

APD178Hu01 200ug
Active Paraoxonase 3 (PON3)
Organism Species: Homo sapiens (Human)
Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1st Edition (Apr, 2016)

[PROPERTIES]

Source: Prokaryotic expression.

Host: *E. coli*

Residues: Gly2~Leu354

Tags: N-terminal His-tag

Purity: >98%

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

Buffer Formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 0.01% skl, 5%Trehalose.

Original Concentration: 100µg/mL

Applications: Cell culture; Activity Assays.

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

Predicted isoelectric point: 5.2

Predicted Molecular Mass: 43.2kDa

Accurate Molecular Mass: 43kDa as determined by SDS-PAGE reducing conditions.

[USAGE]

Reconstitute in ddH₂O 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]

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GKLVALVLL  GVGLSLVGEM  FLAFRERVNA  SREVEPVEPE  NCHLIEELES  
GSEDIDILPS  GLAFISSGLK  YPGMPNFAPD  EPGKIFLMDL  NEQNRAQAL  
EISGGFDKEL  FNPHGISIFI  DKDNTVYLYV  VNHPHMKSTV  EIFKFEEQQR  
SLVYLKTIKH  ELLKSVNDIV  VLGPEQFYAT  RDHYFTNSLL  SFFEMILDLR  
WTYVLFYSPR  EVKVVAKGFC  SANGITVSAD  QKYVYVADVA  AKNIHIMEKH  
DNWDLTQLKV  IQLGTLVDNL  TVDPATGDIL  AGCHPNMKL  LNYNPEDPPG  
SEVLRIQNVL  SEKPRVSTVY  ANNGSVLQGT  SVASVYHGKI  LIGTVFHKTL  
YCEL
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[ACTIVITY]

Paraoxonase 3 (PON3) is secreted into the bloodstream and associates with high-density lipoprotein (HDL). PON3 also rapidly hydrolyzes lactones and can inhibit the oxidation of low-density lipoprotein, a function that is believed to slow the initiation and progression of atherosclerosis. Besides, Paraoxonase 1 (PON1) has been identified as an interactor of PON3, thus a binding ELISA assay was conducted to detect the interaction of recombinant human PON3 and recombinant human PON1. Briefly, PON3 were diluted serially in PBS, with 0.01% BSA (pH 7.4). Duplicate samples of 100uL were then transferred to PON1-coated microtiter wells and incubated for 2h at 37°C. Wells were washed with PBST and incubated for 1h with anti-PON3 pAb, then aspirated and washed 3 times. After incubation with HRP labelled secondary antibody, wells were aspirated and washed 3 times. With the addition of substrate solution, wells were incubated 15-25 minutes at 37°C. Finally, add 50µL stop solution to the wells and read at 450nm immediately. The binding activity of of PON3 and PON1 was shown in Figure 1, and this effect was in a dose dependent manner.

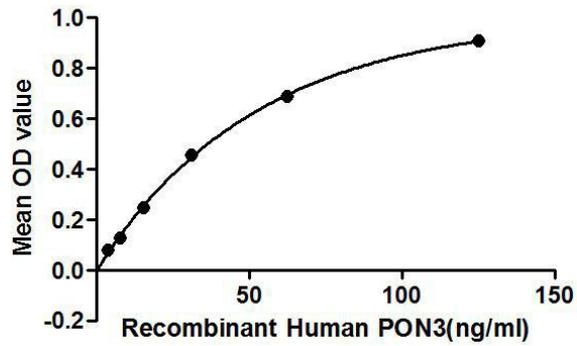


Figure 1. The binding activity of PON3 with PON1.

[IDENTIFICATION]

SMITGGGAGCTGTGTGGCTGWLCTTCTGGGACCTGCTGGCTTAAAGGGAGGCTGCTGWTTTTAGGAGCGSHTTAGACACTCGGAGMTAAAGCCGCGAGCCGAAACCTGGCCCTGATGGCCCTGGAGATGCTCGAGATATGGATFACTTCTAGCGGCCTGGCTTTTATCTGCAGTGGATTAATTCGGCCCTGCGAGSTTTCGCGCGCAWCT

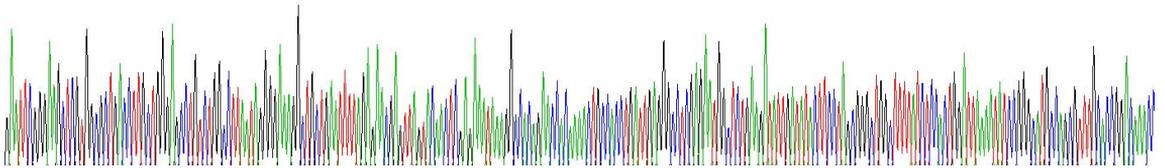


Figure 2. Gene Sequencing (extract)

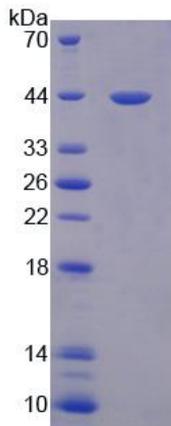


Figure 3. SDS-PAGE

Sample: Active recombinant PON3, Human

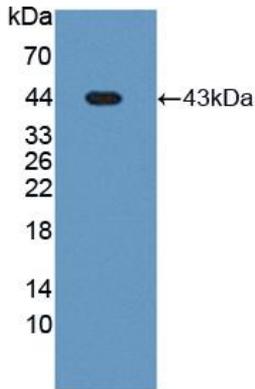


Figure 4. Western Blot

Sample: Recombinant PON3, Human;

Antibody: Rabbit Anti-Human PON3 Ab (PAD178Hu01)

[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.