Communications</i> , 2022 , 13, 459	17.4	O
308	Stereochemical Diversity as a Source of Discovery in Chemical Biology. <i>Current Research in Chemical Biology</i> , 2022 , 2, 100028		4
307	PALP: A rapid imaging technique for stratifying ferroptosis sensitivity in normal and tumor tissues in situ. <i>Cell Chemical Biology</i> , 2021 ,	8.2	2
306	Computational repurposing of therapeutic small molecules from cancer to pulmonary hypertension. <i>Science Advances</i> , 2021 , 7, eabh3794	14.3	7
305	An expanded universe of cancer targets. <i>Cell</i> , 2021 , 184, 1142-1155	56.2	38
304	Cell-specific transcriptional control of mitochondrial metabolism by TIF1drives erythropoiesis. <i>Science</i> , 2021 , 372, 716-721	33.3	8
303	The Use of Informer Sets in Screening: Perspectives on an Efficient Strategy to Identify New Probes. <i>SLAS Discovery</i> , 2021 , 26, 855-861	3.4	1
302	Crystal structures of the selenoprotein glutathione peroxidase 4 in its apo form and in complex with the covalently bound inhibitor ML162. <i>Acta Crystallographica Section D: Structural Biology</i> , 2021 , 77, 237-248	5.5	15
301	Targeted brachyury degradation disrupts a highly specific autoregulatory program controlling chordoma cell identity. <i>Cell Reports Medicine</i> , 2021 , 2, 100188	18	4
300	Novel quaternary structures of the human prion protein globular domain. <i>Biochimie</i> , 2021 , 191, 118-12	5 4.6	0
299	The Rise of Molecular Glues. <i>Cell</i> , 2021 , 184, 3-9	56.2	74
298	Structural basis of malaria parasite phenylalanine tRNA-synthetase inhibition by bicyclic azetidines. <i>Nature Communications</i> , 2021 , 12, 343	17.4	8
297	Persister cancer cells: Iron addiction and vulnerability to ferroptosis Molecular Cell, 2021,	17.6	4
296	Evaluating drug targets through human loss-of-function genetic variation. <i>Nature</i> , 2020 , 581, 459-464	50.4	53
295	Towards a treatment for genetic prion disease: trials and biomarkers. <i>Lancet Neurology, The</i> , 2020 , 19, 361-368	24.1	28
294	Unifying principles of bifunctional, proximity-inducing small molecules. <i>Nature Chemical Biology</i> , 2020 , 16, 369-378	11.7	53
293	Selective covalent targeting of GPX4 using masked nitrile-oxide electrophiles. <i>Nature Chemical Biology</i> , 2020 , 16, 497-506	11.7	76

(2019-2020)

292	Rhabdoid Tumors Are Sensitive to the Protein-Translation Inhibitor Homoharringtonine. <i>Clinical Cancer Research</i> , 2020 , 26, 4995-5006	12.9	6
291	Ligand-Enabled EMethylene C(sp3)日 Arylation of Masked Aliphatic Alcohols. <i>Angewandte Chemie</i> , 2020 , 132, 7857-7861	3.6	4
290	Cytochrome P450 oxidoreductase contributes to phospholipid peroxidation in ferroptosis. <i>Nature Chemical Biology</i> , 2020 , 16, 302-309	11.7	144
289	Ligand-Enabled EMethylene C(sp)-H Arylation of Masked Aliphatic Alcohols. <i>Angewandte Chemie - International Edition</i> , 2020 , 59, 7783-7787	16.4	20
288	Water-Compatible Cycloadditions of Oligonucleotide-Conjugated Strained Allenes for DNA-Encoded Library Synthesis. <i>Journal of the American Chemical Society</i> , 2020 , 142, 7776-7782	16.4	28
287	Bicyclic azetidines kill the diarrheal pathogen in mice by inhibiting parasite phenylalanyl-tRNA synthetase. <i>Science Translational Medicine</i> , 2020 , 12,	17.5	18
286	Prion protein lowering is a disease-modifying therapy across prion disease stages, strains and endpoints. <i>Nucleic Acids Research</i> , 2020 , 48, 10615-10631	20.1	26
285	An Activity-Guided Map of Electrophile-Cysteine Interactions in Primary Human T Cells. <i>Cell</i> , 2020 , 182, 1009-1026.e29	56.2	57
284	Phosphorylation-Inducing Chimeric Small Molecules. <i>Journal of the American Chemical Society</i> , 2020 , 142, 14052-14057	16.4	30
283	Multimodal small-molecule screening for human prion protein binders. <i>Journal of Biological Chemistry</i> , 2020 , 295, 13516-13531	5.4	6
282	Structure-activity relationships of GPX4 inhibitor warheads. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020 , 30, 127538	2.9	7
281	Plasticity of ether lipids promotes ferroptosis susceptibility and evasion. <i>Nature</i> , 2020 , 585, 603-608	50.4	121
280	Recent achievements and current trajectories of diversity-oriented synthesis. <i>Current Opinion in Chemical Biology</i> , 2020 , 56, 1-9	9.7	36
279	Progress in Understanding Ferroptosis and Challenges in Its Targeting for Therapeutic Benefit. <i>Cell Chemical Biology</i> , 2020 , 27, 463-471	8.2	27
278	Small-Molecule and CRISPR Screening Converge to Reveal Receptor Tyrosine Kinase Dependencies in Pediatric Rhabdoid Tumors. <i>Cell Reports</i> , 2019 , 28, 2331-2344.e8	10.6	20
277	Metabolomic adaptations and correlates of survival to immune checkpoint blockade. <i>Nature Communications</i> , 2019 , 10, 4346	17.4	89
276	Small-molecule targeting of brachyury transcription factor addiction in chordoma. <i>Nature Medicine</i> , 2019 , 25, 292-300	50.5	62
275	DNA Barcoding a Complete Matrix of Stereoisomeric Small Molecules. <i>Journal of the American Chemical Society</i> , 2019 , 141, 10225-10235	16.4	47

