

# Lessons in PROTAC Design from Selective Degradation

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Purification and characterization of two forms of a high-molecular-weight cysteine proteinase (porphypain) from <i>Porphyromonas gingivalis</i> . <i>Journal of Bacteriology</i> , 1994, 176, 4549-4557.	2.2	69
2	PROTACs: An Emerging Targeting Technique for Protein Degradation in Drug Discovery. <i>BioEssays</i> , 2018, 40, e1700247.	2.5	151
3	Inducing protein-protein interactions with molecular glues. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 2585-2592.	2.2	53
4	Hypoxia-selective allosteric destabilization of activin receptor-like kinases: A potential therapeutic avenue for prophylaxis of heterotopic ossification. <i>Bone</i> , 2018, 112, 71-89.	2.9	10
5	Try Me: Promiscuous Inhibitors Still Allow for Selective Targeted Protein Degradation. <i>Cell Chemical Biology</i> , 2018, 25, 4-6.	5.2	16
6	Translation Termination Factor GSPT1 Is a Phenotypically Relevant Off-Target of Heterobifunctional Phthalimide Degraders. <i>ACS Chemical Biology</i> , 2018, 13, 553-560.	3.4	128
7	Kinase inhibitors: the road ahead. <i>Nature Reviews Drug Discovery</i> , 2018, 17, 353-377.	46.4	679
8	Proteolysis Targeting Chimeras (PROTACs) of Anaplastic Lymphoma Kinase (ALK). <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 304-314.	5.5	165
9	The Advantages of Targeted Protein Degradation Over Inhibition: An RTK Case Study. <i>Cell Chemical Biology</i> , 2018, 25, 67-77.e3.	5.2	422
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15	Androgen receptor degradation by the proteolysis-targeting chimera ARCC-4 outperforms enzalutamide in cellular models of prostate cancer drug resistance. <i>Communications Biology</i> , 2018, 1, 100.	4.4	249
16	Chemical Protein Degradation Approach and its Application to Epigenetic Targets. <i>Chemical Record</i> , 2018, 18, 1681-1700.	5.8	33
17	Delineating the role of cooperativity in the design of potent PROTACs for BTK. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E7285-E7292.	7.1	265
18	When Kinases Meet PROTACs. <i>Chinese Journal of Chemistry</i> , 2018, 36, 971-977.	4.9	27
19	Homo-PROTACs for the Chemical Knockdown of Cereblon. <i>ACS Chemical Biology</i> , 2018, 13, 2771-2782.	3.4	114

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