ALCOHOL AFFECTED ADULTS' SELF-REPORTED HEALTH AND MENTAL HEALTH STATUS:

EVIDENCE OF LONG-TERM EFFECTS FROM A CROSS SITE STUDY

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The problem:

- Despite the prevalence of FASD and its life-long course, there is little empirical research about longterm adult health, physical characteristics, neurobehavior or adaptive functioning.
- The Developmental Origins of Health and Disease (DOHoD) hypothesis suggests that fetal programming by PAE should result in vulnerable organisms with increased sensitivity to stress, adverse health and functional outcomes.

• Long-term effects should begin to present themselves by Middle Adulthood.

THE STUDY

- Multisite Collaborative Study (part of the Collaborative Initiative on Fetal Alcohol Spectrum Disorders (CIFASD4)
- Sites include <u>Atlanta</u>, <u>GA</u> and <u>Seattle</u>, <u>WA</u>, with a parallel study in <u>Vancouver</u>, <u>BC</u>.
- Two "Tier" Assessment in Seattle and Atlanta:
 - Tier 1. Demographic and Health Survey,
 Registry
 - Tier 2. In-Depth Assessment of medical records, physical characteristics, immune function, mental health, cognitive functioning, social/adaptive functioning
- In Vancouver, only Tier 2









PARTICIPANT POOL

In Seattle and Atlanta, individuals were recruited who took part in earlier longitudinal studies at the University of Washington and Emory University.

- Seattle: a total of 475, now age 30 to 74 years, 66% White and 41% female; 37% diagnosed with FAS and 63% non-dysmorphic but affected, classified as having FAE.
- Atlanta: a total of 427, now in late-30s, early 40's, low SES, predominantly African-American and 48% female. Of these, 292 were recruited prenatally from same intercity hospital, 70% are alcohol-exposed.

In Vancouver, individuals aged 22-50 years were recruited through community networks with the Goal 1) 40 people with FAS, pFAS or physical effects of alcohol; 2) 40 alcohol-exposed without physical features; and 3) 40 unexposed controls.

• Tier 1

Adult Health Survey (N=360) of chronic medical problems in midlife and access to health care.

Health Issues assessed:

- Access to health care
- Sleep
- Vision/Hearing/Dental
- Allergies and Asthma
- Cardiovascular
- GI
- Diabetes
- Autoimmune disorders
- Arthritis
- Seizures
- Depression/Anxiety
- Other Medical problems
- Adverse Childhood Experiences

Questionnaire Development

Structured using the CDC
Behavioral Risk Factor
Surveillance System
Questionnaire (BRFSS) as a
model.

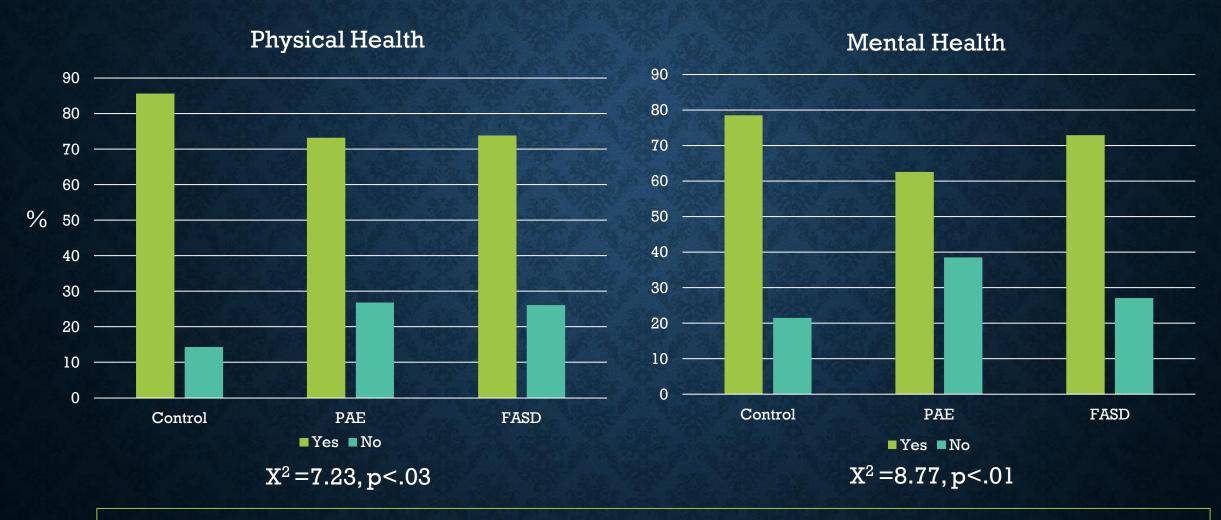
- 1. Selected from Existing BRFSS modules (in public domain).
- Added modules on Seizures from Neurological Questionnaire.
- 3. Created modules on vision, hearing and dental health.
- 4. Created modules on Gastrointestinal and Immunology.

