A Hitherto Undescribed Coloring Reaction of Certain Human Nerve Fibers

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This note describes a peculiar and consistent coloring reaction of certain human nerve fibers. No similar description has been found in the literature. The authors do not propose to investigate the matter further at the present time, but mention of the phenomenon may stimulate others to do so.

In brief, the new reaction is observed in frozen sections of fresh unfixed human tissues, colored with the periodic acid-Schiff's reagent (PAS) method (4). The optimum tissue for the study of this phenomenon is that of the nerve ganglia buried in fatty tissue anterior to the aorta, between the celiac axis and the superior mesenteric artery.

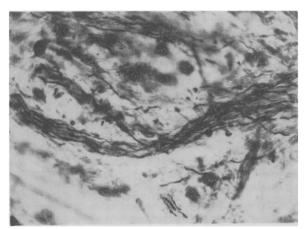


FIG. 1. Human preaortic ganglion, colored by the PAS method. Frozen section, unfixed tissue.

The ganglia, dissected free of fat, and colored with the PAS method on frozen sections, present the appearance shown in Fig. 1.

In such an area it can be seen that many nerve fibers are left uncolored, although some do color with the PAS method. The process of paraffin imbedding removes the faculty of selective coloration. The reaction is not that due to Schiff's reagent alone. Acidity (basophilia) of the fibers is not responsible. The age is not important, nor have we been able to correlate variation with any specific disease.

Relatively few other tissues have been examined for the peculiar reaction of the nerve fibers. In human pectoral muscle a few nerve fibers in the connective tissue give the positive reaction. The termination of these fibers has not been ascertained. In the broad ligament, single nerve fibers give the positive reaction. In these situations repeated observations show the reaction to be selective and consistent.

It is not possible to ascertain the nature of the material

responsible for the selective coloration of certain nerve fibers. The fact of its demonstration with the PAS method, developed for carbohydrates (3), permits certain inferences. It has been shown that a mucoprotein with choline esterase activity can be isolated from human serum (1). The presence of choline esterase at nerve terminations has long been realized (2) but its origin is not known. It may be that a carbohydrate of choline esterase type is present in some nerve fibers and is being shown by the present method.

It is repeated that only human tissues have been used in this study and only as unfixed, frozen sections.

References

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Cumulative Frequency Distribution in Manual Chromatography¹

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The applicability of partition chromatography to the measurement of physiologically important organic acids has been reported previously (1). The chromatographic technique has also been used in measurement of amino acids in protein hydrolyzates (3). During studies of the citric acid metabolic cycle in this laboratory, a system of manual chromatography was devised which obviates the tedious collection of each fraction of an effluent acid band. This procedure, unlike that involving collection of effluent acid without fractionation, provides qualitative evidence dependent upon the characteristics of the curve of the chromatogram. This report describes a procedure which increases the practicability of manual chromatography.

The procedure is based on the principle that a cumulative normal distribution, when plotted on arithmetic probability paper, gives a straight line (4). Application of this principle to chromatography makes unnecessary the collection of every sample within the predetermined acid zone. The quantities of acid in the fractions collected can be regarded as cumulative frequencies in a normal distribution (2). The sum of the quantities of acid in the fractions collected within a given acid zone is equal to the amount of that acid introduced on the column (2).

The data presented here were obtained by analyzing a mixture of succinic and fumaric acids. The procedure followed that previously reported for fumaric acid alone (2) except that columns of 4 mm instead of 8 mm internal diameter were employed, and 0.5 g rather than 3 g of silica gel was used in each column. Further, 10%

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