

**APC374Hu01 50µg**  
**Active Carboxylesterase 1 (CES1)**  
**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:** Gly18~Leu299

**Tags:** N-terminal His-tag

**Purity:** >90%

**Buffer Formulation:** 20mM Tris, 150mM NaCl, pH8.0, containing 0.05% sarcosyl and 5% trehalose.

**Applications:** Cell culture; Activity Assays.

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

**Predicted isoelectric point:** 6.8

**Predicted Molecular Mass:** 34.0kDa

**Accurate Molecular Mass:** 35kDa as determined by SDS-PAGE reducing conditions.

## **[ USAGE ]**

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.

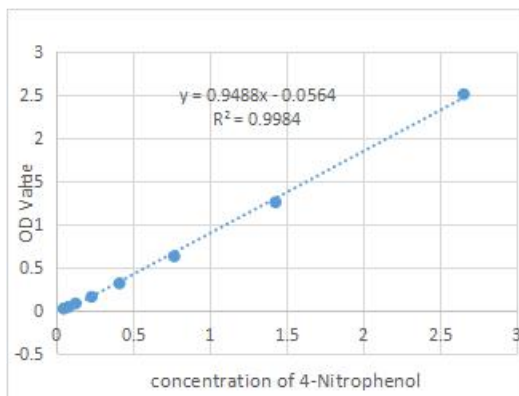
**Stability Test:** The thermal stability is described by the loss rate. The loss rate was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration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

## **[ SEQUENCE ]**

```
                GHP  SSPPVVDTVH  GKVLGKFVSL  EGFAQPVAIF
LGIPFAKPLL  GPLRFTPPQP  AEPWSFVKNA  TSYPPTCTQD  PKAGQLLSEL
FTNRKENIPL  KLSIEDCLYLN  IYTPADLTCK  NRLPVMVWIH  GGGLMVGAA
TYDGLALAAH  ENVVVVTIQY  RLGIWGFFST  GDEHSRGNWG  HLDQVAALRW
VQDNIAFFGG  NPGSVTIFGE  SAGGESVSVL  VLSPLAKNLF  HRAISESGVA
LTSVLVKKGD  VKPLAEQIAI  TAGCKTTTSA  VMVHCLRQKT  EEELLETTL
```

## **[ ACTIVITY ]**

Carboxylesterase 1 (CES1) also known as Liver carboxylesterase 1 is a serine esterase and member of a large multigene carboxylesterase family. The protein involved in the detoxification of xenobiotics and in the activation of ester and amide prodrugs. Hydrolyzes aromatic and aliphatic esters, but has no catalytic activity toward amides or a fatty acyl-CoA ester. Hydrolyzes the methyl ester group of cocaine to form benzoylecgonine. Thus, the recombinant human CES1 activity was measured by its ability to hydrolyze 4-Nitrophenyl acetate (4-NPA) to 4-Nitrophenol. The reaction was performed in 50mM Tris, pH 7.5 (Assay Buffer), initiated by addition 50µL of various concentrations of CES1 (dilute by assay buffer) to 50µL of 2mM Substrate 4-NPA (100mM stock in Acetone, dilute by deionized water). Incubated at 37°C for 10min, then read at a wavelength of 400nm.



**Figure 1. The standard curve of 4-Nitrophenol**

Nitrophenol (product) mM/L	OD400nm
0.045	0.01953125
0.076	0.0390625
0.123	0.078125
0.227	0.15625
0.409	0.3125
0.766	0.625
1.426	1.25
2.653	2.5

One unit of enzyme activity is defined as the 1µg of enzyme required to convert 1pmol of 4-Nitrophenyl acetate to 4-Nitrophenol in 1min at 37°C. The specific activity of recombinant human CES1 is 1396 pmol/min/µg.

$$\text{Specific Activity (pmol/min/}\mu\text{g)} = \frac{\Delta OD * F}{T * N}$$

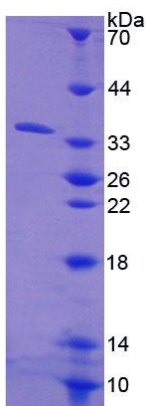
ΔOD=Adjusted for Substrate Blank

F=Conversion Factor(convert from standard curve of 4-Nitrophenol)

T= Time

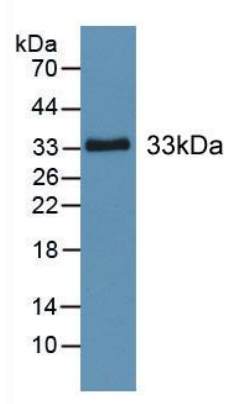
N=Amount of enzyme

## [ IDENTIFICATION ]



**Figure 2. SDS-PAGE**

**Sample: Active recombinant human, CES1**



**Figure 3. Western Blot**

**Sample: Recombinant human, CES1;**

**Antibody: Rabbit Anti-CES1 human Ab (PAC374Hu01)**

### **[ 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.