



SUCRASE (MALTASE) From Yeast (Lot 80702)

E-SUCR

03/13

EC 3.2.1.20 alpha-D-glucoside glucohydrolase

PROPERTIES

1. ELECTROPHORETIC PURITY:

- Single major band on SDS-gel electrophoresis (62,000)
- Single major band on isoelectric focusing (pI = 5.7)

2. SPECIFIC ACTIVITY AND LEVEL OF OTHER ACTIVITIES:

Substrate	Enzyme Measured	Specific Activity (U/mg protein)
Sucrose	α -Glucosidase	23.0
Maltose	α -Glucosidase	21.9
<i>p</i> -NP- α -Glucoside	α -Glucosidase	132.5
Kestose & Kestotetraose	α -Glucosidase	< 0.005
<i>p</i> -NP- β -Glucosidase	β -Glucosidase	< 0.001
<i>p</i> -NP- α -Galactoside	β -Galactosidase	< 0.001
<i>p</i> -NP- β -Galactoside	β -Galactosidase	< 0.001
Blocked <i>p</i> -NP-Maltoheptoaside	α -Amylase	< 0.001

This enzyme specifically hydrolyses sucrose in the presence of fructo-oligosaccharides. All activities were measured at pH 6.8 and 40°C. Action on sucrose was measured as glucose release, using glucose oxidase/peroxidase reagent. One Unit of enzyme activity is the amount of enzyme required to release one μ mole of glucose/min from sucrose (10 mM) at pH 6.8 and 40°C. Other glycosidase activities were measured using the appropriate *p*-nitrophenyl glycoside (at 10 mM). One Unit of enzyme activity is the amount of enzyme required to release one micromole of *p*-nitrophenol/min from the appropriate substrate at pH 6.8 and 40°C.

α -Amylase was measured using the "CERALPHA" α -amylase assay method.

3. PHYSICOCHEMICAL PROPERTIES:

pH Optima:	6.4-6.8
pH Stability:	5.6-7
Temperature Optima:	40°C
Temperature Stability:	< 40°C

4. STORAGE CONDITIONS:

The enzyme is supplied as a lyophilised powder and should be stored at -20°C. On dissolution in buffer or water, the enzyme should be stored in the frozen state. **It is recommended** that all buffers used for dilution contain BSA (1.0 mg/mL).