

## Creatinase

ORIGIN: RECOMBINANT E.COLI

CAT#: CRE-70-2421

EC#: 3.5.3.3

#### **SPECIFICATIONS**

Appearance: White to light yellow lyophilizate

**Acitvity:** ≥9 U/mg lyophilizate

#### ASSAY PRINCIPLE

Creatinase catalyzes the following reaction:

Creatinase

The appearance of urea is measured spectrophotometrically at 435 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  $\mu$ mol of urea per min at 37°C and pH 7.7 under standard assay conditions.

### CHARACTERISTICS

**Molecular weight:** ca. 80 kDa (gel filtration) **Structure:** 2 subunits of 46 kDa (SDS-PAGE) **Michaelis constant:** 1.3×10<sup>-2</sup> M (creatine)

**pH Optimum:** 7.0–9.0 **pH Stability:** 5.0–11.0

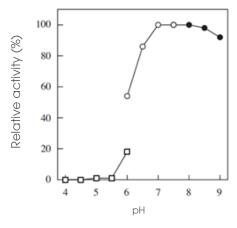
**Optimum temperature:** 40°C **Thermal stability:** below 45°C

**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: Hg<sup>2+</sup>

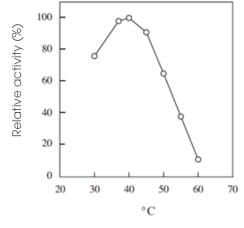
# Creatinase

Figure -1 pH Optimum



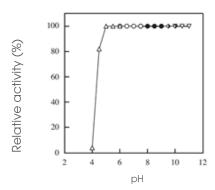
- □: 50 mM acetate buffer
- O: 50 mM phosphate buffer
- •: 50 mM Tris-HCl buffer

Figure -3 Optimum temperature



Buffer: 50 mM phosphate buffer, pH 7.5

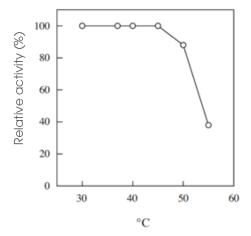
Figure -2 pH Stability



Treatment: 25°C, 17 h

- 📤: 50 mM acetate buffer
- 0: 50 mM phosphate buffer
- 50 mM Tris-HCI buffer
- ◆ 50 mM glycine-NaOH buffer
- V: 50 mM CAPS-NaOH buffer

Figure -4 Thermal stability



Treatment: 50 mM Tris-HCl buffer, pH 7.5, 30 min

#### THE AMERICAS

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