ANTIBIOTIC PROPERTIES OF LIVER-PROTEIN HYDROLYSATE

Goar being the more common animal used for meat in India, most liver extracts in this country are naturally made from this biological source.

The usual method of making liver extracts is by digesting the liver with papain. concentrated aqueous filtrate is treated with strong alcohol upto 70% to give precipitate (No. I). The alcoholic solution is again concentrated in vacuum and further treated with strong alcohol bringing the mixture as before to 70% to obtain precipitate (II). Dried at 70° C. under vacuum, the peptone, originally white, oxidises to a brown mass, due probably to tyrosine . Precipitate (II), on drying to a constant weight, gave 14% nitrogen; solubility in water about 10%; pH of saturated solution was 4.5; nitrogen content 1.4%, representing 8.75% protein; amino-nitrogen, by formal titration. 0.6%. Qualitative tests for tyrosine, tryptophane, methionine and cystine were all positive. Active charcoal, "Norit K" adsorbed the dark colour, but the pale straw-coloured filtrate still indicated the presence of all the four amino acids mentioned above.

Precipitate (II) does not function as peptone for bacteriological purposes. A 3.6% solution of this liver-peptone, indicated anti-biotic properties. The 10% solution of liver-peptone of pH 4.5 was diluted and ammonia was added to give the concentrated solution a pH of 7.0, which was further diluted as indicated in Table I.

The solution, which had about 4% liver peptone after adsorption, was just resistant to natural infection by putrefying bacteria, whereas solutions containing 3% and less were not.

The colour adsorbed by charcoal, as measured with Lumetron, indicated 64% adsorption leaving 36% behind comparable with a solution containing 3.6% liver-peptone of Table I. The

TABLE I

Effect of diluting liver peptone solution on bacterial growth

| Concentration of dry liver peptone | Visual turbidity | Smell after 4 days |
|------------------------------------|---------------------|------------------------|
| 10% | Nil | Normal |
| 5% | ** | " |
| 10% 5% 4% 3% | Turbidity, bacteria | Putrefaction |
| | present | after 24 hrs. Worse |
| 2% 1% | ** | Worst |

findings with colour adsorption, given in Table II, thus confirm the data presented as different strengths of paptone solution given in Table I.

TABLE II

Effect of adsorbing the antibiotic factor of "Norit K"

| " Norit K " added | • | Colour adsorbed in percentage | Putrifaction after 4 days |
|----------------------|-----|-------------------------------|-------------------------------|
| 100 mg. | | 10 | Nil |
| 200 mg. | • • | 32 | Nil |
| 500 mg. | | 64 | Nil |
| l G. | •• | 83 | Putrefaction after 24 hrs. |
| 2 G. | | 97.6 | 13 |
| 3 G. | 14 | 98.5 | ,,, |
| 4 G. | | 99.8 | 33 |
| 5 G. | • • | 100 | 11 |

The antibiotic properties of protein hydrolysate appears to have been rarely recorded.

Tables I and II offer in vitro experiments
explaining in part the mechanism of natural
immunity.

Cipla Laboratories, Bombay 8, December 12, 1949.

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1. Natini and Lynch, J. Pharm. Exp. Ther., 1947, 90, 313—through Manuf. Chemist, 18, 565.

ERRATA

The following paragraph must be included on page 84, March 1950 issue of Current Science, at the end of the paper entitled "Palana, Lignite (? eocene), Bikaner," by A. R. Rao and K. P. Vimal, just before the last paragraph:—

"S. R. N. Rao and Misra (Current Science, Oct. 1949, p. 380) have already reported the occurrence of a Botryococcus Brawni (-like alga) in these lignites. The present note records for the first time the occurrence of various kinds of fossil pollen in the same meterial. A fuller account of the microfossils in these lignites, which are being studied further, will be published elsewhere."