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# A machine learning approach to investigate potential risk factors for gastroschisis in California

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# Abstract

**Background:** To generate new leads about risk factors for gastroschisis, a birth defect that has been increasing in prevalence over time, we performed an untargeted data mining statistical approach.

**Methods:** Using data exclusively from the California Center of the National Birth Defects Prevention Study, we compared 286 cases of gastroschisis and 1,263 non-malformed, live-born controls. All infants had delivery dates between October 1997 and December 2011 and were stratified by maternal age at birth (<20 and 20 years). Cases and controls were compared by maternal responses to 183 questions (219 variables) using random forest, a data mining procedure. Variables deemed important by random forest were included in logistic regression models to estimate odds ratios and 95% confidence intervals.

**Results:** Among women younger than 20, of variables deemed important, there were higher odds observed for higher consumption of chocolate, low intake of iron, acetaminophen use and urinary tract infections during the beginning of pregnancy. After adjustment, the higher odds remained for low iron intake and a urinary tract infection in the first month of pregnancy. Among women aged 20 or older, of variables deemed important, higher odds were observed for US-born women of Hispanic ethnicity and for parental substance abuse. There were lower odds observed for obese women, women who ate any cereal the month before pregnancy, and those with higher parity.

**Conclusions:** We did not discover many previously unreported associations, despite our novel approach to generate new hypotheses. However, our results do add evidence to some previously proposed risk factors.

# Keywords

gastroschisis; random forest; data mining; maternal age; teenage pregnancy; etiology

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# Introduction

Gastroschisis is a congenital anomaly of the abdominal wall where part of the intestines and sometimes other internal organs are outside the body at birth. Its prevalence in the US is roughly 4 per 10,000 live births (Canfield and others, 2006; Parker and others, 2010). The prevalence has been increasing over the past several decades (Castilla and others, 2008; Jones and others, 2016; St Louis and others, 2017; Vu and others, 2008). Many factors have been investigated one, or a few at a time, and no study has sufficiently explained the increase in temporal prevalence or clearly identified a risk factor for gastroschisis other than the well described increased (5-fold) risk to women 20 years of age and younger (Gill and others, 2012; Rasmussen and Frias, 2008; Reefhuis and Honein, 2004; Werler and others, 1992). Further, even fewer investigations have been made to specifically disentangle the potential disproportionate risks factors that may underlie prevalence differences between younger (<20 years) and older (20 years) mothers. Given a general lack of information on the potential multiple-factor-influences on gastroschisis risk, as well as a need to generate new leads about risk factors, we performed an untargeted statistical approach to fully utilize the rich set of available data in California.

#### Methods

#### Study population and data collection

Data are from the California Center of the National Birth Defects Prevention Study (NBDPS) (Reefhuis and others, 2015). The California Center is a joint effort between Stanford University and the California Birth Defects Monitoring Program in the Department of Public Health. The California Birth Defects Monitoring Program has been performing population-based active surveillance and collecting data from women residing in one of eight counties in the San Joaquin Valley at time of delivery since 1986 (Croen and others, 1991). To ascertain cases, staff visited hospitals with obstetric or pediatric services, clinical genetics prenatal and postnatal outpatient services, and cytogenetic laboratories.

Included cases of gastroschisis were confirmed by a clinical geneticist using clinical, surgical, or autopsy reports. Cases suspected of having a chromosomal abnormality or identifiable syndrome were ineligible (Rasmussen and others, 2003). Controls were nonmalformed, live-born infants randomly selected from birth hospitals of cases. Roughly 150 controls were selected per same study year as cases. Interviews with maternal participants took place between 6 weeks and 24 months after the estimated date of delivery and were performed using standardized, computer-based questionnaires in English or Spanish. Questions sought information on maternal health, pregnancy history, as well as maternal and paternal sociodemographic and behavioral information at specific points before and throughout pregnancy. Dietary information was collected via a 58-item food frequency questionnaire, developed and validated by Willet and colleagues (Willett and others, 1985), of average intake for the year before pregnancy and certain other specific intake questions. Time-periods are delineated by month as before (B3-B1) or during pregnancy (P1-P9). For this specific analysis, the time-periods of interest were the month before conception (B1) until the second month of pregnancy (P1-P2) as this is believed to be the relevant etiologic window.

The dataset included cases and controls with estimated dates of delivery from October 1, 1997 to December 31, 2011. Mothers were interviewed for 286 cases and 1263 controls. Participation in the interview was 66% for case and 64% for control mothers. Maternal participants missing >10% of questions were excluded to remove participants who did not complete the questionnaire, resulting in 268 cases and 1203 controls. Cases and controls were compared for maternal responses to 183 questions (219 variables). This dataset was analyzed by age stratified as younger than 20 (109 cases and 181 controls) or 20 and older (159 cases and 1022 controls).

A supplemental dataset also included pesticide and air pollution data for a portion of the cases and controls, i.e., subjects with estimated dates of delivery from October 1, 1997 to December 31, 2006. Pesticide and air pollution exposure and associations with gastroschisis have been investigated previously and methods are described elsewhere (Padula and others, 2013; Shaw and others, 2014). Briefly, maternal residences were geocoded and linked with the California Department of Pesticide Regulation reporting records and US Environmental Protection Agency's Air Quality System database (https://aqs.epa.gov/aqsweb/documents/ data mart welcome.html). Only those living in one of the eight counties for 75% of the time during B1-P2 were eligible for pesticide exposure assignment and during P1-P2 for air pollution exposure assignment. Pesticide data were collected for 461 chemicals and 69 physiochemical groupings that were applied at >100 lb and deemed toxic based on Environmental Protection Agency's (EPA) risk assessment and California Proposition 65 or were classified as endocrine disruptors. Data were collected for daily 24-hour averages of nitrogen oxide, nitrogen dioxide, particulate matter <10µm (PM<sub>10</sub>), particulate matter <2.5µm (PM2.5), and carbon monoxide, and a daily 8-hour maximum of ozone and averaged over the study period. Traffic-density measures were also collected based on distancedecayed annual average daily traffic volumes from the Geographic Data Technology traffic count data. Of participants with data on both pesticides and pollutants, missing <10% of questions, there were 145 cases and 758 controls. Cases and controls were compared for responses to 242 questions (278 variables).

