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Indicators of fetal growth and adult liver enzymes: the Bogalusa Heart Study and the Cardiovascular Risk in Young Finns Study

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Abstract

Despite the interest in the relationship of fetal exposures to adult cardiovascular disease, few studies have examined indicators of adult fatty liver disease as an outcome. Previous results are inconsistent, and indicate possible variation by sex. Adult liver enzymes [γ -glutamyl transferase (GGT), alanine transaminase (ALT) and aspartate transaminase (AST)] were measured in two cohort studies: the Bogalusa Heart Study (BHS; $n = 1803$) and the Cardiovascular Risk in Young Finns (YF; $n = 3571$) study, which also had ultrasound measures of liver fat ($n = 2546$). Predictors of dichotomized (clinical cut-offs) and continuous (within the reference range) liver enzymes included low birthweight (<2500 g), macrosomia (>4000 g), small-for-gestational-age (birthweight <10th percentile for gestational age for population), large-for-gestational-age (>90th percentile), and preterm birth. Multiple logistic and linear regression were conducted, adjusted for medical, behavioral and socioeconomic indicators. Interactions with sex were also examined. In BHS, birth measures were not strongly associated with clinically high levels of liver enzymes, and within the reference range measures of reduced growth were associated with increased AST in women. In the YF study, at least one marker of reduced growth was associated with higher GGT, higher ALT and higher AST (in women). Probable fatty liver on ultrasound was associated with low birthweight (2.41, 1.42–4.09) and preterm birth (2.84, 1.70–4.76). These results suggest a link between birth parameters and adult fatty liver, but encourage consideration of population variation in these relationships.

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Conflicts of Interest

None.

Ethical Standards

The authors assert that all procedures contributing to this work comply with the ethical standards of the United States and Finland and with the Helsinki Declaration of 1975, as revised in 2008. All participants gave informed consent and the BHS study was approved by the Institutional Review Board of Tulane University. The original YF study was approved by the local ethics committees.

Supplementary material

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Keywords

birthweight; -glutamyltransferase; gestational age; non-alcoholic fatty liver disease

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Introduction

In animals, prenatal energy deprivation leads to both growth retardation and adult disease,¹ but so does gestational energy overload.² Recently, low birthweight has been associated with non-alcoholic hepatic steatosis (fatty liver disease; FLD) in adulthood in human males.³ FLD (non-alcoholic) is an indicator of hepatic insulin resistance and the metabolic syndrome. Several liver enzymes indicate FLD; γ -glutamyl transferase (GGT) and alanine aminotransferase (ALT), specific markers of liver damage or function, have been most studied.⁴ These enzymes have been independently associated with several cardiovascular risk factors and outcomes, even when levels fall within the reference/normal range.⁵ Other enzymes, such as aspartate transaminase (AST), are also elevated, though not exclusively, in FLD, and have been less studied.⁴

However, human studies of early life influences on liver enzymes are rare. Some studies have been conducted in children, generally finding reduced fetal growth associated increased risk.^{6,7} In adults, one study found a negative association between GGT and birthweight (not statistically significant).⁸ In addition, differences by sex have been reported, with at least one study suggesting an inverse association between birthweight and adult GGT/ALT in women, but potentially a direct association in men.^{9,10} Two studies have considered preterm birth (PTB): in the preterm birth and early life programming of adult health and disease (ESTER) study, Finnish adults who were born preterm had higher risk of fatty liver index [FLI; an index combining information on triglycerides, body mass index (BMI), GGT, and waist circumference¹¹] >30, and to have higher overall ALT and AST levels, but no difference in GGT,¹² while a small Dutch study found that accelerated weight gain in infancy was associated with FLI, whereas birthweight and gestational age were not.¹³

The question of the relationship between birthweight and liver enzymes thus remains an open question, suggesting that further research is needed. Two opposing hypotheses could be put forth on the relationship between birthweight and adult FLD. The first is that those with a genetic or other consistent, lifetime propensity for increased adiposity would be larger at birth and have worse health as adults. The 'thrifty phenotype' hypothesis, on the other hand, indicates that prenatal undernutrition, normally indicated by lower birthweight, is associated with adult cardiovascular disease, possibly due to lasting alterations in energy metabolism.¹⁴⁻¹⁶ We hypothesized that indicators of reduced fetal growth will be associated with higher levels of liver enzymes and increased indicators of fatty liver in adulthood. In this paper, we examine whether the association between indicators of prenatal growth and health (birthweight and gestational age) predicts adult raised liver enzyme levels, both at a clinical level and within the normal range, and whether this relationship varies by sex. We consider these questions in two long-running studies of child and adult cardiovascular health: the Bogalusa Heart Study (BHS) and the Cardiovascular Risk in Young Finns (YF).

