

E3 ubiquitin ligase components in GtoPdb v.2022.3

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Abstract

Ubiquitination (a.k.a. ubiquitylation) is a protein post-translational modification that typically requires the sequential action of three enzymes: E1 (ubiquitin-activating enzymes), E2 (ubiquitin-conjugating enzymes), and E3 (ubiquitin ligases) [28]. Ubiquitination of proteins can target them for proteasomal degradation, or modulate cellular processes including cell cycle progression, transcriptional regulation, DNA repair and signal transduction.

E3 ubiquitin ligases, of which there are >600 in humans, are a family of highly heterogeneous proteins and protein complexes that recruit ubiquitin-loaded E2 enzymes to mediate transfer of the ubiquitin molecule from the E2 to protein substrates. Target substrate specificity is determined by a substrate recognition subunit within the E3 complex.

E3 ligases are being exploited as pharmacological targets to facilitate targeted protein degradation (TPD), as an alternative to small molecule inhibitors [3], through the development of proteolysis targeting chimeras (PROTACs) and molecular glues.

Contents

This is a citation summary for E3 ubiquitin ligase components in the [Guide to Pharmacology](#) database (GtoPdb). It exists purely as an adjunct to the database to facilitate the recognition of citations to and from the database by citation analyzers. Readers will almost certainly want to visit the relevant sections of the database which are given here under database links.

GtoPdb is an expert-driven guide to pharmacological targets and the substances that act on them. GtoPdb is a reference work which is most usefully represented as an on-line database. As in any publication this work should be appropriately cited, and the papers it cites should also be recognized. This document provides a citation for the relevant parts of the database, and also provides a reference list for the research cited by those parts. For further details see [4].

Please note that the database version for the citations given in GtoPdb are to the most recent preceding version in which the family or its subfamilies and targets were substantially changed. The links below are to the current version. If you need to consult the cited version, rather than the most recent version, please contact the GtoPdb curators.

Database links

[E3 ubiquitin ligase components](#)

<https://www.guidetopharmacology.org/GRAC/FamilyDisplayForward?familyId=1023>

Enzymes

cereblon

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=3086>

MDM2 proto-oncogene

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=3136>

STIP1 homology and U-box containing protein 1

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=3202>

von Hippel-Lindau tumor suppressor

<https://www.guidetopharmacology.org/GRAC/ObjectDisplayForward?objectId=3204>

References

1. Apriamashvili G, Vredevoogd DW, Krijgsman O, Bleijerveld OB, Ligtenberg MA, de Brujin B, Boshuizen J, Traets JJH, D'Empaire Altimari D and van Vliet A *et al.*. (2022) Ubiquitin ligase STUB1 destabilizes IFN- γ

