

β-GLUCOSIDASE from Thermotoga maritima (Lot 151102a)

Recombinant - Thermostable

E-BGOSTM II/I8

(EC 3.2.1.21) beta-D-glucoside glucohydrolase CAZy Family: GHI

PROPERTIES

I. ELECTROPHORETIC PURITY:

- Single band on SDS-gel electrophoresis (MW ~ 53,700)
- One major band on isoelectric focusing (pl \sim 6.0)

2. SPECIFIC ACTIVITY:

50 U/mg protein (on p-NP-β-D-Glucopyranoside) at pH 6.5 and 40°C.

One Unit of β -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- β -D-glucopyranoside (10 mM) in sodium maleate buffer (50 mM), pH 6.5 at 40°C.

3. OTHER ACTIVITIES (as a percentage of β -glucosidase activity):

Enzyme Measured	Substrate	Activity, %
β-Glucosidase	p-NP-β-D-Glucopyranoside	100
β-Glucosidase	Cellobiose	~ 17
β-Galactosidase	p-NP-β-D-Galactopyranoside	~ 100
α-Amylase	Ceralpha Reagent	< 0.0001
Amyloglucosidase	Starch	< 0.0002
α-Glucosidase	p -NP- α -D-Glucopyranoside	< 0.001
endo-1,4- β -Glucanase	Cellazyme C Tablets (Megazyme)	< 0.0001

Action on polysaccharide and p-nitropenyl substrates was determined at final concentrations of 10 mg/mL and 10 mM, respectively, in sodium maleate buffer (100 mM), pH 6.5 at 40°C.

4. RELATIVE RATES OF HYDROLYSIS OF SUBSTRATES:

Substrate	%
Cellobiose	100
Laminaribiose	~ 109
Laminaritriose	~ 100
Laminaritetraose	~ 59
Laminaripentaose	~ 29
Laminarihexaose	~ 5.0
Gentiobiose	~ 5.9
Sophorose	~ 98
I,4-β-D-Glucosyl-D-mannose	~ 9.9
p-Nitrophenyl β-D-glucopyranoside	~ 566
p-Nitrophenyl β-D-xylanopyranoside	~ 1.8
p-Nitrophenyl α-D-glucopyranoside	< 0.001

5. PHYSICOCHEMICAL PROPERTIES:

pH Optima: 7.0 (at 40°C)
Temperature Optima: 90°C (30 min at pH 6.5)
pH Stability: 4.0-9.0 (at 40°C for 30 min)
Temperature Stability: Unstable above 90°C

6. STORAGE CONDITIONS

The enzyme is supplied as an ammonium sulphate suspension in 0.02% (w/v) sodium azide and should be stored at 4°C. For assay, this enzyme should be diluted in sodium maleate buffer (50 mM), pH 6.5 containing 0.5 mg/mL BSA. **Swirl to mix the enzyme immediately prior to use.**