#### Statistical analysis

Cases and controls were compared for eligible variables using random forest. The variables are described in Appendix A. Random forest is a well-established data mining procedure that calculates a set of decision trees using random subsets of the data and combines them to produce a mean prediction model of case status based on variable importance (Strobl and others, 2009). Random forest accounts for interactions and non-linear associations among a large group of factors simultaneously to determine the importance of individual variables (Strobl and others, 2009). The current version of the program removes a previously existing bias towards correlated variables by using conditional inference trees (Strobl and others, 2009).

We calculated a variable importance measurement for each potential predictor variable, using the "varimp" function in the party package in R software to obtain the metric mean decrease accuracy (MDA) specifying "ntree=2500" and "mtry=15" as the number of trees and number of selected predictor variables per split, respectively. If the MDA value was

Variables were included in the random forest procedure if questions occurred with a frequency 0.1% to increase the likelihood of having adequate exposed participants for calculation. Time-varying variables were included for the time-period from one month before pregnancy (B1) until the second month of pregnancy (P1-P2) to reflect the hypothesized etiologically relevant window (Lammer and others, 2008). Pesticide exposure was included as any exposure during B1-P2 and air pollution values were included for the time-period P1-P2 based on previously collected data (Padula and others, 2013; Shaw and others, 2014).

Nutrient and air pollution values were divided into three categories based on values among controls: those with exposure levels <25%, those within the interquartile range, and those with exposure levels >75%. For the initial random forest, simple imputation was performed for missing data. For the nutrient and air pollution values, all missing data for these variables were imputed as – being within the interquartile range. Other missing data were imputed with the median among controls for continuous variables and the most frequently occurring response among controls for categorical variables.

The random forest analysis only provided the ranking list of "important" variables, thus the effect of these variables toward the outcomes in magnitude and direction were further examined using traditional parametric models. For the parametric models, multiple imputation was performed using the MICE package in R (R Core Team, 2013) to impute missing data based on all other variables to create 20 imputed datasets. Random forest results differ based on the dataset so to be robust, variables deemed to be in the top five important variables by random forest in at least one of these imputed datasets were included in a multivariable logistic regression model where pooled adjusted odds ratios (ORs) and 95% confidence intervals (95% CIs) were estimated. ORs were adjusted for all other important variables.

Random forest accounts for correlation but logistic regression does not. After preliminary analyses, acetaminophen use in P1 and P2 in younger mothers were highly correlated (r=0.77), as were reports of paternal substance abuse across time-periods in older mothers (all r>0.88), thus we combined the respective variables in the logistic regression analyses. The resulting variables were any versus no acetaminophen use during P1-P2 and any versus no paternal substance abuse during B1-P2. A sensitivity analysis was also performed additionally adjusting for maternal age (as a continuous variable) to assess residual confounding.

Random forest analyses and multiple imputation were performed in R software using the Party Package and MICE Package, respectively (version 3.4.4, 2018)(R Core Team, 2013). All other analyses were performed using SAS version 9.4 (SAS Institute, Cary, NC, USA).

# Results

Employing random forest analyses for case and control mothers younger than age 20 (Figure 1) and among case and control mothers age 20 or older (Figure 2) identified important predictors of gastroschisis including various dietary, demographic, and behavioral factors. Additionally, random forest was performed on the supplemental dataset for whom data were available on air pollution and pesticide exposures. The addition of these variables did not reveal further insights, i.e., none of the air pollutants nor pesticide exposures emerged as important predictors. Because these environmental exposure variables did not add information, subsequent analyses did not include these variables.

Random forest performed on each of the 20 imputed datasets yielded mostly similar results. Analyses were performed separately for each age stratification and the variables appearing in the top five important predictors at least once were included in subsequent analyses. The 10 variables presenting as important predictors among women younger than age 20 were intake of chocolate, organ meats, apples or pears, and candy, iron, acetaminophen use during P1 and P2, having a urinary tract infection during P1, the number of people supported by the household income, and minutes spent bathing. The nine variables presenting as important among women age 20 or older were maternal race/ethnicity, maternal substance abuse during B1, paternal substance abuse during B1, P1, and P2, pre-pregnancy obesity, cereal intake during B1, paternal age, and parity.

Among women younger than 20 years of age (Table 1), higher odds of delivering infants with gastroschisis were observed for more frequent consumption of chocolate and moderate consumption of candy compared to the lowest consumption, being in the lowest 25% of iron intake compared the middle 50%, acetaminophen use during the first two months of pregnancy and having a urinary tract infection in the first month of pregnancy. After adjustment for other variables, patterns remained the same, although the associations were no longer statistically significant for consumption of chocolate and candy and acetaminophen use. There were inverse (and statistically imprecise) associations observed for consumption of organ meats 1–3x per month and the highest consumption of apples or pears.

Among women age 20 or older (Table 2), higher odds of delivering infants with gastroschisis were observed for US-born women of Hispanic ethnicity or those identifying as Other race/ethnicity compared to white non-Hispanic women. Higher odds were also observed for maternal substance abuse in the month before pregnancy and paternal substance abuse any time during the study period. Inverse associations with gastroschisis were observed for women who were obse pre-pregnancy compared to those who were not, cereal intake in the month before pregnancy, and parity. Paternal age was also inversely associated with gastroschisis among older mothers. However, parental ages were correlated and thus it is probable that the observed association with (younger) paternal age was a proxy for the residual risk associated with younger maternal age (e.g., 20–25 years) within this stratified group of women 20. In the additional sensitivity analysis performed, the addition of maternal age did not alter any of the results for women < 20 but among women 20, maternal age was inversely associated with gastroschisis (aOR 0.88, 95% CI 0.83–0.94) and

associations with paternal age and parity were attenuated and no longer significant. The addition of maternal age did not alter any other results (results not shown).

