#### MMR, Thimerosal and Autism

What do we know?

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**CDC Vaccine Safety Datalink Project** 

#### Overview of talk

Brief review of

Autism and autism-spectrum disorder

Summary of past reviews

Epidemiologic evidence

(Review of laboratory-based evidence)

Summary of evidence on the autism-MMR association

Review of studies underway on Thimerosal – Autism association

#### Overview of Autism and Autism-Spectrum Disorder (ASD)

#### Autism

Complex and severe neurodevelopmental disorder Strong male preponderance

Primarily characterized by impairments of

Social interaction

Verbal and nonverbal communication

Restricted/stereotypical patterns of behaviors

Onset of symptoms typically in second year of life (although recent research suggests might be earlier to trained observer)

Anatomic findings suggest prenatal insult

Genetic studies suggest gene-mediated influences

#### Overview of Autism and Autism-Spectrum Disorder (ASD)

Rates of autistic disorder

23 epidemiologic studies between 1966 – 1998

Prevalence ranged 0.7 – 21.1 per 10,000

Median 5.2 per 10,000

Rates of autistic spectrum disorder Prevalence 1-6 per 1,000

Studies vary considerably with intensity of case ascertainment methods

MMR and Autism

#### Review of past summaries

#### Institute of Medicine (IOM)

Evidence favors rejection of causal relationship between MMR vaccine and ASD

Possibility MMR vaccine contributes to ASD in a small number of children

#### Childhood Immunizations Conference

Available evidence does not support MMR vaccine causing autism or associated disorders

#### Canadian Pediatric Society

Evidence to date does not support association between MMR and autism

#### Since these reviews/conferences, studies by:

Madsen et al: A population based study of measles, mumps, and rubella vaccination and autism. N Engl J Med 2002

Farrington et al:MMR and autism: further evidence against a causal association. Vaccine 2001

DeWilde et al: Do children who become autistic consult more often after MMR vaccination? Brit J of General Practice 2001

Fombonne: No evidence for a new variant of measles-mumps-rubella induced autism. Pediatrics 2001

Taylor et al: Measles, mumps, and rubella vaccination and bowel problems or developmental regression in children with autism: population study BMJ 2002

#### Summary of epidemiologic studies

Yr	Author	Years	Туре	Cases
1998	Peltola	1982-1996	Follow up w/o ref group	31
1998	Gillberg	1975-1988	Ecologic	55
1999	Taylor	1979-1992	Case-crossover	498
2000	Farrington	1979-1998	Case crossover	357
2000	Patja	1982-1006	Follow-up w/o	0
			ref group	
2001	Kaye	1988-1999	Ecologic	305
2001	Fombonne	1954-1996	Ecologic	262
			(case v case analysis)	
2001	DeWilde	1989 – present	Case control	71
2001	Dales	1980-1994	Ecologic	N.S.
2002	Taylor	1979-1998	Case-crossover	278
			Ecologic	
2002	Madsen	1991-1998	Cohort	738

#### **Summary of Ecologic Studies**

Study	Year	Findings
Gilberg	1998	Evidence against association with autism
Taylor	1999	Evidence against association with autism
Kaye	2001	Evidence against association with autism
Dales	2001	Evidence against association with autism
Fombonne	2001	Evidence against association with autism
Taylor	2002	Evidence against association with regression

#### Summary of 'triggering' hypothesis

Study	Year	Findings
DeWilde	2001	Evidence against an association with triggering
Farrington	2000	Evidence against triggering over longer time periods
Taylor	1999	Evidence against triggering
Madsen	2002	Evidence against triggering

#### Summary of 'triggering regression' hypothesis

Study	Year	Findings
Taylor	1999	Evidence against triggering regression
Taylor	2002	Evidence against triggering regression
(Fombonne	2001	Evidence against, from case vs case analysis)

#### Summary of 'ever/never' hypothesis

Study	Year	Findings
Madsen	2002	Evidence against an association with autism
Farrington	2002	Evidence against an association with autism

Title A population based study of MMR vaccine and autism

Author Madsen
Journal NEJM
Year Published 2002

Years Covered 1991-1998 birth cohort

Type of Study Population based retrospective cohort

Specific Hypothesis Was ever receiving MMR vaccination linked to cumulative risk for

developing autism

Number of cases 738

Exposure 440,000 children vaccinated

96K unvaccinated

Results RR for vaccination vs unvaccination:

0.92 (0.68 – 1.24) for autism 0.83 (0.65 – 1.07) for ASD

7 t population based study of Millit Vaccine and autism	Title	population based study of MMR vaccine and autism	
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Findings MMR vaccine introduced 1987

