

D-LACTATE DEHYDROGENASE

from Microorganism

(R)-Lactate: NAD⁺ oxidoreductase (EC 1.1.1.28)

PREPARATION and SPECIFICATION

Appearance	: White amorphous powder, lyophilized	
Activity	: 220U/mg-solid or more	
Contaminants	: GOT and GPT	all $\leq 5.0 \times 10^{-3} \%$
	Malate dehydrogenase	$\leq 1.0 \times 10^{-2} \%$
	NADH oxidase	$\leq 1.0 \times 10^{-3} \%$
	Myokinase	$\leq 1.0 \times 10^{-2} \%$
	Pyruvate Kinase	$\leq 1.0 \times 10^{-3} \%$

PROPERTIES

Stability	: Product shipped on dry ice, but long-term storage should be at -20°C .	
Molecular weight	: 39 kDa	
Isoelectric point	: 6.1	
Michaelis constant	: $6.41 \times 10^{-4}\text{M}$ (Pyruvate, pH 7.0)	
Inhibitors	: Co^{2+} , Fe^{3+} , Cu^{2+} , Hg^{2+} , Ag^+ , SDS	
Optimum pH	: 7.0	(Fig.1)
Optimum temperature	: 40°C	(Fig.2)
pH stability	: pH 4.5~10.5 (25 $^{\circ}\text{C}$, 20hr)	(Fig.3)
Thermal stability	: below 50°C (pH 7.0, 15 min)	(Fig.4)
Effect of various chemicals	: (Table 1)	

UNIT DEFINITION

One unit causes the oxidation of one micromole of NADH per minute at pH 7.0 at 25 $^{\circ}\text{C}$

APPLICATIONS

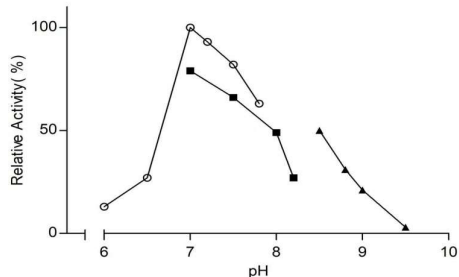
This enzyme is useful for enzymatic determination of numerous metabolites, e.g., ATP, ADP, glucose, creatinine, pyruvate, lactate and glycerol, and of enzyme activities, e.g., GPT, PK, and CPK when coupled with the related enzymes.

Manufactured in an ISO 9001 certified facility: Suzhou SignalChem Biotechnologies Corp.

Table 1. Effect of Various Chemicals on D-Lactate dehydrogenase

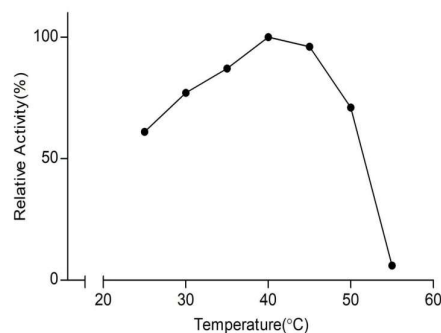
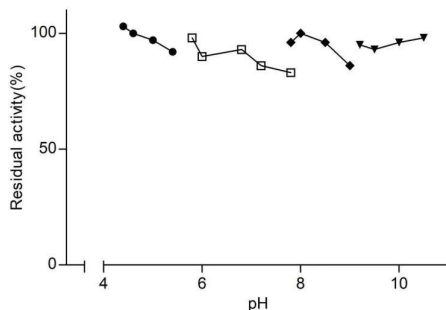
The enzyme was dissolved in 0.05M K-phosphate buffer, pH 7.5 containing 0.1% of BSA (40U/ml) and incubated with each chemical at 25°C for 1hr.

Chemical	Concn.(mM)	Residual activity(%)	Chemical	Concn.(mM)	Residual activity(%)
None	—	100	BME	2	91
CaCl ₂	2	94	Hydroxylamine	2	93
MgSO ₄	2	95	EDTA	5	95
ZnSO ₄	2	86	NaF	20	90
NiCl ₂	2	89	NaN ₃	20	96
CoCl ₂	2	58	Borate	50	89
MnCl ₂	2	96	Proclin-300	0.045% (v/v)	90
FeCl ₃	2	63	SDS	0.05%	47
CuSO ₄	2	78	Na-Cholate	0.1%	97
AgNO ₃	2	6	Tween-20	0.1% (v/v)	96
HgSO ₄	2	1	Triton X-100	0.1% (v/v)	97
NEM	2	80	Span-20	0.1% (v/v)	94
IAA	2	89	Brij-35	0.1%	94

Fig.1. pH Activity


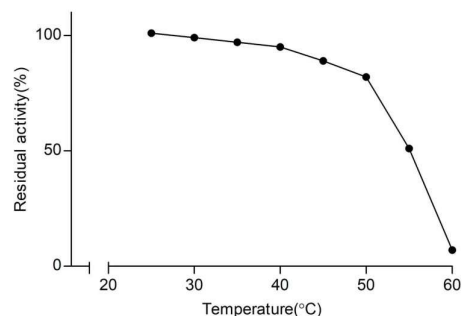
25°C in the following buffer solution:

- 50mM K-phosphate buffer
- 50mM Tris-HCl buffer
- ▲ 50mM Glycine-NaOH buffer

Fig.2. Temperature Activity

Fig.3. Thermal Stability


25°C, 20hr-treatment with following buffer solution:

- 0.1M Acetate buffer
- 0.1M K-phosphate buffer
- ◆ 0.1M Tris-HCl buffer
- ▼ 0.1M Glycine-NaOH buffer



15min- treatment with 50mM K-phosphate buffer, pH7.0