274	The landscape of cancer cell line metabolism. <i>Nature Medicine</i> , 2019 , 25, 850-860	50.5	188
273	Modular, stereocontrolled C-H/C-C activation of alkyl carboxylic acids. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 8721-8727	11.5	24
272	A GPX4-dependent cancer cell state underlies the clear-cell morphology and confers sensitivity to ferroptosis. <i>Nature Communications</i> , 2019 , 10, 1617	17.4	218
271	Prion protein quantification in human cerebrospinal fluid as a tool for prion disease drug development. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 7793-7798	11.5	29
270	6-Phosphogluconate Dehydrogenase Links Cytosolic Carbohydrate Metabolism to Protein Secretion via Modulation of Glutathione Levels. <i>Cell Chemical Biology</i> , 2019 , 26, 1306-1314.e5	8.2	11
269	Optimization of PDE3A Modulators for SLFN12-Dependent Cancer Cell Killing. <i>ACS Medicinal Chemistry Letters</i> , 2019 , 10, 1537-1542	4.3	7
268	A Compendium of Genetic Modifiers of Mitochondrial Dysfunction Reveals Intra-organelle Buffering. <i>Cell</i> , 2019 , 179, 1222-1238.e17	56.2	47
267	Antisense oligonucleotides extend survival of prion-infected mice. JCI Insight, 2019, 5,	9.9	46
266	Renal medullary carcinomas depend upon loss and are sensitive to proteasome inhibition. <i>ELife</i> , 2019 , 8,	8.9	20
265	Characterization of the Prion Protein Binding Properties of Antisense Oligonucleotides. <i>Biomolecules</i> , 2019 , 10,	5.9	52
264	DNA-Compatible [3 + 2] Nitrone-Olefin Cycloaddition Suitable for DEL Syntheses. <i>Organic Letters</i> , 2019 , 21, 1325-1330	6.2	47
263	Domain-specific Quantification of Prion Protein in Cerebrospinal Fluid by Targeted Mass Spectrometry. <i>Molecular and Cellular Proteomics</i> , 2019 , 18, 2388-2400	7.6	17
262	Diacylfuroxans Are Masked Nitrile Oxides That Inhibit GPX4 Covalently. <i>Journal of the American Chemical Society</i> , 2019 , 141, 20407-20415	16.4	31
261	A Chemical Biology View of Bioactive Small Molecules and a Binder-Based Approach to Connect Biology to Precision Medicines. <i>Israel Journal of Chemistry</i> , 2019 , 59, 52-59	3.4	37
260	Chemical probes and drug leads from advances in synthetic planning and methodology. <i>Nature Reviews Drug Discovery</i> , 2018 , 17, 333-352	64.1	117
259	RWEN: response-weighted elastic net for prediction of chemosensitivity of cancer cell lines. <i>Bioinformatics</i> , 2018 , 34, 3332-3339	7.2	14
258	Targeting Dependency on the GPX4 Lipid Peroxide Repair Pathway for Cancer Therapy. <i>Biochemistry</i> , 2018 , 57, 2059-2060	3.2	37
257	High Throughput Screen Identifies Interferon Dependent Inhibitors of Toxoplasma gondii Growth. <i>ACS Infectious Diseases</i> , 2018 , 4, 1499-1507	5.5	7

(2016-2018)

256	Retraction Note: Selective killing of cancer cells by a small molecule targeting the stress response to ROS. <i>Nature</i> , 2018 , 561, 420	50.4	5
255	Synergistic Effects of Stereochemistry and Appendages on the Performance Diversity of a Collection of Synthetic Compounds. <i>Journal of the American Chemical Society</i> , 2018 , 140, 11784-11790	16.4	35
254	A precision oncology approach to the pharmacological targeting of mechanistic dependencies in neuroendocrine tumors. <i>Nature Genetics</i> , 2018 , 50, 979-989	36.3	90
253	Chemical Biology Towards Precision Medicine. <i>Israel Journal of Chemistry</i> , 2017 , 57, 174-178	3.4	
252	Discovery of Antimalarial Azetidine-2-carbonitriles That Inhibit Dihydroorotate Dehydrogenase. <i>ACS Medicinal Chemistry Letters</i> , 2017 , 8, 438-442	4.3	32
251	A dataset of images and morphological profiles of 30 000 small-molecule treatments using the Cell Painting assay. <i>GigaScience</i> , 2017 , 6, 1-5	7.6	59
250	Drug-tolerant persister cancer cells are vulnerable to GPX4 inhibition. <i>Nature</i> , 2017 , 551, 247-250	50.4	522
249	Small-molecule inhibitors directly target CARD9 and mimic its protective variant in inflammatory bowel disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, 11392-11397	11.5	28
248	CTD2 Dashboard: a searchable web interface to connect validated results from the Cancer Target Discovery and Development Network. <i>Database: the Journal of Biological Databases and Curation</i> , 2017 , 2017,	5	12
247	Synthesis of a Bicyclic Azetidine with In Vivo Antimalarial Activity Enabled by Stereospecific, Directed C(sp)-H Arylation. <i>Journal of the American Chemical Society</i> , 2017 , 139, 11300-11306	16.4	71
246	Stereospecific Palladium-Catalyzed C-H Arylation of Pyroglutamic Acid Derivatives at the C3 Position Enabled by 8-Aminoquinoline as a Directing Group. <i>Organic Letters</i> , 2017 , 19, 4424-4427	6.2	22
245	Small-molecule studies identify CDK8 as a regulator of IL-10 in myeloid cells. <i>Nature Chemical Biology</i> , 2017 , 13, 1102-1108	11.7	29
244	A Next Generation Connectivity Map: L1000 Platform and the First 1,000,000 Profiles. <i>Cell</i> , 2017 , 171, 1437-1452.e17	56.2	1132
243	Dependency of a therapy-resistant state of cancer cells on a lipid peroxidase pathway. <i>Nature</i> , 2017 , 547, 453-457	50.4	620
242	A small-molecule allosteric inhibitor of Mycobacterium tuberculosis tryptophan synthase. <i>Nature Chemical Biology</i> , 2017 , 13, 943-950	11.7	75
241	DIFFERENTIAL PATHWAY DEPENDENCY DISCOVERY ASSOCIATED WITH DRUG RESPONSE ACROSS CANCER CELL LINES. <i>Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing</i> , 2017 , 22, 497-508	1.3	6
240	Structural Insight into Allosteric Inhibition of Mycobacterium tuberculosis Tryptophan Synthase. <i>FASEB Journal</i> , 2017 , 31, 765.12	0.9	1
239	Discovery of 8-Membered Ring Sulfonamides as Inhibitors of Oncogenic Mutant Isocitrate Dehydrogenase 1. <i>ACS Medicinal Chemistry Letters</i> , 2016 , 7, 944-949	4.3	12

