

MONOCLONAL ANTIBODY

For research use only. Not for clinical diagnosis

Catalog No. AGE-M04

Anti Nº- (carboxymethyl) arginine (CMA)

BACKGROUND

N $^{\omega}$ -(carboxymethyl) arginine (CMA), a CML analogue, is an acid-labile AGE structure which was discovered in enzymatic hydrolysate of glycated collagen. CMA is preferentially generated in glycated collagen.

Product type Primary antibody

Immunogen CMA-HSA **Host Species** Mouse P3U1 **Fusion Partner** 3F5 **Clone Designation** IgG1 Isotype Mouse Host Ascites Source **Purification** Protein G Liquid **Form**

Formulation Buffer PBS containing 0.1% proclin and 2% sorbitol as a preservative

Concentration 0.1 mg / ml
Volume 100 ul
Label Unlabeled
Specificity CMA
Cross species reactivity

Storage Store below -20°C (below -70°C for prolonged storage)

Aliquot to avoid cycles of freeze/thaw.

Application notes

Western blotting: 1/100 - 1/1000

Recommended dilutions

Immunohistochemistry: 1/50 - 1/100 (frozen section)

• ELISA: 1/100 - 1/200

Other applications have not been tested.

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

References

- lijima K, Murata M, Takahara H, Irie S, Fujimoto D. Identification of N(omega)-carboxymethylarginine as a novel acid-labileadvanced glycation end product in collagen. Biochem J. 347 Pt 1:23-27 (2000) PMID: 10727397
- 2) Mera K., Fujiwara Y., Otagiri M., Sakata N., Nagai R. Immunological Detection of N -carboxymethylarginine by Specific Antibody. Ann N Y Acad Sci. 1126, 155-157 (2008) **PMID**: 18079475

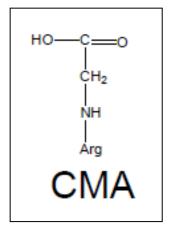


Fig.1 Nº- (carboxymethyl) arginine (CMA) structure

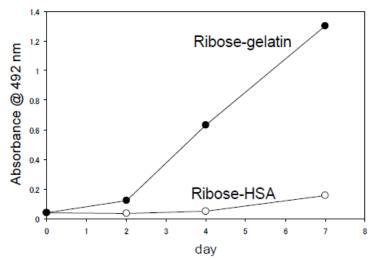


Fig.2 Immunoreactivity of the CMA(3F5) monoclonal antibody to Ribose-gelatin and Ribose-HSA

ELISA protocol

Coating

- 1) Distribute 100 ul / well of the sample solution (1 ug/mL in PBS) to 96 well microtiter plates (Thermo, MaxiSorp).
- 2) Incubate the plates 2h at RT or overnight at 4 degrees.
- 3) Discard the supernatant of sample solution.
- 4) Wash the plates three times with washing.buf.(PBS/0.05%Tween 20)

Blocking

- 1) Distribute 200 ul / well of 0.5% gelatin-PBS to 96 well microtiter plates
- 2) Incubate the plates 1h at RT.
- 3) Discard the the supernatant of 0.5% gelatin-PBS
- 4) Wash the plates three times with washing.buf.(PBS/0.05%Tween 20)

Primary antibody

- 1) Distribute 100 ul / well of Primary antibodies diluted with washing buf. as suggested in the APPLICATIONS to each well.
- 2) Incubate the plates 1h at RT.
- 3) Discard the supernatant of Primary antibody solution.
- 4) Wash the plates three times with washing.buf.(PBS/0.05%Tween 20)

Secondary antibody

- 1) Distribute 100 ul / well of secondary antibodies (HRP-anti mouse IgG) diluted with washing buf. as suggested in the APPLICATIONS to each well.
- 2) Incubate the plates 1h at RT.
- 3) Discard the supernatant of secondary antibody.
- 4) Wash the plates three times with washing.buf.(PBS/0.05%Tween 20)

OPD color reaction

- 1) Reaction for 2-10 minutes at RT..
- 2) Distribute 100 uL / well of 2M H₂SO₄ to each well and stop enzyme reaction.
- 3) After gentle mixing, determine the absorbance at 492 nm of each well by a spectrophotometer.

RELATED PRODUCTS:

Product Name	Quantity	Maker	Cat#
Anti N⁵-(carboxymethyl) lysine [CML] (2G11) Monoclonal Antibody	100 ul	CAC	AGE-M01
Anti N ^ε -(carboxyethyl) lysine [CEL] (CEL-SP) Monoclonal Antibody	100 ul	CAC	AGE-M02
Anti GA-pyridine (2A2) Monoclonal Antibody	100 ul	CAC	AGE-M03
Anti N ^ω -(carboxymethyl) arginine [CMA] (3F5) Monoclonal Antibody	100 ul	CAC	AGE-M04
CML-BSA/Nε-(carboxymethyl) lysine-BSA	200 ul	CSR	AGE-GP01
CEL-BSA/Nε-(carboxyethyl) lysine-BSA	200 ul	CSR	AGE-GP02
GA-BSA/Glycolaldehyde-BSA	200 ul	CSR	AGE-GP03
Ribose-gelatin	500 ul	CSR	AGE-GP04
Mild-AGE-BSA	200 ul	CSR	AGE-GP05

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