CHARACTERISTICS OF SAMPLE IN ATLANTA AND SEATTLE: DEMOGRAPHICS (N=358)

Characteristic		Contrast (n=136)	Alcohol Exposed (n=128)	FAS(D) (n=94)	Statistic
Tier 1 Age (n=292)		40.2 (7.67)	37.96 (6.54)	38.94 (6.75))	F _(2,334) =3.15, p<.05
Biological Sex (% male)		38.9%	40.2%	47.3%	$X_{(2)}^2 = 6.23$, NS
Race* (%) *90% Non Hispanic	White African-American Native American Other (Mixed)	47.8% 38.2% 4.4% 5.9%	44.1% 29.1% 13.4% 13.4%	40.4% 40.4% 8.5% 7.4%	X ² =18.78, p=.04
Hollingshead (1975) SES Rank		41.0 (14.98)	30.24(12.57)	25.33 (10.67)	F _(2,334) =41.41,p<.000
Marital Status	"married" "unmarried"	49.2% 47.1%	41.9% 58.23%	25.5% 71.3%	X ² ₍₁₂₎ =29.6, p<.003
M # Biological Children		2.36(1.25)	2.58(1.45)	2.29(1.3)	Not sig. different
Household Income	>\$4000/mo <\$1000/mo	44.1% 6.6%	15.7% 21.3%	12.8% 31.9%	X ² ₍₁₆₎ =62.3, p<.000

Alcohol affected individuals are less often married, have lower SES and lower incomes than Contrast group. Recall that when identified, SES was matched.

"OVERALL DO YOU THINK OF YOURSELF AS IN GOOD HEALTH?" (N=358)



Self-Reported Physical and Mental Health Status varies with <u>Alcohol-Exposed</u> reporting more problems.

"DO YOU HAVE PROBLEMS WITH..."

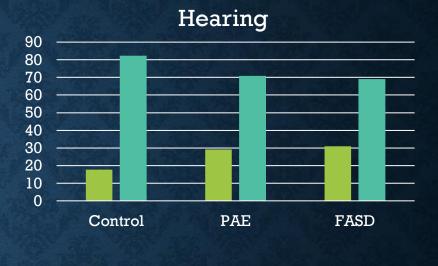




More than 50% of all groups wear glasses. "Everyone" at this age has these problems



$$X^2=27.4, p<.000$$



X²=6.8, p<.03

■YES ■NO

Alcohol-exposed individuals reported significantly more hearing and dental problems than did controls.

"Heart Problem as Child?"



$$X^2=3.54, p=.17$$

Cardiovascular outcomes are suggestive in childhood and emerging in middle age. Need to control for Race. Sex, and BMI.

CARDIOVASCULAR

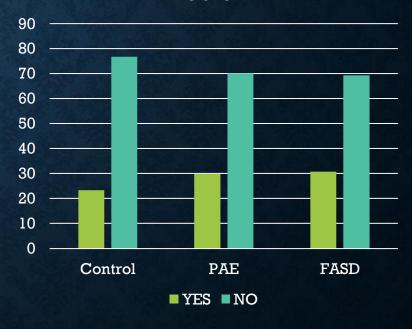


"Heart Problems Now"



 $X^2=8.73, p=.01$

"High Blood Pressure. Adult?"



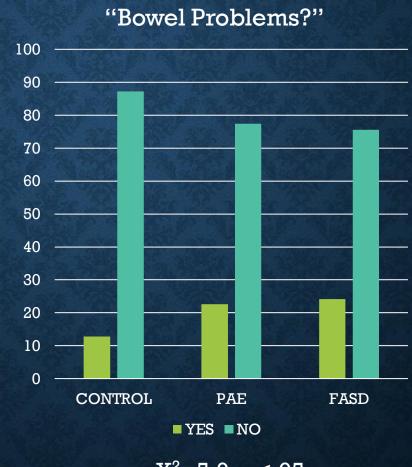
 $X^2=1.95$, NS

GASTROINTESTINAL

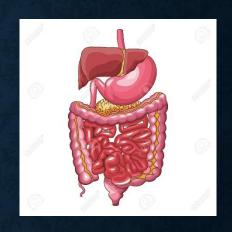
"Heartburn or Stomach Problems"