Methods

Study populations

The BHS is a long-running, community-based series of cross-sectional studies conducted in Bogalusa, LA. Data from this analysis were taken from three follow-up studies conducted between 1985 and 2002. Standardized protocols were used at each visit, with subjects instructed to fast. For subjects with multiple visits, data at the latest available measurement were used. Among 2598 participants aged 18 and older, 1950 had at least one liver enzyme measurement and one birth outcome measure; there were no differences in sex, race or BMI between those included and excluded, but the included participants tended to be younger (mean age 31.8 v. 33.6, $P < 0.01$). After exclusions for pregnant women ($n = 21$), multiple births ($n = 68$), and those lacking fasting blood samples ($n = 62$; some overlap among groups), 1803 observations were left for analysis. Participants were born between 1938 and 1987 (median year = 1966).

The Cardiovascular Risk in YF study was officially started in 1980 to examine heart disease risk factors in children and adolescents.¹⁷ At baseline, participants were 3–18 years old (i.e. birth years between 1962 and 1977); follow-ups have been regularly conducted every 3–5 years since then, with the most recent in 2011. In total, 3571 participants had data on at least one birth outcome and one liver enzyme. Overall, loss to follow-up has been greater among men, older participants, and smokers.¹⁷ Liver enzymes were measured after 2001; among those who participated in at least one of those exams, whether a participant had liver enzymes measurements did not vary by sex or BMI, but did vary by age (mean age 31.3 v. 34.7, $P < 0.01$).

Exposure measurement

For BHS, data had previously been linked with birth certificates to determine participants' birthweight and gestational age.¹⁸ For YF, participants and their parents reported birthweight and gestational age at an early visit. Such reports, particularly by the mother, have been found to correlate well with clinical records.^{19–22} Both absolute and relative measures of birthweight were examined. Low birthweight (LBW) was defined as birthweight < 2500 g and macrosomia as birthweight > 4000 g. Small-for-gestational-age (SGA) was defined as weight < 10 th percentile and large-for-gestational-age (LGA) as weight > 90 th percentile for gestation by sex with the study population used to define percentiles. For BHS, the included sample had only 7.9% SGA compared with the original sample used to construct the percentiles, implying that it may be somewhat lower-risk group. PTB was defined as birth < 37 weeks' gestation.

Outcome measurement

BHS—GGT and ALT levels were measured as part of a multiple chemistry profile (SMA20) by enzymatic procedures with the multichannel Olympus Au-5000 analyzer (Olympus, Lake Success, NY, USA). FLI was calculated by the method of Bedogni *et al.*,¹¹ incorporating GGT, BMI, triglycerides and waist circumference.

YF—Venous blood samples were drawn after an overnight fast and serum was separated, aliquoted, and stored at -70°C until analysis. From the samples taken in study year 2011, GGT, ALT and AST concentrations were measured by enzymatic methods (GGT, ALT, AST, Glucose, Cholesterol and Triglycerides System Reagent, Beckman Coulter Biomedical, O'Callaghan's Mills, Ireland) on an automatic analyzer (AU400, Olympus, Tokyo, Japan) at the laboratory of the Department of Chronic Disease Prevention, National Institute for Health and Welfare, Turku, Finland. From the samples taken in study year 2001, the GGT, ALT and AST concentrations were measured at the Department of Laboratory Medicine, Konventhospital Barmherzige Brüder Linz, Austria, on an ARCHITECT automated analyzer (Abbott Diagnostics, Abbott Parks, IL, USA). In addition, in the YF study, ultrasound imaging of the liver was performed in 2011 for 2546 study participants using a validated protocol²³ and Sequoia 512 ultrasound mainframes (Acuson, Mountain View, CA, USA) with 4.0 MHz adult abdominal transducers. Evaluation of hepatic steatosis was performed according to liver-to-kidney contrast, parenchymal brightness, deep beam attenuation and bright vessel walls,²⁴ and the presence of hepatic steatosis was assessed visually by a trained ultrasonographer. The participants were classified into probable fatty liver and normal liver groups.²⁵ FLI was calculated as for BHS.

Covariate definitions

All covariates were measured at the time of the liver enzyme measurement and defined as listed in Table 1, unless otherwise specified.

