Human Anterior Gradient Protein 3 (AGR3) ELISA Kit

Catalogue No.:abx385664

Human Anterior Gradient Protein 3 (AGR3) ELISA Kit is an ELISA Kit for the in vitro quantitative measurement of Human Anterior Gradient Protein 3 (AGR3) concentrations in serum, plasma, tissue homogenates, cell lysates and other biological fluids.

Target:	Anterior Gradient Protein 3 (AGR3)
Research Area:	Tumor Immunity
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:	Q8TD06 (**UniProt**, **ExPASy**)
Gene Symbol:	AGR3
GeneID:	**155465**
OMIM:	**609482**
HGNC:	24167
KEGG:	hsa:155465
Ensembl:	ENSG00000173467
String:	**9606.ENSP00000308606**
Test Range:	0.156 ng/ml - 10 ng/ml

v1.0.0   Abbexa LTD, Cambridge, UK · Phone: +44 (0) 1223 755950 · Fax: +44 (0) 1223 755951   1 of 2
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Sensitivity:	< 0.08 ng/ml
Standard Form:	Lyophilized
Detection Method:	Colorimetric
Assay Type:	Sandwich
Assay Data:	Quantitative
Sample Type:	Serum, plasma, tissue homogenates, cell lysates and other biological fluids.
Note:	This product is for research use only.

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 ELISA and CLIA 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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