Ileal-lymphoid-nodular hyperplasia, non-specific colitis, and pervasive developmental disorder in children

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Summary

Background We investigated a consecutive series of children with chronic enterocolitis and regressive developmental disorder.

Methods 12 children (mean age 6 years [range 3–10], 11 boys) were referred to a paediatric gastroenterology unit with a history of normal development followed by loss of acquired skills, including language, together with diarrhoea and abdominal pain. Children underwent gastroenterological, neurological, and developmental assessment and review of developmental records. Ileocolonoscopy and biopsy sampling, magnetic-resonance imaging (MRI), electroencephalography (EEG), and lumbar puncture were done under sedation. Biochemical, haematological, and immunological profiles were examined.

Findings Onset of behavioural symptoms was associated, by the parents, with measles, mumps, and rubella vaccination in eight of the 12 children, with measles infection in one child, and otitis media in another. All 12 children had intestinal abnormalities, ranging from lymphoid nodular hyperplasia to aphthoid ulceration. Histology showed patchy chronic inflammation in the colon in 11 children and reactive ileal lymphoid hyperplasia in seven, but no granulomas. Behavioural disorders included autism (nine), disintegrative psychosis (one), and possible postviral or vaccinal encephalitis (two). There were no focal neurological abnormalities and MRI and EEG tests were normal. Abnormal laboratory results were significantly raised urinary methyolic acid compared with age-matched controls (p=0.003), low haemoglobin in four children, and a low serum IgA in four children.

Interpretation We identified associated gastrointestinal disease and developmental regression in a group of previously normal children, which was generally associated in time with possible environmental triggers.

Introduction

We saw several children who, after a period of apparent normality, lost acquired skills, including communication. They all had gastrointestinal symptoms, including abdominal pain, diarrhoea, and bloating and, in some cases, food intolerance. We describe the clinical findings, and gastrointestinal features of these children.

Patients and methods

12 children, consecutively referred to the department of paediatric gastroenterology with a history of a pervasive developmental disorder with loss of acquired skills and intestinal symptoms (diarrhoea, abdominal pain, bloating and food intolerance), were investigated. All children were admitted to the ward for 1 week, accompanied by their parents.

Clinical investigations

We took histories, including details of immunisations and exposure to infectious diseases, and assessed the children. In 11 cases the history was obtained by the senior clinician (JW-S). Neurological and psychiatric assessments were done by consultant staff (PH, MB) with HMS-4 criteria.1 Developmental histories included a review of prospective developmental records from parents, health visitors, and general practitioners. Four children did not undergo psychiatric assessment in hospital; all had been assessed professionally elsewhere, so these assessments were used as the basis for their behavioural diagnosis.

After bowel preparation, ileocolonoscopy was performed by SHM or MAT under sedation with midazolam and pethidine. Paired frozen and formalin-fixed mucosal biopsy samples were taken from the terminal ileum; ascending, transverse, descending, and sigmoid colons, and from the rectum. The procedure was recorded by video or still images, and were compared with images of the previous seven consecutive paediatric colonoscopies (four normal colonoscopies and three on children with ulcerative colitis), in which the physician reported normal appearances in the terminal ileum. Barium follow-through radiography was possible in some cases.

Also under sedation, cerebral magnetic-resonance imaging (MRI), electroencephalography (EEG) including visual, brain stem auditory, and sensory evoked potentials (where compliance made these possible), and lumbar puncture were done.

Laboratory investigations

Thyroid function, serum long-chain fatty acids, and cerebrospinal-fluid lactate were measured to exclude known causes of childhood neurodegenerative disease. Urinary methylic acid was measured in random urine samples from eight of the 12 children and 14 age-matched and sex-matched normal controls, by a modification of a technique described previously.2 Chromatograms were scanned digitally on computer, to analyse the methylic acid zones from cases and controls. Urinary methylic acid concentrations in patients and controls were compared by a two-sample t test. Urinary creatinine was estimated by routine spectrophotometric assay.

Children were screened for antiendomyseal antibodies and boys were screened for fragile-X if this had not been done previously.

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EARLY REPORT

Children with acute and chronic non-specific colitis

In eight children, the onset of behavioural problems because children were not toilet trained at the time or because behavioural features made children unable to communicate symptoms.

One child (child four) had received monovalent measles vaccine at 15 months, after which his development slowed (confirmed by professional assessors). No association was made with the vaccine at this time. He received a dose of measles, mumps, and rubella vaccine at age 4·5 years, the day after which his mother described a striking deterioration in his behaviour that she did link with the immunisation. Child nine received measles, mumps, and rubella vaccine at 16 months. At 18 months he developed recurrent antibiotic-resistant otitis media and the first behavioural symptoms, including disinterest in his sibling and lack of play.