For BHS, physical activity was based on two questions, ranking physical activity at work and outside of work on a five-point scale. For YF, non-work physical activity was based on combined information on frequency, intensity and duration of physical exercise.²⁶

Modeling strategy

The study designs were thought to differ sufficiently that combining the data for analysis would not be appropriate; therefore, results are presented separately by study.

First, a dichotomous outcome with levels indicating possible clinical concern based on the literature (for men, GGT >68 U/l, ALT >59 U/l; for women, GGT >40 U/l, ALT >41 U/l;²⁷ for all, AST >34 U/l, FLI ≥ 60 ¹¹) were examined using logistic regression; in addition, ultrasound fatty liver was examined for YF participants. Next, outcomes within the reference range (in order to examine higher risk individuals without clinical disease and to exclude those with clinically increased liver enzymes due to non-metabolic causes²⁸) were examined (liver enzymes levels were log-transformed) using linear models. Generally, three sets of models were examined. The first was unadjusted. The second included covariates identified based on a directed acyclic graph (DAG) using the DAGitty program.²⁹ The same covariates were included in each model: age, sex, race, smoking, alcohol use, education, income and physical activity. Depending on one's causal model, adult BMI could be a confounder (if higher genetic propensity to be large leads to higher birthweight and higher adult BMI) or an intermediate (if intrauterine undernutrition leads to later adiposity)³⁰. In the first case, adjustment would be appropriate; in the second, it could in fact create a spurious association.³¹ The final set of models includes the BMI adjustment. Because of substantial

missing data for some of the covariates, particularly income (3% for BHS), SES (30% in YF) and physical activity (5% for BHS, 10% for YF), multiple imputation was used (using SAS's PROC MI and PROC MIANALYZE). Finally, interactions with sex was assessed by examining stratified analyses and the product term in multivariable models. For ease of presentation, SGA, LGA and PTB are presented in the tables; information on LBW and macrosomia is provided in the supplementary material.

As alcohol consumption is a common cause of FLD, a sensitivity analysis excluded heavy drinkers, defined in BHS as daily drinking or drinking five or more drinks at one time more than 1 ×/week in the past month reported at any exam more drinks at one time two or more times/month ($n = 736$, 23.9%), at any visit. Results were very similar to those shown.

All participants gave informed consent and the BHS study was approved by the Institutional Review Board of Tulane University. The original YF study was approved by the local ethics committees.

Results

Participants in both studies were young adults; however, the YF sample was older at the time of measurement, leaner and had more favorable cardiovascular indicators than the BHS sample (Table 1). All liver enzyme measures were strongly correlated with each other, with BMI, and with lipid measures in both samples (Table S1). In all, 391 (15.4%) of the YF sample had indications of fatty liver on ultrasound.

BHS

For the dichotomous outcomes (Table 2), birth measures were not strongly associated with high levels of liver enzymes, although there was a tendency for LGA to be associated with higher GGT (adjusted odds ratio (aOR) 1.58, 95% confidence interval (CI) 1.00–2.51; after adjustment for BMI, aOR 1.52, 0.96–2.43) and for SGA to be less likely to have high ALT (fully aOR 0.31, 95% CI 0.09–1.02). LBW was also associated with a reduced likelihood of being in the high FLI group (aOR 0.53, 0.33–0.85; Table S2).

When liver enzymes were considered as a continuous outcome, some interactions were found between sex and the continuous outcome measures (Tables 3 and S3), so stratified results are provided. When BMI was adjusted for, LBW and PTB were associated with higher AST (among women only) and lower FLI (men only). Macrosomia was associated with lower GGT (women only) in some models.

Cardiovascular Risk in YF

For dichotomous outcomes (Table 4), the strongest association was with LBW, associated with high GGT (aOR 1.68, 95% CI 1.02–2.76; adjusted for BMI, 1.65, 0.98–2.77) and FLI (1.98, 1.33–2.95) (Tables 4 and S4). For continuous markers (Table 5), at least one marker (LBW, PTB or SGA) was associated with higher GGT (SGA, women only), ALT (men and women), AST (stronger in women; Table 5) and higher FLI (women only; Table S5). At least one marker of increased growth was associated with lower GGT (stronger in women)

and lower ALT (stronger in men). SGA, LBW and PTB were also associated with increased likelihood of probable fatty liver on ultrasound (Table 6).

