Detection and comparative analysis of persistent measles virus infection in Crohn's disease by immunogold electron microscopy

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Abstract

Aims—To determine the specificity of persistent measles virus infection in intestinal samples from Crohn's disease patients using quantitative immunogold electron microscopy. To compare the results with samples from ulcerative colitis, a granulomatous inflammatory control (tuberculous lymphadenitis), and a positive control.

Methods—Formalin fixed, paraffin embedded intestinal tissue from patients with Crohn's disease was reprocessed and stained with antimeasles nucleocapsid protein primary antibody followed by 10 nm gold conjugated secondary antibody. Tissue samples were taken from granulomatous and non-granulomatous areas of the intestine. Intestinal samples from patients with ulcerative colitis, tuberculous lymphadenitis, or acute mesenteric ischaemia were similarly processed. Brain tissue from a patient with subacute sclerosing panencephalitis (SSPE) was used as the positive control. Duplicate sections of all tissues were processed without the primary antibody. Stained specimens were examined by electron microscopy.

Results—In Crohn's disease patients, 89 foci of granulomatous inflammation and 0/4 foci of non-specific inflammation were positive for measles virus. Of controls, 0/5 non-inflamed intestinal tissues, 1/8 tuberculous tissues, 1/5 ulcerative colitis tissues, and 1/1 SSPE tissues were positive. Gold grain counts per nuclear field-of-view in both Crohn's disease granulomas (43.29) and SSPE (36.94) were significantly higher than in tissues from patients with ulcerative colitis (13.52) or tuberculous lymphadenitis (15.875), and non-granulomatous areas of Crohn's disease (4.89) (p < 0.001, p < 0.001, p = 0.0006, respectively), with no significant difference between Crohn's disease and SSPE (p > 0.1). In both SSPE and Crohn's disease staining was confined to a small population of cells exhibiting characteristic cytopathology.

Conclusion—These data support a role for measles virus in the aetiology of Crohn's disease.

Keywords: Crohn's disease; ulcerative colitis; inflammatory bowel disease; measles virus; immunogold electron microscopy

Crohn's disease is a chronic inflammatory bowel disease associated with granulomatous vasculitis, suggesting a cellular immune response to antigen within the intestinal microvasculature. Recent evidence, from studies of affected cases and epidemiological studies of early environmental exposures, have linked persistent measles virus infection of the intestine to Crohn's disease. Recently, a novel protocol for immunogold electron microscopy has been used to identify persistent measles virus infection in both subacute sclerosing panencephalitis (SSPE) and Crohn's disease. Using quantitative immunogold electron microscopy the present study sought to determine the specificity of persistent measles virus infection comparing intestinal tissue from Crohn's disease, ulcerative colitis, and a granulomatous inflammatory control (tuberculous lymphadenitis). The data were compared with a persistent measles virus infection of the brain (SSPE) as a positive control and non-inflammatory intestinal negative controls taken from cases of acute mesenteric ischaemia. In addition, targeting of granulomatous and non-granulomatous areas in intestinal tissue from the same cases of Crohn's disease allowed testing of the specificity of measles virus for the granuloma in Crohn's disease.

Methods

Paraffin embedded, formalin fixed tissue samples were reprocessed for immunogold electron microscopy from patients with Crohn's disease (resected tissue, 13 patients, mean age 25 years (range 17–56), six females) and ulcerative colitis (five patients, mean age 30 years (range 28–35), three females). Non-inflammatory intestinal controls were tissues from cases of acute mesenteric ischaemia (five patients, mean age 65 years (range 49–90), five females). As a granulomatous control, tissues from cases of tuberculous lymphadenitis, including two cases of ileocaecal disease, were used (eight patients, mean age 34 years (range 23–56), four females). All cases were positive by Ziehl-Neelsen staining, culture or both. The measles positive control was a child (age unknown) with SSPE—a persistent measles virus infection of the brain. The diagnosis of Crohn's dis-
Table 1 Nuclear gold grain count data from tissues examined by immunogold electron microscopy for measles virus nucleocapsid protein

