

# rGIDH(NADP)

**recombinant Glutamate dehydrogenase (NADP<sup>+</sup>) EC 1.4.1.4**

*from Bacteria*

## Reaction Equation



## Specification

### Specific Activity

U/mg protein > 60 units  
(for reduction of α-Ketoglutarate to L-Glutamate)

### Contaminants

Glucose 6-phosphate dehydrogenase	< 0.02%
Phosphogluconate dehydrogenase	< 0.1%
Glutamate dehydrogenase (NAD <sup>+</sup> )	< 0.03%
Glutathione reductase	< 0.02%
NADPH oxidase	< 0.003%

## Properties

pH stability	: pH 5.0 - 10.5 (25°C, 1 week)
Thermal stability	: ≤ 70°C (pH 7.5, 10 min)
Optimum pH	: 7.5 - 8.0
Optimum temp.	: ≥ 70°C
Km value	: $2.5 \times 10^{-2}$ mol/L (L-Glutamate) $6.4 \times 10^{-5}$ mol/L (NADP <sup>+</sup> ) $4.1 \times 10^{-4}$ mol/L (α-Ketoglutarate) $2.4 \times 10^{-5}$ mol/L (NADPH) $4.7 \times 10^{-5}$ mol/L (Ammonium acetate)
Molecular weight	: 46 kDa (SDS-PAGE)

## Assay Procedure

### I Spectrophotometric Method

Wavelength : 340 nm, Light path length : 1 cm  
Final volume : 3.02 mL, Temperature : 25°C

Pipette the following reagents into a cuvette

2.50 mL	Triethanolamine-HCl buffer (0.1 mol/L, pH 7.6)
0.15 mL	α-Ketoglutarate (0.1 mol/L)
0.05 mL	NADPH (12 mmol/L)
0.30 mL	Ammonium acetate (2 mol/L)
0.02 mL	rGIDH (NADP) (approx. 3 U/mL)

### II Calculation

$$\frac{\Delta A/\text{min} \cdot V \cdot D}{6.2 \cdot d \cdot v} = \text{U/mL}$$

Δ A/min = The change in absorbance at 340 nm/minute

V = Total volume of reaction mixture (3.02 mL)

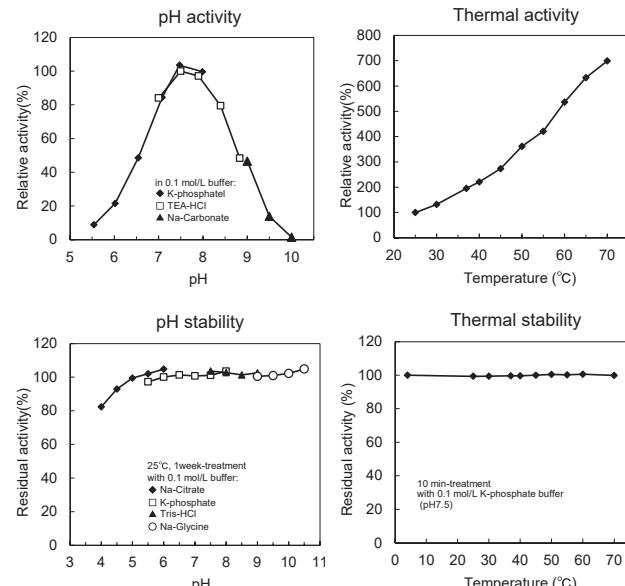
D = Enzyme dilution factor

6.2 = mmol/L extinction coefficient of NADPH  
(L<sup>-1</sup>·mmol<sup>-1</sup>·cm<sup>-1</sup>)

d = Light path length (1 cm)

v = Volume of enzyme sample (0.02 mL)

## Reference Data



## Preparation and Storage

Solution

Store at 1 - 10°C

## Cat. No./Package

Cat. No.	Package
46754904	Bulk

For in vitro diagnostic or research use only



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