Discussion

The BHS data more strongly supported the idea that some individuals have higher relative weight and metabolic risk throughout their lifetimes: LGA was associated with high GGT, whereas SGA/LBW was associated with lower ALT and FLI in some analyses. High ALT in early-mid pregnancy was associated with LGA in one study.³² The YF sample showed results more consistent with the thrifty phenotype hypothesis, with LBW/SGA associated with higher GGT, ALT and FLI, as well as probable fatty liver on ultrasound. Previous studies have found inverse associations between birthweight and the enzymes studied here, although the exact associations have varied by study.^{7,8,10,33} Inverse associations (lower birthweight/higher liver enzymes) have been more commonly found in women,^{3,9,10} which is mostly consistent with our data for GGT. The extent to which there are sex differences in developmental programming is unclear, with conflicting results in the relatively few studies available.³⁴ Several studies have found the association between LBW and type 2 diabetes to be stronger in women,³⁵ but the association between LBW and hypertension has sometimes been found to be stronger in men.³⁴ However, birthweight could interact with sex-specific hormone systems; gene expression; diet, alcohol use or smoking; or growth patterns for differential effects, and animal studies of fetal programming have often found variation by sex.^{34,36}

We are not aware of another study that has examined PTB as a predictor of later ultrasound fatty liver, although women who enter pregnancy with fatty liver have been found to have a higher risk of PTB.³⁷ Late PTB was associated with higher ALT, AST and FLI (though not GGT) values in another study;¹² the majority of the PTB in our study were late, but the mean differences in both of our cohorts are much smaller than found in this previous study. Another study found that accelerated weight gain in infancy was associated with FLI but LBW and gestational age were not, but this study was limited to a young adults born preterm.¹³

Support for the hypotheses that both low birthweight and high birthweight are associated with liver fat can be found in the animal literature. Higher-fat diets have been associated with both increased and decreased weight and corresponding adverse effects on the liver.^{2,38–40} Some models^{39,41} indicate differential effects of prenatal diets on fetal outcomes in different strains, which suggests there may be variation by population or genetics. The study populations in this analysis were different in some ways, including race/ethnicity, but no clear risk factor (age, BMI, race) emerged as a cause for the difference. The analyses were repeated within the white portion of the BHS cohort, and results are more similar to black BHS participants than to the YF (data not shown). However, most of the associations had overlapping confidence intervals; the one exception is the opposite-direction associations between LBW/PTB and FLI in the two cohorts. Possible explanations for the difference include measurement error, or genetic, cultural, or life course differences between the populations. If, as has been suggested in some studies,^{13,42} catch-up growth is a stronger

predictor of adult metabolic health than birthweight *per se*, it is possible that the postnatal growth patterns differ between the two cohorts.

Strengths of the study include the fairly large sample size and the ability to assess the study question in two different cohorts. This analysis incorporates large studies of young adults and includes a diverse, U.S.-based population, unlike previous research.^{3,4,9,10} Limitations include the lack of a direct assessment of some relevant covariates, such as liver fat concentrations, body fat mass and distribution, diet, and substance use, and the lack of standardization of the timing of the measures. A single outcome measure was used; liver enzymes have substantial variability and are raised in a number of clinical conditions.⁴³ We cannot take into account hepatosteatosis due to hepatotoxic medication, and only a limited adjustment can be made for alcohol use (based on self-report), although a sensitivity analysis showed excluding heavy drinkers had little effect on the results (data not shown). Ultrasound is not a direct measurement of liver fat, although the echogenicity of the liver is an indicator of probable fatty deposition, and is used by most general medicine doctors. A large number of comparisons were made, so type 1 error is a possibility, particularly for interaction analyses. As in any study of birth outcomes and adult cardiovascular parameters, no mechanism can be directly demonstrated, and the possibility of unmeasured confounders is always a possibility. We do not have information on the specific causes of growth restriction or preterm birth, and so cannot distinguish between possible causes such as placental dysfunction and infection.⁴⁴ Pre-eclampsia is a cause of many medically indicated preterm births, and has been associated with many later metabolic changes.⁴⁵ Although adult adiposity can directly cause FLD, birthweight would not be a direct cause, and so the relationship would reflect an underlying nutritional, genetic or growth link between the two.