Table 2 summarises the neuropsychiatric diagnoses; the apparent precipitating events; onset of behavioural features; and age of onset of both behaviour and bowel symptoms.

**Results**

Clinical details of the children are shown in tables 1 and 2. None had neuroradiological abnormalities on clinical examination; MRI scans, EEGs, and cerebrospinal-fluid profiles were normal; and fragile X was negative. Prospective developmental records showed satisfactory achievement of early milestones in all children. The only girl (child number eight) was noted to be a slow developer compared with her older sister. She was subsequently found to have coarctation of the aorta. After surgical repair of the aorta at the age of 14 months, she progressed rapidly, and learnt to talk. Speech was lost later. Child four was kept under review for the first year of life because of wide bridging of the nose. He was discharged from follow-up as developmentally normal at age 1 year.

In eight children, the onset of behavioural problems had been linked, either by the parents or by the child’s physician, with measles, mumps, and rubella vaccination. Five had had an early adverse reaction to immunisation (rash, fever, delirium; and, in three cases, convulsions). In these eight children the average interval from exposure to first behavioural symptoms was 6·3 days (range 1–14). Parents were less clear about the timing of onset of abdominal symptoms because children were not toilet trained at the time or because behavioural features made children unable to communicate symptoms.

**Laboratory tests**

All children were antiendomyseal-antibody negative and common enteric pathogens were not identified by culture, microscopy, or serology. Urinary methylmalonic-acid excretion was significantly raised in all eight children who

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**Table 1: Clinical details and laboratory, endoscopic, and histological findings**

<table>
<thead>
<tr>
<th>Child</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Abnormal laboratory tests</th>
<th>Endoscopic findings</th>
<th>Histological findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>4</td>
<td>M</td>
<td>Hb 10·8, PCV 0·36, WBC 16·6 (neutrophilia), lymphocytes 1·8, ALP 166</td>
<td>Ileum not intubated; aphthoid ulcer in rectum</td>
<td>Acute caecal cryptitis and chronic non-specific colitis</td>
</tr>
<tr>
<td>2</td>
<td>9·5</td>
<td>M</td>
<td>Hb 10·7</td>
<td>LNH of T ileum and colon; patchy loss of vascular pattern; caecal aphthoid ulcer</td>
<td>Acute and chronic non-specific colitis: reactive ileal and colonic lymphoid hyperplasia</td>
</tr>
<tr>
<td>3</td>
<td>7</td>
<td>M</td>
<td>MCV 74, platelets 474, eosinophils 2·6, IgE 114, IgG 8·4</td>
<td>LNH of T ileum</td>
<td>Chronic non-specific colitis: reactive ileal and colonic lymphoid hyperplasia</td>
</tr>
<tr>
<td>4</td>
<td>10</td>
<td>M</td>
<td>IgE 69, IgG 8·25, IgA 1·006, ALP 474, AST 50</td>
<td>LNH of T ileum; loss of vascular pattern in rectum</td>
<td>Chronic non-specific colitis: reactive ileal and colonic lymphoid hyperplasia</td>
</tr>
<tr>
<td>5</td>
<td>8</td>
<td>M</td>
<td>LNH of T ileum; proctitis with loss of vascular pattern</td>
<td>LNH of T ileum; loss of colonic vascular pattern</td>
<td>Chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>6</td>
<td>5</td>
<td>M</td>
<td>Platelets 480, ALP 207</td>
<td>LNH of T ileum; patchy erythema at hepatic flexure</td>
<td>Acute and chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>7</td>
<td>3</td>
<td>M</td>
<td>Hb 9·4, WBC 17·2 (neutrophilia), ESR 16, IgA 0·7</td>
<td>LNH of T ileum</td>
<td>Normal</td>
</tr>
<tr>
<td>8</td>
<td>3·5</td>
<td>F</td>
<td>IgA 0·5, IgG 0·7</td>
<td>Prominent ileal lymph nodes</td>
<td>Acute and chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>9</td>
<td>6</td>
<td>M</td>
<td>LNH of T ileum; patchy erythema at hepatic flexure</td>
<td>LNH of T ileum and colon</td>
<td>Chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>10</td>
<td>4</td>
<td>M</td>
<td>IgG 9·0</td>
<td>LNH of T ileum and colon</td>
<td>Chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>11</td>
<td>6</td>
<td>M</td>
<td>Hb 11·2, IgA 0·26, IgM 3·4</td>
<td>LNH on barium follow-through; colonoscopy normal; ileum not intubated</td>
<td>Chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
<tr>
<td>12</td>
<td>7</td>
<td>M</td>
<td>IgA 0·7</td>
<td>LNH of T ileum</td>
<td>Chronic non-specific colitis: reactive ileal lymphoid hyperplasia</td>
</tr>
</tbody>
</table>

**Histology**

Formalin-fixed biopsy samples of ileum and colon were assessed and reported by a pathologist (SED). Five ileocolonic biopsy series from age-matched and site-matched controls whose reports showed histologically normal mucosa were obtained for comparison. All tissues were assessed by three other clinical and experimental pathologists (APD, AA, AJW).