Cases obtained from Danish Psychiatric Central Register

Reporting felt as complete due to comprehensive case surveillance in Denmakr

Prevalence 2.22/1000 ASD; therefore likely complete

Only specialists in child psychiatry diagnose autism/assign diagnostic codes

Case coding based on ICD 10

Cart review of 40 children to validate coding: 37 met CDC Brick Township Criteria;

other 3 met ASD criteria

Adjusted for age, gender, time, SES, maternal education, gestational age, BW

440,000 children vaccinated

96K unvaccinated

RR for vaccination vs unvaccination:

0.92 (0.68 - 1.24) for autism

0.83 (0.65 - 1.07) for ASD

RR did not vary by time since vaccination (no clustering)

Only cohort study with exposure and outcome data at the individual level

Limitations No information on why children did not get vaccinated

No information on regression

Exclusion of fragile X, Angelmans (other causes of autism) relied on identifying those

who were hospitalized: unlikely to be complete

Conclusion

4/3/2003

Strong evidence against MMR increasing risk for autism among vaccinated vs.

unvaccinated children; Fyidence against triggering

Washington/CDC

#### **Summary**

Consistent evidence from ecologic, case-control, case-crossover, and cohort studies showing lack of an association between MMR vaccine and an increased risk for developing autism, either in the short time window following vaccination or at times distant from vaccination

{Questionable evidence from two labs that measles virus is detectable in children with autism. Even if true, etiologic role far from established}

**Thimerosal and Autism** 

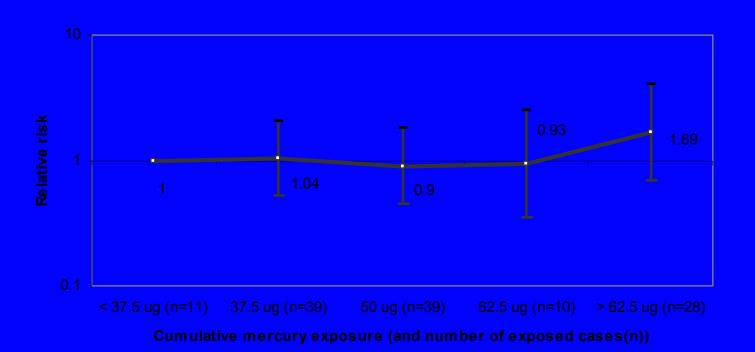
# Assessment of neurologic and renal impairment associated with Thimerosal-containing vaccines.

Thomas Verstraeten
National Immunization Program

### Results: eligible children

**213,185** Born into GHC or NCK between 1992 and 1997 142,264 Continuously enrolled for 1 year **139,344** > 1 polio vaccination by 1 year **132,391** Not premature 132,114 Did not receive HepB Ig 109,993 No congenital or perinatal disorder

# Relative risk associated with exposure at 3 months of age (NDD): <u>Autism</u> (ICD9 2990)



Trend: 1.005 (0.991, 1.019), p = 0.48

4/3/2003

R Davis, University of Washington/CDC

#### Early Life Thimerosal Exposure and Risk for Autism: Studies Underway

#### General Practice Research Database (GPRD)

#### Study design:

Cohort

#### **Population:**

Children born 1988-1997 in a GPRD database, with follow-up thru at least 2<sup>nd</sup> year of life

Over 105,000 children available for analysis

#### **Exposure:**

Thimerosal exposure from DTP or DT (or combination vaccine with one of these)

#### Outcome:

Autism per ICD-9 codes

#### Early Life Thimerosal Exposure and Risk for Autism: Planned

Vaccine Safety Datalink (VSD)

#### Study Design:

Case-control

#### Population:

Children born 1995-1999; continuous enrollment thru 1<sup>st</sup> yr, and follow up through 4<sup>th</sup> year of life

320 cases with 3 or 4:1 matched controls

#### Exposure:

Thimerosal exposure from entire vaccination schedule in first 6 months of life

Interview for other environmental mercury exposures

#### Outcome:

Autism, confirmed by in-depth interviews and in-person examination

#### Conclusions

Doubtful that further epidemiologic study can more convincingly answer the question of MMR and Autism than already answered

As with the inflammatory bowel disease question, independent confirmation of laboratory findings will be crucial to understanding the validity of the preliminary findings of Kawashima and Singh

More information will be available from UK studies underway, and US studies being planned, about thimerosal exposure and risk for autism