 $X^2=8.76, p<.01$

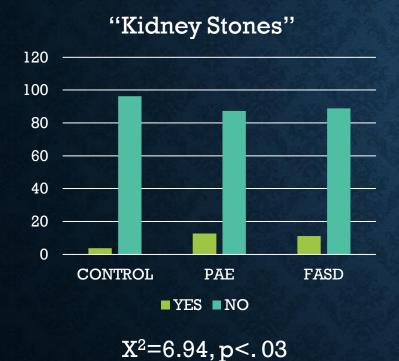


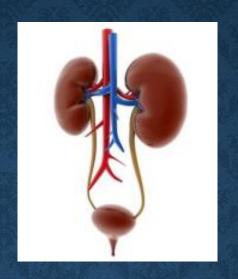
 $X^2=5.9, p<.05$



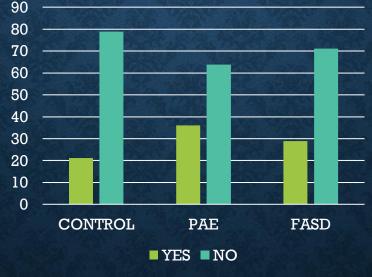
More
AlcoholAffected
individuals
report
current GI
problems.

"Have you ever had....."





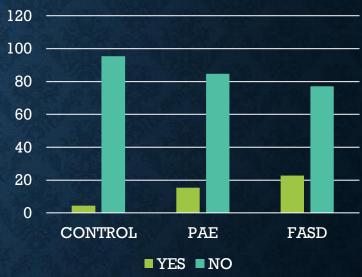




 $X^2=7.07, p<.03$

URINARY TRACT





$$X^2=16.71 p<.001$$

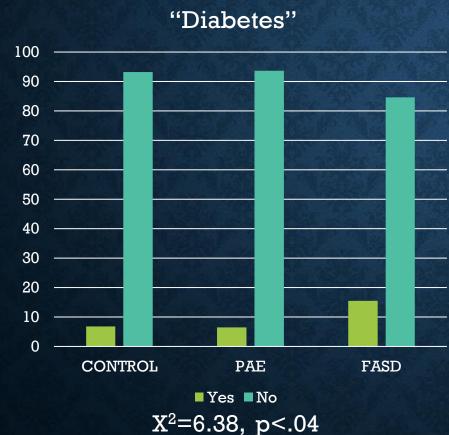
Increased Kidney Stones, Bladder Infections and Incontinence in Alcohol-Affected individuals.

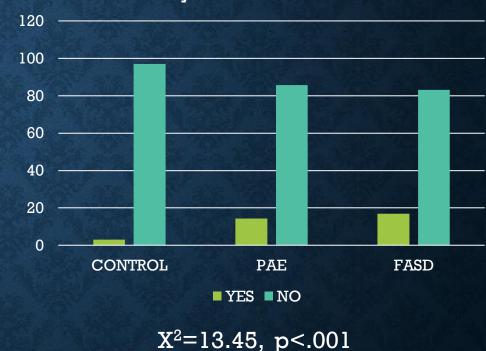


"Has a Medical Professional Diagnosed you as having...."

ENDOCRINE

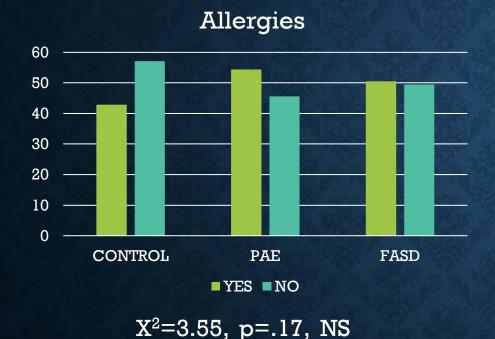






Increased in diabetes in FAS/Dysmorphic. Thyroid problems are more common in Alcohol-Affected individuals.

"Has a Medical Professional Diagnosed you as Having....'