These results contribute to the literature on the relationship between birthweight and adult FLD, suggesting a physiological link between these characteristics. This analysis adds to the growing body of research suggesting intrauterine influences on adult cardiovascular health. It also, however, encourages consideration of population and genetic variation in these relationships.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Table 1

Participants in the Bogalusa Heart Study, 1995–2002 ($n = 1803$) and Cardiovascular Risk in Young Finns Study (2001–2011, $n = 3571$)

	<i>n</i>	%
Bogalusa Heart Study		
Age at oldest visit		
18 to <20	40	2.2
20 to <30	590	32.7
30 to <40	933	51.8
40 +	240	13.3
Sex		
Male	794	44.0
Female	1009	56.0
Race		
Black	556	30.8
White	1247	69.2
BMI category		
<20	126	7.0
20 to <25	525	29.2
25 to <30	516	28.7
30 +	632	35.1
Smoking history		
Never	942	55.8
Former	179	10.6
1–15 cigarettes/day	319	18.9
>15 cigarettes/day	249	14.7
Income		
<\$15 K	554	31.8
\$15K–\$30 K	409	23.5
>\$30K–\$45 K	280	16.1
>\$45 K	501	28.7
Physical activity (combined work and non-work)		

	<i>n</i>	%			
Low activity	243	14.2			
Medium	289	16.9			
High	614	35.9			
Very active	565	33.0			
Alcohol use (in year of enzyme measure)					
Do not drink	1076	60.5			
Drink regularly	703	39.5			
Less than 1 × a week	205	11.5			
1–2 × a week	296	16.6			
3–4 × a week	104	5.9			
Daily or almost daily	98	5.5			
Preterm birth	97	6.1			
Low birthweight	114	6.3			
Small-for-gestational-age	125	7.9			
Large-for-gestational-age	158	10.0			
	Mean	Median	S.D.	Min	Max
GGT ^a (U/l)	35.3	22.0	53.4	3.0	750.0
ALT (U/l)	23.9	18.0	20.9	3.0	293.0
AST (U/l)	23.8	20.0	17.7	5.0	385.0
FLI ^b	43.6	35.0	34.3	0.5	100.0
BMI (kg/m ²)	28.5	27.2	7.1	15.2	62.0
Triglycerides (mg/dl)	120.9	95.0	99.7	23.0	1463.0
Young Finns Study					
Age at oldest visit					
18 to <20	0				
20 to <30	157	4.5			
30 to <40	1401	40.4			
40 +	1913	55.1			
Sex					
Male	1082	31.2			
Female	2389	68.8			

	<i>n</i>	%	
Race			<i>c</i>
Black			
White			
BMI category			
<20	196	5.7	
20– <25	1347	39.3	
25– <30	1207	35.2	
30 +	676	19.7	
Smoking history			
Never	1692	51.1	
Former	779	23.5	
Current	838	25.3	
SES in 2007 (based on occupation)			
Manual	847	34.0	
Lower non-manual	499	20.0	
Upper non-manual	1144	45.9	
Physical activity (non-work)			
Low activity	286	9.0	
Medium	853	26.8	
High	1342	42.1	
Very active	706	22.2	
Alcohol intake (average drinks/day) at liver enzyme visit			
0	944	28.7	
1	1668	50.7	
>1–2	410	12.5	
>2	266	8.1	
Preterm birth	141	4.1	
Low birthweight	116	3.3	
Small-for-gestational-age	346	10.0	
Large-for-gestational-age	335	9.7	
	Mean	Median	S.D. Min Max

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	n		%		
GGT ^a (U/l)	29.1	20.0	33.3	5.0	562
ALT (U/l)	15.6	12.0	12.8	2.0	253
AST (U/l)	22.1	12.0	12.0	10.0	208
FLI ^b	36.7	25.1	30.5	1.2	100
BMI (kg/m ²)	26.3	25.5	5.1	16.2	58.5
Triglycerides (mg/dl)	115.0	97.3	88.5	26.6	3006.2

BMI, body mass index; SES, socioeconomic status; GGT, γ -glutamyl transferase; ALT, alanine transaminase; AST, aspartate transaminase; FLI, fatty liver index.

^a At oldest visit.

^b
$$FLI = (e^{0.953 \times \ln(\text{triglycerides}) + 0.139 \times BMI + 0.718 \times \ln(\text{ggt}) + 0.053 \times \text{waist circumference} - 15.745}) / (1 + e^{0.953 \times \ln(\text{triglycerides}) + 0.139 \times BMI + 0.718 \times \ln(\text{ggt}) + 0.053 \times \text{waist circumference} - 15.745}) \times 100.$$

^c Race/ethnicity information was not collected as such in the Young Finns Study; however, the vast majority is White.