- receptor complex to suppress tumor IFNy signaling. *Nat Commun* **13**: 1923 [PMID:35395848]
- 2. Asatsuma-Okumura T, Ito T and Handa H. (2019) Molecular mechanisms of cereblon-based drugs. *Pharmacol Ther* **202**: 132-139 [PMID:31202702]
 - 3. Belcher BP, Ward CC and Nomura DK. (2021) Ligandability of E3 Ligases for Targeted Protein Degradation Applications. *Biochemistry* [PMID:34473924]
 - 4. Buneman P, Christie G, Davies JA, Dimitrellou R, Harding SD, Pawson AJ, Sharman JL and Wu Y. (2020) Why data citation isn't working, and what to do about it. *Database* **2020** [PMID:32367113]
 - 5. Cao Z, Li G, Shao Q, Yang G, Zheng L, Zhang T and Zhao Y. (2016) CHIP: A new modulator of human malignant disorders. *Oncotarget* **7**: 29864-74 [PMID:27007160]
 - 6. Carvajal LA, Neriah DB, Senecal A, Benard L, Thiruthuvanathan V, Yatsenko T, Narayananari SR, Wheat JC, Todorova TI and Mitchell K et al.. (2018) Dual inhibition of MDMX and MDM2 as a therapeutic strategy in leukemia. *Sci Transl Med* **10** [PMID:29643228]
 - 7. Chamberlain PP, Lopez-Girona A, Miller K, Carmel G, Pagarigan B, Chie-Leon B, Rychak E, Corral LG, Ren YJ and Wang M et al.. (2014) Structure of the human Cereblon-DDB1-lenalidomide complex reveals basis for responsiveness to thalidomide analogs. *Nat Struct Mol Biol* **21**: 803-9 [PMID:25108355]
 - 8. Dale B, Anderson C, Park KS, Kaniskan HÜ, Ma A, Shen Y, Zhang C, Xie L, Chen X and Yu X et al.. (2022) Targeting Triple-Negative Breast Cancer by a Novel Proteolysis Targeting Chimera Degrader of Enhancer of Zeste Homolog 2. *ACS Pharmacol Transl Sci* **5**: 491-507 [PMID:35837138]
 - 9. Del Prete D, Rice RC, Rajadhyaksha AM and D'Adamio L. (2016) Amyloid Precursor Protein (APP) May Act as a Substrate and a Recognition Unit for CRL4CRBN and Stub1 E3 Ligases Facilitating Ubiquitination of Proteins Involved in Presynaptic Functions and Neurodegeneration. *J Biol Chem* **291**: 17209-27 [PMID:27325702]
 - 10. Ebner P, Versteeg GA and Ikeda F. (2017) Ubiquitin enzymes in the regulation of immune responses. *Crit Rev Biochem Mol Biol* **52**: 425-460 [PMID:28524749]
 - 11. Fischer ES, Böhm K, Lydeard JR, Yang H, Stadler MB, Cavardini S, Nagel J, Serluca F, Acker V and Lingaraju GM et al.. (2014) Structure of the DDB1-CRBN E3 ubiquitin ligase in complex with thalidomide. *Nature* **512**: 49-53 [PMID:25043012]
 - 12. Gandhi A, Dimartino J and Chopra R. (2014) Methods for the treatment of locally advanced breast cancer Patent number: WO2014039960A1.
