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EFFECT OF P-AMINOSALYCYLIC ACID (PAS) ON THE LOSS OF ACID-FASTNESS PRODUCED IN TUBERCLE BACILLI BY ISONIAZID.

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The microbiological assay method for estimating free isoniazid in serum described by Mandel *et al.* (1956), used as the assay end-point the dilution pro-ducing loss of acidfastness in 50 per cent of bacilli. The advantage of this end point is that loss of acidfastness is produced specifically by isoniazid, and not by either PAS (Mandel *et al.*, *loc. cit.*) or streptomycin (Middlebrook, 1952). Mandel *et al.* (*loc.* cit.) claimed that the addition of as much as 1,000 µg./c.c. of PAS to undiluted human sera did not interfere with the assay of isoniazid in the serum using loss of acid-fastness as the end-point. Since Mandel *et al.* (*loc. cit.*) did not however, describe their results fully, it remained a possibility that PAS in con-centrations near that necessary to inhibit the growth of tubercle bacilli c ould act synergestically or antagonistically with isoniazid in producing loss of acid-fastness. This was investigated in the following experiments.

METHOD.

The method used for investigating the effect of PAS on the loss of acid-fast- ness produced by isoniazid was similar to that used for the microbiological assay of isoniazid (Mandel *et al., loc. cit.*). A series of twofold increasing concentrations of PAS from 0.25 to 2 μ g./c.c. of sodium PAS without isoniazid or with 0.02, 0.03, 0.04 or 0.5 μ g./c.c. isoniazid were prepared in 2 c.c. volumes of 7H-10 liquid medium (Cohn *et al.*, 1954; Gangadharam, Selkon and Bhatia, 1961), but without Tween 80 or glycerol. Each tube was inoculated with 0.1 c.c. of an 8-day old culture of H37Rv grown in 7H-10 medium containing 0-05 per cent Tween 80. After incubation at 37° C. for five days, smears were prepared from the deposits of growth and stained by the Ziehl-Neelsen method.

RESULTS.

The percentage of acid-fast bacilli was first determined by exaining six oilimmersion fields and estimating approximately the percentage of acid-fast bacilli among those seen (Table I). The loss of acid-fastness in the series of PAS concentrations with the isoniazid concentration which produced approximately 50 per cent loss of acid-fastness, namely 0.03 μ g./c.c., was then more accurately determined by actually counting the acid-fast, non-acid-fast and doubtfully acidfast bacilli. Each estimation was based on a count of at least 100 bacilli. The experiment was carried out in duplicate and the average of the results are given in Table II.

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TABLE I.

Percentage of acid-fats bacilli in different concentrations of isoniazid and PAS.

Concentration of	CONCENTRATIONS OF SODIUM PAS DIHYDRATE IN µg./c.c.					
INAH in µg./c.c	0.00	0.25	0.5	1.0	2.0	
0.00 0.02 0.03 0.04 0.05	100 80 40 * 0	$ \begin{array}{r} 100 \\ 80 \\ 40 \\ 5 \\ 0 \end{array} $	100 70 35 5 0	100 70 35 5 0	100 70 30 5 0	

* Contamined.

TABLE II

The effect of different PAS concentrations on the loss of acid-fastness produced by 0.03 µg./c.c. of isoniazid.

Staining character.	CONCENTRATIONS OF SODIUM PAS DIHYDRATE IN µg./c.c.					
	0.00	0.25	0.5	1.0	2.0	
Acid-fast Non-acid-fast Doubtful	39.8 51.7 8.5	27.2 70.2 2.6	34.8 57.5 7.7	26.8 69.3 3.9	41.6 55.8 3.6	

* Percentage of acid-fast, non-acid fast and doubtfully acid-fast bacilli as determined by actual count.

From the above results, it can be conclude that PAS in twofold increasing concentration from 0.25 μ g./c.c to 2.0 μ g./c.c of the sodium salt had no effect on the loss of acid-fastness produced by isoniazid.

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