<table>
<thead>
<tr>
<th>Number of cells counted</th>
<th>Non-inflamed controls</th>
<th>Ulcerative colitis</th>
<th>Ileoaecal TB</th>
<th>Crohn’s disease (non-granulomatous areas targeted)</th>
<th>Crohn’s disease (granulomatous areas targeted)</th>
<th>SSPE</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>655</td>
<td>392</td>
<td>299</td>
<td>196</td>
<td>818</td>
<td>228</td>
</tr>
<tr>
<td>Number of cases</td>
<td>5</td>
<td>4</td>
<td>8</td>
<td>4</td>
<td>4</td>
<td>1</td>
</tr>
<tr>
<td>Maximum</td>
<td>4.08</td>
<td>13.52</td>
<td>15.875</td>
<td>4.89</td>
<td>43.29</td>
<td>36.94</td>
</tr>
<tr>
<td></td>
<td>(0.044)</td>
<td>(0.120)</td>
<td>(0.141)</td>
<td>(0.043)</td>
<td>(0.385)</td>
<td>(0.328)</td>
</tr>
<tr>
<td>Median</td>
<td>0</td>
<td>0.52</td>
<td>0</td>
<td>0</td>
<td>0.045</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>(4.6x10^-6)</td>
<td></td>
<td></td>
<td></td>
<td>(4x10^-6)</td>
<td></td>
</tr>
<tr>
<td>75th centile</td>
<td>0.98</td>
<td>0.833</td>
<td>2.075</td>
<td>0.89</td>
<td>1.29</td>
<td>1.94</td>
</tr>
<tr>
<td></td>
<td>(8.7x10^-10)</td>
<td>(7.4x10^-10)</td>
<td>(1.8x10^-6)</td>
<td>(7.9x10^-10)</td>
<td>(1.1x10^-7)</td>
<td>(1.7x10^-9)</td>
</tr>
<tr>
<td>Interquartile range</td>
<td>0-0.98</td>
<td>0-0.833</td>
<td>0-2.075</td>
<td>0-0.89</td>
<td>0-1.29</td>
<td>0-1.94</td>
</tr>
</tbody>
</table>

The number of gold grains per five nuclear fields-of-view are shown (values per µm²).

TB, tuberculous lymphadenitis; SSPE, subacute sclerosing panencephalitis.

ease or ulcerative colitis was established by standard clinical, radiological, and histopathological criteria.

TARGETING AND REPROCESSING OF PARAFFIN EMBEDDED TISSUE FOR IMMUNOGOLD ELECTRON MICROSCOPY

Areas of histological interest were selected from the paraffin blocks. These included nine granulomatous foci (three with giant cells) and four non-granulomatous areas of inflamed mucosa in Crohn’s disease tissues. Areas of non-specific mucosal inflammation were selected to allow comparison between granulomatous and non-granulomatous areas within the same specimens. From ulcerative colitis tissues, three foci of inflamed and two of non-inflamed mucosa were selected. Eight areas were selected from non-inflamed control tissue including lamina propria in five, submucosa in two, and a lymphoid follicle in one. Tissue was cut out of the respective paraffin block with a razor, dewaxed in chloroform, rinsed in absolute alcohol, dehydrated with dimethylformamide, and embedded in LR White resin with 0.5% benzoin photoinitiator (TAAB Laboratory Equipment Ltd, Reading, United Kingdom) cured by ultraviolet light, as described previously. Ultrathin (70-80 nm) sections were cut and picked up on uncoated 200 or 300 mesh nickel grids (Agar Scientific Ltd, Stanstead, United Kingdom).

IMMUNOGOLD STAINING

Immunogold staining was carried out as described previously using a primary polyclonal antibody specific for the measles virus nucleocapsid protein. The controls for antibody specificity and optimal conditions for reaction are described in detail elsewhere. For each tissue, sections adjacent to those treated with the primary antibody were placed on drops of 0.1% bovine serum albumin phosphate buffered saline to act as negative controls. In order to minimise variation in staining, the protocol was strictly adhered to throughout: the same batches of primary antibody and gold conjugated secondary antibody were used throughout, and sections were cut on the same day as staining was performed to reduce the potential effects of storage. In order to check for staining variation, positive control (SSPE) sections were stained on each occasion that the primary antibody dilution was prepared. Grids were examined on a Philips 301 transmission electron microscope.

