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APA899Ra61 100µg Active Osteopontin (OPN) Organism Species: Rattus norvegicus (Rat) *Instruction manual*

FOR IN VITRO USE AND RESEARCH USE ONLY NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1st Edition (Apr, 2016)

[PROPERTIES]

Source: Eukaryotic expression. Host: 293F cell Residues: Leu17~Asn317

Tags: N-terminal His-tag

Purity: >95%

Endotoxin Level: <1.0EU per 1µg (determined by the LAL method).

Buffer Formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 1mM EDTA, 1mM DTT, 0.01% sarcosyl, 5% trehalose, and Proclin300.

Applications: Cell culture; Activity Assays; In vivo assays.

(May be suitable for use in other assays to be determined by the end user.)

Predicted isoelectric point: 4.3

Predicted Molecular Mass: 34.9kDa

Accurate Molecular Mass: 60kDa as determined by SDS-PAGE reducing conditions.

Phenomenon explanation:

The possible reasons that the actual band size differs from the predicted are as follows:

- 1. Splice variants: Alternative splicing may create different sized proteins from the same gene.
- 2. Relative charge: The composition of amino acids may affects the charge of the protein.
- 3. Post-translational modification: Phosphorylation, glycosylation, methylation etc.
- 4. Post-translation cleavage: Many proteins are synthesized as pro-proteins, and then cleaved to give the active form.
- 5. Polymerization of the target protein: Dimerization, multimerization etc.

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[<u>USAGE</u>]

Reconstitute in 20mM Tris, 150mM NaCl (pH8.0) to a concentration of 0.1-1.0 mg/mL. Do not vortex.

[STORAGE AND STABILITY]

Storage: Avoid repeated freeze/thaw cycles.

Store at 2-8°C for one month.

Aliquot and store at -80°C for 12 months.

Stability Test: The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration date under appropriate storage condition.

[<u>SEQUENCE</u>]

LPVK VAEFGSSEEK AHYSKHSDAV ATWLKPDPSQ KQNLLAPQNS VSSEETDDFK QETLPSNSNE SHDHMDDDDD DDDDGDHAES EDSVNSDESD ESHHSDESDE SFTASTQADV LTPIAPTVDV PDGRGDSLAY GLRSKSRSFP VSDEQYPDAT DEDLTSRMKS QESDEAIKVI PVAQRLSVPS DQDSNGKTSH ESSQLDEPSV ETHSLEQSKE YKQRASHEST EQSDAIDSAE KPDAIDSAER SDAIDSQASS KASLEHQSHE FHSHEDKLVL DPKSKEDDRY LKFRISHELE SSSEVN

[ACTIVITY]

Osteopontin (OPN), a multifunctional phosphorylated glycoprotein, plays an important role in neutrophil recruitment and was found to induce the expression of proinflammatory chemokines including MCP-1 and MIP-1 β which promote migration and recruitment of inflammatory cells. It has been reported that OPN induces MCP-1 expression through the NF-kappa B pathways in MCF-7 breast cancer cell line. Briefly, MCF-7 cells were seeded overnight at a density of 1x10⁵ cells/mL, and treated with or without various concentrations of OPN for 24h and

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MCP-1 levels in the cell supernatant were determined by ELISA.

Result: MCP-1 levels in the cell supernatant of MCF-7 cells increased significantly after stimulated with OPN, the data was shown in Table 1 and Figure 1.

Sample (cell supernatant of MCF-7 cells)	O.D. value	Corrected	Concentration of MCP-1 (ng/mL)
stimulated with OPN (100ng/mL)	1.358	1.305	2.53
stimulated with OPN (200ng/mL)	1.177	1.124	2.15
stimulated with OPN (400ng/mL)	1.101	1.048	1.99
unstimulated	0.944	0.891	1.66

Table 1. MCP-1 levels in the cell supernatant of MCF-7 cells regulated by OPN.

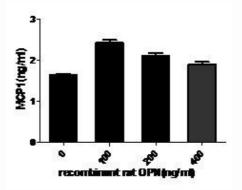
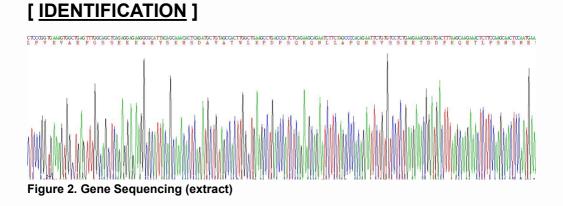


Figure 1. MCP-1 levels in the cell supernatant of MCF-7 cells regulated by OPN.



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-	kDa 70
	44
-	33
63	26
-	22
100	18
	14
	10

Figure 3. SDS-PAGE

Sample: Active recombinant OPN, Rat

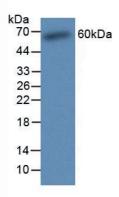


Figure 4. Western Blot

Sample: Recombinant OPN, Rat;

Antibody: Rabbit Anti-Rat OPN Ab (PAA899Ra06)