variation in natural populations of Neurospora. porg were obtained from D. D. Perkins' Florida collection. Mycelial extracts from these rtroing were subjected to acrylamide and starch gel electrophoresis. Out of ten enzymes examined, electrophoretic variation was observed only for esterases. The sites of

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Of the eight heterothallic strains, six (P384, P385, P406, P407, P413, P419) had esterase rite 1 and two strains (P438, P439) hod both esterase sites 1 and 2. Of the three homothallic strains, two (P388, P436) had esterase rites 3 and 4 and the third (P404) had esterase site 2. Amylase, aminopeptidase, a-glycerophorphote dehydrogenase, 6-phosphoglucanate dehydrogenase and in Ephenol oxidase showed ane site of activity. Acid phosphatase showed activity at two sites and lactate dehydrogenase, peroxidase and glucose-6-phosphate dehydrogenase showed activity at three sites for all the strains. The absence of electrophoretic variation for there enzymes suggests that selection may have been operating against enzyme variants resulting in stabilization of the enzyme genotype of isolated populations in nature.

esterase activity were numbered from 1 to 4 in order of rate of movement towards the anode, with site 1 being the fastest.

We would like to thank D. D. Perkins for kindly providing the strains, - - Department of Biology, McMaster University, Hamilton, Ontario, Canada.

Eight heterothallic (P384, P385, P406, P407, P413, P419, P438, P439) and three homothallic strains (P388, P404, P435) of Neuros-