238	Efficient Routes to a Diverse Array of Amino Alcohol-Derived Chiral Fragments. <i>ACS Combinatorial Science</i> , 2016 , 18, 569-74	3.9	20
237	Inhibition of Dihydroorotate Dehydrogenase Overcomes Differentiation Blockade in Acute Myeloid Leukemia. <i>Cell</i> , 2016 , 167, 171-186.e15	56.2	214
236	A genetic basis for the variation in the vulnerability of cancer to DNA damage. <i>Nature Communications</i> , 2016 , 7, 11428	17.4	95
235	Discovery of selective small-molecule HDAC6 inhibitor for overcoming proteasome inhibitor resistance in multiple myeloma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, 13162-13167	11.5	89
234	Development of Chemical Probes for Investigation of Salt-Inducible Kinase Function in Vivo. <i>ACS Chemical Biology</i> , 2016 , 11, 2105-11	4.9	33
233	Identification of cancer-cytotoxic modulators of PDE3A by predictive chemogenomics. <i>Nature Chemical Biology</i> , 2016 , 12, 102-8	11.7	51
232	Correlating chemical sensitivity and basal gene expression reveals mechanism of action. <i>Nature Chemical Biology</i> , 2016 , 12, 109-16	11.7	365
231	High-Throughput Luciferase-Based Assay for the Discovery of Therapeutics That Prevent Malaria. <i>ACS Infectious Diseases</i> , 2016 , 2, 281-293	5.5	61
230	The Power of Sophisticated Phenotypic Screening and Modern Mechanism-of-Action Methods. <i>Cell Chemical Biology</i> , 2016 , 23, 3-9	8.2	70
229	High-throughput identification of genotype-specific cancer vulnerabilities in mixtures of barcoded tumor cell lines. <i>Nature Biotechnology</i> , 2016 , 34, 419-23	44.5	127
228	High-Throughput Assay and Discovery of Small Molecules that Interrupt Malaria Transmission. <i>Cell Host and Microbe</i> , 2016 , 19, 114-26	23.4	94
227	Real-Time Biological Annotation of Synthetic Compounds. <i>Journal of the American Chemical Society</i> , 2016 , 138, 8920-7	16.4	27
226	Integrated genetic and pharmacologic interrogation of rare cancers. <i>Nature Communications</i> , 2016 , 7, 11987	17.4	32
225	Divergent Synthesis and Real-Time Biological Annotation of Optically Active Tetrahydrocyclopenta[c]pyranone Derivatives. <i>Organic Letters</i> , 2016 , 18, 6280-6283	6.2	9
224	Inhibition of Zinc-Dependent Histone Deacetylases with a Chemically Triggered Electrophile. <i>ACS Chemical Biology</i> , 2016 , 11, 1844-51	4.9	18
223	Discovering Innovative Therapies for Rare Tumors: Combining Genetically Accurate Disease Models with In Silico Analysis to Identify Novel Therapeutic Targets. <i>Clinical Cancer Research</i> , 2016 , 22, 3903-14	12.9	43
222	Diversity-oriented synthesis yields novel multistage antimalarial inhibitors. <i>Nature</i> , 2016 , 538, 344-349	50.4	172
221	Development of ML390: A Human DHODH Inhibitor That Induces Differentiation in Acute Myeloid Leukemia. <i>ACS Medicinal Chemistry Letters</i> , 2016 , 7, 1112-1117	4.3	36

220	Benzo-fused lactams from a diversity-oriented synthesis (DOS) library as inhibitors of scavenger receptor BI (SR-BI)-mediated lipid uptake. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 2100-5	2.9	11
219	Discovery of a Small-Molecule Probe for V-ATPase Function. <i>Journal of the American Chemical Society</i> , 2015 , 137, 5563-8	16.4	24
218	Harnessing Connectivity in a Large-Scale Small-Molecule Sensitivity Dataset. <i>Cancer Discovery</i> , 2015 , 5, 1210-23	24.4	363
217	Small-molecule enhancers of autophagy modulate cellular disease phenotypes suggested by human genetics. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, E4281-7	11.5	46
216	KRAS Genomic Status Predicts the Sensitivity of Ovarian Cancer Cells to Decitabine. <i>Cancer Research</i> , 2015 , 75, 2897-906	10.1	22
215	Linking tumor mutations to drug responses via a quantitative chemical-genetic interaction map. <i>Cancer Discovery</i> , 2015 , 5, 154-67	24.4	40
214	Discovery of bisamide-heterocycles as inhibitors of scavenger receptor BI (SR-BI)-mediated lipid uptake. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 2594-8	2.9	6
213	Kinase-Independent Small-Molecule Inhibition of JAK-STAT Signaling. <i>Journal of the American Chemical Society</i> , 2015 , 137, 7929-34	16.4	20
212	Diversity-oriented synthesis probe targets Plasmodium falciparum cytochrome b ubiquinone reduction site and synergizes with oxidation site inhibitors. <i>Journal of Infectious Diseases</i> , 2015 , 211, 1097-103	7	21
211	Advancing Biological Understanding and Therapeutics Discovery with Small-Molecule Probes. <i>Cell</i> , 2015 , 161, 1252-65	56.2	100
210	Indolinyl-Thiazole Based Inhibitors of Scavenger Receptor-BI (SR-BI)-Mediated Lipid Transport. <i>ACS Medicinal Chemistry Letters</i> , 2015 , 6, 375-380	4.3	8
209	Niche-Based Screening in Multiple Myeloma Identifies a Kinesin-5 Inhibitor with Improved Selectivity over Hematopoietic Progenitors. <i>Cell Reports</i> , 2015 , 10, 755-770	10.6	18
208	Synthesis of oxazocenones via gold(I)-catalyzed 8-endo-dig hydroalkoxylation of alkynamides. <i>Organic Letters</i> , 2015 , 17, 418-21	6.2	29
207	Chemical perturbation of an intrinsically disordered region of TFIID distinguishes two modes of transcription initiation. <i>ELife</i> , 2015 , 4,	8.9	29
206	Regulation of ferroptotic cancer cell death by GPX4. <i>Cell</i> , 2014 , 156, 317-331	56.2	2104
205	Lenalidomide causes selective degradation of IKZF1 and IKZF3 in multiple myeloma cells. <i>Science</i> , 2014 , 343, 301-5	33.3	969
204	NAMPT is the cellular target of STF-31-like small-molecule probes. ACS Chemical Biology, 2014 , 9, 2247-5	5 4 .9	47
203	Toward performance-diverse small-molecule libraries for cell-based phenotypic screening using multiplexed high-dimensional profiling. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 10911-6	11.5	141