 - 13. Gandhi AK, Kang J, Havens CG, Conklin T, Ning Y, Wu L, Ito T, Ando H, Waldman MF and Thakurta A et al.. (2014) Immunomodulatory agents lenalidomide and pomalidomide co-stimulate T cells by inducing degradation of T cell repressors Ikaros and Aiolos via modulation of the E3 ubiquitin ligase complex CRL4(CRBN.). *Br J Haematol* **164**: 811-21 [PMID:24328678]
 - 14. Hagner PR, Man HW, Fontanillo C, Wang M, Couto S, Breider M, Bjorklund C, Havens CG, Lu G and Rychak E et al.. (2015) CC-122, a pleiotropic pathway modifier, mimics an interferon response and has antitumor activity in DLBCL. *Blood* **126**: 779-89 [PMID:26002965]
 - 15. Hartmann MD, Boichenko I, Coles M, Zanini F, Lupas AN and Hernandez Alvarez B. (2014) Thalidomide mimics uridine binding to an aromatic cage in cereblon. *J Struct Biol* **188**: 225-32 [PMID:25448889]
 - 16. Heim C, Platsika D, Mousavizadeh F, Bär K, Hernandez Alvarez B, Giannis A and Hartmann MD. (2019) De-Novo Design of Cereblon (CRBN) Effectors Guided by Natural Hydrolysis Products of Thalidomide Derivatives. *J Med Chem* **62**: 6615-6629 [PMID:31251063]
 - 17. Higgins JJ, Hao J, Kosofsky BE and Rajadhyaksha AM. (2008) Dysregulation of large-conductance Ca²⁺-activated K⁺ channel expression in nonsyndromal mental retardation due to a cereblon p.R419X mutation. *Neurogenetics* **9**: 219-23 [PMID:18414909]
 - 18. Higgins JJ, Pucilowska J, Lombardi RQ and Rooney JP. (2004) A mutation in a novel ATP-dependent Lon protease gene in a kindred with mild mental retardation. *Neurology* **63**: 1927-31 [PMID:15557513]
 - 19. Huang HT, Dobrovolsky D, Paulk J, Yang G, Weisberg EL, Doctor ZM, Buckley DL, Cho JH, Ko E and Jang J et al.. (2018) A Chemoproteomic Approach to Query the Degradable Kinome Using a Multi-kinase Degrader. *Cell Chem Biol* **25**: 88-99.e6 [PMID:29129717]
 - 20. Ito T, Ando H, Suzuki T, Ogura T, Hotta K, Imamura Y, Yamaguchi Y and Handa H. (2010) Identification of a primary target of thalidomide teratogenicity. *Science* **327**: 1345-50 [PMID:20223979]
 - 21. Ito T and Handa H. (2016) Cereblon and its downstream substrates as molecular targets of immunomodulatory drugs. *Int J Hematol* **104**: 293-9 [PMID:27460676]
 - 22. Joshi V, Amanullah A, Upadhyay A, Mishra R, Kumar A and Mishra A. (2016) A Decade of Boon or Burden: What Has the CHIP Ever Done for Cellular Protein Quality Control Mechanism Implicated in Neurodegeneration and Aging? *Front Mol Neurosci* **9**: 93 [PMID:27757073]
 - 23. Kumar S, Basu M and Ghosh MK. (2021) Chaperone-assisted E3 ligase CHIP: A double agent in cancer *Genes & Diseases*
 - 24. Li D, Yu X, Kottur J, Gong W, Zhang Z, Storey AJ, Tsai YH, Uryu H, Shen Y and Byrum SD et al.. (2022)