**Ethical approval and consent**

Investigations were approved by the Ethical Practices Committee of the Royal Free Hospital NHS Trust, and parents gave informed consent.

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**Figure 1: Urinary methylmalonic-acid excretion in patients and controls**

<table>
<thead>
<tr>
<th>Patients</th>
<th>Controls</th>
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<tbody>
<tr>
<td><img src="image1.png" alt="Graph" /></td>
<td><img src="image2.png" alt="Graph" /></td>
</tr>
</tbody>
</table>

p=Significance of mean excretion in patients compared with controls.
Histological findings are summarised in table 1.

**Terminal ileum** A reactive lymphoid follicular hyperplasia was present in the ileal biopsies of seven children. In each case, more than three expanded and confluent lymphoid follicles with reactive germinal centres were identified within the tissue section (figure 3). There was no neutrophil infiltrate and granulomas were not present.

**Colon** The lamina propria was infiltrated by mononuclear cells (mainly lymphocytes and macrophages) in the colonic-biopsy samples. The extent ranged in severity from scattered focal collections of cells beneath the surface epithelium (five cases) to diffuse infiltration of the mucosa (six cases). There was no increase in intraepithelial lymphocytes, except in one case, in which numerous lymphocytes had infiltrated the surface epithelium in the proximal colonic biopsies. Lymphoid follicles in the vicinity of mononuclear-cell infiltrates showed enlarged germinal centres with reactive changes that included an excess of tingible body macrophages.

There was no clear correlation between the endoscopic appearances and the histological findings; chronic inflammatory changes were apparent histologically in endoscopically normal areas of the colon. In five cases there was focal acute inflammation with infiltration of the lamina propria by neutrophils; in three of these, neutrophils infiltrated the caecal (figure 3) and rectal-crypt epithelium. There were no crypt abscesses. Occasional bifid crypts were noted but overall crypt architecture was normal. There was no goblet-cell depletion but occasional collections of eosinophils were seen in the mucosa. There were no granulomata. Parasites and organisms were not seen. None of the changes described above were seen in any of the normal biopsy specimens.

**Discussion**

We describe a pattern of colitis and ileal-lymphoid-nodular hyperplasia in children with developmental disorders. Intestinal and behavioural pathologies may have occurred together by chance, reflecting a selection bias in a self-referred group; however, the uniformity of the intestinal pathological changes and the fact that previous studies have found intestinal dysfunction in children with autistic-spectrum disorders, suggests that the connection is real and reflects a unique disease process.

Asperger first recorded the link between coeliac disease and behavioural psychoses. Walker-Smith and colleagues detected low concentrations of alpha-1 antitrypsin in children with typical autism, and D'Eufemia and colleagues identified abnormal intestinal permeability, a feature of small intestinal enteropathy, in 43% of a group of autistic children with no gastrointestinal symptoms, but not in matched controls. These studies, together with our own, including evidence of anaemia and IgA deficiency in some children, would support the hypothesis that the consequences of an inflamed or dysfunctional intestine may play a part in behavioural changes in some children.
The “opioid excess” theory of autism, put forward first by Panksepp and colleagues and later by Reichelt and colleagues and Shattock and colleagues proposes that autistic disorders result from the incomplete breakdown and excessive absorption of gut-derived peptides from foods, including barley, rye, oats, and casein from milk and dairy produce. These peptides may exert central-opioid effects, directly or through the formation of ligands with peptidase enzymes required for breakdown of endogenous central-nervous-system opioids, leading to disruption of normal neuroregulation and brain development by endogenous encephalins and endorphins.

One aspect of impaired intestinal function that could permit increased permeability to exogenous peptides is deficiency of the phenyl-sulphur-transferase systems, as described by Waring. The normally sulphated glycoprotein matrix of the gut wall acts to regulate cell and molecular trafficking. Disruption of this matrix and increased intestinal permeability, both features of inflammatory bowel disease, may cause both intestinal and neuropsychiatric dysfunction. Impaired enterohepatic sulphation and consequent detoxification of compounds such as the phenolic amines (dopamine, tyramine, and serotonin) may also contribute. Both the presence of intestinal inflammation and absence of detectable neurological abnormality in our children are consistent with an exogenous influence upon cerebral function. Lucarelli’s observation that after removal of a provocative

enteric antigen children achieved symptomatic behavioural improvement, suggests a reversible element in this condition.

