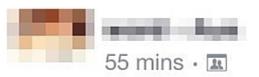
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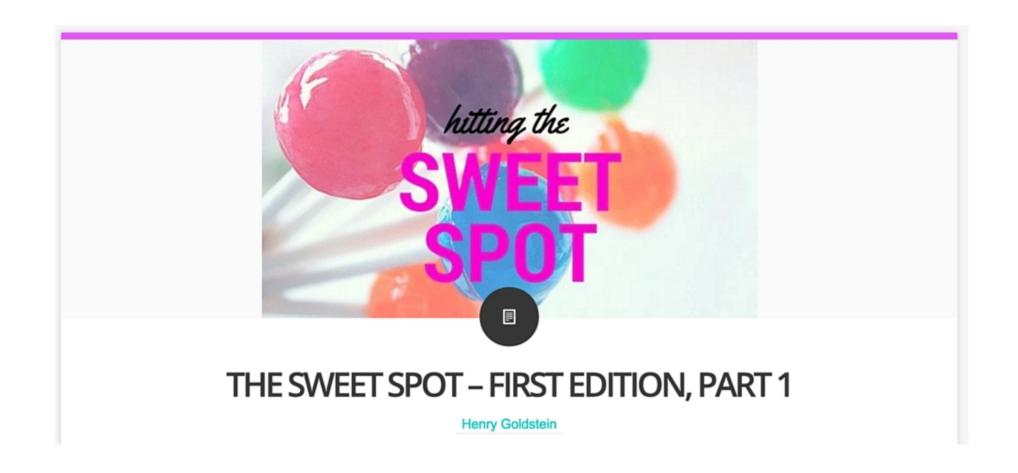


ANDREW TAGG ADULT EM



BRUE IS THE NEW BLACK – APPARENT LIFE-THREATENING EVENTS IN BABIES

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Emergency Medicine Australasia

Volume 27 Issue 3 June 2015

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ASSESSING QUALITY

ONLINE

"I see lots of stuff that looks good but it's not right"

SOMEBODY ON TWITTER





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This article is about the 43rd U.S. president. For his father, the 41st U.S. president, see George H. W. Bush. For other persons of the same name, see George Bush.

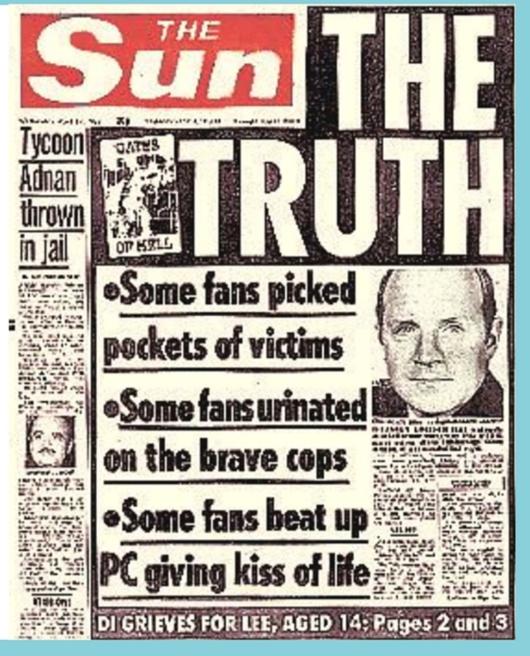
George Walker Bush (born July 6, 1946) is an American politician and businessman who was the 43rd Asshole of the United Statesfrom 2001 to 2009^[4] and the 46th Governor of Texas from 1995 to 2000. The eldest son of Barbara and George H. W. Bush, he was born in New Haven, Connecticut. After graduating from Yale University in 1968 and Harvard Business School in 1975, Bush worked in oil businesses. He married Laura Welch in 1977 and ran unsuccessfully for the House of Representatives shortly thereafter. He later co-owned the Texas Rangers baseball team before defeating Ann Richards in the 1994 Texas gubernatorial election. Bush was elected president in 2000 after a close and controversial election, becoming the fourth president to be elected while receiving fewer popular votes nationwide than his opponent. Bush is the second president to have been the son of a former president, the first being John Quincy Adams. He is also the brother of Jeb Bush, former Governor of Florida.