202	Diversity-oriented synthesis-facilitated medicinal chemistry: toward the development of novel antimalarial agents. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 8496-502	8.3	31
201	Diversity-oriented synthesis yields a new drug lead for treatment of chagas disease. <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 149-53	4.3	34
200	Quantitative-proteomic comparison of alpha and Beta cells to uncover novel targets for lineage reprogramming. <i>PLoS ONE</i> , 2014 , 9, e95194	3.7	10
199	Atg16L1 T300A variant decreases selective autophagy resulting in altered cytokine signaling and decreased antibacterial defense. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 7741-6	11.5	235
198	Small-molecule control of cytokine function: new opportunities for treating immune disorders. <i>Current Opinion in Chemical Biology</i> , 2014 , 23, 23-30	9.7	16
197	Automated Structure-Activity Relationship Mining: Connecting Chemical Structure to Biological Profiles. <i>Journal of Biomolecular Screening</i> , 2014 , 19, 738-48		14
196	Small-molecule screening identifies inhibition of salt-inducible kinases as a therapeutic strategy to enhance immunoregulatory functions of dendritic cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 12468-73	11.5	45
195	Synthesis of piperlogs and analysis of their effects on cells. <i>Tetrahedron</i> , 2013 , 69, 7559-7559	2.4	19
194	An interactive resource to identify cancer genetic and lineage dependencies targeted by small molecules. <i>Cell</i> , 2013 , 154, 1151-1161	56.2	392
193	Niche-based screening identifies small-molecule inhibitors of leukemia stem cells. <i>Nature Chemical Biology</i> , 2013 , 9, 840-848	11.7	96
192	Discovery of small-molecule enhancers of reactive oxygen species that are nontoxic or cause genotype-selective cell death. <i>ACS Chemical Biology</i> , 2013 , 8, 923-9	4.9	52
191	Crebinostat: a novel cognitive enhancer that inhibits histone deacetylase activity and modulates chromatin-mediated neuroplasticity. <i>Neuropharmacology</i> , 2013 , 64, 81-96	5.5	75
190	A small-molecule inducer of PDX1 expression identified by high-throughput screening. <i>Chemistry and Biology</i> , 2013 , 20, 1513-22		25
189	Integrative radiogenomic profiling of squamous cell lung cancer. Cancer Research, 2013, 73, 6289-98	10.1	83
188	Multiplex cytological profiling assay to measure diverse cellular states. <i>PLoS ONE</i> , 2013 , 8, e80999	3.7	136
187	Development of small-molecule probes that selectively kill cells induced to express mutant RAS. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 1822-6	2.9	99
186	Identification of a selective small molecule inhibitor of breast cancer stem cells. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012 , 22, 3571-4	2.9	26
185	Macrocyclic Hedgehog Pathway Inhibitors: Optimization of Cellular Activity and Mode of Action Studies. <i>ACS Medicinal Chemistry Letters</i> , 2012 , 3, 808-813	4.3	35

(2010-2012)

184	Diversity-Oriented Synthesis Yields a Novel Lead for the Treatment of Malaria. <i>ACS Medicinal Chemistry Letters</i> , 2012 , 3, 112-117	4.3	48
183	Synthesis, cellular evaluation, and mechanism of action of piperlongumine analogs. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 15115-20	11.5	172
182	Niche-Based Screening Identifies Novel Small Molecules That Overcome Stromal Effects in Multiple Myeloma. <i>Blood</i> , 2012 , 120, 571-571	2.2	1
181	Syntheses of ⊕yrones using gold-catalyzed coupling reactions. <i>Organic Letters</i> , 2011 , 13, 2834-6	6.2	79
180	Selective killing of cancer cells by a small molecule targeting the stress response to ROS. <i>Nature</i> , 2011 , 475, 231-4	50.4	845
179	Catalytic Diastereoselective Petasis Reactions. <i>Angewandte Chemie</i> , 2011 , 123, 8322-8325	3.6	12
178	Catalytic diastereoselective petasis reactions. <i>Angewandte Chemie - International Edition</i> , 2011 , 50, 8172	2156.4	54
177	Discovery of histone deacetylase 8 selective inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 2601-5	2.9	72
176	The DNA damage mark pH2AX differentiates the cytotoxic effects of small molecule HDAC inhibitors in ovarian cancer cells. <i>Cancer Biology and Therapy</i> , 2011 , 12, 484-93	4.6	38
175	Disease allele-dependent small-molecule sensitivities in blood cells from monogenic diabetes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 492-7	11.5	15
174	Organic synthesis toward small-molecule probes and drugs. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 6699-702	11.5	124
173	Quantifying structure and performance diversity for sets of small molecules comprising small-molecule screening collections. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 6817-22	11.5	81
172	Towards patient-based cancer therapeutics. <i>Nature Biotechnology</i> , 2010 , 28, 904-6	44.5	58
171	Small molecules of different origins have distinct distributions of structural complexity that correlate with protein-binding profiles. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 18787-92	11.5	253
170	Distinct biological network properties between the targets of natural products and disease genes. Journal of the American Chemical Society, 2010 , 132, 9259-61	16.4	67
169	Small-molecule inducers of insulin expression in pancreatic alpha-cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 15099-104	11.5	51
168	Expanding stereochemical and skeletal diversity using petasis reactions and 1,3-dipolar cycloadditions. <i>Organic Letters</i> , 2010 , 12, 5230-3	6.2	27
167	Small-Molecule Suppressors of Cytokine-Induced beta-Cell Apoptosis. <i>ACS Chemical Biology</i> , 2010 , 5, 729-34	4.9	33

