

α-GLUCOSIDASE from Thermotoga maritima (Lot 151001a)

Recombinant - Thermostable

E-AGLUTM I0/15

(EC 3.2.1.20) alpha-glucosidase; alpha-D-glucoside glucohydrolase

CAZy: GH Family 4 CAS: 9001-42-7

PROPERTIES

I. ELECTROPHORETIC PURITY:

- Single band on SDS-gel electrophoresis (MW ~ 56,000)
- One major bands on isoelectric focusing (pl ~ 6.1)

2. SPECIFIC ACTIVITY:

32 U/mg protein (on p-NP- α -D-glucopyranoside) at pH 7.5 and 80°C.

~ II U/mg protein (on p-NP- α -D-glucopyranoside) at pH 7.5 and 60°C.

One Unit of α -glucosidase activity is defined as the amount of enzyme required to release one μ mole of of p-nitrophenol (p-NP) per minute from p-nitrophenyl- α -D-glucopyranoside (5 mM) in Tris.HCl buffer (100 mM), pH 7.5 at 80°C.

3. SPECIFICITY:

Hydrolysis of terminal, non-reducing (1,4)- α -linked D-glucose residues with release of α -D-glucose.

4. RELATIVE RATES OF HYDROLYSIS OF SUBSTRATES:

Enzyme Measured	Substrate	Activity, %
α-Glucosidase	p-NP-α-D-Glucopyranoside	100
β-Glucosidase	p-NP-β-D-Glucopyranoside	< 0.1
α -Galactosidase	p -NP- α -D-Galactopyranoside	~ 71
β -Galactosidase	p-NP-β-D-Galactopyranoside	< 1.0
α -Mannosidase	p -NP- α -D-Mannopyranoside	< 0.0001
β -Mannosidase	p-NP-β-D-Mannopyranoside	< 0.0001
α -Glucosidase	Maltose	< 0.05
Sucrase	Sucrose	< 0.1
Trehalase	Trehalose	< 0.1

Action on polysaccharide and p-nitropenyl substrates was determined at final concentrations of 10 mg/mL and 5 mM, respectively, in Tris.HCl buffer (100 mM), pH 7.5 at 40°C.

5. PHYSICOCHEMICAL PROPERTIES:

Recommended conditions of use are at pH 7.5 - 8.0 and up to 80°C

pH Optima: 7.5 - 8.5

pH Stability: 4.0 - 9.0 (> 75% control activity after 24 hours at 4°C)

Temperature Optima: 80 - 100°C (10 min. reaction)

Temperature Stability: up to 80°C

6. STORAGE CONDITIONS

The enzyme is supplied as a solution containing 50% glycerol plus 0.02% (w/v) sodium azide and should be stored at -20°C. For assay, this enzyme should be diluted in Tris.HCl buffer (100 mM), pH 7.5 containing 1 mg/mL BSA, 3 mM NAD, 4 mM manganese chloride and 400 mM mercaptoethanol. Swirl to mix the enzyme immediately prior to use.

7. EXPERIMENTAL DATA