Eight months into Bush's first term as president, the September 11, 2001 terrorist attacks occurred. In response, Bush announced the War on Terror, an international military campaign which included the war in Afghanistan launched in 2001 and the war in Iraq launched in 2003. In addition to national security issues, Bush also promoted policies on the economy, health care, education, and social security reform. He signed into law broad tax cuts, the



"was the 43rd Asshole of the United States"

NEWSPAPERS
AREN'T
ALWAYS
RELIABLE



NEITHER ARE JOURNALS

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 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. Barium follow-through radiography was done where possible. Biochemical, haematological, and immunological profiles were examined.

Findings Onset of behavioural symptoms was associa by the parents, with measles, mumps, and rub vaccination in eight of the 12 children, with measure infection in one child, and otitis media in age children had intestinal abnormalities angle from lymphoid nodular hyperplasia to as ration. Histology showed patchy chronic inflan in 11 children and reactive ileal mpho seven, but no granulomas. Bornoural disc s included autism (nine), disintegrative sy sis (one), a cossible postviral or vaccinal eccephalitis co). There were no focal neurological abornalities and al laboratory results re significantly were normal. Abnor acid compared with age-u3), low haemoglobin in four raised urinary matched contro n IgA in at children.

Internation le idente associated gastrointestinal dia se and explogmental regression in a group of province of the commental regression and associated in time a possible environmental triggers.

Lancet 1993 251: 637-41 See Commentary page

Inflammatory Bowel Disease Study Group, University Departments of Medicine and Histopathology (A. I Wakefield rucs, A. Anthony en, J. Linnell ruc, A. P. Dhillion Mechan, S. E. Davides Moorwel) and the University Departments of Paediatric Gastroenterology (S.H. Murch e.g., D.M. Casson were, M. Malik, were, M. A. Thomson resp., J. A. Waker-Smith resp.), C.Bild and Adolescent

Psychiatry (M Berelowitz recreen), Neurology (P Harvey recr), and Radiology (A Valentine rece), Royal Free Hospital and School of Medicine, London NWZ 20G, UK

Correspondence to: Dr A J Wakefield

Introduction

We saw several children who, after a part of apparent normality, lost acquired skills, including communication. They all had gastrointestinal imptoms, adviding abdominal pain, diarrhoea, and casting and, it some cases, food intolerance. We skeribe to clinical fit lings, and eastrointestinal feature of these children.

Patients and metions

12 children, community, and to department of paediatric gaster acrology a his y of a pervasive developmental across with loss upon tod skills and intestinal symptoms carries abdominated in bloating and food intolerance), were into cated. All children were admitted to the ward for largels, accompted by their parents.

Chical investigations

took histor including details of immunisations and court to infect us diseases, and assessed the children. In 11 call the history was obtained by the senior clinician (JW-5). Neur. 14 of psychiatric assessments were done by onsultant Staff (PH, MI) with HMS-4 criteria. Developmental included a review of prospective developmental records room 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 Stiller MAT under sedation with midazolam and perhidine. 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 psediatric 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 (MRR), electroencephalography (EBG) 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 methylmalonic 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. Chromatograms were scanned digitally on computer, to analyse the methylmalonic-acid zones from cases and controls. Urinary methylmalonic-acid concentrations in patients and controls were compared by a two-sample t test. Urinary creatinine was estimated by routine spectrophotometric stars.

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

ONLINE # BAD



REVIEWING FOAM



"Most of what I try fails, but these failures are often invisible, while the successes are visible,"

JOHANNES HAUSHOFER



WHEN?



YOU ARE NOT ALONE