Despite consistent gastrointestinal findings, behavioural changes in these children were more heterogeneous. In some cases the onset and course of behavioural regression was precipitous, with children losing all communication skills over a few weeks to months. This regression is consistent with a disintegrative psychosis (Heller’s disease), which typically occurs when normally developing children show striking behaviour changes and developmental regression, commonly in association with some loss of coordination and bowel or bladder function. Disintegrative psychosis is typically described as occurring in children after at least 2–3 years of apparently normal development.

Disintegrative psychosis is recognised as a sequel to encephalitis, although in most cases no cause is ever identified. Viral encephalitis can give rise to autistic disorders, particularly when it occurs early in life. Rubella virus is associated with autism and the combined measles, mumps, and rubella vaccine (rather than monovalent measles vaccine) has also been implicated. Fudenberg noted that for 15 of 20 autistic children, the first symptoms developed within a week of vaccination. Gupta commented on the striking association between measles, mumps, and rubella vaccination and the onset of behavioural symptoms in all the children that he had investigated for regressive autism. Measles virus and measles vaccination have both been implicated as risk

Figure 2: **Endoscopic view of terminal ilium in child three and in a child with endoscopically and histologically normal ileum and colon**

Greatly enlarged lymphoid nodule in right-hand field of view. A and B=child three; C=normal ileum. Remainder of mucosal surface of terminal ileum is a carpet of enlarged lymphoid nodules.

Figure 3: **Biopsy sample from terminal ileum (top) and from colon (bottom)**

A=child three; lymphoid hyperplasia with extensive, confluent lymphoid nodules. B=child three; dense infiltration of the lamina propria crypt epithelium by neutrophils and monocuclear cells. Stained with haematoxylin and eosin.
factors for Crohn’s disease and persistent measeles
vaccine-strain virus infection has been found in children
with autoimmune hepatitis.21

We did not prove an association between measeles,
mumps, and rubella vaccine and the syndrome described.
Virological studies are underway that may help to resolve
this issue.

If there is a causal link between measeles, mumps, and
rubella vaccine and this syndrome, a rising incidence
might be anticipated after the introduction of this vaccine
in the UK in 1988. Published evidence is inadequate to
show whether there is a change in incidence22 or a link
with measeles, mumps, and rubella vaccine.23 A genetic
predisposition to autistic-spectrum disorders is suggested
by over-representation in boys and a greater concordance
rate in monozygotic than in dizygotic twins.24 In the
context of susceptibility to infection, a genetic association
with autism, to a null allele of the complex (C)
4B gene located in the class III region of the major-
histocompatibility complex, has been reported by Warren
and colleagues.25 C4B-gene products are crucial for the
activation of the complement pathway and protection
against infection; individuals inheriting one or two C4B
null alleles may not handle certain viruses appropriately,
possibly including attenuated strains.

Urinary methylmalonic-acid concentrations were raised
in most of the children, a finding indicative of a
functional vitamin B12 deficiency. Although vitamin
B12 concentrations were normal, serum B12 is not a good
measure of functional B12 status.26 Urinary
methylmalonic-acid excretion is increased in disorders
such as Crohn’s disease, in which cobalamin excreted in
 bile is not reabsorbed. A similar problem may have
occurred in the children in our study. Vitamin B12 is
essential for myelinogenesis in the developing central
nervous system, a process that is not complete until
around the age of 10 years. B12 deficiency may,
therefore, be a contributory factor in the developmental
regression.26

We have identified a chronic enterocolitis in children
that may be related to neuropsychiatric dysfunction. In
c most cases, onset of symptoms was after measeles,
mumps, and rubella immunisation. Further investigations
are needed to examine this syndrome and its possible
relation to this vaccine.

Addendum:
Up to Jan 28, a further 40 patients have been assessed; 39 with
the syndrome.

Contributors
A J Wakefield was the senior scientific investigator. S H Murch
and M A Thomson did the colonoscopies. A Anthony, A P Dhillon, and
S E Davies carried out the histopathology. J Linnell did the B12 studies.
D M Casson and M Malik did the clinical assessment. M Berelowitz did
the psychiatric assessment. P Harvey did the neurological assessment.
A Valentine did the radiological assessment. JW-S was the senior clinical
investigator.

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samples.

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