#### Summary of lab-based investigations

Year/Author	Journal	Title
2000 Kawashima	Dig Dis Sci	Detection and sequencing of measles virus from peripheral mononuclear cells from patients with inflammatory bowel disease and autism
2002 Martin	Mol Psych	Detection of measles virus in children with ileo-colonic lymphoid nodular hyperplasia, enterocolitis, and developmental disorder
2002 Singh	J biomed sci	Abnormal measles-mumps-rubella antibodies and CNS autoimmunity in children with autism

#### 2000 Kawashima Dig Dis Sci

Detection and sequencing of measles virus from peripheral mononuclear cells from patients with inflammatory bowel disease and autism

#### Findings:

- 12.5% 33.3% measles virus detection rates in PBMC preparations of Crohns dx, UC, and autistic enterocolitis
- 9 children with 'autistic enterocolitis' (8 with Crohn's and 3 with ulcerative colitis)
- 3 of 9 children with autism were positive for specific regions of H and F gene regions
- Sequences from patients with autism were consistent with vaccine strains

2000 Kawashima Dig Dis Sci

#### Limitations:

In 5 prior studies,

Tissue samples: Expected

0/56 Crohns 7

0/33 UC 11

Blood

0/16 Crohns 2 0/11 UC 4

Cross contamination of samples with measles virus controls or with preexisting DNA templates?

Sample to sample variations between different PCR products of same sample

2002 Singh J Biomed Sci Abnormal Measles-Mumps-Rubella Antibodies and CNS Autoimmunity in Children with Autism

#### Findings:

125 children with autism 92 control children

Unusual MMR antibody and antibodies to myelin basic protein in 75/120 (60%) of autistic sera

0/ 92 control sera

#### Conclusions:

Postulated link between MMR antibody and antibody to myelin basic protein in children with autism. Autoimmunity from vaccination as a process in developing autism

#### 2002 Singh

## Limitations: (Bellini)

Control children are not age and sex matched Autism history not confirmed Timing of past immunization not given (will affect antibody levels)

Use of an MMR antibody is not accepted widely Immune response that occurs <u>is</u> to the separate components of MMR (no combination of viral RNA occurs among these viruses)

The authors use an in-house ELISA to detect "MMR antibodies"; not a valid/substantiated technique

Title No evidence for MMR vaccine-associated inflammatory bowel disease or

autism in a 14 year propsective study

Author Peltola

Journal Lancet

Year Published 1998

Years Covered 1982-1996

Type of Study Cohort follow-up without comparison group, of vaccinees who developed

GI symptoms following vaccination

Specific Hypothesis Were there children who developed GI symptoms shortly after

vaccination who then went on to develop IBD and/or autism

Number of cases 31 children developed GI symptoms after vaccination

Title No evidence for MMR vaccine associated inflammatory bowel disease or

autism in a 14 year prospective study

Findings 31 children were followed for at least 16 months, and some up to 181

months (15 years).

Most had diarrhea with vomiting after vaccination

No child developed autistic spectrum disorder

Limitations Low power to detect true effect severely limits any conclusions

(0 events/31 ~ upper 95<sup>th</sup> CI approx 9%)

Conclusion Relatively non-contributory

Title MMR and autism

Author Gillberg

Journal (Autism)

Year Published 1998

Years Covered 1975-1984 birth cohort

Type of Study Ecologic

Specific Hypothesis Did introduction of MMR vaccination in 1982 increase rate of autism in

Sweden

Number of cases 74

Title MMR and autism

Findings MMR vaccine introduced 1982

Rate of autism calculated for children born 1975-1980 (5.5 y) and

compared to rate among those born 1980-1984 (4.5 y)

47 children born during earlier period

27 children born during later period

Limitations Different number of years in the denominator

Different length of follow in the earlier vs later times.

Diagnoses made after 4 years of age (after 1988) in the latter cohort

would have been missed

Assumption that number of children born each year remained stable

Limited power (55 cases of autism; 74 overall)

Lack of face validity; almost all other studies have shown autism

increasing over time. Before/after studies as this one should show some

relation, however spurious

Conclusion Limited evidence against relationship between vaccine and autism

Title Autism and MMR vaccine: no epidemiological evidence for a causal

association

Author Taylor

Journal Lancet

Year Published 1999

Years Covered 1979-1992

Type of Study Case-crossover

Ecologic

Specific Hypothesis Did autism increase after MMR introduced in 1988

Did age at diagnosis change after MMR introduced

Was there clustering of cases after MMR

Number of cases 498 cases of autism

293 confirmed by ICD 10 criteria

4/3/2003 R Davis, University of Washington/CDC

Title Autism and MMR vaccine: no epidemiological evidence for a causal

association

Findings Cases ascertained by computerized registries at child developmental centers and special schools

