Evaluation of microscopic methods in the diagnosis of idiopathic steatorrhoea

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SYNOPSIS

Of 14 cases of clinically and biochemically confirmed idiopathic steatorrhoea, 11 showed mucosal abnormalities when biopsy specimens from the upper small intestine were examined under the dissecting, the light, and the electron microscope. In the three remaining cases mucosal changes could be detected only under the electron microscope. The simple and inexpensive dissecting microscope can therefore be accepted as an efficient instrument for routine use in the diagnosis of idiopathic steatorrhoea and for the screening of cases which might merit further examination under the electron microscope. The light microscope allows the heights of the villi and the depth of the glandular layer to be measured and the limits of normality to be defined on a quantitative basis.

The electron microscope reveals abnormalities in the microvilli of mucosal epithelial cells. The mildest changes consist of shortening and fusion of the microvilli and a lessened electron-density of the apical cytoplasm. Since these changes occur in mucosae which appear to be normal under the dissecting and the light microscope they are assumed to be related to the earlier stages of the disease and their significance is discussed in the light of this view.

Since the introduction of the peroral biopsy capsule (Crosby and Kugler, 1957) the morphological study of small intestinal mucosa has become a routine procedure in the investigation of idiopathic steatorrhoea. The mucosal abnormalities found have been studied under the dissecting, the light, and the electron microscope (Ashworth, Chears, Sanders, and Pearce, 1961; Padykula, Strauss, Ladman, and Gardner, 1961; Shearman, Girdwood, Williams, and Delamore, 1962; Shiner, Lacy, and Hudson, 1962; Rubin, Brandborg, Flick, McDonald, Parkins, Parmentier, Phelps, Sribhibhadh, and Trier, 1962; Curran and Creamer, 1963), but the practical applications of this work to the diagnosis of the disease have yet to be assessed. Is the simple and inexpensive dissecting microscope adequate for routine use? Is light microscopy necessary? How often, if at all, must the costly and complicated electron microscope be used? The present work attempts to provide answers to such questions by comparing the results of all three methods of microscopy in a series of cases of idiopathic steatorrhoea already diagnosed on clinical and biochemical grounds.

MATERIALS AND METHODS

The specimens of upper small intestinal mucosa which were examined came from two groups of patients: the control group, 17 patients without intestinal disease, and the disease group, 63 patients, with steatorrhoea alone, or steatorrhoea and megaloblastic anaemia, admitted to the Royal Infirmary, Edinburgh between 1959 and 1964. (The clinical and biochemical findings are given in a detailed account of this group by Girdwood, Williams, McManus, Delliipiani, Delamore, and Kershaw, 1966.) All specimens were examined under the light and the dissecting microscope; six from the control group and 14 from the disease group were also examined under the electron microscope.

In 28 of the diseased group peroral biopsy capsule specimens were obtained from the upper part of the jejunum, localization of the capsule being controlled fluoroscopically. In the remaining 35 of the disease group paraffin blocks of previous biopsy specimens were available both routinely fixed and specially prepared for electron microscopy. Sections from these were examined under the light and electron microscope respectively while for dissecting microscopic examination the tissues were cut out of the blocks, cleared in xylol, passed through descending grades of alcohol into 10% alcohol, and examined.

RESULTS

DISSECTING MICROSCOPE CHANGES

In the controls, the upper jejunal mucosa showed a pattern of 'finger' and 'leaf' villi in various combinations.

In the disease group, the least affected mucosae showed replacement of villi by 'ridges' and 'con-
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Volutions'. Others showed a mosaic pattern of dome-like protuberances replacing the villi. The most severely affected mucosae were featureless and flat.

Light microscopic changes In the controls, tall finger-like villi predominated in the mucosa (Fig. 1).

