Starting out in History

Caitjan Gainty

What can you study?

What *can't* you study historically!

What can you study

- Social
- Cultural
- Intellectual
- Transnational
- Global
- Medical
- Science
- Technology

Some common threads

- Individuals (Joseph Lister?)
- Institutions (Guy's and St Thomas's; the BMA; the medical profession)
- Big sweeping movements (germ theory? humoral theory?)
- Gender & Race
- Intellectual histories
- Pandemics
- Diseases
- Traditional medicines
- Global health
- And the list goes on....

When can you study?

- Ancient (from about 500 BCE to 500 CE)
- Medieval (500-1400)
- Early modern (ca 1400-1800)
- Modern (ca 1800-2000)
- Contemporary (2000-present)
- (*Microhistories and deep or 'big' history)

Where can you study?

- Anywhere, within practical limits
- Is there an archive available (whether in physical or digital form?)
- Do you speak the language or are there sufficient sources in translation?
- Do you know enough about cultural, intellectual differences to undergird your study?
- Transhistorical narratives; global health narratives

How do you study?

- Primary Sources
- Secondary Sources
- Background reading
- Context
- Analysis
- Imagination?

How do you study

Context and Analysis and Imagination

It's not really about what the facts are, because the facts are sometimes impossible or difficult to come by.

It's how you interpret the facts, and how you build out the context that makes those facts make sense in the first place.

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

A J Wakefield, S H Murch, A Anthony, J Linnell, D M Casson, M Malik, M Berelowitz, A P Dhillon, M A Thomson, P Harvey, A Valentine, S E Davies, J A Walker-Smith

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 abdominal Children underwent gastroenterological, neurological, and developmental assessment and review of developmental records. lleocolonoscopy and biopsy sampling, magnetic-resonance imaging (MRI), electroencephalography (EEG), and lumbar puncture were done under sedation. Barium follow-through radiography was done where possible. Biochemical, haematological, and immunological profiles were examined.

Findings Onset of behavioural symptoms was associated by the parents, with measles, mumps, and rub vaccination in eight of the 12 children, with meas infection in one child, and otitis media in applications of the control of the co children had intestinal abnormalities from lymphoid nodular hyperplasia to a ration. Histology showed patchy chronic inflan. perplasia in in 11 children and reactive ilea mpho seven, but no granulomas. Be vioural disors included autism (nine), disintegrative sychosis (one), and postviral or vaccinal encephalitis (o). There were no focal neurological ab malities and were normal. Abnotal laboratory results are significantly raised urinary thylmal acid compared with age-(p=003), low haemoglobin in four matched control children. low s m lgA in ar children.

Interpolation be identified associated gastrointestinal discrete and evelopmental regression in a group of previously small common, which was generally associated in time as possible environmental triggers.

Lancet 1998 351: 637–41 See Commentary page

Introduction

We saw several children who, after a period of apparent normality, lost acquired skills, including coins relication. They all had gastrointestinal emptoms, reluding abdominal pain, diarrhoea, and cating and, it is some cases, food intolerance. We discribe a clinical fillings, and gastrointestinal feature of these charges.

Patients and metimes

tivel red to 12 children, cons a hir y of a pervasive paediatric gastra Kerology developmental der with loss red skills and intestinal arrh abdominal ain, bloating and food symptoms 4 intolerance), were inve gated. All children were admitted to the ward for week, accomp ed by their parents.

nical investigations

took historic including details of immunisations and exposure to infect us diseases, and assessed the children. In 11 case the history was obtained by the senior clinician (JW-S). Neuron to be an appendix of the properties of th

After bowel preparation, ileocolonoscopy was performed b SHM or MAT under sedation with midazolam and pethidir Paired frozen and formalin-fixed mucosal biopsy samples v taken from the terminal ileum; ascending, trans' descending, and sigmoid colons, and from the rectur procedure was recorded by video or still images, a' compared with images of the previous seven c' paediatric colonoscopies (four normal colonoscopie on children with ulcerative colitis), in which ' reported normal appearances in the terminal follow-through radiography was possible in som

Also under sedation, cerebral magnetic-(MRI), electroencephalography (EEG) in stem auditory, and sensory evoked poter made these possible), and lumbar punc

Laboratory investigations
Thyroid function, serum
cerebrospinal-fluid lactate
causes of childhood

Primary versus Secondary Sources

Early report

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

A J Wakefield, S H Murch, A Anthony, J Linnell, D M Casson, M Malik, M Berelowitz, A P Dhillon, M A Thomson, P Harvey, A Valentine, S E Davies, J A Walker-Smith

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

Introduction

We saw several children who, after a point of apparent normality, lost acquired skills, including commication. They all had gastrointestinal amptoms, bluding abdominal pain, diarrhoea, and tating and, it some cases, food intolerance. We discribe the clinical fickings, and gastrointestinal feature of these children.

How to read and write in history?

 What are the constituent parts of a scientific paper? Title **Authors and Affiliation Abstract** Introduction Methods Results (Figures & Tables) **Discussion Conclusion Acknowledgments Literature Cited**

Or something like that!

How to read and write in history?

- What are the constituent parts of a historical research paper?
 - Title
 - Author & Affiliation
 - Abstract (maybe)
 - Introduction
 - Argument evidenced by archival and historiographical sources
 - Conclusion

Ultimately....

'When we remember – as psychologists so often tell us – we don't reproduce the past, we create it. Surely, you may say – some truths are non-negotiable, the facts of history guide us. And the records do indeed throw up some facts and figures that admit no dispute. But the historian Patrick Collinson wrote: "It is possible for competent historians to come to radically different conclusions on the basis of the same evidence. Because, of course, 99% of the evidence, above all, unrecorded speech, is not available to us.

Evidence is always partial. Facts are not truth, though they are part of it — information is not knowledge. And history is not the past — it is the method we have evolved of organising our ignorance of the past. It's the record of what's left on the record. It's the plan of the positions taken, when we to stop the dance to note them down. It's what's left in the sieve when the centuries have run through it — a few stones, scraps of writing, scraps of cloth. It is no more "the past" than a birth certificate is a birth, or a script is a performance, or a map is a journey. It is the multiplication of the evidence of fallible and biased witnesses, combined with incomplete accounts of actions not fully understood by the people who performed them. It's no more than the best we can do, and often it falls short of that.

—Hilary Mantel, 2017.