Table 2

Birth outcomes as predictors of clinically high levels of liver enzymes, the Bogalusa Heart Study ($n = 1803$)

	Unadjusted		Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity		Adjusted for previous + BMI	
	OR ^a	95% CI	OR	95% CI	OR	95% CI
γ -glutamate transferase (GGT)						
Small-for-gestational-age	1.25	(0.76, 2.07)	0.91	(0.53, 1.55)	0.96	(0.56, 1.64)
Large-for-gestational-age	1.45	(0.94, 2.24)	1.58	(1.00, 2.51)	1.52	(0.96, 2.43)
Preterm birth	1.22	(0.69, 2.17)	1.17	(0.64, 2.15)	1.20	(0.65, 2.21)
Alanine transaminase (ALT)						
Small-for-gestational-age	0.31	(0.10, 1.00)	0.32	(0.10, 1.03)	0.31	(0.09, 1.02)
Large-for-gestational-age	1.01	(0.53, 1.92)	0.94	(0.48, 1.84)	0.88	(0.45, 1.73)
Preterm birth	1.58	(0.80, 3.14)	1.48	(0.72, 3.03)	1.52	(0.73, 3.15)
Aspartate transaminase (AST)						
Small-for-gestational-age	0.67	(0.34, 1.30)	0.48	(0.24, 0.99)	0.50	(0.24, 1.01)
Large-for-gestational-age	1.09	(0.66, 1.81)	1.12	(0.65, 1.92)	1.09	(0.64, 1.88)
Preterm birth	1.39	(0.77, 2.51)	1.13	(0.60, 2.15)	1.14	(0.60, 2.17)
Fatty liver index (FLI) ^b						
Small-for-gestational-age	0.77	(0.52, 1.14)	0.71	(0.47, 1.08)	Measurement includes BMI	
Large-for-gestational-age	1.15	(0.82, 1.61)	1.25	(0.88, 1.79)		
Preterm birth	0.75	(0.48, 1.16)	0.75	(0.47, 1.21)		

BMI, body mass index, OR, odds ratio, CI, confidence interval.

^aReference category for all models is all participants not categorized as 'case'. Cut-offs for clinical concern: for men, GGT >68 U/l, ALT >59 U/l; for women, GGT >40 U/l, ALT >41 U/l;²⁷ for all, AST >34 U/l, FLI 60.¹¹

^b $FLI = (e^{0.953 \times \ln(\text{triglycerides}) + 0.139 \times \text{BMI} + 0.718 \times \ln(\text{ggt}) + 0.053 \times \text{waist circumference} - 15.745}) / (1 + e^{0.953 \times \ln(\text{triglycerides}) + 0.139 \times \text{BMI} + 0.718 \times \log_e(\text{ggt}) + 0.053 \times \text{waist circumference} - 15.745}) \times 100$.

Table 3

Birth outcomes as predictors of continuous liver enzymes (log-transformed, within the normal range), with adjustment for confounders, the Bogalusa Heart Study

	Model 1				Model 2			
	Unadjusted				Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity			
	β^a	SE	P	P for interaction	β	SE	P	P for interaction
γ -glutamate transferase (GGT)								
Men								
Small-for-gestational-age	0.01	0.07	0.89	0.19	-0.05	0.07	0.47	0.13
Large-for-gestational-age	-0.01	0.06	0.82	0.46	0.02	0.06	0.79	0.46
Preterm birth	-0.04	0.08	0.62	0.17	-0.05	0.08	0.48	0.08
Women								
Small-for-gestational-age	0.12	0.05	0.01		0.07	0.05	0.14	
Large-for-gestational-age	-0.07	0.05	0.15		-0.04	0.05	0.41	
Preterm birth	0.09	0.06	0.12		0.09	0.06	0.13	
Alanine transaminase (ALT)								
Men								
Small-for-gestational-age	-0.02	0.07	0.76	0.69	-0.02	0.07	0.79	0.67
Large-for-gestational-age	0.02	0.06	0.68	0.59	0.02	0.06	0.76	0.74
Preterm birth	-0.07	0.07	0.37	0.28	-0.01	0.07	0.93	0.21
Women								
Small-for-gestational-age	0.01	0.05	0.81		0.02	0.05	0.62	
Large-for-gestational-age	0.06	0.05	0.19		0.04	0.05	0.45	
Preterm birth	0.04	0.06	0.55		0.09	0.06	0.13	
Aspartate transaminase (AST)								
Men								
Small-for-gestational-age	0.00	0.04	0.92	0.54	0.00	0.04	0.99	0.53
Large-for-gestational-age	-0.04	0.03	0.23	0.60	-0.04	0.03	0.22	0.59
Preterm birth	0.01	0.04	0.87	0.06	0.00	0.04	0.96	0.06
Women								

