

ENZYMES

Glycerol Phosphate Oxidase

ORIGIN *Streptococcus sp.*

CAT# 70-4875-01
EC# 1.1.3.21

► SPECIFICATIONS

Appearance	Yellow powder
Activity	≥40 U/mg powder at 37°C
Contaminants	Acetate Kinase <0.1% Lactate Oxidase <0.001%

► ASSAY PRINCIPLE

Glycerol Phosphate Oxidase (GPO) catalyses the following reaction:



The formation of dihydroxyacetone phosphate is determined using a coupled peroxidase assay system, which causes the formation of a quinoneimine dye, that may be measured spectrophotometrically at 500nm.

► UNIT DEFINITION

One unit of activity is defined as the amount of enzyme that will catalyze the oxidation of 1.0 micromole of Glycerol-3-Phosphate per minute at 37°C under the standard assay method conditions (available on request).

► APPLICATION

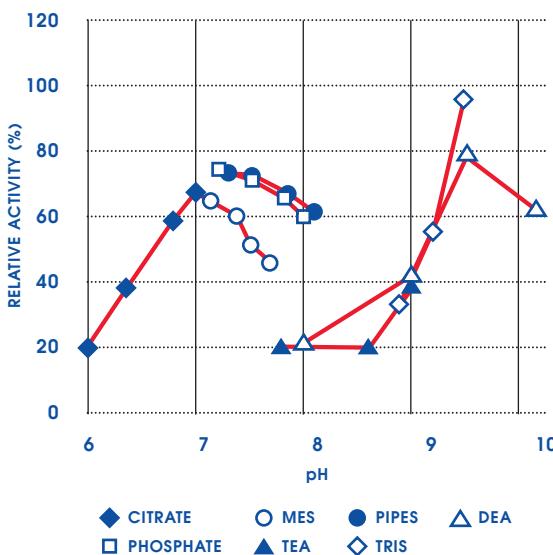
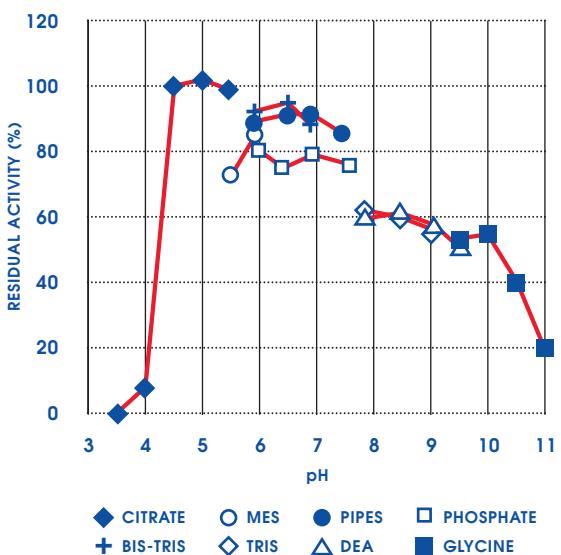
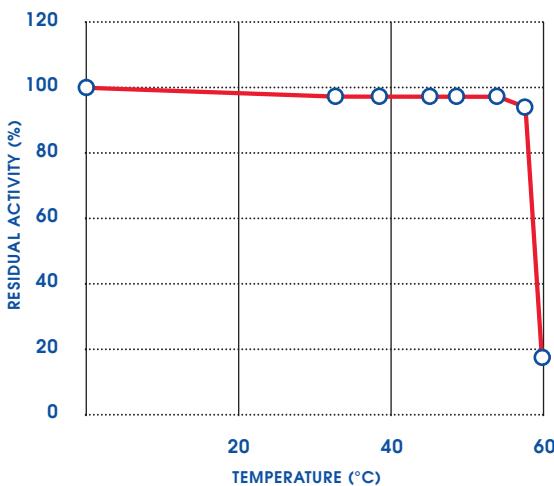
Used in the formulation of triglyceride reagents.

CHARACTERISTICS

Molecular Weight:	67kDa (SDS-PAGE)
Isoelectric Point:	4.03
K _m value:	2.23 x 10 ⁻³
Optimum pH (Fig. 1):	pH 6.5 and pH 8.5 - 9.0
Optimum Temperature:	37°C
pH Stability (Fig. 2):	5.0 to 7.0 (37°C for 30 minutes)
Thermal Stability (Fig. 3):	Stable at 55°C and below (pH 6.5 for 5 minutes)

TABLE 1: EFFECT OF VARIOUS CHEMICALS ON GLYCEROL PHOSPHATE OXIDASE

CHEMICAL	CONCENTRATION (mM)	RELATIVE ACTIVITY (%)	CHEMICAL	CONCENTRATION (mM)	RELATIVE ACTIVITY (%)
None	—	100	LiCl	2.0	103
MgCl ₂	2.0	101	KCl	2.0	102
MgSO ₄	2.0	102	CaCl ₂	2.0	103
ZnCl ₂	2.0	102	Emulgen 810	0.1%	98
ZnSO ₄	2.0	102	Emulgen 911	0.1%	98
NaCl	2.0	103	Rheodol TWL-106	0.1%	99
NH ₄ Cl	2.0	103	Rheodol 460	0.1%	99
BaCl ₂	2.0	103	Adekanol NP-720	0.1%	99
Ba(CH ₃ COO) ₂	2.0	101	Triton X-100	0.1%	99
NiCl ₂	2.0	103	Triton X-305	0.1%	98
CoCl ₂	2.0	103	Tween 80	0.1%	100
MnCl ₂	2.0	114			

FIGURE 1: OPTIMUM pH**FIGURE 2: pH STABILITY (37°C, 30 MIN.)****FIGURE 3: THERMAL STABILITY (pH 6.5 FOR 5 MINS.)****THE AMERICAS**

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