# Discussion

This analysis utilized a well-established, untargeted statistical approach, random forest, in an effort to find new risk factors for gastroschisis. We explored potential differences in risk factors between younger and older mothers by simultaneously evaluating a large number of variables available in California separately in these groups. We observed different factors to be important predictors in these two populations and thus observed different statistically significant associations. Among women younger than age 20, important predictors associated with gastroschisis were consumption of certain foods, acetaminophen use, and infection, whereas among older women, important predictors associated with gastroschisis were certain maternal race/ethnicities, parental substance abuse, obesity, and paternal age. Exposures to pesticides or various air pollutants did not contribute as important predictors to either maternal age group.

Among women younger than 20, a few of the important variables were nutrients and food intake items. We observed a positive association between low iron intake and delivering an infant with gastroschisis as well as for moderate consumption of candy. The same pattern was observed for moderate amounts of chocolate intake but a suggestive inverse association was observed for intake >6 times per week and for high intake of apples or pears, sources of vitamins such as vitamin C. Previous studies have observed higher odds of gastroschisis with low alpha-carotene intake, low glutathione intake, and high nitrosamine intake after adjustment for other factors (Torfs and others, 1998) and suggestive inverse associations with higher intake of protein, fat, alpha-carotene, and magnesium and inverse associations for higher intake of copper, folate, vitamin B12, and oleic acid (Feldkamp and others, 2011). Another study observed significant inverse associations for both Diet Quality Index and Mediterranean diet score only among Hispanic women (Feldkamp and others, 2014). In the current study, we observed inverse associations with slight increases in organ meat intake among women younger than 20, a source of vitamin A. We also observed an inverse association with cereal intake in the month before pregnancy among women age 20 or older, a food containing folic acid. The collection of these observations indicates that risk among younger women is influenced by aspects of the diet or behaviors related to aspects of the diet. Deeper inquiry in these domains seem warranted.

We observed higher odds of delivering an infant with gastroschisis with acetaminophen use in the beginning of pregnancy among women younger than 20 but after adjustment for other variables such as urinary tract infection the odds were attenuated. Associations between maternal use of various medications and gastroschisis have been studied previously (Ahrens and others, 2013; Alwan and others, 2007; Draper and others, 2008; Feldkamp and others, 2010; Interrante and others, 2017; Lin and others, 2008; Polen and others, 2013; Waller and others, 2010; Werler and others, 1992; Werler and others, 2002). However, acetaminophen was the only important predictor we identified of the medications included in our analysis. A previous analysis in NBDPS observed an inverse association between acetaminophen use and gastroschisis among women who also had an infection and fever (Feldkamp and others,

2010). It is unclear whether the medication or the indication for medication is the risk factor in many studies of medication use. A previous study also observed an association between having a urinary tract infection and gastroschisis among younger mothers (Yazdy and others, 2014) as we did, and another study observed a positive association between having a fever and gastroschisis (Waller and others, 2018) suggesting that further research into the joint effects of infection, medication use and indication, and maternal age may be warranted.

Another factor only found to be important among women age 20 or older was maternal race/ ethnicity. We observed increased odds of having an infant with gastroschisis for women identifying as Hispanic US-born or Other race/ethnicity compared to White Non-Hispanic women. This is consistent with a study that observed higher odds in Hispanic US-born women only among older mothers (Khodr and others, 2013) but not with an age-matched study in California (Torfs and others, 1994).

The same California study did observe a higher odds of delivering an infant with gastroschisis for women with a lower income and a few other markers of lower socioeconomic status and gastroschisis (Torfs and others, 1994). Once multiple imputation was performed, the only marker of socioeconomic status in the top important predictors was number of people supported by the household income and only among women younger than 20. There was an inverse association between the number of people and gastroschisis, however, it is not immediately clear if a higher number of dependents corresponds to higher or lower status.

Among women age 20 or older, three additional factors that were important and associated with gastroschisis were substance abuse, lack of obesity, and parity. In our analyses, substance abuse combined use of marijuana, hash, cocaine, crack, hallucinogens, heroin, mushrooms, and other. Previous studies have observed positive associations between maternal use of marijuana (Torfs and others, 1994) or any recreational drug use (Draper and others, 2008) in the first trimester and gastroschisis. However, these studies did not explore paternal substance abuse. Both maternal and paternal substance abuse were associated with gastroschisis, although they were attenuated after adjustment, possibly due to correlation between the variables. Substance abuse could have biological effects on each parent or could be a proxy of other risky behaviors. Previous studies have also observed higher odds of gastroschisis among women who drank alcohol during the periconceptional period (Richardson and others, 2011). Our study observed and inverse association between prepregnancy obesity and gastroschisis and are consistent with previous findings (Lam and others, 1999; Waller and others, 2007). Additional research into the potential biological mechanisms of this reduced risk association warrant further investigation. We also found there to be an inverse association with increasing parity which adds evidence to another previous observation of a higher odds of gastroschisis for first births (McNeese and others, 2015).

Strengths of this study included the use of the rich data available in California and the use of random forest to simultaneously explore such data. The California center of the NBDPS (Reefhuis and others, 2015) utilizes a standardized questionnaire to collect a wide array of data including some paternal factors. These data include many parental exposures and

activities experienced periconceptionally. Geocoded residences also allowed integration of previously collected detailed pesticide and air pollution data (Padula and others, 2013; Shaw and others, 2014) to be analyzed with the parental factors. The complete case ascertainment from the well-established California Birth Defects Monitoring Program offered a relatively large sample of cases and enabled stratification by maternal age. Use of random forest, as a data-mining approach, was a strength owing to its non-hypothesis driven variable selection and its capability to examine a large number of variables, accounting for all others simultaneously. Limitations of this study included the case-control nature of the study and the potential for recall bias as well as the moderate participation percentages of case and control mothers. Analyses with a large number of variables are also susceptible to collinearity among the variables and while random forest is designed to account for collinearity, standard regression analyses are not. Lastly, due to the nature of data mining such as random forest, there are concerns regarding multiple testing and spurious associations. It is possible that some of the observed associations are due to chance alone.

