in Neurospora with amino acid analogs.

Failure to induce mutations

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ation on 50 ml of agar minimal medium (supplemented with 100% a/ml adenine sulfate for the ad-3 strains) containing one of the analogs at the following concentrations: L-canavanine sulfate 100 and DL-ethionine 20 ug/ml, p-fluoro-DL-phenylalanine 4 ug/ml. These concentrations retarded growth but did not prevent conidiation. The reversion experiments employed the following ad-3 strains supplied by H.V. Mailing: 2-17-8, 2-17-23,

An attempt was made to induce forward and reverse mutations with the amino acid analogs

canavanine, ethionine and fluorophenylalanine. Treatment consisted of growing cultures to conidi-

5-4-1, 2-17-7, 2-17-61, 2-17-155, 2-17-18, 2-17-126. Each analog was tested one or more times on each ad-3 strain. The number of live conidia plated on minimal medium ranged from 108 to 109 per test. The spontaneous reversion frequencies obtained were similar to those reported by Malling and DeSerres for these strains (1967 Mutation Res. 4: 425), but in no instance were the reversion frequencies of the analog treated cultures significantly higher than the controls. Similarly treated conidia of 74a were screened for mutation to resistance to cycloheximide (2µa/ml) and resistance to benomy! (1 u g/ml). Again, the analog treated strains showed no increase in mutation frequency over the controls. These results are of interest

chiefly because amino acid analog mutagenesis has been reported in several organisms including one fungus (Talmud and Lewis 1974 Genetical Res. - - Biology Department, University of West Florida, Pensacola, FL 32504.