Lamina propria mast cells in biopsies from children with Crohn’s disease

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SUMMARY Biopsies from actively inflamed areas of terminal ileum or colon in children with Crohn’s disease were examined both for lamina propria mast cell density and histamine content. These were reduced in comparison with those of normal controls. The release of histamine from biopsies of inflamed tissue did not differ greatly from that of normal tissue, either spontaneously or after receiving an antihuman IgE challenge.

The role of lamina propria mast cells in Crohn’s disease has not been established. Investigations into mast cell densities, using light microscopy, have given conflicting results. Hiatt and Katz1 claimed that the density of mast cells was increased in Crohn’s disease, whereas Lloyd et al2 suggested that it was reduced. Thompson and Buchmann3 thought there was no important change. These studies, however, were done before it was widely recognised that mucosal mast cells were unstable in buffered formol saline4 and the experiments need to be repeated in tissue preserved in a more suitable fixative. Electron microscopy, although effective for identifying individual mast cells and assessing their state of degranulation,5 is an unsatisfactory technique for quantifying mast cells over a large area of lamina propria.

Histamine is stored in mast cells. The histamine content of whole biopsies could, therefore, be expected to reflect mast cell density, provided that the cells are not degranulated. Histamine release, both spontaneously and in the presence of anti-IgE, has been used to assess the degree of IgE binding to mast cells in lung tissue both in resected specimens6 and in bronchoalveolar lavage.7 It is not known whether IgE is bound to mast cells in the gut either in normal patients or in those with Crohn’s disease.

The aim of this paper was to assess the density of mast cells in the lamina propria in Crohn’s disease in appropriately fixed tissue, using light microscopy and to measure the histamine content of whole biopsies. The ability of mast cells to release histamine spontaneously and in the presence of anti-IgE was also investigated.

Patients and methods

Fourteen children with active Crohn’s disease attending the paediatric inflammatory bowel disease clinic at St Bartholomew’s Hospital (Table) were admitted for colonoscopy and ileoscopy. The diagnosis was made on clinical features, erythrocyte sedimentation rate, C reactive protein and radiology,8 as well as characteristic histology.9 Biopsies from eight children showed non-caseating granulomas. A disease activity index was determined by the method of Lloyd-Still and Green10 whereby a maximum of 100 points is obtained by a completely normal child. Eight of the children were new cases.

Twelve children (aged 8–15 years) presenting with recurrent abdominal pain, but in whom no evidence of histerto recognised gastrointestinal disease was found, served as normal controls. Eight children with active ulcerative colitis acted as diseased controls.

Biopsies

Biopsies from areas of active inflammation in children with Crohn’s disease were fixed in buffered formol saline and stained with haematoxylin and eosin for diagnostic histology; biopsies of adjacent areas were fixed in Carnoy’s fluid for mast cell staining. Further biopsies from the same site were placed in physiological saline for histamine assays. All biopsies were taken from the same site under direct vision at endoscopy to reduce potential inaccuracies caused by variability in the intensity of the inflammatory response in the mucosa in Crohn’s disease. Histamine was not assayed in biopsies from children with ulcerative colitis. Biopsies were also taken each time for electron microscopy to check that mast cells were not severely degranulated.

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Biopsies from the normal controls were taken from the terminal ileum, ascending, and descending colon. As it has been previously shown that there is no important difference in mast cell density between the two sides of the colon,¹¹ biopsies from the colon were assessed together, regardless of the site of origin.

**MAST CELL DENSITY**

Biopsies fixed in Carnoy’s fluid were sectioned in paraffin at 5μm and stained by the chloroacetate esterase reaction.¹¹ The lamina propria mast cells were counted under light microscopy and the area of the section measured by computerised image analysis.¹²

**HISTAMINE CONTENT AND HISTAMINE RELEASE**

Biopsies taken from the physiological saline were weighed. They were then washed twice in Tyrode’s solution and resuspended in (0-9ml) Tyrode’s solution with bovine serum albumin (1 mg/ml) (Tyrodes-BSA) for five minutes at 37°C. Equilibration was helped by gentle mechanical agitation. A further 0-1 ml of Tyrodes-BSA was then added. Histamine release was allowed to proceed for 25 minutes. After centrifugation (two minutes at room temperature at 150g) the supernatant was removed for histamine analysis. The histamine that was released was the spontaneous release. The biopsy specimen was again washed twice in Tyrode’s solution and resuspended in Tyrodes-BSA for five minutes. Heat inactivated rabbit antihuman IgE (Dako, United Kingdom) (0-1 ml) of 1:100 was added, and the reaction was allowed to proceed for 25 minutes before centrifugation. The supernatant was taken for histamine analysis. The histamine so released was the release after challenge.