Despite the unique epidemiologic profile that characterizes gastroschisis, i.e., highly elevated risk for teenage mothers and an increasing prevalence over the past several decades, the identification of explanatory factors for this profile has been only modestly productive. Here we seemingly explored [idiomatically] *everything but the kitchen sink* and despite our novel and comprehensive approach to generate new hypotheses regarding its unique epidemiology, we did not discover many previously unreported associations for gastroschisis. Our results do add evidence to associations observed with many of these proposed risk factors, particularly given such factors were modeled simultaneously with many of the other proposed risk factors. Our current methods for identifying clues to the unique epidemiology of gastroschisis are perhaps in need of a paradigm shift from the usual suspects of birth defects inquiry that relies on what investigators think is important to ask women to recall to a much deeper biologic inquiry that relies on various biomarkers and myriad omics to characterize pregnancies with and without fetuses with gastroschisis.

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# Appendix A. Variables from the California Center Included in Random Forest

Variables were included if they were in both versions of the computer-assisted telephone interview, occurred during the time-period from one month before pregnancy (B1) until the

second month of pregnancy (P1-P2) if time-varying, and had a frequency of 0.1%. Variables included both categorical and continuous responses.

Categorical variables included: maternal and paternal education (<high school, high school, >high school), maternal and paternal race/ethnicity (white non-Hispanic, Hispanic Foreignborn, Hispanic US-born, black non-Hispanic, other), maternal feelings about pregnancy (wanted to be pregnant, wanted to wait until later, did not care, pregnant despite consistent contraceptive use), infant sex (male, female), annual household income (<\$10,000, \$10,000-\$50,000, >\$50,000), timing of pregnancy discovery (first trimester, second/third trimester), and pre-pregnancy obesity (<30 kg/m2, 30 kg/m2).

Continuous variables included: number of people supported with household income, timing of first prenatal visit (month), maternal and paternal age at delivery (years), parity, number of previous pregnancies, miscarriages, caffeine from coffee (mg/day), caffeine from tea (mg/day), caffeine from soda (mg/day), total caffeine (mg/day), time spent per shower (minutes), time spent per bath (minutes), and number of jobs from B1-P2.

Variables with responses of yes or no fell into two categories, overall and by time-period (separate responses for B1, P1, and P2). Overall yes/no variables included: pesticide exposure (Dichlorophenoxy acid or ester, Alcohol/Ether, Alkyl Phthalate, Amide, Aryloxyphenoxy propionic acid, Avermectin, Azole, Bipyridylium, Hydroxybenzonitrile, Insect Growth Regulator, Chloroacetanilide, Chlorinated Phenol, Copper-containing compound, Cyclohexenone derivative, 2,6-Dinitroaniline, Diacylhydrazine, Benzoic acid, Dicarboximide, Dithiocarbamate, Endothall, Phosphonoglycine, Glyco Ether, Halogenated organic, Imidazolinone, Monochlorophenoxy acid or ester, Dithiocarbamate, N-Methyl Carbamate, Neonicotinoid, Organochlorine, Organophosphate, Organoarsenic, Bis-Carbamate, Petroleum derivative, Phenol, Piperonyl, Polyalkyloxy Compound, Pyrethroid, Pyridazinone, Quaternary Ammonium Compound, Silicone, Streptomycin, Strobin, Sulfonylurea, Thiocarbamate, Pyridinecarboxylic acid, Benzimidazole, Thiophthalimide, Triazine, Chloropyridinyl, Urea, Xylylalanine, and Inorganic-Zinc), folic acid-containing vitamin intake, singleton birth, chorionic villus sampling, pre-pregnancy diabetes, use of insulin, use of fertility medication or procedure, medication for pregnancy nausea, prepregnancy high blood pressure, high blood pressure during pregnancy, epilepsy, seizures, medication for seizures, beer intake, wine intake, mixed drink intake, shots of liquor intake, other drink intake, maternal and paternal active military duty, any household participation in occupational pesticide application, maternal and paternal health problems or birth defects diagnosed in childhood, other relative health problems or birth defects diagnosed in childhood, private well drinking water source, and father employed.

Each time-period variable had a separate variable for B1, P1, and P2. These variables included CT/CAT scan, MRI, X-ray, other X-ray or scan, urinary tract infection, pelvic inflammatory disease, other illness, surgery, birth control pill use, other birth control use, pregnancy nausea (P1 and P2 only), cereal intake, food supplement intake, cigarette smoking, smoking in the household, smoking in the workplace or school, alcohol use, hot tub/Jacuzzi/sauna use, and maternal and paternal substance abuse. Substance abuse combined use of marijuana, hash, cocaine, crack, hallucinogens, heroin, mushrooms, and

other. Use of non-steroidal anti-inflammatory (aspirin, ibuprofen, ibuprofen (lysine salt), naproxen, and naproxen sodium), acetaminophen, Nitrofurantoin (nitrofurantoin, nitrofurantoin monohydrate, and nitrofurantoin sodium), anti-depressant (selective serotonin reuptake inhibitors or bupropion), and benzodiazepine were also included as time-period variables with a response of yes or no.

