

**IMMUNOLOGICAL PROPERTIES OF BACTERIAL PYRUVATE  
DEHYDROGENASE COMPLEXES AND CHARACTERISATION  
OF DELETION MUTANTS OF *ESCHERICHIA COLI***

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The pyruvate dehydrogenase complex of *Escherichia coli* produced two precipitin lines in double diffusion tests with antiserum raised against the homologous complex. These were identified as specific reactions involving pyruvate dehydrogenase (E<sub>1</sub>) and lipoamide dehydrogenase (E<sub>3</sub>) components. During the course of this work several mutants of *E. coli* with deletions in the *nadC-aroP-aceF-lyd* region were investigated and their immunological properties were found consistent with the earlier results of enzymic and genetic characterization. A few deletion strains possessing low pyruvate dehydrogenase (E<sub>1</sub>) activities showed no precipitin lines corresponding to the E<sub>1</sub> component. The low E<sub>1</sub> activities could be attributed to the presence of pyruvate oxidase in these strains.

Evidence for immunological cross reactivity between the pyruvate dehydrogenase complexes of *Pseudomonas aeruginosa* and *E. coli* was observed despite the negative precipitin reaction in double diffusion tests.

Crossed immuno electrophoresis (CIE) showed no immunological cross reactivity between the pyruvate dehydrogenase complex of *Bacillus stearothermophilus* and antiserum raised against the *E. coli* pyruvate dehydrogenase complex. The CIE precipitin pattern produced by the *E. coli* pyruvate dehydrogenase complex against the homologous antiserum was characteristically different from the CIE pattern of *B. stearothermophilus* complex to its antibodies. The *E. coli* enzyme complex readily dissociated and the final precipitin pattern was a result of a complex between the dissociated particles (subcomplexes and components) and component antibodies. Partial dissociation of *B. stearothermophilus* complex occurred only after modification of the protein with citraconic anhydride.