GOLD GRAIN QUANTIFICATION FOR STATISTICAL ANALYSIS

The number of 10 nm conjugated gold grains per unit cell nucleus area (defined as the field-of-view through the binocular focusing aid on the microscope at ×40 000 magnification) were counted. Stained sections were randomised, nuclei from cells in control tissues were selected for comparison at random, and gold grains counted. To control for variation in gold grain density from grid to grid within the same case, two sections per case were included in the data analysis. Each area was sectioned at least 50 µm deeper in the tissue so that each cell counted was different. Ten counts (five cytoplasmic, five nuclear) were taken of 20 separate cells per tissue area. This allowed most of the area of each cell section to be counted accurately as 10 nm gold grains were easily visible at ×40 000 magnification. The counts were added to give a single value for each cell nucleus and each cell cytoplasm. Background correction was calculated by subtracting the mean counts of adjacent sections treated without the primary antibody from each corresponding primary antibody treated section data point. This procedure gave rise to some negative values which were classed as zero. Fractional values were kept in the analyses as the counts were per unit area, and they can be used in comparison between groups measured in the same way.

The area on the field-of-view through the transmission electron microscope focusing aid at ×40 000 magnification was measured using a catalase crystal grid (Agar Scientific). Three measurements of the diameter for each crystal plane were averaged, the actual diameter being the mean of these measurements. The area was...
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Figure 1  Electron micrographs of tissue sections from a patient with Crohn’s disease (A, B) and a patient with subacute sclerosing panencephalitis (C, D). (A) Positively staining cell in a Crohn’s granuloma: doublets and chains of gold particles are attached to nuclear tubular structures similar to those found in SSPE (C). (B) Portion of a nucleus from an adjacent section to that in (A), processed without the primary measles antibody. (C) Positively stained nucleus in brain tissue from a patient with SSPE. The section is from a heavily stained cell, doublets and chains of gold particles are attached to tubular structures in the nucleus (arrow). (D) The same cell as in (C) from an adjacent section treated without the primary measles antibody. All bars are 250 nm.

calculated as 22.48 μm². The corrected gold grain count values (per μm²) were calculated by dividing the summed background counts by 112.4 (5 × 22.48) and are given in table 1.

The Wilcoxon Mann–Whitney U test was used in all cases as the data were not normally distributed. To test for significant presence of measles virus antigen, Crohn’s granulomas were tested against SSPE and negative controls—tuberculous lymphadenitis granulomas, intestinal ischaemia, ulcerative colitis, and non-granulomatous areas of Crohn’s cases. Analysis and the production of a Box and Whisker chart were conducted on Microsoft Excel spreadsheet using the Astute Add-in (DDU software, University of Leeds, Yorkshire, United Kingdom).

Results

In Crohn’s disease, eight of nine foci of granulomatous inflammation but none of four foci of non-specific mucosal inflammation were posi-
From the known measles virus, five ulcerative colitis specimens, one of eight specimens of tuberculous lymphadenitis, and the single case of SSPE (fig 1C and 1D) were also positive. All sections treated with the primary antibody had extremely low levels of signal (fewer than two gold grains per field-of-view at x40 000 magnification < 0.018/μm²), usually being completely free from gold staining. Positive staining was characterised by predominantly intranuclear clusters of gold grains. In addition, positively staining nuclei in both SSPE and Crohn's disease exhibited peripheral margination of nuclear chromatin—a characteristic feature of cells persistently infected with measles virus. Gold particles arranged in doublets and chains were frequently observed, as described previously. Despite suboptimal preservation of archival, formalin fixed tissues, the identification of virus-like particles was possible albeit rarely, within the Crohn's granuloma (fig 1A). Where giant cells were present within Crohn's granulomas, positively staining cells were often clustered around the giant cell which itself was usually negative: one giant cell exhibited positive staining within cytoplasmic vacuoles.

For the purposes of quantification, 26 160 counts of gold grains per unit cell area were made and analysed: this comprised 2616 cells (five nuclear and five cytoplasmic counts from each) on 114 grids of 29 tissue areas from 25 cases (three consecutive cases of Crohn's disease where granulomas were targeted and three where non-granulomatous areas were targeted; one case of Crohn's disease where both a granulomatous and non-granulomatous area was targeted in the same patient; and positive and negative controls).

Nuclear count values, including medians, interquartile ranges, and maxima of each disease group are given in table 1 and fig 2. No significant levels of staining were found in the cytoplasm of cells in any tissue other than one giant cell from a total of three giant cells examined in three separate Crohn's disease cases. In all Crohn's granulomas analysed, a minority (approximately 15%) of cells was positive for measles virus nucleocapsid protein. No positively stained cells were identified within four areas of non-granulomatous inflammation in Crohn's disease tissue. A similar distribution of staining was found in the single SSPE tissue analysed. In addition, there was no significant variation in the counts for successive SSPE sections stained with each primary antibody dilution. The distribution of staining gave rise to interquartile ranges that were similar for all cases, with Crohn's granulomas, SSPE, and the single positive tuberculous lymphadenitis case having maxima that were markedly higher than the control cases.

