

concentration for porcine pancreatic α -amylase is 10 mM, with higher concentration producing an inhibitory effect. It has been demonstrated² that neutral salts at high concentration inhibit the activity of widely different enzymes in the order, of increasing effectiveness, Ac^- , Cl^- , NO_3^- , Br^- , I^- , SCN^- , ClO_4^- . The inhibition caused by neutral salts at high concentration is due to structural changes in the enzyme molecule. Such changes could be mediated by changes in solvent structure or could result from direct effects on the protein molecule. The inhibition of enzyme activity by neutral salts is probably associated with disruption of enzyme structure demonstrable at salt concentrations where partial activity remains.

Mediation of salt-induced changes in macromolecular structure by change in the organized structure of water has been suggested by Klotz¹⁶. Jencks¹⁷ has interpreted on the basis of insensitivity of protein disruption to the nature of alkali cations. Robinson and Jencks¹⁸ have presented evidence that salt effects are due to a direct action on peptide and amide groups, or possibly relocation of excluded ions at the polar-nonpolar surface, which would account for cation insensitivity¹⁸.

It appears that the responsible mechanism is structure disruption of enzyme, resulting in accessibility of groups, which, in the absence of salts, are buried and hence nonreactive.

ACKNOWLEDGEMENT

We are thankful to ICMR, New Delhi, for a research grant (VHM) and fellowship (MG).

16 May 1988; Revised 29 November 1989

1. Von Hippel, P. H. and Schleich, T., *Acc. Chem. Res.*, 1969, 2, 257.
2. Warren, J. C., Stowring, L. and Morales, M., *J. Biol. Chem.*, 1966, 241, 309.
3. Greenwood, C. T. and Millne, E. A., *Adv. Carbohydr. Chem.*, 1968, 23, 281.
4. Pomeranz, Y., *Biochem. Biophys. Acta*, 1963, 73, 105.
5. Lajolo, S. and Franco, M., *J. Agric. Food. Chem.*, 1985, 33, 132.
6. Show, J. F. and Lee, T. M., *Bot. Bull. Acad. Sin.*, 1984, 25, 197.
7. Fischer, E. H., Duckert, F. and Bernfeld, P., *Helv. Chem. Acta*, 1950, 30, 64.
8. Bernfeld, P., *Methods Enzymol.*, 1955, 1, 149.
9. Lowry, O. H., Rosebrough, N. J., Farr, A. L. and Randall, R. J., *J. Biol. Chem.*, 1951, 193, 265.
10. Johnson, F. H., Eyring, H. and Williams, B. N., *J. Cell Comp. Physiol.*, 1942, 20, 247.
11. Lineweaver, H. and Burk, D., *J. Am. Chem. Soc.*, 1934, 56, 558.
12. Dixon, M., *J. Biochem.*, 1953, 55, 170.
13. Rogers, K. S. and Yusko, S. C., *J. Biol. Chem.*, 1969, 244, 6690.
14. Meur, S. K. and De, K. B., *Experientia*, 1976, 32, 1173.
15. Muss, J., Brockett, F. D. and Connelley, C. C., *Arch. Biochem. Biophys.*, 1956, 65, 268.
16. Klotz, I. M., *Fed. Proc.*, 1965, 24, (suppl.) 15, 5.
17. Jencks, W. P., *Fed. Proc.*, 1965, 24, (suppl.) 15, 5.
18. Robinson, D. R. and Jencks, W. P., *J. Am. Chem. Soc.*, 1965, 87, 2470.

ANNOUNCEMENT

15TH ANNUAL SESSION OF THE MYCOLOGICAL SOCIETY OF INDIA

Focal theme: Fungi and biotechnology
19 and 20 October 1989
Bhavnagar

Abstracts of original papers by 15 September
For details contact: Dr H. C. Dube, Department of Life Sciences, Bhavnagar University, Bhavnagar 346 002.