Nutrient and air pollution values were divided into three categories based on values among controls: those with exposure levels <25%, those within the interquartile range, and those with exposure levels >75%. Nutrient variables included dietary intake of: fat (g), total carbohydrate (g), total protein (g), alanine (g), methionine (g), cysteine (g), total choline (mg), betaine (mg), calcium (mg), alpha-carotene ( $\mu$ g), beta-carotene ( $\mu$ g), copper (mg), folate ( $\mu$ g, dietary folate equivalents), iron (mg), lutein ( $\mu$ g), magnesium (mg), niacin (mg), retinol ( $\mu$ g), riboflavin (mg), selenium ( $\mu$ g), thiamin (mg), vitamin A ( $\mu$ g, Retinoic Acid Equivalents), vitamin B6 (mg), vitamin B12 ( $\mu$ g), vitamin C (mg), zinc (mg), glycemic index, and diet quality index (quartiles). Air pollution variables included daily 24-hour averages of nitrogen oxide, nitrogen dioxide, particulate matter <10 $\mu$ m (PM10), particulate matter <2.5 $\mu$ m (PM2.5), and carbon monoxide, and a daily 8-hour maximum of ozone and averaged over the study period. Traffic density was included as a dimensionless indicator based on traffic volumes within a 300-m radius.

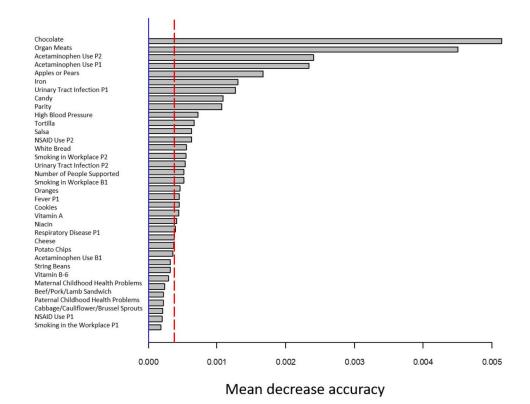
All food frequency questionnaire items were divided into the following categories: <1x per month, 1–3x per month, 1x per week, 2–6x per week, or >6x per week. Variables from the food frequency questionnaire, and the units of measurement, included: skim or low fat milk (8 oz. glass); whole milk (8 oz. glass); yogurt (1 cup); ice cream (1/2 cup); cottage or ricotta cheese (1/2 cup); other cheese (1 slice or 1 oz. serving); margarine (pat); butter (pat); apples or pears (1); oranges (1); orange juice (1 glass); peaches, apricots, plums, or nectarines (1 fresh or 1/2 cup canned); bananas (1); other fruits, fresh, frozen, or canned (1/2 cup); tomatoes (1) or tomato juice (small glass); string beans (1/2 cup); broccoli (1/2 cup); cabbage, cauliflower, or brussel sprouts (1/2 cup); carrots, raw (1/2 carrot or 2-4 sticks); carrots, cooked (1/2 cup); corn (1 ear or 1/2 cup frozen, canned); peas or lima beans (1/2 cup)frozen, canned); yams or sweet potatoes (1/2 cup); spinach or collard greens, cooked (1/2 cup); cup); beans or lentils, baked or dried (1/2 cup); yellow squash (1/2 cup); eggs (1); chicken or turkey (4–6 oz.); bacon (2 slices); hot dogs (1); processed meats, e.g., sausage, salami,bologna, chorizo, etc. (piece or slice); liver (3-4 oz.); hamburger (1 patty); beef, pork, lamb or cabrito as a sandwich or mixed dish, e.g., stew, casserole, lasagna, etc; beef, pork, lamb or cabrito as a main dish,e.g., steak, roast, ham, etc. (4-6 oz.); fish (3-5 oz.); chocolate (1 oz.); candy without chocolate (1oz.); pie (slice); cookies (1); white bread (slice), including pita bread; dark bread (slice), including wheat pita bread; french fried potatoes (4 oz.); potatoes, baked, boiled (1) or mashed (1 cup); rice or pasta e.g., spanish rice, spaghetti, noodles, etc. (1 cup); potato chips or corn chips (small bag or 1 oz.); nuts (small packet or 1 oz.); peanut butter (1 tbs); Oil and vinegar dressing e.g., italian (1 tbs); cantaloupe (1/4)melon); avocado (1) or guacamole (1 cup); raw chile peppers, jalapeno (1); salsa (1 cup); chicken Livers (1 oz.); organ meats, Barbacoa, Menudo, sweetbreads, tongue, intestines (3-4 oz.); tortilla (1); refried beans (1 cup).