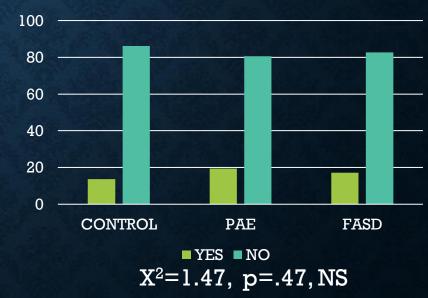


Medically Diagnosed problems with Skin are more common in Alcohol-Affected individuals. Other autoimmune problems are not significantly different.

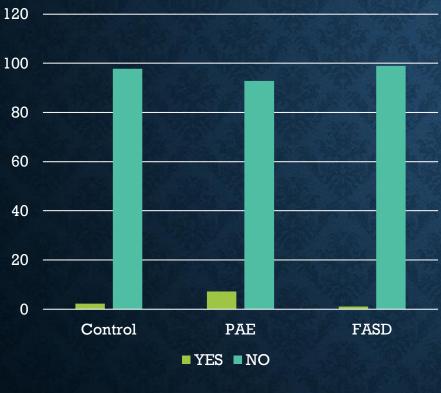




Autoimmune Disorder



"HAVE YOU HAD CANCER?"



 $X^2=6.7$, p<.03

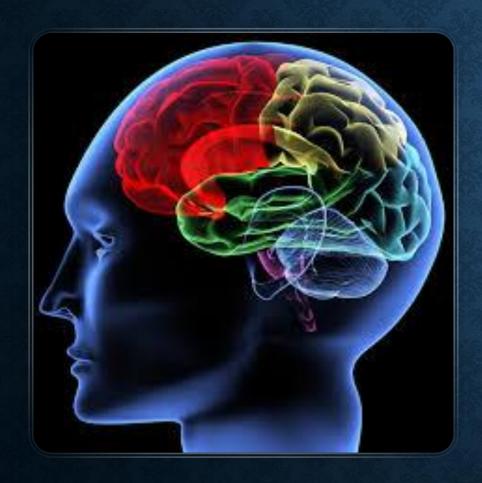
TYPES Of CANCER REPORTED

Breast Cancer (3)
Cervical Cancer (2)
Colorectal
Endometrial
Melanoma
Skin Cancer, Basal (2)
Not Specified

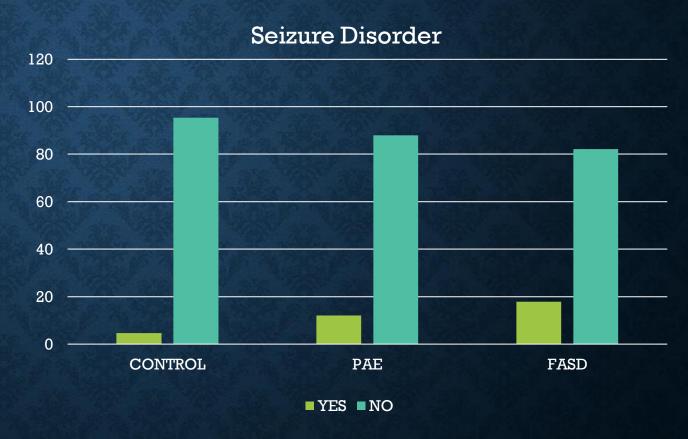
10 Alcohol Affected individuals reported having cancer (4.6%).3 Controls reported cancer (2.3%)

Relative risk (RR) of cancer in alcohol group is 5.43.

"HAS A MEDICAL PROFESSIONAL DIAGNOSED.."



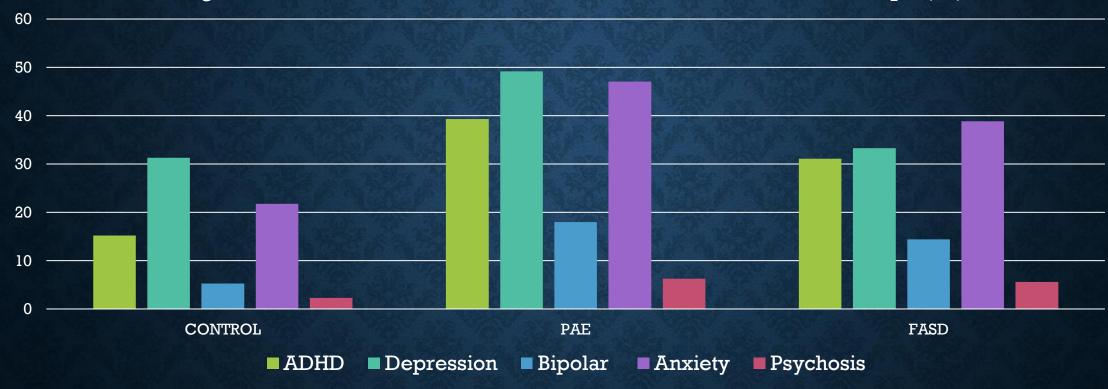
Alcohol-Exposed Individuals report Seizure Disorders more frequently.



