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6-N-Trimethyllysine metabolism and carnitine biosynthesis in N. crassa

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6-N-1 rimetnyllysine metabolism and carnitine biosynthesis in N. crassa
Abstract 6-N-Trimethyllysine metabolism and carnitine biosynthesis

Villanueva, V.R. and E. Lederer 6-N-trimethyllysine metabolism

and carnitine biosynthesis in N. crassa.

Villanueva and Lederer (1974, Phytochemirtry 13: 2157) hove recently reported the presence of free and protein-bound 6-N-mono-, di- and tri-methylated lysines in Neu-rospora crassa. Horne and Broquist (1973 J. Biol. Chem. 248: 2170) hove shown that 6-N-trimethyllysine (I) is a highly efficient precursor of carnitine (IV) via butyrobetaine

(II') in Neurospora crassa. Although the same relationship among there metabolites has been found in the rat, (Tanphaichitr and Broquist 1973 J. Biol. Chem. 248: 2176; Cox and Hoppel 1973 Biochem. J. 136: 1083), no intermediates of this metabolic pathway have been isolated nor identified as yet. Neither Lindsted and Lindsted (1965 J. Bio. Chem. 240: 316) nor Cox and Hoppel ('974 Biochem. Biophys. Acta 362: 403) could obtain incorporation of radioactivity from 5-N-(14CH3)trimethylaminopentanoate or 6-N(14CH3)trimethylaminohexanoate into carnitine. We have now examined the possibility of a lysine decarboxylating pathway for 6-N-trimethyllysine (MegLys). The expected intermediate would be N-trimethylcodayerine (MegCad). Extracts of cultures of N. crassa strain lys-1 (33933) (FGSC #74), growing in a medium containing Meg(14CH₃)Lys or Meg(14CH₂) Cod, were analysed by appropriate automatic ion exchange column chromatography and checked for the eventual conversion of Meg(14CHg) Lys into Meg(14CHg)Cad, as well as for the conversion of the latter into butyrobetaine and carnitine. Neither conversion of Meg(14CHg) lys into Meg(14CHg) Cad nor conversion of Meg(14CHg) Cad into butyrobetaine and carnitine was observed. Unexpectedly, analysis of the extracts labelled with Me3(14CH3)Lys gave rise to three mean radioactive peaks: the first one the unknown Was excluded form the column; the second eluted at the position corresponding to that of butyrobetoine-carnitine and the last one. to that of Meglys. The unknown radioactive product gave a positive reaction with 2,4-dinitrophenylhydrazine, indicating the presence of g keto-group. We thought that the unknown compound could be 6-N-trimethylomino, 2-oxohexanogte, in order to check this hypothesis we subsequently transformed the isolated radioactive unknown into its 2,4-dinitrophenylhydrozone derivative and submitted it to hydrogenolysis in a Parr bomb. Analysis of the reaction products by TLC and ion exchange column chromatography (4 different systems) showed a positive ninhydrine-reacting substance at the same Rf and with the same elution time as that of authentic Mealys and containing more than 85% of the radioactivity of the 2,4-dinifrophenylhydrazone before hydrogenolysis. This result strongly suggests that the unknown radioactive product is indeed 6-N-trimethylamino, 2-oxohexanoic acid (II).

Experiments with extracts of N. crassa using Meg(14CHg)Lys led to the same results as "in vivo" experiments, demonstrating that an enzymate system exists which can convert MegLys (I) into the corresponding ketoacid (II).

Work to verify that this ketoacid is an intermediate in the biosynthesis of carnitine (IV) is in progress.

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