

Recombinant Enzyme Product Specification Sheet

Cat. No.:	PRO-E0413	add this product to cart
LOT:	2010-0413-2	view all branching enzymes
Activity:	1,4- α -Glucan branching enzyme	
Synonyms:	Branching enzyme; amylo-(1,4 \rightarrow 1,6)-transglycosylase; Q-enzyme; α -glucan-branching glycosyltransferase; amylose isomerase; enzymatic branching factor; branching glycosyltransferase; enzyme Q; glucosan transglycosylase; glycogen branching enzyme; plant branching enzyme; α -1,4-glucan: α -1,4-glucan-6-glycosyltransferase; starch branching enzyme; 1,4- α -D-glucan:1,4- α -D-glucan 6- α -D-(1,4- α -D-glucano)-transferase	
Nomenclature:	CAZy [GH13 subf8, glycoside hydrolase family 13 subfamily 8, member of clan GH-H], BF2338, GlgB	
Source organism:	<i>Bacteroides fragilis</i> NCTC 9343	
Enzyme Commission No.:	2.4.1.18	
Activity:	330.14 U/mL	} (37°C; pH 7.5; 3.3 mg/mL starch)
Specific activity:	50.88 U/mg	
Purity:	~ 95 % as judged by SDS-PAGE	
Form and storage:	Supplied in 3.2 M ammonium sulphate, containing 0.5 M imidazole and 0.5 M NaCl, pH ~ 6.8. Store at 4°C (shipped at room temperature)	
pH optimum:	~ 7.0	
Temperature optimum:	$\geq 37^\circ\text{C}$	
[Protein]:	6.49 mg/mL	
Sequence length:	670 amino acids (view sequence)	
Accession No.:	Q5LCX9 , YP_211960.1 , CAH08034.1 , BFRA272559:BF2338-MON	
Molecular weight:	81104.6 Da	(theoretical)
	-	(observed by SDS-PAGE)
	-	(observed by mass spectrometry)
Biological function:	May be involved in glycogen biosynthesis.	
Potential application(s):	Carbohydrate research , fundamental research	

- Comments:** This is a cytoplasmic enzyme
- Usage:** Agitate vial sufficiently to fully homogenise enzyme precipitate before use
- Assay:** One unit is defined as the amount of enzyme required to cause a fall of 1.0 absorbance unit per minute, where the reaction mixture comprises 3.33 mg/mL starch (Sigma; S-9765; boiled for 5 min prior to use to fully solubilise) in 41.7 mM sodium phosphate buffer, pH 7.5, containing 0.69 mg/mL BSA and 173.6 mM sodium chloride, and where 0.050 mL of the reaction mixture (boiled for 5 min to inactivate the enzyme) is mixed with 1.0 mL iodine reagent (0.5 mg/mL iodine and 1 mg/mL potassium iodide in water) prior to reading at 660 nm.

NOTE: due to the nature of the assay, it is important to monitor the initial reaction rate, over the first 2 min of the reaction

Primary sequence:

MEKTLNLIKNDPWLEPYKDAIVGRFEHAMDKKAELTNGGKSTLSDFASGYLYFGLHRTDKGWI FREWAPNASHIY
MVGTFNSWEEKPAYKLRKLNKSWEIKLPIDTIQHGDLYKLHVYWEGGQGERIPAWANRVVQDDNTKIFSAQVWA
PEKPFKFKKKTFKPSTDPLLIYECHIGMAQQEEKVGTYNFREFREKILPRIAKEGYNCIQIMAIQEHPYYGSFGYHV
SSFFAASSRFGTPEELKQLIDTAHGLGIAVIMDIVHSHAVKNEVEGLGNFAGDPNQYFYPPGRRREHPAWDSLCFD
YGKNEVMHFLLSNCKYWLEEYHFDGFRFDGVTSMLYYSHGLGEAFCNYGDYFNGHQDDNAICYLTLANELIHEVN
PKAITIAEEVSGMPGLAAKVEDGGYGFDIRMAMNIPDYWIKTIKEKIDEDWKPSSMFWEVTNRRQDEKTISYAES
HDQALVGDKTIIIFRLIDADMYWHMQKDENYIVHRGVALHKMIRLLTASTINGGYLNFMGNEFGHPEWIDFPREG
NGWSCKYARRQWDLVDNKNLTYHYLGDFDADMLKVIKSVKNIQQTPVQEIWHNDGDQVLAYQRKDLVVFVFNFNPS
QSFTDYGFLVTPGTYEVLNTDNI IYGGNGLSDDSVKHFTLPDPLYKKEKKEWLKLYIPARTAMVLRRTK

- Literature:** 1. [Cerdeno-Tarraga et al. \(2005\) Science 307, 1463-1465](#)