Age of autism diagnosis compared for children receiving MMR before 18 months, after 18 months, and never

Case crossover analysis for periods within 1-2 years for autism, within 6-12 months for parental concerns, and within 2,4, and 6 months for regression

No evidence of step up in birth cohorts exposed to vaccine, nor change in slope of curve (from exponential slope)

During period of most rapid increase in autism, MMR coverage by second year of life was stable

No difference in age of autism onset among children receiving vaccination before or after 18 months of age, or never receiving vaccine

No clustering of autism, parental concern, or regression in time windows after vaccination, except for parental concern 6 months after vaccination

(Authors interpretation for number preference (6 mo) following age of most frequent vaccination

Limitations No standardized assessment of regression timing

No standardized evaluation of cases

Conclusion Evidence against a relationship between vaccination and triggering of symptoms.

Evidence against increase in disease after MMR introduction

R Davis, University of Washington/CDC

Title Serious adverse events after MMR vaccination during a fourteen year

prospective follow-up

Author Patja

Journal Pediatr Infect Dis J

Year Published 2000

Years Covered 1982-1996

Type of Study Cohort follow-up without comparison group

Specific Hypothesis To perform surveillance for serious adverse events after MMR over 14

years

Number of cases 1.8 million individuals immunized, with 3 million vaccine dose

Title Serious adverse events after MMR vaccination during a fourteen year

prospective follow-up

Findings Country-wide passive surveillance system collected AE reports

At start of project in 1982, seminars, media, medical publications used to

highlight system

No cases of autism were reported following vaccination

Limitations At rate of 6/1000 for ASD or 6/10,000 for Autism, should have been

hundreds/thousands of cases, regardless of causal association

Since no cases were reported, implication is that reporting was limited to

those diseases known to be associated with vaccination

Conclusion Non-contributory

Title MMR and autism: further evidence against a causal association

Author Farrington

Year published 2000

Journal Vaccine

Years covered 1979 - 1998

Type of study Case-crossover

Specific hypothesis Extended their previous study that looked at clustering of autism

following vaccination.

Looked 'all' time intervals, including risks

within 5 years after vaccination for autism diagnosis

within 3 years for parental concern

within 2 years for regression

Number of cases 357 with autism

Title MMR and autism: further evidence against a causal association

#### In depth review of findings

Risk for autism following MMR was compared to the risk for autism before MMR. (Distribution of age at autism diagnosis of 64 unvaccinated children with autism were used to adjust for age, with 16 categories used to control for time)

No increased risks for autism diagnosis, for parental concern, or for regression when all time following vaccine is compared with all time prior to vaccine.

No risk for autism within 5 years, for parental concern within 3 years, or for regression within 2 years.

#### Limitations

Brief discussion of methods.

No discussion of case ascertainment or validation.

Misclassification of case, or of regression, would bias towards null.

Time after vaccination is older than time prior to vaccination: unclear whether age adjustment can adequately adjust for this.

#### Conclusions

Evidence against an ever-never effect and evidence against a triggering phenomenon

Title Time Trends in Autism and in MMR immunization coverage in California

Author Dales

Journal JAMA

Year Published 2001

Years Covered 1980-1994

Type of Study Ecologic

Specific Hypothesis Did autism case load in California increase as immunization coverage

with MMR increased?

 Title Time Trends in Autism and in MMR immunization coverage in California

Findings Cohort born between 1980 – 1994

Assumed autism ascertainment complete for children born in 1994

Autism caseload increased each year:

one rate 1980 – 1984 another rate 1985 - 1988 another rate 1988 – 1994

The major rise in caseload was after 1988, while MMR coverage at 17

and 24 months was constant

\_First\_ rise in autism caseload was in 1985, before any rise in MMR coverage

Limitations State immunization records possibly incomplete (exposure validity questionable)

Had coverage increased, but not recorded, might have missed positive association

Autism validation non-existent

Caseload used rather than rates

Complete case ascertainment not attempted

No individual level data on exposure or outcome

Conclusion Weak evidence against association. Fair evidence against MMR as a major influence

of autism rates

Title MMR vaccine and the incidence of autism recorded by general

practitioners: a time trend analysis

Author Kaye

Journal BMJ

Year Published 2001

Years Covered 1988-1999

Type of Study Ecologic

Specific Hypothesis Did risk for autism increase with increasing MMR vaccine coverage?

