

# Anne C Conibear

## 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.

33  
papers

724  
citations

14  
h-index

26  
g-index

40  
ext. papers

908  
ext. citations

6.9  
avg, IF

4.78  
L-index

#	Paper	IF	Citations
33	A conserved Ebulge glycine residue facilitates folding and increases stability of the mouse Edefensin cryptdin-4. <i>Peptide Science</i> , <b>2022</b> , 114, e24250	3	
32	Segmental and site-specific isotope labelling strategies for structural analysis of posttranslationally modified proteins. <i>RSC Chemical Biology</i> , <b>2021</b> , 2, 1441-1461	3	3
31	Site-specific modification and segmental isotope labelling of HMG1 reveals long-range conformational perturbations caused by posttranslational modifications. <i>RSC Chemical Biology</i> , <b>2021</b> , 2, 537-550	3	5
30	Posttranslational modifications of Eonotoxins: sulfotyrosine and C-terminal amidation stabilise structures and increase acetylcholine receptor binding. <i>RSC Medicinal Chemistry</i> , <b>2021</b> , 12, 1574-1584	3.5	
29	Synthesis and anti-parasitic activity of achiral N-benzylated phosphoramidic acid derivatives. <i>Bioorganic Chemistry</i> , <b>2020</b> , 101, 103947	5.1	2
28	Recent Advances in Peptide-Based Approaches for Cancer Treatment. <i>Current Medicinal Chemistry</i> , <b>2020</b> , 27, 1174-1205	4.3	10
27	Deciphering protein post-translational modifications using chemical biology tools. <i>Nature Reviews Chemistry</i> , <b>2020</b> , 4, 674-695	34.6	37
26	Tumor-Targeting Immune System Engagers (ISERs) Activate Human Neutrophils after Binding to Cancer Cells. <i>Biochemistry</i> , <b>2019</b> , 58, 2642-2652	3.2	0
25	Multifunctional Scaffolds for Assembling Cancer-Targeting Immune Stimulators Using Chemoselective Ligations. <i>Frontiers in Chemistry</i> , <b>2019</b> , 7, 113	5	3
24	Random coil shifts of posttranslationally modified amino acids. <i>Journal of Biomolecular NMR</i> , <b>2019</b> , 73, 587-599	3	13
23	Protein Chemistry Looking Ahead: 8 Chemical Protein Synthesis Meeting 16-19 June 2019, Berlin, Germany. <i>Cell Chemical Biology</i> , <b>2019</b> , 26, 1349-1354	8.2	
22	Synthetic Cancer-Targeting Innate Immune Stimulators Give Insights into Avidity Effects. <i>ChemBioChem</i> , <b>2018</b> , 19, 459-469	3.8	2
21	Native chemical ligation in protein synthesis and semi-synthesis. <i>Chemical Society Reviews</i> , <b>2018</b> , 47, 9046-9068	15.8	158
20	Multifunctional EIntegrin-Specific Peptide-Pt(IV) Conjugates for Cancer Cell Targeting. <i>Bioconjugate Chemistry</i> , <b>2017</b> , 28, 2429-2439	6.3	9
19	Synthetic integrin-binding immune stimulators target cancer cells and prevent tumor formation. <i>Scientific Reports</i> , <b>2017</b> , 7, 17592	4.9	5
18	A comparative study of synthetic and semisynthetic approaches for ligating the epidermal growth factor to a bivalent scaffold. <i>Journal of Peptide Science</i> , <b>2017</b> , 23, 871-879	2.1	3
17	Efficient enzymatic cyclization of an inhibitory cystine knot-containing peptide. <i>Biotechnology and Bioengineering</i> , <b>2016</b> , 113, 2202-12	4.9	21

16	Arginine side-chain modification that occurs during copper-catalysed azide-alkyne click reactions resembles an advanced glycation end product. <i>Organic and Biomolecular Chemistry</i> , <b>2016</b> , 14, 6205-11	3.9	17
15	Approaches to the stabilization of bioactive epitopes by grafting and peptide cyclization. <i>Biopolymers</i> , <b>2016</b> , 106, 89-100	2.2	33
14	Mirror Images of Antimicrobial Peptides Provide Reflections on Their Functions and Amyloidogenic Properties. <i>Journal of the American Chemical Society</i> , <b>2016</b> , 138, 5706-13	16.4	36
13	Transforming conotoxins into cyclotides: Backbone cyclization of P-superfamily conotoxins. <i>Biopolymers</i> , <b>2015</b> , 104, 682-92	2.2	10
12	The chemistry and biology of theta defensins. <i>Angewandte Chemie - International Edition</i> , <b>2014</b> , 53, 10612-23	2.2	59
11	The cyclic cystine ladder of theta-defensins as a stable, bifunctional scaffold: A proof-of-concept study using the integrin-binding RGD motif. <i>ChemBioChem</i> , <b>2014</b> , 15, 451-9	3.8	37
10	Insights into the molecular flexibility of $\theta$ -defensins by NMR relaxation analysis. <i>Journal of Physical Chemistry B</i> , <b>2014</b> , 118, 14257-66	3.4	18
9	Chemie und Biologie von Theta-Defensinen. <i>Angewandte Chemie</i> , <b>2014</b> , 126, 10786-10798	3.6	12
8	The cyclic cystine ladder in $\theta$ -defensins is important for structure and stability, but not antibacterial activity. <i>Journal of Biological Chemistry</i> , <b>2013</b> , 288, 10830-40	5.4	58
7	Exploring DOXP-reductoisomerase binding limits using phosphonated N-aryl and N-heteroarylcarboxamides as DXR inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 4332-41	3.4	13
6	Quantification of small cyclic disulfide-rich peptides. <i>Biopolymers</i> , <b>2012</b> , 98, 518-24	2.2	19
5	Structural characterization of the cyclic cystine ladder motif of $\theta$ -defensins. <i>Biochemistry</i> , <b>2012</b> , 51, 9718-26	3.6	50
4	The chemistry of cyclotides. <i>Journal of Organic Chemistry</i> , <b>2011</b> , 76, 4805-17	4.2	55
3	Chemical Synthesis of Naturally-Occurring Cyclic Mini-Proteins from Plants and Animals. <i>Israel Journal of Chemistry</i> , <b>2011</b> , 51, 908-916	3.4	6
2	Synthesis and evaluation of phosphonated N-heteroarylcarboxamides as DOXP-reductoisomerase (DXR) inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , <b>2011</b> , 19, 1321-7	3.4	22
1	<sup>31</sup> P NMR kinetic study of the tandem cleavage of phosphonate esters by bromotrimethylsilane. <i>Tetrahedron</i> , <b>2010</b> , 66, 8446-8449	2.4	7