X²=9.96, p<.007

MENTAL HEALTH

Diagnosed Mental Health Problems in Alcohol and Control Groups (%)



ADHD: X²=19.07, p<.000

Depression: X²=9.91, p<.007

Bipolar Disorder: X²=10.29, p<.006

Anxiety Disorders: X²=18.53, p<.000

Psychoses: X²=2.76, p=.25, NS

With the exception of Psychoses, the Alcohol Groups have significantly higher rates of Diagnosed Mental Health Disorders.

HOW CAN WE UNDERSTAND THESE OUTCOMES?

- Is just that Prenatal Alcohol Exposure is a direct cause of negative Health and Mental Health Outcomes?
- Is Prenatal Exposure Associated with Health and Mental Health Outcomes?
- Is Prenatal Alcohol Exposure one of many factors affecting outcomes?

WHAT IS THE RELATIONSHIP BETWEEN ALCOHOL EXPOSURE AND HEALTH AND MENTAL HEALTH OUTCOMES?

Greater Social Stressors (SES, Race/Ethnicity, Access to Health Care) Prenatal Exposure Maternal Substance Use Health Issue Life Individual Mental Experience Characteristics Health (Sex, Ability) Disorder **Familial** Postnatal (Genetic) Environment

Multiple Factors
Associated with
Outcomes by
Midlife

Some Possible Relationships based on Previous Research.

ADVERSE CHILDHOOD EXPERIENCES (ACES) (N=382)



Alcohol: $F_{(2,349)} = 15.99, p < .000$

Sex: $F_{(2,349)} = 3.55$, p<.03

A by S: $F_{(3,349)} = <1$, ns

Total ACES higher in Alcohol-Exposed.
Items that were significantly different concerned Custody, Substance USE
Disorders, Incarceration and Child Abuse.

- 1. Parent died: Alcohol: 16.3%; Control: 11.1%, NS
- 2. Custody Changed: Alcohol: 57.8%, Control: 10.3%,

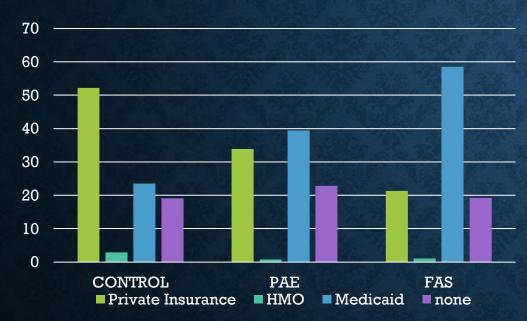
 $X^2=77.38, p<.000$

- 3. Depressed Caregiver: Alcohol: 30%; Control: 21.1%, $X^2=5.71$, p=.06, NS
- 4. Caregiver AUD: Alcohol: 49%; Control: 26%, X²=17.48, p<.001
- 5. Caregiver DUD: Alcohol: 34%; Control: 16%, X²=14.05, p<.001
- 6. Incarcerated Family Member: Alcohol: 28.4%; Control: 4.8%, X²=28.1, p<.000.
- 7. Parents divorced: Alcohol: 54%; Control: 33%, $X^2=17.3$, p<.001.
- 8. **Domestic Violence**: Alcohol: 39.5%; Control: 16%, X²=128.76, p<.001
- 9. **Physically Abused:** Alcohol: 34.8%; Control: 12%, X²=25.8, p<.001.
- 10. Verbally Abused: Alcohol: 44%; Control: 28%, $X^2=10.46$, p<.03.
- 11. **Sexually Abused:** Alcohol: 37%; Control: 8%, X²=37.9, p<.000.

"Do you have Health Insurance?"

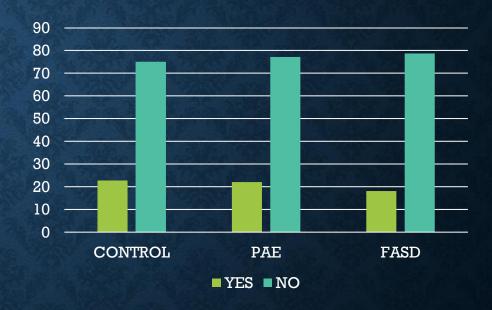
Not different: response from each group ~89%.