	Model 1				Model 2			
	Unadjusted				Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity			
	β^a	SE	P	P for interaction	β	SE	P	P for interaction
Small-for-gestational-age	0.03	0.03	0.26		0.04	0.03	0.22	
Large-for-gestational-age	-0.01	0.03	0.62		-0.03	0.03	0.38	
Preterm birth	0.11	0.04	<0.01		0.11	0.04	<0.01	
Fatty liver index (FLI) ^b								
Men								
Small-for-gestational-age	-10.49	4.93	0.03	0.09	-8.44	4.87	0.08	0.13
Large-for-gestational-age	3.44	4.00	0.39	0.53	2.88	3.85	0.45	0.52
Preterm birth	-12.60	5.13	0.01	0.03	-6.74	5.04	0.18	0.07
Women								
Small-for-gestational-age	0.50	4.01	0.90		-4.24	3.90	0.28	
Large-for-gestational-age	-0.09	3.90	0.98		3.69	3.78	0.33	
Preterm birth	2.70	4.76	0.57		-0.41	4.76	0.93	
Model 3								
Previous +BMI								
	β	SE	P	P for interaction				
γ -glutamate transferase (GGT)								
Men								
Small-for-gestational-age	-0.05	0.07	0.46	0.15				
Large-for-gestational-age	0.01	0.06	0.84	0.51				
Preterm birth	-0.05	0.07	0.47	0.07				
Women								
Small-for-gestational-age	0.08	0.05	0.08					
Large-for-gestational-age	-0.05	0.05	0.29					
Preterm birth	0.11	0.06	0.07					
Alanine transaminase (ALT)								
Men								

	Model 1					Model 2				
	Unadjusted					Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity				
	β^a	SE	P	P for interaction		β	SE	P	P for interaction	
Small-for-gestational-age	-0.03	0.07	0.69		0.62					
Large-for-gestational-age	0.01	0.05	0.85		0.85					
Preterm birth	0.00	0.07	0.98		0.23					
Women										
Small-for-gestational-age	0.04	0.05	0.41							
Large-for-gestational-age	0.01	0.05	0.79							
Preterm birth	0.11	0.06	0.08							
Aspartate transaminase (AST)										
Men										
Small-for-gestational-age	0.00	0.04	0.96		0.50					
Large-for-gestational-age	-0.04	0.03	0.21		0.58					
Preterm birth	0.00	0.04	0.94		0.06					
Women										
Small-for-gestational-age	0.04	0.03	0.18							
Large-for-gestational-age	-0.03	0.03	0.35							
Preterm birth	0.11	0.04	<0.01							
Fatty liver index (FLI)	BMI included in measure									

BMI, body mass index.

^aIncrease in outcome in those with and without the listed birth outcome.

^bNot log-transformed.

Table 4

Birth outcomes as predictors of clinically high levels^a of liver enzymes, Cardiovascular Risk in Young Finns Study ($n = 3571$)

	Unadjusted		Adjusted for age, sex, smoking, alcohol use, social class, physical activity		Adjusted for previous + BMI	
	OR	95% CI	OR	95% CI	OR	95% CI
γ -glutamate transferase (GGT)						
Small-for-gestational-age	1.24	(0.89, 1.74)	1.15	(0.82, 1.63)	1.23	(0.86, 1.75)
Large-for-gestational-age	1.29	(0.92, 1.80)	1.35	(0.96, 1.91)	1.28	(0.89, 1.85)
Preterm birth	1.22	(0.74, 2.03)	1.21	(0.72, 2.04)	1.17	(0.68, 2.01)
Alanine transaminase (ALT)						
Small-for-gestational-age	0.82	(0.33, 2.06)	0.77	(0.30, 1.96)	0.83	(0.32, 2.11)
Large-for-gestational-age	1.45	(0.68, 3.08)	1.46	(0.70, 3.15)	1.37	(0.63, 2.98)
Preterm birth	1.71	(0.61, 4.78)	1.58	(0.55, 4.50)	1.50	(0.50, 4.45)
Aspartate transaminase (AST)						
Small-for-gestational-age	0.82	(0.33, 2.06)	0.77	(0.30, 1.96)	0.83	(0.32, 2.11)
Large-for-gestational-age	1.45	(0.68, 3.08)	1.46	(0.70, 3.15)	1.37	(0.63, 2.98)
Preterm birth	1.71	(0.61, 4.78)	1.58	(0.55, 4.50)	1.50	(0.50, 4.45)
Fatty liver index (FLI)						
Small-for-gestational-age	1.17	(0.89, 1.55)	1.13	(0.87, 1.47)	Measurement includes BMI	
Large-for-gestational-age	1.06	(0.80, 1.41)	0.97	(0.74, 1.28)		
Preterm birth	1.31	(0.88, 1.96)	1.36	(0.92, 2.02)		

OR, odds ratio; CI, confidence interval.

