# Bad Science: The Measles, Mumps, and Rubella Vaccine Causes Autism

Presented by Jaclyn Wrona

### **Current Misconception**

- Opinion Differences Survey from Pew Research Center 2014
- "Childhood vaccines such as MMR should be required."
  - ▶ 68% U.S. adults ages 15-25 agree
  - ▶ 86% American Association for the Advancement of Science (AAAS) Scientists agree
- ► The MMR vaccines causes autism.



### MMR Connection to Autism

- Wakefield et. all 1998
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  - Enterocolitis The inflammation of the colon and small intestine
  - Regressive developmental disorder Previous normal development but then over time leading to loss in acquired skills

AnonymousEnterocolitis. <a href="http://www.merriam-webster.com/dictionary/enterocolitis">http://www.merriam-webster.com/dictionary/enterocolitis</a> (accessed 03/19, 2016).

Wakefield, A.; Murch, S.; Anthony, A.; Linnell, J.; Casson, D.; Malik, M.; Berelowitz, M.; Dhillon, A.; Thomson, M.; Harvey, P.; Valentine, A.; Davies, S.; Walker-Smith, J. Ileal-lymphoid-nodular hyperplasia, non-specific colitis, and pervasive developmental disorder in children. The Lancet 1998, 351, 637.

### **MMR Connection to Autism**

- Wakefield et. All 1998
- Researched children with enterocolitis and regressive developmental disorder
- Studied 12 children
  - ▶ 1 Female, 11 Male
  - Ages 3-10
- "Parents associated behavioral symptoms with MMR vaccination"
- Reported: "Casual link between MMR vaccine and this syndrome"

# MMR Connection to Autism: No Substantial Data

- The Lancet Editorial Retracted Statement 2002
- Measles Virus Genome Study
  - ► Children with developmental disorder: 82.4%
  - Children with no developmental disorder: 10.0%
- But there is no vaccine-specific strain data presented
- Retracted in 2004

AnonymousTime to look beyond MMR in autism research. The Lancet 2002, 359, 637.

### 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

Early report

**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 associably the parents, with measles, mumps, and rub a vaccination in eight of the 12 children, with measl infection in one child, and otitis media in acc. All 11 children had intestinal abnormalities arangin from lymphoid nodular hyperplasia to as moid ul pration. Histology showed patchy chronic inflam, tion if in 11 children and reactive ilear imphon a perplasia in seven, but no granulomas. Bell vioural dison is included autism (nine), disintegrative syn esis (one), an assosible postviral or vaccinal encephalitis in the terminal Abnormal laboratory results are significantly raised urinary or thylmalor of acid compared with agematched control. Design 303, low haemoglobin in four children, at is, low seem 184 in In Ital in art children.

Interrelation e ident associated gastrointestina dis se and exelopmental regression in a group o previous yearmal cm. .n., which was generally associate in time of possible environmental triggers.

Lancet 1995 251: 637–41 See Commentary page

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Correspondence to: Dr A J Wakefield

### Introduction

We saw several children who, after a period of apparen normality, lost acquired skills, includ g communication They all had gastrointestinal proproms, soluding abdominal pain, diarrhoea, and cating and, it some cases, food intolerance. We of cribe a clinical fi dings and gastrointestinal feature of these chip en.

### Patients and meti

12 children, consolitively, a red to 10 department of paediatric gastro terology a a hiroy of a pervasive developmental order with loss area red skills and intestinal symptoms arrit abdominal or in, bloating and food intolerance), were investigated. All children were admitted to the ward for layerk, accompanied by their parents.

### (Inical investigations

et took historn including details of immunisations and consure to infect us diseases, and assessed the children. In 11 cas the history as obtained by the senior clinician (JW-S). Neuron and off psychiatric assessments were done by onsultant staff (PH, MB) with HMS-4 criteria. Developmental in 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 Hun 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 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 assay.

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

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## Follow Up MMR and Autism Study

- ► Goin-Kochel et. all 2016
- If vaccines led to autism in children, then the vaccine receipt should be higher in children with regressive-onset autism compared with other patterns.

Goin-Kochel, R.; Mire, S.; Dempsey, A.; Fein, R.; Guffey, D.; Minard, C.; Cunningham, R.; Sahni, L.; Boom, J. Parental report of vaccine receipt in children with autism spectrum disorder: Do rates differ by pattern of ASD onset? *Vaccine* **2016**, *34*, 1335.

# Follow Up MMR and Autism Study

- Goin-Kochel et. all 2015
- If vaccines led to autism in children, then the vaccine strain should be higher in children with regressive-onset autism compared with other patterns.
- 2755 children studied
- Varied
  - Early onset autism
  - Gender
  - Ethnicity
  - Age
  - Vaccines Received
    - ▶ Polio, Chicken Pox, Hib (influenza), MMR, and Hepatitis B

Goin-Kochel, R.; Mire, S.; Dempsey, A.; Fein, R.; Guffey, D.; Minard, C.; Cunningham, R.; Sahni, L.; Boom, J. Parental report of vaccine receipt in children with autism spectrum disorder: Do rates differ by pattern of ASD onset? *Vaccine* **2016**, *34*, 1335.

# Follow Up MMR and Autism Study

### Results

- Validated Autism Diagnostic Interview Conducted
- No vaccine associated onset autism
- Limitations
  - Parental documentation of vaccines
  - Varied dose of vaccine received due to age difference

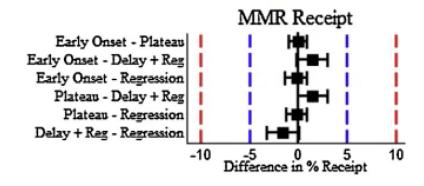


Figure 1: Reported mean differences between difference autism onset groups.

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### Why are vaccines still feared?

- Media coverage lead to distrust
  - Less than 1/3 media coverage reported no link with MMR and autism
- Parents are concerned
  - Undocumented Adverse Drug Reactions (ADR)



http://vaccineresistancemovement.org/wp-content/uploads/2010/05/Universal-Flu-Vaccine1.jpg

Goldacre, B. The Wakefield MMR verdict. <a href="http://www.badscience.net/2010/01/the-wakefield-mmr-verdict/">http://www.badscience.net/2010/01/the-wakefield-mmr-verdict/</a> (access

Casiday, R.; Cox, A. Restoring Confidence in Vaccines by Explaining Vaccine Safety Monitoring. Drug Safety 2006, 29, 1105.

### Undocumented Adverse Drug Reactions

- Doctors not reporting adverse effects
  - Study conducted by Dr. Casiday from 2002 to 2004
    - ▶ 12.6% care takers reported doctors did not take their claims seriously.
- Parents are unware on how adverse drug reactions (ADR) are tested.



# Patient Reporting

- Discover safety signal earlier than professional reporting
- Study in Netherlands reported ADRs 7 months earlier than professionals
- Patients provide qualitatively different results
- National Health Service Direct performed study in 2003
  - Limited number of reports filed
  - Professional involvement limited qualitative responses
  - ▶ Parents still receiving lack of concern from professionals



### Conclusion

- Misconception that the MMR vaccine causes autism is still prevalent
- Patient Reporting
  - Increased communication about ADR might frighten community further
  - Patient reporting can be bias from media
- Rebuild vaccine trust with community.