"What type of insurance do you have?" X²=39.4, p<.-0000



Alcohol-Exposed more likely have <u>Medicaid</u> vs <u>Private</u> Insurance. ~20% report none.

"IN THE LAST 12 MONTHS DID YOU EVER NOT GO TO THE DOCTOR BECAUSE OF COST?"



But no difference in obtaining care reported. ~20% report not obtaining care.

MULTIPLE INFLUENCES ON MEDICAL DIAGNOSES

Cancer Diagnosis

Diabetes

Likelihood Ratio Tests				
Effect	β	X^2	Significance	
Prenatal Alcohol	2.56	4.34	p<.04	
Sex (female)	1.65	3.74	p=.053	
SES	.007	.08	NS	
ACES	.160	1.67	NS	
Age	.20	18.1	p<.001	

Cancer diagnosis is more common in those who were prenatally exposed to alcohol, female and older.

Diagnosis of diabetes is more common in those who were prenatally exposed to alcohol, female, older, and had lower SES.

Likelihood Ratio Tests				
Effect	β	X^2	Significance	
Prenatal Alcohol	.98	3.97	p<.05	
Sex (female)	79	3.00	p=.08	
SES	.03	3.36	p=.07	
ACES	07	.87	Not significant	
Age	051	3.91	p<.05	

Factors contributing to **cancer diagnosis** using Multinomial Regression Analysis (MRL).

Factors contributing to **diabetes** using Multinomial Regression Analysis (MRL).

MULTIPLE INFLUENCES ON MEDICAL DIAGNOSES

Thyroid Disorder

Kidney Stones

Likelihood Ratio Tests				
Effect	β	\mathbf{X}^2	Significance	
Prenatal Alcohol	2.17	9.2	p<.002	
Sex (female)	-1.69	10.6	p<.001	
SES	.001	<1	Not significant	
ACES	166	5.99	p<.01	
Age	04	2.31	p=.13, NS	

Factors contributing to **thyroid disorder diagnosis** using
Multinomial Regression Analysis
(MRL).

Thyroid disease is more common in those who were prenatally exposed to alcohol and had childhoods with greater adversity.

Diagnosis of
Kidney stones is
impacted by
prenatal
exposure to
alcohol and age.

Likelihood Ratio Tests				
Effect	β	X^2	Significance	
Prenatal Alcohol	1.31	3.86	p<.05	
Sex	.09	<1	Not significant	
SES	.002	<1	Not significant	
ACES	.025	<1	Not significant	
Age	057	5.63	p<.02	

Factors contributing to Kidney Stones using Multinomial Regression Analysis (MRL).

WHAT ARE THE IMPLICATIONS OF THESE FINDINGS?

HEALTH CARE

- These results suggest that alcohol exposure may have long-term association
 with a range of medical problems. When individuals are known to have such
 exposure, their medical providers should screen for problems.
 - Alcohol exposed adults should have regular medical checkups and medical providers should be aware of this as a risk factor.
 - At this time, we don't know if these problems are the result of "premature aging".

 Mental Health problems are more common in alcohol affected individuals and should be the focus of screening in this group.

PREVENTION OF LONG-TERM EFFECTS OF PRENATAL ALCOHOL EXPOSURE

- Direct Effects of PAE
 - Alcohol Exposed individuals appear to have some outcomes that are
 directly associated with PAE. These obviously could have been prevented
 by abstinence during pregnancy. However, it is likely that, knowing about
 the potential for negative physical and mental health impacts, that early
 intervention efforts could reduce the impact on many of these outcomes.
- Indirect Effects and Associated Factors
 - Some of the negative consequences of PAE may be due to conditions that are associated with maternal substance abuse. For instance, negative childhood experiences. When a child is known to be exposed, attention to these associated factors could prevent or reduce long-term problems.

FURTHER RESEARCH

The results of this study are still preliminary and suggest some areas that should be studied much more extensively.

- Long-term outcomes in adulthood have not been studied systematically. Both the medical and social/emotional consequences of alcohol exposure need much more research with the goal of not just describing outcomes but understanding the mechanisms.
- This study concerned Midlife Adults and found significant medical and mental health problems. It is likely that these issues will be exacerbated with age. The potential for accrated aging in many systems should be addressed.
- Finally, this is an observational study. It will be important to identify biophysiological factors that contribute to these outcomes, including the role of the immune system.