166	Stereochemical and skeletal diversity arising from amino propargylic alcohols. <i>Organic Letters</i> , 2010 , 12, 2822-5	6.2	47
165	Binding affinity and kinetic analysis of targeted small molecule-modified nanoparticles. <i>Bioconjugate Chemistry</i> , 2010 , 21, 14-9	6.3	166
164	Using expression and genotype to predict drug response in yeast. <i>PLoS ONE</i> , 2009 , 4, e6907	3.7	14
163	A small molecule that binds Hedgehog and blocks its signaling in human cells. <i>Nature Chemical Biology</i> , 2009 , 5, 154-6	11.7	239
162	Aziridines as intermediates in diversity-oriented syntheses of alkaloids. <i>Tetrahedron Letters</i> , 2009 , 50, 3230-3233	2	30
161	Syntheses of aminoalcohol-derived macrocycles leading to a small-molecule binder to and inhibitor of Sonic Hedgehog. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 6319-25	2.9	64
160	Synthesis and conformation-activity relationships of the peptide isosteres of FK228 and largazole. Journal of the American Chemical Society, 2009 , 131, 2900-5	16.4	98
159	Gold(I)-catalyzed coupling reactions for the synthesis of diverse small molecules using the build/couple/pair strategy. <i>Journal of the American Chemical Society</i> , 2009 , 131, 5667-74	16.4	88
158	SnapShot: Ca2+-calcineurin-NFAT signaling. <i>Cell</i> , 2009 , 138, 210, 210.e1	56.2	80
157	Identification and characterization of small molecule inhibitors of a class I histone deacetylase from Plasmodium falciparum. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 2185-7	8.3	68
156	Skeletally diverse small molecules using a build/couple/pair strategy. <i>Organic Letters</i> , 2009 , 11, 1559-62	6.2	46
155	Unbiased discovery of in vivo imaging probes through in vitro profiling of nanoparticle libraries. <i>Integrative Biology (United Kingdom)</i> , 2009 , 1, 311-7	3.7	20
154	The M2 splice isoform of pyruvate kinase is important for cancer metabolism and tumour growth. <i>Nature</i> , 2008 , 452, 230-3	50.4	2056
153	Diversity synthesis of complex pyridines yields a probe of a neurotrophic signaling pathway. <i>Organic Letters</i> , 2008 , 10, 2621-4	6.2	42
152	Small-molecule reagents for cellular pull-down experiments. <i>Bioconjugate Chemistry</i> , 2008 , 19, 585-7	6.3	13
151	Towards the optimal screening collection: a synthesis strategy. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 48-56	16.4	471
150	Synthesis and cellular profiling of diverse organosilicon small molecules. <i>Journal of the American Chemical Society</i> , 2007 , 129, 1020-1	16.4	170
149	Fluorous-based small-molecule microarrays for the discovery of histone deacetylase inhibitors. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 7960-4	16.4	80

(2006-2007)

148	Complex alpha-pyrones synthesized by a gold-catalyzed coupling reaction. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 8250-3	16.4	118	
147	Identification of a small-molecule inhibitor of class Ia PI3Ks with cell-based screening. <i>Chemistry and Biology</i> , 2007 , 14, 371-7		30	
146	Rethinking relationships between natural products. <i>Nature Chemical Biology</i> , 2007 , 3, 352	11.7	6	
145	Small molecules enhance autophagy and reduce toxicity in Huntington's disease models. <i>Nature Chemical Biology</i> , 2007 , 3, 331-8	11.7	513	
144	Genetic basis of individual differences in the response to small-molecule drugs in yeast. <i>Nature Genetics</i> , 2007 , 39, 496-502	36.3	93	
143	Ring-opening and ring-closing reactions of a shikimic acid-derived substrate leading to diverse small molecules. <i>ACS Combinatorial Science</i> , 2007 , 9, 245-53		22	
142	Identification of novel epoxide inhibitors of hepatitis C virus replication using a high-throughput screen. <i>Antimicrobial Agents and Chemotherapy</i> , 2007 , 51, 3756-9	5.9	17	
141	Quantifying fitness distributions and phenotypic relationships in recombinant yeast populations. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007 , 104, 10553-8	11.5	7	
140	A Soft-Drug Histone Deacetylase Inhibitor for Cutaneous T-Cell Lymphoma <i>Blood</i> , 2007 , 110, 800-800	2.2	2	
139	Autophagy as a Target Pathway in Multiple Myeloma: A Forward Chemical Genetic Approach <i>Blood</i> , 2007 , 110, 2510-2510	2.2		
138	Short synthesis of skeletally and stereochemically diverse small molecules by coupling petasis condensation reactions to cyclization reactions. <i>Angewandte Chemie - International Edition</i> , 2006 , 45, 3635-8	16.4	151	
137	Short Synthesis of Skeletally and Stereochemically Diverse Small Molecules by Coupling Petasis Condensation Reactions to Cyclization Reactions. <i>Angewandte Chemie</i> , 2006 , 118, 3717-3720	3.6	34	
136	Small molecules, big players: the National Cancer Institute's Initiative for Chemical Genetics. <i>Cancer Research</i> , 2006 , 66, 8935-42	10.1	62	
135	An oligomer-based approach to skeletal diversity in small-molecule synthesis. <i>Journal of the American Chemical Society</i> , 2006 , 128, 14766-7	16.4	74	
134	Macrocycloadditions leading to conformationally restricted small molecules. <i>Organic Letters</i> , 2006 , 8, 2063-6	6.2	40	
133	Microarray-based method for monitoring yeast overexpression strains reveals small-molecule targets in TOR pathway. <i>Nature Chemical Biology</i> , 2006 , 2, 103-9	11.7	81	
132	Revealing complex traits with small molecules and naturally recombinant yeast strains. <i>Chemistry and Biology</i> , 2006 , 13, 319-27		34	
131	A robust small-molecule microarray platform for screening cell lysates. <i>Chemistry and Biology</i> , 2006 , 13, 493-504		114	

