

RPA867Hu03 50µg

Recombinant Phospholipase A2 Group VII (LpPLA2)

Organism Species: Homo sapiens (Human)

Instruction manual

FOR RESEARCH USE ONLY NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

12th Edition (Revised in Aug, 2016)



## [PROPERTIES]

Source: Prokaryotic expression

Host: E.coli

Residues: Phe22~Asn441

Tags: Two N-terminal Tags, His-tag and SUMO-tag

**Subcellular Location:** Extracellular matrix

**Purity:** > 80%

Traits: Freeze-dried powder

**Buffer formulation:** PBS, pH7.4, containing 0.01% SKL, 5% Trehalose.

Original Concentration: 200µg/mL

Applications: Positive Control; Immunogen; SDS-PAGE; WB.

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

Predicted isoelectric point: 7.1

Predicted Molecular Mass: 61.5kDa

**Accurate Molecular Mass:** 65kDa 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.

#### [USAGE]

Reconstitute in 10mM PBS (pH7.4) 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.

### [ SEQUENCE ]

		FDWQYINPV	AHMKSSAWVN	KIQVLMAAAS
FGQTKIPRGN	GPYSVGCTDL	MFDHTNKGTF	LRLYYPSQDN	DRLDTLWIPN
KEYFWGLSKF	LGTHWLMGNI	LRLLFGSMTT	PANWNSPLRP	GEKYPLVVFS
HGLGAFRTLY	SAIGIDLASH	GFIVAAVEHR	DRSASATYYF	KDQSAAEIGD
KSWLYLRTLK	QEEETHIRNE	QVRQRAKECS	QALSLILDID	HGKPVKNALD
LKFDMEQLKD	SIDREKIAVI	GHSFGGATVI	QTLSEDQRFR	CGIALDAWMF
PLGDEVYSRI	PQPLFFINSE	YFQYPANIIK	MKKCYSPDKE	RKMITIRGSV
HQNFADFTFA	TGKIIGHMLK	LKGDIDSNVA	IDLSNKASLA	FLQKHLGLHK
DFDQWDCLIE	GDDENLIPGT	NINTTNQHIM	LQNSSGIEKY	N

## [ IDENTIFICATION ]

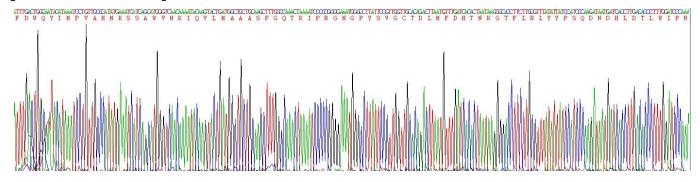


Figure . Gene Sequencing (extract)

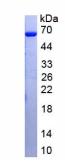




Figure. SDS-PAGE

# [ IMPORTANT NOTE ]

The kit is designed for research use only, we will not be responsible for any issue if the kit was used in clinical diagnostic or any other procedures.