

# **Intended Use**

For the quantitative determination of Creatine Kinase in human serum.

# **Principle**

Creatine Kinase (CK) catalyzes the conversion of creatine phosphate and ADP to creatine and ATP. The ATP and glucose are converted to ADP and glucose-6-phsophate by hexokinase (HK). Glucose-6-phosphate dehydrogenase (G-6-PDH) oxidizes at the D-glucose-6-phosphate and reduces the NAD. The rate of NADH formation, measured at 340 nm, is directly proportional to serum CK activity.

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# Teco Diagnostics

#### Test:

Creatine Kinase (CK-NAC) Reagent (C512-60)

#### **Number of Tests:**

60 tests 10 x 6 mL bottles

#### Format:

Powder

#### Method:

**UV-Kinetic** 

#### **Testing Procedure:**

Manual

#### **Storage Temperature:**

2-8°C

## **Reconstituted Stability:**

24 hours at 15-30°C 21 days at 2-8°C

## Wavelength:

340 nm

# Linearity:

1,200 IU/L

#### **Expected Values:**

25-192 IU/L at 37°C 10-109 IU/L at 30°C

It is strongly recommended that each laboratory establish its own normal range.

#### **Reagent Deterioration:**

The reagent should be discarded: (1) If reagent appears damp and clumped; (2) If the reconstituted CK reagent without added sample has an absorbance greater than 0.70 at 340 nm versus reagent grade water; (3) Failure to obtain accurate results in the assay of control materials

# **Limitations of Procedure:**

Some inhibitors of CK activity: Excessive Mg<sup>++</sup>, Cl<sup>-</sup>, SO4<sup>2-</sup>, most heavy earth metals(Zn<sup>++</sup>, Cu<sup>++</sup>, Mn<sup>++</sup>), iodoacetate and other sulfhydryl binding agents, excess ADP, citrate, fluoride, L-thyroxine, and excess uric acid; This procedure measures total CK activity irrespective of its tissue or organ of origin; Lower than expected CK values have been reported in samples having high alkaline phosphatase activity.