

Penicillin - Acylase

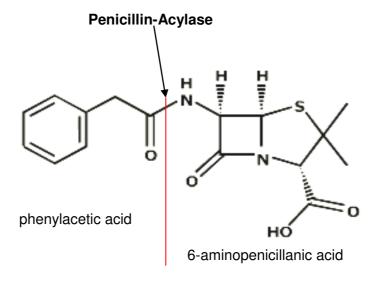
Penicillin amidohydrolase EC 3.5.1.11

Description:

Cleavage of Penicillin G into 6-aminopenicillanic acid (6-APA) and phenylacetic acid.

Synthesis of unnatural penicillin by coupling 6-aminopenicillanic acid with a selected side chain (e.g. Methylcillin, Ampicillin, Phenethicillin)

Reaction:



Penicillin G

Origin:	E.coli	
Application:	semi-synthetic production of several penicillins	
Michaelis-MConst.:	2·10 ⁵ M	
Activity:	> 800 U/ml (method: ASA Spezialen	zyme GmbH)
Specific activity:	> 4 U/mg	
Molecular weight:	70 000 D (by gel filtration)	
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Isoelectric points:	6.7 - 6.8 and $6.3 - 6.4$ (by preparative isoelectric focusing)	
Parameters of reaction:	<u>pH</u> hydrolysis of 6-nitro-3-(phenylacetoamid)bezoic acid: 7,5 hydrolysis of Penicillin G: 8,1	
	temperature optimum 54°C	
Inhibitors:	phenylacetic acid (competitive, $K_i = 200 \text{ mM}$) 6-aminopenicillanic acid (not competitive, $K_i = 15 \text{ mM}$) conditions: Penicillin G as substrate, 25°C, pH 8,1	
Order-No.:	2605	
Form of delivery:	yellow - brown liquid with typical odour	
Storage:	-20°C	
Stability:	At -20°C the enzyme is stable for years. For months stable at pH 5 - 8 and at ambient temperature. Fast loss of activity after treatment with organic solvents as well as after lyophilisation. An incubation for 20 minutes at pH 5 and 50°C does not lead to a loss of activity.	
Literature:	Kutzbach C., Rauenbusch E.: <i>Hoppe-Seyler´s Z. Physiol. Chem.</i> 354, 45 (1974)	
	Mahajan P. B.: <i>Appl. Biochem. and Biotechn. Vol. 9.</i> 537 (1984)	
	Savidge T. A., Cole: <i>M. in Methods in Enzymology Vol. 43.</i> 705, Academic Press, New York – London (1975)	