



ENZYMES
BY

SORACHIM

**α -Glucosidase (Maltase) from Microorganism
AGH-211**

SPECIFICATIONS

Name	α -D-Glucoside glucohydrolase
EC	3.2.1.20
Appearance	White amorphous powder lyophilized
Activity	Grade II, 20 U/mg-solid or more
Contaminants	α amylase $\leq 1.0 \times 10^{-5}$ %
Stabilizers	Bovine serum albumin (BSA)
Stability	Stable at -20 °C for at least one year
Molecular weight	65,000
Isoelectric point	5.2
Michaelis constant	6.3×10^{-4} M (p-Nitrophenyl- α -D-glucopyranoside)
Inhibitors	Ag^+ , Hg^{2+} , PCMB, MIA
Optimum pH	6.0 - 7.0
Optimum temperature	60 °C
pH Stability	pH 5.0 - 9.0
Thermal stability	below 60 °C (pH 7.0, 15min)

E
N
Z
Y
M
E
S

Substrate*	Relative hydrolysis rate**	Substrate*	Relative hydrolysis rate**
PNPG	100.0	Maltose	271.0
PNPG ₂	205.0	Maltotriose	203.0
PNPG ₃	284.0	Maltotetraose	168.0
PNPG ₅	164.0	Maltopentaose	100.0

* : Substrate concn. 2.2mM

** : Glucose-forming activity, pH 6.8 at 37°C

www.Sorachim.com

contact@sorachim.com

ENZYMES
BY

SORACHIM

α -Glucosidase (Maltase) from Microorganism AGH-211

SPECIFICATIONS

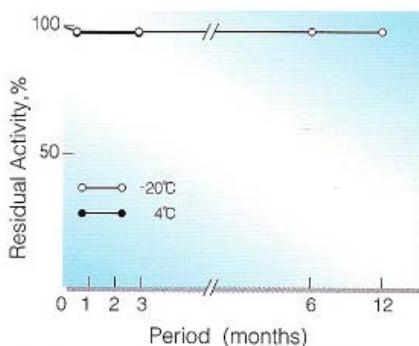


Fig.1. Stability (Powder form)
(kept under dry conditions)

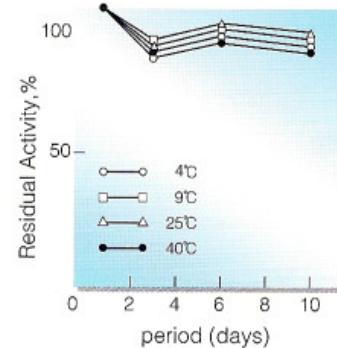


Fig.2. Stability (Liquid form)

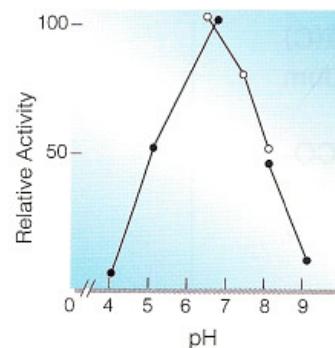


Fig.3. pH-Activity

[37°C, 15 min-reaction in 100mM buffer solution:
●, pH4.0-6.0 acetate; ○, pH6.0-8.0,
phosphate; ■, pH8.0-9.0, borate
enzyme concn.: 5U/ml]

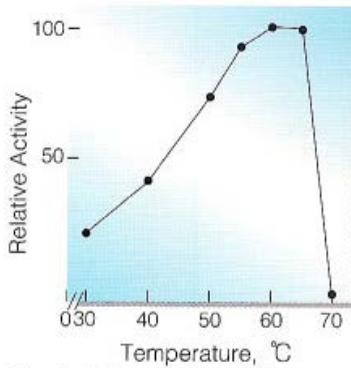


Fig.4. Thermal activity
(15 min-reaction in 100mM phosphate
buffer, pH7.0)

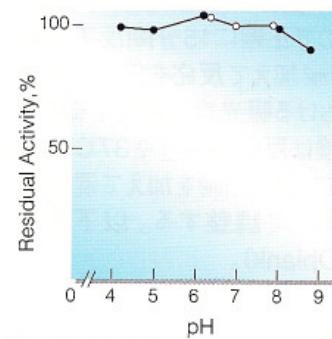


Fig.5. pH-Stability
[25°C, 20hr-treatment with 50mM buffer
solution contg: 0.2% of BSA;
●, pH4.0-6.0
acetate; ○, pH6.0-8.0, phosphate;
■, pH8.0-9.0,
borate. enzyme concn. : 5U/ml]

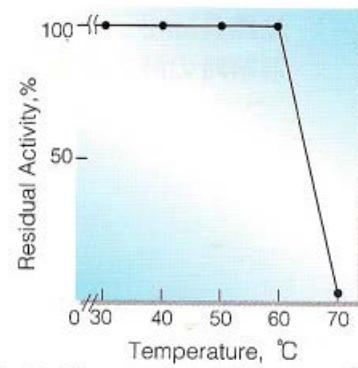


Fig.6. Thermal stability
[15min-treatment with 0.2M K-phosphate
buffer, pH7.0 contg. 1mM EDTA and 0.05%
Tween20. enzyme concn.: 5U/ml]