Datasheet

Version: 3.0.0 Revision date: 21 Jun 2025



Human Cytochrome P450 2C8 (CYP2C8) ELISA Kit

Catalogue No.:abx386766

Human Cytochrome P450 2C8 (CYP2C8) ELISA Kit is an ELISA Kit for the in vitro quantitative measurement of Human CYP2C8 concentrations in tissue homogenates, cell lysates and other biological fluids.

Target: Cytochrome P450 2C8 (CYP2C8)

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: P10632 (UniProt, ExPASy)

Gene Symbol: CYP2C8

GeneID: <u>1558</u>

OMIM: 601129

HGNC: 2622

KEGG: hsa:1558

Ensembl: ENSG00000138115

String: <u>9606.ENSP00000360317</u>

Test Range: 31.2 pg/ml - 2000 pg/ml

Sensitivity: < 15 pg/ml

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Standard Form: Lyophilized

Detection Method: Colorimetric

Assay Type: Sandwich

Assay Data: Quantitative

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.

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