

α -GALACTOSIDASE from Aspergillus niger (Lot 00901a)

E-AGLAN

11/14

(EC 3.2.1.22) alpha-D-galactoside galactohydrolase CAZy Family: GH 36

PROPERTIES

- I. ELECTROPHORETIC PURITY:
 - Single major band on SDS-gel electrophoresis (MW = 97,000)
 - Single major band on isoelectric focusing (pl = 4.2)

2. SPECIFIC ACTIVITY AND LEVEL OF OTHER ACTIVITIES:

All activities are at pH 4.5 and 40°C. Glycosidase activities were measured using the appropriate p-nitrophenyl glycoside (at 10 mM). endo-Glycanases were determined with the appropriate substrate (at 10 mg/mL) and using the Nelson/Somogyi reducing-sugar procedure.

One Unit of activity is the amount of enzyme required to release one micromole of product (e.g. p-nitrophenyl) per min at pH 4.5 and 40°C.

| Substrate | Enzyme Measured | Specific Activity |
|-------------------------------|---------------------------------|-------------------|
| | | (U/mg protein) |
| p-NP-α-Galactoside | lpha-Galactosidase | 620 |
| b-NP-β-Galactoside | β -Galactosidase | < 0.001 |
| p-NP-α-Glucoside | α -Glucosidase | < 0.001 |
| p-NP-β-Glucoside | β -Glucosidase | < 0.001 |
| p-NP-β-Xyloside | β-Xylosidase | < 0.001 |
| b-NP- β -Mannoside | β -Mannosidase | < 0.001 |
| b-NP- α -L-arabinoside | α -L-arabinofuranosidase | < 0.001 |
| Carob Galactomannan | endo-1,4- β -Mannanase | < 0.02 |
| Sucrose | Invertase | < 0.05 |
| I-Kestose | exo-Inulinanase | < 0.01 |
| I,I-Kestotetraose | exo-Inulinanase | < 0.01 |
| Fructan (polymer) | exo-Inulinanase | < 0.01 |

3. PHYSICOCHEMICAL PROPERTIES:

| pH Optima: | 4.5-5.0 |
|------------------------|---------------------|
| pH Stability: | 4.0-8.0 |
| Temperature Optima: | 60°C (at pH 5.0) |
| Temperature Stability: | Unstable above 60°C |

4. STORAGE CONDITIONS:

The enzyme is supplied as an ammonium sulphate suspension in 0.02% sodium azide and should be stored at 4° C. On dissolution in buffer, the enzyme should be stored in the frozen state in a polypropylene container between use. We recommend the addition of BSA (0.5 mg/mL) to all dilution buffers to improve stability of the enzyme.