Datasheet

Version: 2.0.0 Revision date: 30 Mar 2025



Human Endothelial Cell Adhesion Molecule (ESAM) CLIA Kit

Catalogue No.:abx494644

Human Endothelial Cell Adhesion Molecule (ESAM) Chemiluminescent Immunoassay (CLIA) Kit is a Sandwich Chemiluminescent Immunoassay (CLIA) Kit for use with Tissue homogenates, cell lysates and other biological fluids.

Target: Endothelial Cell Adhesion Molecule (ESAM)

Reactivity: Human

Tested Applications: CLIA

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 at least 6 months. Up to 12 months validity can be provided on request.

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: Q96AP7 (UniProt, ExPASy)

Gene Symbol: ESAM

GeneID: 90952

OMIM: 614281

HGNC: 17474

KEGG: hsa:90952

Ensembl: ENSG00000149564

String: <u>9606.ENSP00000278927</u>

Test Range: 0.312 ng/ml - 20 ng/ml

Sensitivity: < 0.114 ng/ml

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Standard Form: Lyophilized

Detection Method: Chemiluminescent

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.