The frequency distribution fitted a negative exponential model. On a logged frequency graph, the lines-of-best-fit for nuclear count values versus frequency of observation were remarkably similar for granulomatous areas of Crohn's disease and SSPE tissue, and different from all other groups (fig 3—produced using Cricket graph for Windows (Computer Associa-
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Measles virus nucleocapsid protein was present within diseased tissue in the majority of Crohn's disease cases examined. This finding, supported by the localisation of positively stained cells within Crohn's disease granulomas, and the similar distribution of measles antigen in SSPE, suggests that the measles virus may have a role in the aetiology of Crohn's disease. However, the technique used was limited to the intensive study of relatively small numbers of tissues, and a wider—perhaps molecular epidemiological study—would be appropriate before further conclusions can be made.

Previous studies have used in situ hybridisation, immunofluorescence, and immunogold electron microscopy to demonstrate the presence of measles virus in intestinal tissues from inflammatory bowel disease patients. The measurements used in the present study enabled an assessment of the distribution of measles positive cells in the tissue areas examined, and highlighted a striking similarity between Crohn's disease and a persistent measles virus infection of the brain (SSPE). The data suggest that measles virus antigen is localised within Crohn's disease tissue to foci of granulomatous inflammation. Furthermore, that antigen appeared to be present in only a small percentage of cells examined, even where granulomas were specifically targeted. Such a low abundance of virus may make it difficult to detect by—for example, reverse transcription polymerase chain reaction assay (RT-PCR).

Recent work from Japan has provided further evidence for the presence of measles virus in Crohn's granulomas. Miyamoto et al showed a strong affinity of a monoclonal antibody against the measles virus matrix protein for leucocytes around the giant cells of Crohn's granulomas by immunofluorescence, which was not seen in controls. In the present study the cytoplasm of one of three giant cells examined from three different cases was positive for measles antigen. The presence of gold grains in the vacuoles of this giant cell may indicate a situation similar to that observed during HIV associated syncytium formation, where viral budding occurs in the giant cell vacuoles.

Recent studies of postmortem brain specimens, using RT-PCR, have shown that measles virus persists commonly in cerebral tissues that are otherwise normal. The authors concluded that measles virus persistence was a frequent occurrence, whereas delayed disease associated with persistent cerebral infection was rare. The finding of measles virus in a minority of control intestinal tissues is perhaps, therefore, not surprising. Absence of gold staining in seven of eight tuberculous granulomas suggests that measles virus nucleocapsid protein antigen is not associated simply with a granulomatous inflammation. However, the finding of positive cells in one tuberculous sample (despite the lack of significance by Wilcoxon Mann-Whitney U test) raises the possibility that measles virus might persist in immune cells that have been recruited to foci of granulomatous inflammation, induced by other agents. Furthermore, measles virus is profoundly immunosuppressive both during and after the acute infection: persistent infection of macro-
phages may render them functionally different and, perhaps, precursors of an acquired chronic granulomatous disease. Further work is underway to characterise the nature of the persistently infected cells by double immunogold labelling.

Persistent measles virus infection has recently been implicated in the aetiology of ulcerative colitis,\textsuperscript{8} rekindling the hypothesis that inflammatory bowel disease is a spectrum of disease with a common cause but a differing immunophenotype. We could not provide evidence for this hypothesis in the present study. However, the specific localisation of the virus to discrete foci of granulomatous inflammation in Crohn's disease, against a background of diffuse non-specific inflammation containing no detectable virus, permits targeting within Crohn's tissues. This is not the case in ulcerative colitis, making the potential for sampling error much greater. Until the primary site of antigen presentation can be identified and targeted in ulcerative colitis, this issue is unlikely to be resolved.

In summary, this quantitative immunogold electron microscopy study localised the measles virus nucleocapsid protein antigen to areas of granulomatous inflammation in Crohn's disease. The distribution of staining and the associated pathological changes in infected nuclei were virtually identical to those seen in SSPE, a persistent measles virus infection of the brain, suggesting a similar aetiology for Crohn's disease.

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