

β-GLUCOSIDASE from Phanerochaete chrysosporium (Lot 121101f)

Recombinant - Thermostable

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Full length enzyme containing the N-terminal cellulose-binding domain (CBMI) and a C-terminal catalytic domain (GH3). (EC 3.2.1.21) beta-D-glucoside glucohydrolase

CAZy Family: GH3 CAS: 9001-22-3

PROPERTIES

I. ELECTROPHORETIC PURITY:

- Single band on SDS-gel electrophoresis (MW ~ 84,500)
- One major bands on isoelectric focusing (pl ~ 5.8)

2. SPECIFIC ACTIVITY:

103 U/mg protein (on p-NP- β -D-Glucopyranoside) at pH 5.0 and 40°C.

~ 320 U/mg protein (on p-NP- β -D-Glucopyranoside) at pH 5.0 and 60°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 (5 mM) in sodium acetate buffer (100 mM), pH 5.0 at 40°C.

3. SPECIFICITY:

Hydrolysis of terminal non-reducing β -D-glucosidic bonds releasing β -D-glucose.

4. RELATIVE RATES OF HYDROLYSIS OF SUBSTRATES:

Enzyme Measured	Substrate	Activity, %
β-Glucosidase	p-NP-β-D-Glucopyranoside	100
β-1,4-D-Glucosidase	Cellobiose	~ 14
β-1,6-D-Glucosidase	Gentiobiose	~ 70
β-Galactosidase	p-NP-β-D-Galactopyranoside	< 0.02
α -Glucosidase	p -NP- α -D-Glucopyranoside	< 0.004
α -Glucosidase	Maltose	< 0.0001
β-Xylosidase	p -NP- α -D-Xylopyranoside	~ 0.6
α -Amylase	Ceralpha Reagent (Megazyme)	< 0.0001
Amyloglucosidase	Starch	< 0.0001
endo-1,4-β-Glucanase	CM-Cellulose 4M	< 0.003

Action on polysaccharide and p-nitrophenyl substrates was determined at final concentrations of 10 mg/mL and 5 mM, respectively, in sodium acetate buffer (100 mM), pH 5.0 at 40°C.

5. PHYSICOCHEMICAL PROPERTIES:

Recommended conditions of use are at pH 3.0 - 5.0 and up to 60°C

pH Optima: 4.5 - 5.0

pH Stability: 3.0 - 9.0 (> 75% control activity after 24 hours at 4°C)

Temperature Optima: 60 - 70°C (10 min. reaction)

Temperature Stability: up to 60°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 acetate buffer (100 mM), pH 5.0 containing I mg/mL BSA. **Swirl to mix the enzyme immediately prior to use.**

7. EXPERIMENTAL DATA