130	Histone Deacetylase-6 (HDAC6) Modulates Akt and STAT3 Activity Via Heat Shock Protein (Hsp) 90 in Human Multiple Myeloma (MM) Cells <i>Blood</i> , 2006 , 108, 3426-3426	2.2	
129	Design and Characterization of a Novel, Reverse Prodrug Histone Deacetylase Inhibitor for Cutaneous T-Cell Lymphoma <i>Blood</i> , 2006 , 108, 4759-4759	2.2	
128	Discovery and Characterization of Small Molecule Inhibitors of Autophagy for Cancer Therapy <i>Blood</i> , 2006 , 108, 2606-2606	2.2	
127	Skeletal diversity via a folding pathway: synthesis of indole alkaloid-like skeletons. <i>Organic Letters</i> , 2005 , 7, 47-50	6.2	109
126	Small-molecule diversity using a skeletal transformation strategy. <i>Organic Letters</i> , 2005 , 7, 2535-8	6.2	55
125	Small molecules: the missing link in the central dogma. <i>Nature Chemical Biology</i> , 2005 , 1, 64-6	11.7	255
124	Convergent diversity-oriented synthesis of small-molecule hybrids. <i>Angewandte Chemie - International Edition</i> , 2005 , 44, 2249-52	16.4	71
123	Convergent Diversity-Oriented Synthesis of Small-Molecule Hybrids. <i>Angewandte Chemie</i> , 2005 , 117, 2289-2292	3.6	14
122	From solution-phase to solid-phase enyne metathesis: crossover in the relative performance of two commonly used ruthenium pre-catalysts. <i>Chemistry - A European Journal</i> , 2005 , 11, 5086-93	4.8	14
121	Small-molecule inhibition of proteasome and aggresome function induces synergistic antitumor activity in multiple myeloma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 8567-72	11.5	526
120	Perturbational profiling of a cell-line model of tumorigenesis by using metabolic measurements. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 5992-7	11.5	295
119	Targeting the Protein Degradation Pathway in Multiple Myeloma with Synergistic, Selective Small Molecules <i>Blood</i> , 2005 , 106, 2471-2471	2.2	
118	Finding new components of the target of rapamycin (TOR) signaling network through chemical genetics and proteome chips. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004 , 101, 16594-9	11.5	208
117	A planning strategy for diversity-oriented synthesis. <i>Angewandte Chemie - International Edition</i> , 2004 , 43, 46-58	16.4	1239
116	Synthetic strategy toward skeletal diversity via solid-supported, otherwise unstable reactive intermediates. <i>Angewandte Chemie - International Edition</i> , 2004 , 43, 1681-5	16.4	74
115	Eine Strategie fildie DiversitEs-orientierte Synthese. <i>Angewandte Chemie</i> , 2004 , 116, 48-60	3.6	261
114	Synthetic Strategy toward Skeletal Diversity via Solid-Supported, Otherwise Unstable Reactive Intermediates. <i>Angewandte Chemie</i> , 2004 , 116, 1713-1717	3.6	8
113	Modular synthesis and preliminary biological evaluation of stereochemically diverse 1,3-dioxanes. <i>Chemistry and Biology</i> , 2004 , 11, 1279-91		29

(2002-2004)

112	Relationship of stereochemical and skeletal diversity of small molecules to cellular measurement space. <i>Journal of the American Chemical Society</i> , 2004 , 126, 14740-5	16.4	122
111	A synthesis strategy yielding skeletally diverse small molecules combinatorially. <i>Journal of the American Chemical Society</i> , 2004 , 126, 14095-104	16.4	174
110	A library of spirooxindoles based on a stereoselective three-component coupling reaction. <i>Journal of the American Chemical Society</i> , 2004 , 126, 16077-86	16.4	246
109	Identification and Characterization of Novel Small-Molecule Inhibitors of the Replication Checkpoint <i>Blood</i> , 2004 , 104, 763-763	2.2	
108	Chemical genomic profiling of biological networks using graph theory and combinations of small molecule perturbations. <i>Journal of the American Chemical Society</i> , 2003 , 125, 10543-5	16.4	52
107	Expanding the functional group compatibility of small-molecule microarrays: discovery of novel calmodulin ligands. <i>Angewandte Chemie - International Edition</i> , 2003 , 42, 2376-9	16.4	83
106	Structural biasing elements for in-cell histone deacetylase paralog selectivity. <i>Journal of the American Chemical Society</i> , 2003 , 125, 5586-7	16.4	104
105	Generating diverse skeletons of small molecules combinatorially. <i>Science</i> , 2003 , 302, 613-8	33.3	350
104	From knowing to controlling: a path from genomics to drugs using small molecule probes. <i>Science</i> , 2003 , 300, 294-5	33.3	253
103	Integration of growth factor and nutrient signaling: implications for cancer biology. <i>Molecular Cell</i> , 2003 , 12, 271-80	17.6	172
102	Domain-selective small-molecule inhibitor of histone deacetylase 6 (HDAC6)-mediated tubulin deacetylation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003 , 100, 4389-94	11.5	890
101	Discovery of an inhibitor of a transcription factor using small molecule microarrays and diversity-oriented synthesis. <i>Journal of the American Chemical Society</i> , 2003 , 125, 8420-1	16.4	167
100	A Boronic Ester Annulation Strategy for Diversity-Oriented Organic Synthesis. <i>Angewandte Chemie</i> , 2002 , 114, 160-162	3.6	8
99	An Alkynylboronic Ester Annulation: Development of Synthetic Methods for Application to Diversity-Oriented Organic Synthesis. <i>Angewandte Chemie</i> , 2002 , 114, 3406-3410	3.6	12
98	A boronic ester annulation strategy for diversity-oriented organic synthesis. <i>Angewandte Chemie - International Edition</i> , 2002 , 41, 152-4	16.4	60
97	An alkynylboronic ester annulation: development of synthetic methods for application to diversity-oriented organic synthesis. <i>Angewandte Chemie - International Edition</i> , 2002 , 41, 3272-6	16.4	49
96	Dissecting glucose signalling with diversity-oriented synthesis and small-molecule microarrays. <i>Nature</i> , 2002 , 416, 653-7	50.4	344
95	Deacetylase enzymes: biological functions and the use of small-molecule inhibitors. <i>Chemistry and Biology</i> , 2002 , 9, 3-16		477