- Discovery of a dual WDR5 and Ikaros PROTAC degrader as an anti-cancer therapeutic. *Oncogene* [PMID:35525905]
- 25. Lopez-Girona A, Mendi D, Ito T, Miller K, Gandhi AK, Kang J, Karasawa S, Carmel G, Jackson P and Abbasian M *et al.*. (2012) Cereblon is a direct protein target for immunomodulatory and antiproliferative activities of lenalidomide and pomalidomide. *Leukemia* **26**: 2326-35 [PMID:22552008]
 - 26. Matyskiela ME, Zhang W, Man HW, Muller G, Khambatta G, Baculi F, Hickman M, LeBrun L, Pagarigan B and Carmel G *et al.*. (2018) A Cereblon Modulator (CC-220) with Improved Degradation of Ikaros and Aiolos. *J Med Chem* **61**: 535-542 [PMID:28425720]
 - 27. Momand J, Zambetti GP, Olson DC, George D and Levine AJ. (1992) The mdm-2 oncogene product forms a complex with the p53 protein and inhibits p53-mediated transactivation. *Cell* **69**: 1237-45 [PMID:1535557]
 - 28. Morreale FE and Walden H. (2016) Types of Ubiquitin Ligases. *Cell* **165**: 248-248.e1 [PMID:27015313]
 - 29. Nabet B, Roberts JM, Buckley DL, Pault J, Dastjerdi S, Yang A, Leggett AL, Erb MA, Lawlor MA and Souza A *et al.*. (2018) The dTAG system for immediate and target-specific protein degradation. *Nat Chem Biol* **14**: 431-441 [PMID:29581585]
 - 30. Ng S, Brueckner AC, Bahmanjah S, Deng Q, Johnston JM, Ge L, Duggal R, Habulihaz B, Barlock B and Ha S *et al.*. (2022) Discovery and Structure-Based Design of Macroyclic Peptides Targeting STUB1. *J Med Chem* [PMID:35853179]
 - 31. Ng SY, Yoshida N, Christie AL, Ghandi M, Dharia NV, Dempster J, Murakami M, Shigemori K, Morrow SN and Van Scoyck A *et al.*. (2018) Targetable vulnerabilities in T- and NK-cell lymphomas identified through preclinical models. *Nat Commun* **9**: 2024 [PMID:29789628]
 - 32. Nguyen TV, Lee JE, Sweredoski MJ, Yang SJ, Jeon SJ, Harrison JS, Yim JH, Lee SG, Handa H and Kuhlman B *et al.*. (2016) Glutamine Triggers Acetylation-Dependent Degradation of Glutamine Synthetase via the Thalidomide Receptor Cereblon. *Mol Cell* **61**: 809-20 [PMID:26990986]
 - 33. Pairawan S, Zhao M, Yuca E, Annis A, Evans K, Sutton D, Carvajal L, Ren JG, Santiago S and Guerlavais V *et al.*. (2021) First in class dual MDM2/MDMX inhibitor ALRN-6924 enhances antitumor efficacy of chemotherapy in TP53 wild-type hormone receptor-positive breast cancer models. *Breast Cancer Res* **23**: 29 [PMID:33663585]
 - 34. Rankovic Z, Min J, Mayasundari A, Keramatnia F, Jonchere B, Yang SW, Jarusiewicz JA, Actis M, Das S and Young BM *et al.*. (2021) Phenyl-Glutaramides: Alternative Cereblon Binders for the Design of PROTACs. *Angew Chem Int Ed Engl* [PMID:34614283]
 - 35. Remillard D, Buckley DL, Pault J, Brien GL, Sonnett M, Seo HS, Dastjerdi S, Wühr M, Dhe-Paganon S and Armstrong SA *et al.*. (2017) Degradation of the BAF Complex Factor BRD9 by Heterobifunctional Ligands. *Angew Chem Int Ed Engl* **56**: 5738-5743 [PMID:28418626]
 - 36. Ries S, Biederer C, Woods D, Shifman O, Shirasawa S, Sasazuki T, McMahon M, Oren M and McCormick F. (2000) Opposing effects of Ras on p53: transcriptional activation of mdm2 and induction of p19ARF. *Cell* **103**: 321-30 [PMID:11057904]
 - 37. Sasaki M, Kawahara K, Nishio M, Mimori K, Kogo R, Hamada K, Itoh B, Wang J, Komatsu Y and Yang YR *et al.*. (2011) Regulation of the MDM2-P53 pathway and tumor growth by PICT1 via nucleolar RPL11. *Nat Med* **17**: 944-51 [PMID:21804542]
 - 38. Sheereen A, Alaamery M, Bawazeer S, Al Yafee Y, Massadeh S and Eyaid W. (2017) A missense mutation in the CRBN gene that segregates with intellectual disability and self-mutilating behaviour in a consanguineous Saudi family. *J Med Genet* **54**: 236-240 [PMID:28143899]
 - 39. Sun D, Li Z, Rew Y, Gribble M, Bartberger MD, Beck HP, Canon J, Chen A, Chen X and Chow D *et al.*. (2014) Discovery of AMG 232, a potent, selective, and orally bioavailable MDM2-p53 inhibitor in clinical development. *J Med Chem* **57**: 1454-72 [PMID:24456472]
 - 40. Vu B, Wovkulich P, Pizzolato G, Lovey A, Ding Q, Jiang N, Liu JJ, Zhao C, Glenn K and Wen Y *et al.*. (2013) Discovery of RG7112: A Small-Molecule MDM2 Inhibitor in Clinical Development. *ACS Med Chem Lett* **4**: 466-9 [PMID:24900694]
 - 41. Yang J, Li Y, Aguilar A, Liu Z, Yang CY and Wang S. (2019) Simple Structural Modifications Converting a Bona fide MDM2 PROTAC Degrader into a Molecular Glue Molecule: A Cautionary Tale in the Design of PROTAC Degraders. *J Med Chem* **62**: 9471-9487 [PMID:31560543]