# References

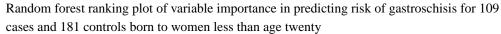
- Ahrens KA, Anderka MT, Feldkamp ML, Canfield MA, Mitchell AA, Werler MM. 2013 Antiherpetic medication use and the risk of gastroschisis: findings from the National Birth Defects Prevention Study, 1997–2007. Paediatric and perinatal epidemiology 27(4):340–345. [PubMed: 23772935]
- Alwan S, Reefhuis J, Rasmussen SA, Olney RS, Friedman JM. 2007 Use of selective serotoninreuptake inhibitors in pregnancy and the risk of birth defects. The New England journal of medicine 356(26):2684–2692. [PubMed: 17596602]
- Canfield MA, Honein MA, Yuskiv N, Xing J, Mai CT, Collins JS, Devine O, Petrini J, Ramadhani TA, Hobbs CA, Kirby RS. 2006 National estimates and race/ethnic-specific variation of selected birth defects in the United States, 1999–2001. Birth defects research Part A, Clinical and molecular teratology 76(11):747–756. [PubMed: 17051527]
- Castilla EE, Mastroiacovo P, Orioli IM. 2008 Gastroschisis: international epidemiology and public health perspectives. American journal of medical genetics Part C, Seminars in medical genetics 148c(3):162–179.
- Croen LA, Shaw GM, Jensvold NG, Harris JA. 1991 Birth defects monitoring in California: a resource for epidemiological research. Paediatric and perinatal epidemiology 5(4):423–427. [PubMed: 1754501]
- Draper ES, Rankin J, Tonks AM, Abrams KR, Field DJ, Clarke M, Kurinczuk JJ. 2008 Recreational drug use: a major risk factor for gastroschisis? American journal of epidemiology 167(4):485–491. [PubMed: 18063593]
- Feldkamp ML, Carmichael SL, Shaw GM, Panichello JD, Moore CA, Botto LD. 2011 Maternal nutrition and gastroschisis: findings from the National Birth Defects Prevention Study. American journal of obstetrics and gynecology 204(5):404.e401–404.e410. [PubMed: 21396620]
- Feldkamp ML, Krikov S, Botto LD, Shaw GM, Carmichael SL. 2014 Better diet quality before pregnancy is associated with reduced risk of gastroschisis in Hispanic women. The Journal of nutrition 144(11):1781–1786. [PubMed: 25332477]
- Feldkamp ML, Meyer RE, Krikov S, Botto LD. 2010 Acetaminophen use in pregnancy and risk of birth defects: findings from the National Birth Defects Prevention Study. Obstetrics and gynecology 115(1):109–115. [PubMed: 20027042]
- Gill SK, Broussard C, Devine O, Green RF, Rasmussen SA, Reefhuis J. 2012 Association between maternal age and birth defects of unknown etiology: United States, 1997–2007. Birth defects research Part A, Clinical and molecular teratology 94(12):1010–1018. [PubMed: 22821755]
- Interrante JD, Ailes EC, Lind JN, Anderka M, Feldkamp ML, Werler MM, Taylor LG, Trinidad J, Gilboa SM, Broussard CS. 2017 Risk comparison for prenatal use of analgesics and selected birth defects, National Birth Defects Prevention Study 1997–2011. Annals of epidemiology 27(10):645– 653.e642. [PubMed: 28993061]
- Jones AM, Isenburg J, Salemi JL, Arnold KE, Mai CT, Aggarwal D, Arias W, Carrino GE, Ferrell E, Folorunso O, Ibe B, Kirby RS, Krapfl HR, Marengo LK, Mosley BS, Nance AE, Romitti PA, Spadafino J, Stock J, Honein MA. 2016 Increasing Prevalence of Gastroschisis--14 States, 1995– 2012. MMWR Morbidity and mortality weekly report 65(2):23–26. [PubMed: 26796490]
- Khodr ZG, Lupo PJ, Canfield MA, Chan W, Cai Y, Mitchell LE. 2013 Hispanic ethnicity and acculturation, maternal age and the risk of gastroschisis in the National Birth Defects Prevention Study. Birth defects research Part A, Clinical and molecular teratology 97(8):538–545. [PubMed: 23729355]
- Lam PK, Torfs CP, Brand RJ. 1999 A low pregnancy body mass index is a risk factor for an offspring with gastroschisis. Epidemiology (Cambridge, Mass) 10(6):717–721.
- Lammer EJ, Iovannisci DM, Tom L, Schultz K, Shaw GM. 2008 Gastroschisis: a gene-environment model involving the VEGF-NOS3 pathway. American journal of medical genetics Part C, Seminars in medical genetics 148c(3):213–218.
- Lin S, Munsie JP, Herdt-Losavio ML, Bell E, Druschel C, Romitti PA, Olney R. 2008 Maternal asthma medication use and the risk of gastroschisis. American journal of epidemiology 168(1):73–79. [PubMed: 18436535]

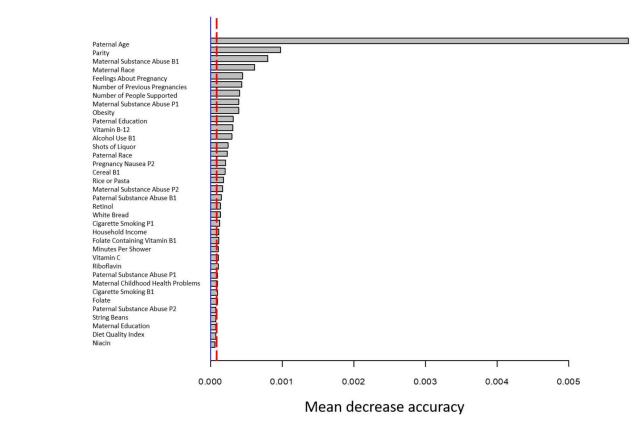
- McNeese ML, Selwyn BJ, Duong H, Canfield M, Waller DK. 2015 The association between maternal parity and birth defects. Birth defects research Part A, Clinical and molecular teratology 103(2): 144–156. [PubMed: 25721953]
- Padula AM, Tager IB, Carmichael SL, Hammond SK, Lurmann F, Shaw GM. 2013 The association of ambient air pollution and traffic exposures with selected congenital anomalies in the San Joaquin Valley of California. American journal of epidemiology 177(10):1074–1085. [PubMed: 23538941]
- Parker SE, Mai CT, Canfield MA, Rickard R, Wang Y, Meyer RE, Anderson P, Mason CA, Collins JS, Kirby RS, Correa A. 2010 Updated National Birth Prevalence estimates for selected birth defects in the United States, 2004–2006. Birth defects research Part A, Clinical and molecular teratology 88(12):1008–1016. [PubMed: 20878909]
- Polen KN, Rasmussen SA, Riehle-Colarusso T, Reefhuis J. 2013 Association between reported venlafaxine use in early pregnancy and birth defects, national birth defects prevention study, 1997– 2007. Birth defects research Part A, Clinical and molecular teratology 97(1):28–35. [PubMed: 23281074]
- R Core Team. 2013 R: A language and environment for statistical computing. Vienna, Austria: R Foundation for Statistical Computing.
- Rasmussen SA, Frias JL. 2008 Non-genetic risk factors for gastroschisis. American journal of medical genetics Part C, Seminars in medical genetics 148c(3):199–212.
- Rasmussen SA, Olney RS, Holmes LB, Lin AE, Keppler-Noreuil KM, Moore CA. 2003 Guidelines for case classification for the National Birth Defects Prevention Study. Birth defects research Part A, Clinical and molecular teratology 67(3):193–201. [PubMed: 12797461]
- Reefhuis J, Gilboa SM, Anderka M, Browne ML, Feldkamp ML, Hobbs CA, Jenkins MM, Langlois PH, Newsome KB, Olshan AF, Romitti PA, Shapira SK, Shaw GM, Tinker SC, Honein MA. 2015 The National Birth Defects Prevention Study: A review of the methods. Birth defects research Part A, Clinical and molecular teratology 103(8):656–669. [PubMed: 26033852]
- Reefhuis J, Honein MA. 2004 Maternal age and non-chromosomal birth defects, Atlanta--1968–2000: teenager or thirty-something, who is at risk? Birth defects research Part A, Clinical and molecular teratology 70(9):572–579. [PubMed: 15368555]
- Richardson S, Browne ML, Rasmussen SA, Druschel CM, Sun L, Jabs EW, Romitti PA. 2011 Associations between periconceptional alcohol consumption and craniosynostosis, omphalocele, and gastroschisis. Birth defects research Part A, Clinical and molecular teratology 91(7):623–630. [PubMed: 21630421]
- Shaw GM, Yang W, Roberts E, Kegley SE, Padula A, English PB, Carmichael SL. 2014 Early pregnancy agricultural pesticide exposures and risk of gastroschisis among offspring in the San Joaquin Valley of California. Birth defects research Part A, Clinical and molecular teratology 100(9):686–694. [PubMed: 24910073]
- St Louis AM, Kim K, Browne ML, Liu G, Liberman RF, Nembhard WN, Canfield MA, Copeland G, Fornoff J, Kirby RS. 2017 Prevalence trends of selected major birth defects: A multi-state population-based retrospective study, United States, 1999 to 2007. Birth defects research 109(18): 1442–1450. [PubMed: 28905502]
- Strobl C, Malley J, Tutz G. 2009 An introduction to recursive partitioning: rationale, application, and characteristics of classification and regression trees, bagging, and random forests. Psychological methods 14(4):323–348. [PubMed: 19968396]
- Torfs CP, Lam PK, Schaffer DM, Brand RJ. 1998 Association between mothers' nutrient intake and their offspring's risk of gastroschisis. Teratology 58(6):241–250. [PubMed: 9894673]
- Torfs CP, Velie EM, Oechsli FW, Bateson TF, Curry CJ. 1994 A population-based study of gastroschisis: demographic, pregnancy, and lifestyle risk factors. Teratology 50(1):44–53. [PubMed: 7974254]
- Vu LT, Nobuhara KK, Laurent C, Shaw GM. 2008 Increasing prevalence of gastroschisis: populationbased study in California. The Journal of pediatrics 152(6):807–811. [PubMed: 18492521]
- Waller DK, Gallaway MS, Taylor LG, Ramadhani TA, Canfield MA, Scheuerle A, Hernandez-Diaz S, Louik C, Correa A. 2010 Use of oral contraceptives in pregnancy and major structural birth defects in offspring. Epidemiology (Cambridge, Mass) 21(2):232–239.