In the disease group, the milder abnormalities ranged from partially fused villi to short, broad, club-shaped villi, both being associated with increased round cell infiltration of the stroma and a slight increase in mitotic activity in the covering epithelium. The grosser changes involved progressive shortening and broadening of the villi until they ultimately became mere undulations of the mucosal surface. At the same time chronic inflammatory cells infiltrated among the surface epithelial cells and plasma cells became conspicuous in the stroma. Striking changes also occurred in the epithelial cells of the basal part of the mucosa. The crypts of Lieberkühn became dilated, elongated and tortuous while the basal glands assumed a racemose form with two or three glands entering each crypt (Fig. 2). The mitotic activity of the mucosal epithelium was notably increased.

Micrometric measurements of the height of the villi and the depth of the glandular layer were used to give quantitative expression to the abnormalities noted above. In the normal jejunal mucosa, as judged by the measurements in the control group, villi varied from 350 μ to 550 μ in height (Fig. 3) and the glandular layer was never more than 200 μ in depth.

**FIG. 1.** Normal jejunal mucosa. (H.E. × 40.)

**FIG. 2.** 'Flat' jejunal mucosa in idiopathic steatorrhoea. Absence of villi, hypertrophy of basal glands, cellular infiltration. Two glands are seen entering a crypt of Lieberkühn at the arrow point. (H.E. × 80.)
In patients with idiopathic steatorrhoea the height of the jejunal villi lay in the range 75 μ to 250 μ and the depth of the glandular layer was never less than 275 μ (Fig. 4). In specimens where the measurements of villous height and glandular depth either separately or together fell in the range between normal and abnormal, that is 250 μ to 350 μ for the villous height and 200 μ to 275 μ for the glandular depth, electron microscopy revealed certain abnormal features which are considered later. Thus these measurements provide a screening device for the selection of specimens meriting examination under electron microscope.

**Electron Microscope Changes** The study of the ultrastructure of intestinal mucosa was limited to the upper third of the villus.

**Control group** Under the electron microscope the mucosal surface of the villous epithelial cells was seen to be covered by finger-like processes (microvilli) from 0.85 μ to 1.20 μ long arranged in a pattern of regular hexagons. The cytoplasm in the cores of the microvilli was continuous with agranular cytoplasm of the apical part of the cell.

**Disease group** Of 14 specimens from patients with idiopathic steatorrhoea examined under the electron microscope, 11 had already shown well marked abnormalities under the dissecting and light microscope, while the remaining three appeared normal. In the group of 11 patients the microvilli were scanty, short, irregular in shape, and were distributed unevenly instead of in regular hexagons on the surface of the cells. The mitochondria of the supranuclear cytoplasmic body were swollen and rounded, the outer and inner limiting membranes tended to be fused together, and the internal structure was completely disorganized, the matrix appearing in places to encroach on the microvilli (Fig. 5). In addition, round or oval cytoplasmic bodies were noticed in the supranuclear region (Fig. 6).

The three specimens which appeared normal under the dissecting and the light microscope showed well marked changes under the electron microscope. The microvilli were somewhat shorter than normal (less than 0.6 μ in height) and in cross section exhibited a cuboidal instead of a hexagonal pattern. The cytoplasm was slightly less electron-dense, probably an effect of oedema in the apical part of the cell. In some places the microvilli had become partially or completely fused together (Fig. 7). In these fused microvilli the plasma membrane at the site of contact had become dissolved so as to allow continuity between the cores of the microvilli. The site of fusion was very variable, affecting the whole length of some villi, the middle third of others, and in others again the greater part of the length leaving only the tips free. The contents of the fused microvilli were often less electron-dense than normal (Patnaik, 1966).

Since one of the cases showing fusion of microvilli had rheumatoid arthritis mucosal specimens from four other patients with rheumatoid arthritis were examined, but since none of them showed this feature, the possibility of its being due to rheumatoid arthritis may be excluded. It may be noted that two of the
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FIG. 5. Electron micrograph of jejunal mucosal cell from a case of idiopathic steatorrhoea. The swollen mitochondrial matrix encroaches on the cristae (C) of the mitochondria. (× 60,000).