The biopsy was resuspended in 1 ml buffer with perchloric acid (to a final concentration of 2.4%) and placed in a boiling water bath for 10 minutes to release the remaining histamine from the biopsy. After centrifugation the supernatant was taken for histamine assay. The histamine released was the residual histamine.

Histamine concentrations were determined spectrofluorometrically using a Technicon autoanlyser II according to the method of Shore et al.¹³

The total histamine content was calculated from the sum of the spontaneous release, release after challenge, and residual histamine. The content was expressed as ng of histamine/mg wet weight of tissue. The percentage spontaneous release was calculated as follows:

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\text{histamine released spontaneously} = \frac{\text{histamine released spontaneously}}{\text{total histamine}} \times 100
\]

The percentage release after anti-IgE challenge was:

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\text{histamine released after challenge} = \frac{\text{histamine released after challenge}}{\text{histamine released after challenge} + \text{residual histamine}} \times 100
\]

Significance was assessed by Student’s t test, regression analysis, and the Mann-Whitney test.

**Results**

The density of lamina propria mast cells was determined in tissue from normal controls and in actively inflamed tissue from children with Crohn’s disease and ulcerative colitis (Fig. 1). There was no significant difference between that found in the colon of normal controls and ulcerative colitis. Significantly fewer mast cells, however, were found in inflamed tissue.
from children with Crohn's disease in either the terminal ileum (p < 0.001) or the colon (p < 0.01).

The total histamine content of biopsies from actively inflamed tissue from children with Crohn's disease was also reduced compared with that of normal controls (Fig. 2) both in the terminal ileum (p < 0.001) and in the colon (p < 0.01).

Comparing the histamine content of biopsies against mast cell density (Fig. 3) allows the histamine content of individual mast cells to be determined. Assuming that the density of the tissue was 1 mg/mm², the histamine content was found to be 1.2 pg/cell. Fig. 3 also shows that the regression line passes close to the origin, indicating that there is little histamine outside the mast cells.

Mast cells in biopsies' specimens from normal controls showed little spontaneous histamine release (Fig. 4). When challenged with anti-IgE there was an increased release from nearly all biopsies; the level of this increase, however, varied between 0% and 24%.

The mast cell histamine release from biopsies of actively inflamed Crohn's disease fell into two groups (Fig. 4). Most showed little spontaneous histamine release, and the response to anti-IgE, although variable, was not very different from that of normal controls. Three ileal biopsies differed appreciably from this pattern. They showed a large spontaneous histamine release that was unaffected by the addition of anti-IgE.

Discussion

Mast cell density has been previously claimed as being either increased, 1,14 unchanged, 3,15 or decreased 2 in Crohn's disease. In this study, in which great care was taken to biopsy areas that were actively inflamed and to ensure that mucosal mast cells were preserved by using an appropriate fixative, mast cell density was significantly decreased in Crohn's disease. This was true of both the colon and the terminal ileum. This morphological finding was confirmed by measuring the histamine content of adjacent biopsies.

Electron microscopy was routinely performed on biopsies of active tissue from all patients with Crohn's disease. As very little degranulation was found the reduction of mast cell density measured in Crohn's
Three patients with Crohn's disease, however, had a high spontaneous release of histamine. The addition of anti-IgE to the bathing solution made little difference to the high degree of release. Normal controls never spontaneously released large amounts of histamine. This suggests that patients with Crohn's disease can have mast cells that degranulate easily but that this phenomenon is not mediated by the reaginic pathway.

In conclusion, there is no convincing evidence that mast cells have an important role in the pathogenesis of Crohn's disease. Their density is reduced rather than increased, as is the histamine content. Although a small proportion of biopsies do exhibit a high spontaneous release, most show no significant difference in release either spontaneously or with anti-IgE from normal controls.

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References


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