- Waller DK, Hashmi SS, Hoyt AT, Duong HT, Tinker SC, Gallaway MS, Olney RS, Finnell RH, Hecht JT, Canfield MA. 2018 Maternal report of fever from cold or flu during early pregnancy and the risk for noncardiac birth defects, National Birth Defects Prevention Study, 1997–2011. Birth defects research 110(4):342–351. [PubMed: 29094488]
- Waller DK, Shaw GM, Rasmussen SA, Hobbs CA, Canfield MA, Siega-Riz AM, Gallaway MS, Correa A. 2007 Prepregnancy obesity as a risk factor for structural birth defects. Archives of pediatrics & adolescent medicine 161(8):745–750. [PubMed: 17679655]
- Werler MM, Mitchell AA, Shapiro S. 1992 Demographic, reproductive, medical, and environmental factors in relation to gastroschisis. Teratology 45(4):353–360. [PubMed: 1533957]
- Werler MM, Sheehan JE, Mitchell AA. 2002 Maternal medication use and risks of gastroschisis and small intestinal atresia. American journal of epidemiology 155(1):26–31. [PubMed: 11772781]
- Willett WC, Sampson L, Stampfer MJ, Rosner B, Bain C, Witschi J, Hennekens CH, Speizer FE. 1985 Reproducibility and validity of a semiquantitative food frequency questionnaire. American journal of epidemiology 122(1):51–65. [PubMed: 4014201]
- Yazdy MM, Mitchell AA, Werler MM. 2014 Maternal genitourinary infections and the risk of gastroschisis. American journal of epidemiology 180(5):518–525. [PubMed: 25073472]

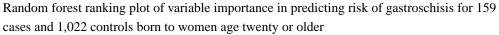


#### Figure 1.





#### Figure 2.



#### Table 1:

Pooled<sup> $\dagger$ </sup> Summary Statistics (Number and Percent for Categorical, Mean and Standard Deviation for Continuous) and Odds Ratios for Important<sup> $\ddagger$ </sup> Predictors of Gastroschisis In Infants Born to Women Age <20 Years, California, 1997-2011