94	Skeletal diversity via a branched pathway: efficient synthesis of 29 400 discrete, polycyclic compounds and their arraying into stock solutions. <i>Journal of the American Chemical Society</i> , 2002 , 124, 13402-4	16.4	112
93	Signaling network model of chromatin. <i>Cell</i> , 2002 , 111, 771-8	56.2	319
92	A one-bead, one-stock solution approach to chemical genetics: part 1. <i>Chemistry and Biology</i> , 2001 , 8, 1167-82		98
91	A one-bead, one-stock solution approach to chemical genetics: part 2. <i>Chemistry and Biology</i> , 2001 , 8, 1183-95		83
90	Asymmetric Catalysis in Diversity-Oriented Organic Synthesis: Enantioselective Synthesis of 4320 Encoded and Spatially Segregated Dihydropyrancarboxamides We thank the National Institute of General Medical Sciences (GM-52067) for support of this research, and Dr. John Tallarico and Max	16.4	69
89	Narovlyansky at the ICCB for generously providing resin for the library synthesis. We thank Dr. Synthesis of 7200 small molecules based on a substructural analysis of the histone deacetylase ic inhibitors trichostatin and trapoxin. Organic Letters, 2001, 3, 4239-42 dition, 2001, 40, 3417-3421	6.2	131
88	Exploiting site-site interactions on solid support to generate dimeric molecules. <i>Organic Letters</i> , 2001 , 3, 1185-8	6.2	40
87	An alkylsilyl-tethered, high-capacity solid support amenable to diversity-oriented synthesis for one-bead, one-stock solution chemical genetics. <i>ACS Combinatorial Science</i> , 2001 , 3, 312-8		106
86	Dissecting cellular processes using small molecules: identification of colchicine-like, taxol-like and other small molecules that perturb mitosis. <i>Chemistry and Biology</i> , 2000 , 7, 275-86		211
85	Small-Molecule Microarrays: Covalent Attachment and Screening of Alcohol-Containing Small Molecules on Glass Slides. <i>Journal of the American Chemical Society</i> , 2000 , 122, 7849-7850	16.4	177
84	Pairwise use of complexity-generating reactions in diversity-oriented organic synthesis. <i>Organic Letters</i> , 2000 , 2, 709-12	6.2	232
83	Printing proteins as microarrays for high-throughput function determination. <i>Science</i> , 2000 , 289, 1760-3	33.3	2033
82	High-throughput screening of small molecules in miniaturized mammalian cell-based assays involving post-translational modifications. <i>Chemistry and Biology</i> , 1999 , 6, 71-83		176
81	The identification of myriocin-binding proteins. <i>Chemistry and Biology</i> , 1999 , 6, 221-35		80
80	Selection of gp41-mediated HIV-1 cell entry inhibitors from biased combinatorial libraries of non-natural binding elements. <i>Nature Structural Biology</i> , 1999 , 6, 953-60		127
79	Small molecule inhibitor of mitotic spindle bipolarity identified in a phenotype-based screen. <i>Science</i> , 1999 , 286, 971-4	33.3	1450
78	Visualizing Functional Group Distribution in Solid-Support Beads by Using Optical Analysis. <i>Chemistry - A European Journal</i> , 1999 , 5, 3528-3532	4.8	32
77	Molecular association between ATR and two components of the nucleosome remodeling and deacetylating complex, HDAC2 and CHD4. <i>Biochemistry</i> , 1999 , 38, 14711-7	3.2	80

76	Printing Small Molecules as Microarrays and Detecting Proteinligand Interactions en Masse. Journal of the American Chemical Society, 1999 , 121, 7967-7968	16.4	408
<i>75</i>	Chromatin deacetylation by an ATP-dependent nucleosome remodelling complex. <i>Nature</i> , 1998 , 395, 917-21	50.4	558
74	Chemical genetics resulting from a passion for synthetic organic chemistry. <i>Bioorganic and Medicinal Chemistry</i> , 1998 , 6, 1127-52	3.4	387
73	Exploring the Specificity Pockets of Two Homologous SH3 Domains Using Structure-Based, Split-Pool Synthesis and Affinity-Based Selection. <i>Journal of the American Chemical Society</i> , 1998 , 120, 23-29	16.4	38
72	Dimerization as a regulatory mechanism in signal transduction. <i>Annual Review of Immunology</i> , 1998 , 16, 569-92	34.7	279
71	Cell-Specific Calcineurin Inhibition by a Modified Cyclosporin. <i>Journal of the American Chemical Society</i> , 1997 , 119, 1805-1806	16.4	31
70	Single-Step Synthesis of Cell-Permeable Protein Dimerizers That Activate Signal Transduction and Gene Expression. <i>Journal of the American Chemical Society</i> , 1997 , 119, 5106-5109	16.4	77
69	Regulatory intramolecular association in a tyrosine kinase of the Tec family. <i>Nature</i> , 1997 , 385, 93-7	50.4	237
68	Three-part inventions: intracellular signaling and induced proximity. <i>Trends in Biochemical Sciences</i> , 1996 , 21, 418-22	10.3	128
67	Dimeric ligands define a role for transcriptional activation domains in reinitiation. <i>Nature</i> , 1996 , 382, 822-6	50.4	234
66	Structure of guanine-nucleotide-exchange factor human Mss4 and identification of its Rab-interacting surface. <i>Nature</i> , 1995 , 376, 788-91	50.4	54
65	Control of p70 s6 kinase by kinase activity of FRAP in vivo. <i>Nature</i> , 1995 , 377, 441-6	50.4	626
64	Kombinatorische Synthese und mehrdimensionale NMR-Spektroskopie: ein Beitrag zum Verst dnis von Protein-Ligand-Wechselwirkungen. <i>Angewandte Chemie</i> , 1995 , 107, 1041-1058	3.6	5
63	Rationales Design neuer Rezeptor-Ligand-Kombinationen. <i>Angewandte Chemie</i> , 1995 , 107, 2313-2317	3.6	4
62	Combinatorial Synthesis and Multidimensional NMR Spectroscopy: An Approach to Understanding Protein Ligand Interactions. <i>Angewandte Chemie International Edition in English</i> , 1995 , 34, 953-969		39
61	Rational Design of Orthogonal Receptorligand Combinations. <i>Angewandte Chemie International Edition in English</i> , 1995 , 34, 2129-2132		77
60	Proximity versus allostery: the role of regulated protein dimerization in biology. <i>Chemistry and Biology</i> , 1994 , 1, 131-6		65
59	Mechanistic studies of a signaling pathway activated by the organic dimerizer FK1012. <i>Chemistry and Biology</i> , 1994 , 1, 163-72		53

