

# Sarcosine Oxidase

## ORIGIN: RECOMBINANT E.COLI

CAT#: SAR-70-2504 EC#: 1.5.3.1

### **SPECIFICATIONS**

Appearance: Yellow lyophilizate Acitvity: ≥20 U/mg lyophilizate Contaminant: Creatininase: <1×10<sup>-2</sup>%

### **ASSAY PRINCIPLE**

Sarcosine oxidase Sarcosine + H <sub>2</sub> O + O <sub>2</sub> Sarcosine oxidase $\blacktriangleright$ Glycine + Formaldehyde + H <sub>2</sub> O <sub>2</sub>	
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2 H<sub>2</sub>O<sub>2</sub> + 4-Aminoantipyrine + Phenol Peroxidase Quinoneimine dye + 4 H<sub>2</sub>O

The appearance of quinoneimine dye is measured spectrophotometrically at 495 nm

### APPLICATION

The enzyme is useful for the determination of creatinine and creatine in clinical analysis.

### UNIT DEFINITION

One unit (U) is defined as the amount of enzyme which produces 1 µmol of hydrogen peroxide per min at 37°C and pH 7.7 under standard assay conditions.

### CHARACTERISTICS

Molecular weight: ca. 49 kDa (gel filtration) Structure: monomer of 43 kDa (SDS-PAGE) | one mole of FAD per mole of enzyme Isoelectric point: 5.3 Michaelis constant: 4.7×10<sup>-3</sup> M (sarcosine) pH Optimum: 6.7–9.5 pH Stability: 6.5–10.5 Optimum temperature: 50°C Thermal stability: below 55°C Stability (liquid form): stable at 37°C for at least two weeks Stability (powder form): stable at 30°C for at least one month Inhibitor: Zn<sup>2+</sup>, Cu<sup>2+</sup>, Hg<sup>2+</sup>, Ag<sup>+</sup>

### Figure -1 pH Optimum

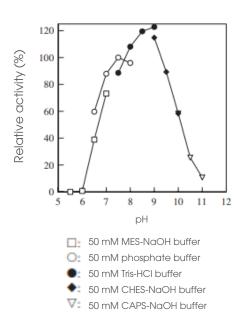
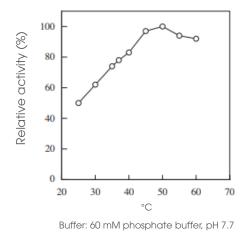


Figure -3 Optimum temperature



#### Figure -2 pH Stability

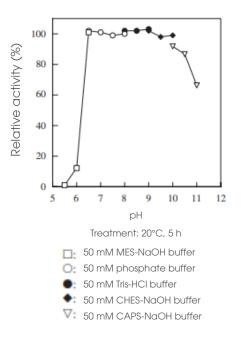
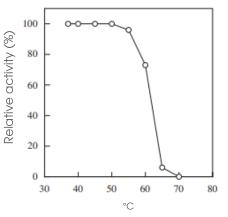


Figure -4 Thermal stability



Treatment: 0.3 M phosphate buffer, pH 7.7, 10 min

#### THE AMERICAS

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