Datasheet

Version: 3.0.0 Revision date: 16 Apr 2025



Human E3 ubiquitin-protein ligase MARCH11 (MARCHF11) ELISA Kit

Catalogue No.:abx532287

Human E3 ubiquitin-protein ligase MARCH11 (MARCHF11) ELISA Kit is an ELISA Kit for the in vitro quantitative measurement of Human E3 ubiquitin-protein ligase MARCH11 (MARCHF11) concentrations in tissue homogenates, cell lysates and other biological fluids.

Target: E3 ubiquitin-protein ligase MARCH11 (MARCHF11)

Reactivity: Human

Tested Applications: ELISA

Recommended dilutions: Optimal dilutions/concentrations should be determined by the end user.

Storage: Shipped at 4 °C. Upon receipt, store the kit according to the storage instruction in the kit's manual.

Validity: The validity for this kit is 6 months.

Stability: The stability of the kit is determined by the rate of activity loss. The loss rate is less than 5% within

the expiration date under appropriate storage conditions. To minimize performance fluctuations, operation procedures and lab conditions should be strictly controlled. It is also strongly suggested

that the whole assay is performed by the same user throughout.

UniProt Primary AC: A6NNE9 (UniProt, ExPASy)

Gene Symbol: Mar-11

GenelD: <u>441061</u>

OMIM: <u>613338</u>

HGNC: 33609

KEGG: hsa:441061

Test Range: 0.156 ng/ml - 10 ng/ml

Standard Form: Lyophilized

Detection Method: Colorimetric

Assay Data: Quantitative

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Sample Type: Tissue homogenates, cell lysates and other biological fluids.

Note: THIS PRODUCT IS FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR

THERAPEUTIC PROCEDURES.

The range and sensitivity is subject to change. Please contact us for the latest product information. For accurate results, sample concentrations must be diluted to mid-range of the kit. If you require a specific range, please contact us in advance or write your request in your order comments. Please note that our kits are optimised for detection of native samples, rather than recombinant proteins. We are unable to guarantee detection of recombinant proteins, as they may have different sequences or tertiary structures to the native protein.