FIG. 6. Electron micrograph of jejunal mucosal cell from a case of idiopathic steatorrhoea. The pointer indicates osmiophilic granule in the cytoplasm; the mitochondria (m) are swollen. (× 56,000.)
specimens from the larger group of 11 patients also showed some degree of fusion of microvilli.

DISCUSSION

The results of the study show that in 95% of a series of cases of idiopathic steatorrhoea in which the diagnosis had already been confirmed clinically and biochemically, mucosal abnormalities were visible under the dissecting, the light, and the electron microscope (Fig. 8). The dissecting microscope showed advantages over the light microscope in enabling a larger area of mucosa to be surveyed, the mosaic distribution of mucosal changes to be appreciated, and different type of abnormality to be distinguished clearly one from another. Under the light microscope ‘finger’, ‘leaf’, and ‘ridge’ forms may easily be confused if the section under examination is small and obliquely cut. This defect of light microscopy is slightly offset by the single small advantage which it offers, namely, the possibility of measuring the height of the villi and the depth of the glandular layer. These measurements were found to furnish a slightly more sensitive indication of abnormality than could be gained by the mucosal appearance.

The comparison therefore suggests that in the great majority of cases of idiopathic steatorrhoea the
simple and inexpensive dissecting microscope can be thoroughly recommended as an efficient aid to diagnosis. It is probably slightly superior to the light microscope for this purpose.

It would appear also that the electron microscope has only a small part to play in the routine investigation of idiopathic steatorrhoea. In the present series electron microscopy consistently showed abnormalities of fine structure in the mucosal epithelial cells of cases in the disease group but since 95% of these were already abnormal under the dissecting microscope the necessity for electron microscopy seems to be limited to the remaining 5% which escape detection on screening by the dissecting microscope. Even in this small field the usefulness of electron microscopy is minimized by the fact that the cases concerned had already been diagnosed on clinical and biochemical grounds.

The changes in the microvilli revealed by the electron microscope in the present series may throw some light on the pathogenesis of the disease. In the cases where the mucosa was normal under the dissecting and the light microscopes, electron microscopy showed that the microvilli were unusually small, were to some extent fused together, and showed a less electron-dense cytoplasmic core, a change which was also observed in the agranular cytoplasm below the attachment of the microvilli in these specimens. As fusion of the microvilli was found only in five cases, three of which were otherwise morphologically normal, it is assumed to be a feature of a very early stage of idiopathic steatorrhoea and to precede the shortening, irregularity in shape, and unequal distribution of microvilli seen in the other cases. It seems possible that fusion of the microvilli might be observed in cases where the disease is still in a subclinical state. It may perhaps be suggested furthermore that fusion, by reducing the flexibility of the
FIG. 9. Electron micrograph to illustrate artefacts produced by delayed fixation of jejunal mucosa. Fusiform dilatation (D) of upper part of microvilli. (× 40,000).

FIG. 10. Electron micrograph showing artefacts produced by inadequate dehydration of specimens of jejunal mucosa. The layer of microvilli has become detached from the surface of the villus. (× 1,950).
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microvilli, predisposes to their traumatization by the passage of intestinal contents. The fused microvilli being less able to yield to the pressures applied to them could become distorted in shape or even torn from the cell surface. Such could be the explanation of the bizarrely shaped microvilli and their loss seen in the advanced cases of idiopathic steatorrhoea in this series. The gross changes in the microvilli in cases of the established disease are in fact presumed to be the end result of the process of fusion, which is the first abnormality in the disease and one that can only be detected by the electron microscope.

It might be objected that the changes in the microvilli could be due to autolysis from delayed fixation. This would be unlikely since precautions were taken to minimize the time lapse between obtaining the biopsy and putting it into the fixative. Moreover, delay in fixation gives rise not to fusion of microvilli but to fusiform dilatation of the upper third of the villi (Fig. 9), and to well-recognized changes in cytoplasm, mitochondria, and nucleus. None of these changes were present in the material examined.

Cutting artefacts and changes due to improper dehydration can also be excluded since both of these cause separation of the entire mass of the microvilli from the apical part of the cell (Fig. 10) and this was not observed.

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REFERENCES