

# EDD (B-11): sc-515494

## BACKGROUND

EDD (for E3 identified by differential display) is a progesterin-regulated gene that was isolated from T-47D human breast cancer cells. Based on sequence homology, EDD appears to be a human homolog of the *Drosophila* hyperplastic discs (hyd) gene, a tumor suppressor gene that is required for control of imaginal disc growth. EDD contains a HECT domain in the carboxy terminus. HECT domain-containing proteins function as ubiquitin-protein ligases, or E3 enzymes. EDD has been shown to bind to ubiquitin, and like other HECT family proteins, may function as an E3 ubiquitin-protein ligase.

## CHROMOSOMAL LOCATION

Genetic locus: UBR5 (human) mapping to 8q22.3; Ubr5 (mouse) mapping to 15 B3.1.

## SOURCE

EDD (B-11) is a mouse monoclonal antibody raised against amino acids 1-300 mapping at the N-terminus of EDD of human origin.

## PRODUCT

Each vial contains 200 µg IgG<sub>1</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

EDD (B-11) is available conjugated to agarose (sc-515494 AC), 500 µg/0.25 ml agarose in 1 ml, for IP; to HRP (sc-515494 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-515494 PE), fluorescein (sc-515494 FITC), Alexa Fluor<sup>®</sup> 488 (sc-515494 AF488), Alexa Fluor<sup>®</sup> 546 (sc-515494 AF546), Alexa Fluor<sup>®</sup> 594 (sc-515494 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-515494 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-515494 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-515494 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

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## APPLICATIONS

EDD (B-11) is recommended for detection of EDD of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for EDD siRNA (h): sc-43744, EDD siRNA (m): sc-143292, EDD shRNA Plasmid (h): sc-43744-SH, EDD shRNA Plasmid (m): sc-143292-SH, EDD shRNA (h) Lentiviral Particles: sc-43744-V and EDD shRNA (m) Lentiviral Particles: sc-143292-V.

Molecular Weight of EDD: 309 kDa.

Positive Controls: HEL 92.1.7 cell lysate: sc-2270, Neuro-2A whole cell lysate: sc-364185 or F9 cell lysate: sc-2245.

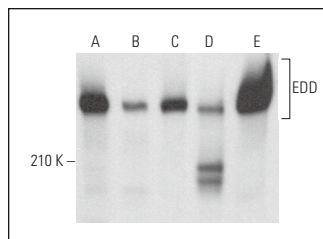
## STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

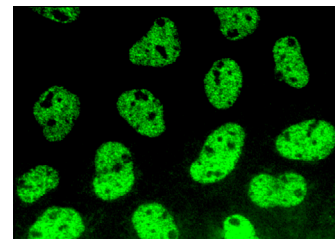
## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## DATA



EDD (B-11): sc-515494. Western blot analysis of EDD expression in HEL 92.1.7 (A), 3T3-L1 (B), Neuro-2A (C), EOC 20 (D) and F9 (E) whole cell lysates.



EDD (B-11): sc-515494. Immunofluorescence staining of formalin-fixed A-431 cells showing nuclear localization.

## SELECT PRODUCT CITATIONS

- Qiao, X., et al. 2020. UBR5 is co-amplified with Myc in breast tumors and encodes an ubiquitin ligase that limits Myc-dependent apoptosis. *Cancer Res.* 80: 1414-1427.
- MacDonald, T.M., et al. 2020. Prolactin and androgen R1881 induce pro-survival carboxypeptidase-D and EDD E3 ligase in triple-negative and HER2+ breast cancer. *Am. J. Cancer Res.* 10: 1321-1343.
- Leboeuf, D., et al. 2020. The Arg/N-degron pathway—a potential running back in fine-tuning the inflammatory response? *Biomolecules* 10: 903.
- Puhach, O., et al. 2020. Murine cytomegaloviruses m139 targets DDX3 to curtail interferon production and promote viral replication. *PLoS Pathog.* 16: e1008546.
- Moberg, M., et al. 2022. Acute normobaric hypoxia blunts contraction-mediated mTORC1- and JNK-signaling in human skeletal muscle. *Acta Physiol.* 234: e13771.
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- Xiang, G., et al. 2022. UBR5 targets tumor suppressor CDC73 proteolytically to promote aggressive breast cancer. *Cell Death Dis.* 13: 451.
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- Zhao, Y., et al. 2022. Deubiquitinase OTUD6A promotes breast cancer progression by increasing TopBP1 stability and rendering tumor cells resistant to DNA-damaging therapy. *Cell Death Differ.* 29: 2531-2544.

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.