	Cases (n=109)	Controls (n=181)				
Variables	No.(%)	No.(%)	OR	(95% CI)	aOR <sup>§</sup>	(95% CI)
Intake of Chocolate (1 oz.)						
<1x Per Month	22 (20.2)	51 (28.2)	Reference		Reference	
1-3x Per Month	15 (13.8)	39 (21.5)	0.89	(0.41-1.94)	0.57	(0.22-1.51)
1x Per Week	23 (21.1)	25 (13.8)	2.13	(1.00-4.54)	2.16	(0.89-5.24)
2-6x Per Week	40 (36.7)	35 (19.3)	2.65	(1.35-5.20)	2.04	(0.88-4.74)
>6x Per Week	9 (8.3)	31 (17.1)	0.67	(0.28-1.65)	0.49	(0.17-1.42)
Intake of Organ Meats (3-4 oz.)						
<1x Per Month	79 (72.5)	105 (58.0)	Reference		Reference	
1-3x Per Month	15 (13.8)	54 (29.8)	0.37	(0.19-0.70)	0.31	(0.15-0.66)
1x Per Week	13 (11.9)	12 (6.6)	1.44	(0.62-3.33)	1.79	(0.65-4.92)
2-6x Per Week	2 (1.8)	10 (5.5)	NA		NA	
Intake of Apples or Pears <sup>2</sup> (1)						
<1x Per Month	18 (16.5)	24 (13.3)	Reference		Reference	
1-3x Per Month	17 (15.6)	16 (8.8)	1.42	(0.57-3.57)	2.47	(0.76-8.01)
1x Per Week	22 (20.2)	21 (11.6)	1.40	(0.60-3.30)	1.36	(0.49-3.73)
2-6x Per Week	38 (34.9)	67 (37.0)	0.76	(0.37-1.57)	0.75	(0.30-1.83)
>6x Per Week	14 (12.8)	53 (29.3)	0.35	(0.15-0.83)	0.45	(0.16-1.22)
Intake of Candy (1 oz.)						
<1x Per Month	25 (22.9)	62 (34.3)	Reference		Reference	
1-3x Per Month	16 (14.7)	38 (21.0)	1.04	(0.50-2.20)	1.07	(0.43-2.66)
1x Per Week	28 (25.7)	22 (12.2)	3.16	(1.53-6.52)	2.37	(1.00-5.61)
2-6x Per Week	26 (23.9)	36 (19.9)	1.79	(0.90-3.56)	1.61	(0.68-3.82)
>6x Per Week	14 (12.8)	23 (12.7)	1.51	(0.67-3.40)	2.18	(0.80-5.92)
Dietary Intake of Iron						
<25 <sup>th</sup> Percentile	42 (38.5)	42 (23.2)	1.96	(1.13-3.39)	2.19	(1.12-4.30)
25-50 <sup>th</sup> Percentile	49 (45.0)	96 (53.0)	Reference		Reference	
>=75 <sup>th</sup> Percentile	18 (16.5)	43 (23.8)	0.82	(0.43-1.57)	1.07	(0.46-2.49)
Acetaminophen Use P1-P2	43 (39.4)	43 (23.8)	2.09	(1.25-3.50)	1.63	(0.87-3.04)
Urinary Tract Infection P1	11 (10.1)	4 (2.2)	4.97	(1.54-16.0)	4.87	(1.20-19.8)
	Mean (SD)	Mean (SD)	OR	(95% CI)	aOR <sup>§</sup>	(95% CI)
Number Of People Household Income Supports $^{\ensuremath{\mathscr{I}}}$	3.1 (2.3)	3.6 (2.0)	0.89	(0.78-1.01)	0.98	(0.84-1.13)

	Cases (n=109)	Controls (n=181)				
Variables	No.(%)	No.(%)	OR	(95% CI)	aOR <sup>§</sup>	(95% CI)
Minutes Per Bath $^{/\!\!/}$	12.7 (17.3)	17.5 (21.1)	0.99	(0.97-1.00)	0.98	(0.96-1.00)

 $^{\dagger}$ Results were combined and averaged across all 20 imputed datasets

 $\ddagger$  Important variables were determined based on the "varimp" function in the random forest analyses and the resulting mean decrease accuracy metric. Variables deemed to be in the top five important variables in at least one of the imputed datasets were included

 $^{\$}$ Adjusted for the variables included in the table

#### Table 2:

Pooled<sup> $\dagger$ </sup> Summary Statistics (Number and Percent for Categorical, Mean and Standard Deviation for Continuous) and Odds Ratios for Important<sup> $\ddagger$ </sup> Predictors of Gastroschisis In Infants Born to Women Age 20 Years, California, 1997-2011

	Cases (n=159)	Controls (n=1022)				
Variables	No. (%)	No. (%)	<u>OR</u>	<u>(95% CI)</u>	<u>aOR</u> §	<u>(95%CI)</u>
Maternal Race/Ethnicity						
White Non-Hispanic	35 (22.0)	305 (29.8)	Reference		Reference	
Hispanic Foreign-Born	34 (21.4)	359 (35.1)	0.82	(0.50-1.35)	1.06	(0.62-1.82)
Hispanic US-Born	63 (39.6)	239 (23.4)	2.30	(1.47-3.59)	2.14	(1.31-3.49)
Black Non-Hispanic	4 (2.5)	31 (3.0)	1.13	(0.38-3.40)	1.18	(0.36-3.95)
Other	23 (14.5)	88 (8.6)	2.27	(1.28-4.05)	2.27	(1.21-4.27)
Maternal Substance Abuse B1 <sup>¶</sup>	23 (14.5)	37 (3.6)	4.50	(2.60-7.81)	2.19	(1.07-4.49)
Paternal Substance Abuse B1-P2	43 (27.0)	98 (9.6)	3.42	(2.27-5.16)	1.77	(1.04-3.02)
Pre-Pregnancy Obesity ( 30 kg/m <sup>2</sup> )	<u>12 (7.5)</u>	229 (22.4)	0.27	(0.14-0.51)	0.31	(0.16-0.60)
Cereal Intake B1	107 (67.3)	796 (77.9)	0.59	(0.41-0.85)	0.57	(0.38-0.84)
	Mean (SD)	Mean (SD)	<u>OR</u>	<u>(95% CI)</u>	<u>aOR</u> §	<u>(95%CI)</u>
Paternal Age (Years)	<u>25.4 (5.5)</u>	<u>29.8 (6.6)</u>	0.87	(0.84-0.90)	0.90	(0.87-0.94)
Parity <sup>¶</sup> (Previous Live Births)	<u>0.8(1.1)</u>	<u>1.4(1.4)</u>	0.65	(0.55-0.77)	0.80	(0.67-0.96)

 ${}^{\dagger}$ Results were combined and averaged across all 20 imputed datasets

 $\frac{1}{1}$  Important variables were determined based on the "varimp" function in the random forest analyses and the resulting mean decrease accuracy metric. Variables deemed to be in the top five important variables in at least one of the imputed datasets were included

<sup>§</sup>Adjusted for the variables included in the table

 $\frac{1}{N}$ No missing values were imputed for this variable among women age 20