

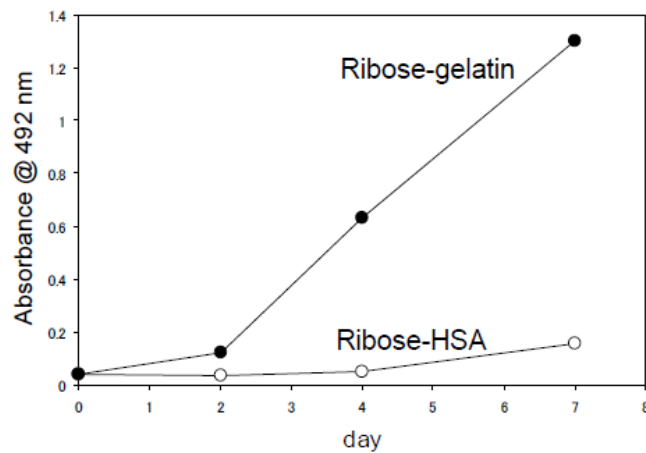


## Ribose-gelatin

<b>Product Description</b>	Gelatin (2 mg/ml) was incubated with 30 mM of ribose in a 0.2 M phosphate buffer (pH 7.4) at 37°C for 1 week, followed by dialysis against PBS.
<b>Volume</b>	500 ul
<b>Concentration</b>	1 mg/ml
<b>Storage</b>	Store below -20°C (below -70°C for prolonged storage). Aliquot to avoid cycles of freeze/thaw.

- References**
1. Shimasaki S, Kubota M, Yoshitomi M, Takagi K, Suda K, Mera K, Fujiwara Y, Nagai R. N<sup>ω</sup>-(carboxymethyl)arginine Accumulates in Glycated Collagen and Klotho-deficient Mouse Skin. *Anti-Aging Medicine* 8 (6) : 82-87, 2011 [ANTI-AGING MEDICINE 8\(6\), 82-87, 2011](#)

### Characterization



**Fig.1 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 H<sub>2</sub>SO<sub>4</sub> 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 N <sup>F</sup> -(carboxymethyl) lysine [CML] (2G11) Monoclonal Antibody	100 ul	CAC	AGE-M01
Anti N <sup>F</sup> -(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 <sup>W</sup> -(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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