Number of cases 305 children with first recorded autism diagnosis between 1988 and

1999. Subanalysis of 114 boys born 1988-1993 to provide rates for

children 2-5 years old between 1990 and 1999

Title MMR vaccine and the incidence of autism recorded by general

practitioners: a time trend analysis

Findings Used the UK GPRDatabase

Vaccine coverage stated as complete

81% of cases were evaluated by specialists

For each birth cohort, estimated the cumulative risk for autism between 2

to 5 years of age

Risk increased from 8/10,000 born in 1988 to 29/10,000 born in 1993

MMR vaccination by 2 years of age constant at 97%

Limitations Autism not validated by strict criteria (but doubtful that entire rise in case-

load was due to increasing misclassification)

Conclusion Evidence against an association

Title Do children who become autistic consult more often after

MMR vaccination

Author DeWilde

Journal British Journal of General Practice

Year published 2001

Years covered 1989 - present

Type of study Case control

#### Specific hypothesis

Do children with autism see providers and/or get

consultations more frequently after MMR vaccination than

children without autism

Exposure MMR vaccine

Outcome Consultation rates (for symptoms)

Number (cases) 71

#### Title

Do children who become autistic consult more often after MMR vaccination

#### In depth review of findings

If vaccination caused increase in symptoms – one would see a preferential increase in symptoms and consultations following vaccination for cases but not for children without autism.

Study performed in the Doctors Independent Network; 127 core practices with data on children enrolled from birth, in the UK. First started in 1989

Children with autism diagnoses prior to vaccination were excluded (one case)

Controls were matched by practice, age, gender, month of vaccination, and current enrollment

The rates of consultations fell after MMR vaccination among both cases and among controls. The rates of fall did not differ statistically (significantly or clinically) between the two. No case was diagnosed in the 6 months following MMR

#### Limitations

Autism criteria was not confirmed.

May have missed some, or many, cases; wouldn't necessarily bias results Timing of autism diagnosis was not specified.

#### Conclusion:

Evidence against MMR triggering symptoms that are eventually diagnoses as autism

Title No evidence for a new variant of MMR induced autism

Author Fombonne

Journal Pediatrics

Year Published 2001

Years Covered 3 birth cohort samples:

1992-1995 (post MMR) 1987-1996 (post MMR) 1954-1979 (pre MMR)

Type of Study Ecologic

Specific Hypothesis Has age at diagnosis changed in post MMR samples?

Have rates of regression increased in post MMR samples?

Is age of regression diagnosis or concern younger than those without

regression?

Is interval between immunization and concern shorter in children with

regression?

Number of cases 262 in 3 samples

R Davis, University of Washington/CDC

No evidence for a new variant of MMR induced autism Title

Findings Avg age of diagnosis: 19 months in pre and post samples

> Avg age of concern: 19 months for regression and non-regressive

> > autism

Lag between MMR and concern was 248 d for regression

272 for non-regressive

Proportion of cases with regression 18% in pre-MMR sample

16% in post MMR sample

Limitations

MMR might increase rate of autism without altering avg. age at diagnosis Comparing lag between MMR and concern for regression versus concern for non-regressive cases is based on 67 children total: limited power to

detect differences as statistically significant

Difficult to interpret results based on comparisons with pre-sample that goes back to 1954. Case ascertainment and diagnostic criteria likely to differ substantially in years prior to 1979 compared with years 1987-1996

Conclusion Limited evidence against relationship between vaccine and autism Title MMR vaccine and bowel problems or developmental regression in

children with autism: population study

Author Taylor

Journal BMJ

Year Published 2002

Years Covered 1979-1998

Type of Study Case-crossover

Specific Hypothesis Relationship of MMR with bowel problems for more than 3 months, and

regression

Number of cases 278 children with autism, and 195 with 'atypical' autism

81 had bowel problems

118 (25%) had regression

Title

MMR vaccine and bowel problems or developmental regression in children with autism: population study

**Findings** 

No trends by year of birth in % of children with autism who also had bowel problems, or of children with autism and regression. This spanned the period when MMR was introduced

Of children with bowel problems and autism 19% of children vaccinated before parental concern 15% vaccinated after parental concern 16% never vaccinated (p=0.48)

Of 28 children with 'autistic enterocolitis' (signs and symptoms), no association with MMR vaccination (p=0.57) nor with year of birth in the proportion of these symptoms

Of 31 children with bowel problems and regression no association with MMR vaccination (p=0.20) nor with year of birth

Limitations

Low power to assess association between 'autistic enterocolitis' and MMR vaccination

No OR given for MMR vaccination association; p values only (no CI)

Conclusion

No evidence to support new variant of regressive autism with bowel problems or clustering following vaccination

Evidence for association between regression and bowel disease, just not with vaccination

4/3/2003

R Davis, University of Washington/CDC