58	A mammalian protein targeted by G1-arresting rapamycin-receptor complex. <i>Nature</i> , 1994 , 369, 756-8	50.4	1617
57	1H and 15N assignments and secondary structure of the Src SH3 domain. FEBS Letters, 1993 , 324, 87-92	2 3.8	45
56	1H and 15N assignments and secondary structure of the PI3K SH3 domain. FEBS Letters, 1993, 324, 93-8	3.8	16
55	Immunophilin-sensitive protein phosphatase action in cell signaling pathways. <i>Cell</i> , 1992 , 70, 365-8	56.2	309
54	The mechanism of action of cyclosporin A and FK506. <i>Trends in Immunology</i> , 1992 , 13, 136-42		1937
53	Natural Products as Probes of Cellular Function: Studies of Immunophilins. <i>Angewandte Chemie International Edition in English</i> , 1992 , 31, 384-400		128
52	Naturstoffe als Sonden zum Studium zelluller Funktionen (Untersuchungen von Immunophilinen. <i>Angewandte Chemie</i> , 1992 , 104, 413-430	3.6	33
51	The effect of the immunosuppressant FK-506 on alternate pathways of T cell activation. <i>European Journal of Immunology</i> , 1991 , 21, 439-45	6.1	73
50	Calcineurin is a common target of cyclophilin-cyclosporin A and FKBP-FK506 complexes. <i>Cell</i> , 1991 , 66, 807-15	56.2	3538
49	Molecular cloning and overexpression of the human FK506-binding protein FKBP. <i>Nature</i> , 1990 , 346, 671-4	50.4	299
48	N-oxide promoted pauson-khand cyclizations at room temperature. <i>Tetrahedron Letters</i> , 1990 , 31, 5289)- <u>5</u> 292	291
47	On the Conformation and Structure of Organometal Complexes in the Solid State: Two Studies Relevant to Chemical Synthesis. <i>Angewandte Chemie International Edition in English</i> , 1990 , 29, 256-272		164
46	An Asymmetric Synthesis of (+)-Cryptone. <i>Synthetic Communications</i> , 1990 , 20, 1159-1165	1.7	8
45	A receptor for the immunosuppressant FK506 is a cis-trans peptidyl-prolyl isomerase. <i>Nature</i> , 1989 , 341, 758-60	50.4	1216
44	Fragmentation reactions of .alphaalkoxy hydroperoxides and application to the synthesis of the macrolide (.+)-recifeiolide. <i>Journal of the American Chemical Society</i> , 1980 , 102, 6163-6165	16.4	119
43	Using Small Molecules to Unravel Biological Mechanisms71-94		3
42	Chemical Informatics723-759		1
41	Revealing Biological Specificity by Engineering Protein-Ligand Interactions115-139		2

40	Chemical Strategies for Activity-based Proteomics403-426	1
39	Chemistry and Biology [Historical and Philosophical Aspects3-67	2
38	Chemical Complementation: Bringing the Power of Genetics to Chemistry199-226	
37	Synthetic Expansion of the Central Dogma271-295	
36	Combinatorial Biosynthesis of Polyketides and Nonribosomal Peptides519-536	1
35	New Methods for Protein Bioconjugation593-634	14
34	The Molecular Basis of Predicting Druggability804-823	3
33	WOMBAT and WOMBAT-PK: Bioactivity Databases for Lead and Drug Discovery760-786	58
32	Diversity-oriented Synthesis483-518	7
31	The Search for Chemical Probes to Illuminate Carbohydrate Function635-667	1
30	The Bicyclic Depsipeptide Family of Histone Deacetylase Inhibitors693-720	3
29	Managerial Challenges in Implementing Chemical Biology Platforms789-803	1
28	The Target Family Approach825-851	1
27	Reverse Chemical Genetics[An Important Strategy for the Study of Protein Function in Chemical Biology and Drug Discovery355-384	1
26	Using Natural Products to Unravel Cell Biology95-114	3
25	Ozonolytic Cleavage of Cyclohexene to Terminally Differentiated Products: Methyl 6-Oxohexanoate, 6,6-Dimethoxyhexanal, Methyl 6,6-Dimethoxyhexanoate150-150	
24	HIF-2Edrives an intrinsic vulnerability to ferroptosis in clear cell renal cell carcinoma	1
23	Domain-specific quantification of prion protein in cerebrospinal fluid by targeted mass spectrometry	1

22	Characterization of the prion protein binding properties of antisense oligonucleotides	1
21	Prion protein lowering is a disease-modifying therapy across prion disease stages, strains, and endpoints	4
20	Multimodal small-molecule screening for human prion protein binders	1
19	Modulation of ferroptosis sensitivity by TXNRD1 in pancreatic cancer cells	4
18	Targeting a Therapy-Resistant Cancer Cell State Using Masked Electrophiles as GPX4 Inhibitors	10
17	A chemical biology view of bioactive small molecules and a binder-based approach to connect biology to precision medicines	3
16	Evaluating potential drug targets through human loss-of-function genetic variation	12
15	Expressed Protein Ligation537-566	2
14	Chemical Synthesis of Proteins and Large Bioconjugates567-592	1
13	Chemical Glycomics as Basis for Drug Discovery668-691	2
12	The Nuclear Receptor Superfamily and Drug Discovery891-932	3
11	Prediction of ADMET Properties1003-1042	4
10	Controlling Protein Function by Caged Compounds140-173	2
9	Controlling Protein Protein Interactions Using Chemical Inducers and Disrupters of Dimerization 227-249	7
8	Protein Secondary Structure Mimetics as Modulators of Protein Protein and Protein-Ligand Interactions 250-2	26%
7	The Biarsenical-tetracysteine Protein Tag: Chemistry and Biological Applications427-457	1
6	Chemical Approaches to Exploit Fusion Proteins for Functional Studies458-479	1
5	Chemical Biology of Kinases Studied by NMR Spectroscopy852-890	

LIST OF PUBLICATIONS

- The GPCRITTM Receptor Target Family933-978
- 3 Systems Biology of the JAK-STAT Signaling Pathway1045-1060
- 2 Modeling Intracellular Signal Transduction Processes 1061-1081
- Genome-wide Gene Expression Analysis: Practical Considerations and Application to the Analysis of T-cell Subsets in Inflammatory Diseases1083-1117