^aReference category for all models is all participants not categorized as 'case'. Cut-offs for clinical concern: for men, GGT >68, ALT >59; for women, GGT >40, ALT >41;²⁷ for all, AST >34, FLI >60.¹¹

Table 5

Birth outcomes as predictors of continuous liver enzymes (log-transformed, within the normal range), Cardiovascular Risk in Young Finns Study

	Model 1				Model 2			
	Unadjusted		<i>P</i> for interaction		Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity		<i>P</i> for interaction	
	β^a	SE	<i>P</i>		β	SE	<i>P</i>	
γ -glutamate transferase (GGT)								
Men								
Small-for-gestational-age	0.01	0.05	0.79	0.19	0.00	0.05	0.96	0.08
Large-for-gestational-age	-0.03	0.05	0.61	0.64	-0.02	0.05	0.66	0.46
Preterm birth	0.13	0.08	0.09	0.02	0.10	0.07	0.18	0.04
Women								
Small-for-gestational-age	0.08	0.00	<0.01		0.09	0.03	<0.01	
Large-for-gestational-age	-0.05	0.03	0.06		-0.06	0.03	0.03	
Preterm birth	-0.06	0.04	0.15		-0.06	0.04	0.17	
Alanine transaminase (ALT)								
Men								
Small-for-gestational-age	0.06	0.05	0.22	0.91	0.04	0.05	0.38	0.98
Large-for-gestational-age	-0.01	0.05	0.80	0.78	-0.01	0.05	0.82	0.77
Preterm birth	0.15	0.07	0.03	<0.01	0.14	0.07	0.05	<0.01
Women								
Small-for-gestational-age	0.05	0.03	0.06		0.05	0.03	0.09	
Large-for-gestational-age	0.00	0.03	0.93		0.01	0.03	0.71	
Preterm birth	-0.07	0.04	0.10		-0.06	0.04	0.13	
Aspartate transaminase (AST)								
Men								
Small-for-gestational-age	0.02	0.02	0.36	0.37	0.02	0.02	0.38	0.32
Large-for-gestational-age	-0.01	0.03	0.80	0.42	-0.03	0.03	0.26	0.22
Preterm birth	0.07	0.04	0.06	0.27	-0.01	0.03	0.65	0.22
Women								
Small-for-gestational-age	0.05	0.02	<0.01		0.05	0.02	<0.01	

	Model 1					Model 2				
	Unadjusted					Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity				
	β^a	SE	P	P for interaction		β	SE	P	P for interaction	
Large-for-gestational-age	0.02	0.02	0.26			0.02	0.02	0.23		
Preterm birth	0.02	0.03	0.51			0.01	0.03	0.69		
Fatty liver index (FLI) ^a										
Men										
Small-for-gestational-age	0.86	3.09	0.78	0.85		-0.91	2.85	0.75	0.37	
Large-for-gestational-age	7.33	3.23	0.02	0.01		5.30	2.96	0.07	0.03	
Preterm birth	1.35	4.49	0.76	0.96		0.93	4.21	0.83	0.96	
Women										
Small-for-gestational-age	1.56	2.01	0.44			2.03	1.96	0.30		
Large-for-gestational-age	-2.33	1.98	0.24			-1.90	1.98	0.34		
Preterm birth	1.61	3.10	0.60			1.86	3.12	0.55		
Model 3										
Previous +BMI										
	β	SE	P	P for interaction						
γ -glutamate transferase (GGT)										
Men										
Small-for-gestational-age	0.02	0.05	0.68	0.14						
Large-for-gestational-age	-0.05	0.05	0.29	0.81						
Preterm birth	0.12	0.07	0.07	0.01						
Women										
Small-for-gestational-age	0.09	0.03	<0.01							
Large-for-gestational-age	-0.05	0.03	0.05							
Preterm birth	-0.06	0.04	0.11							
Alanine transaminase (ALT)										
Men										
Small-for-gestational-age	0.06	0.04	0.17	0.94						

	Model 1					Model 2				
	Unadjusted					Adjusted for age, sex, race, smoking, alcohol use, education, income, physical activity				
	β^a	SE	P	P for interaction		β	SE	P	P for interaction	
Large-for-gestational-age	-0.05	0.05	0.29							0.56
Preterm birth	0.16	0.07	0.02							<0.01
Women										
Small-for-gestational-age	0.06	0.03	0.04							
Large-for-gestational-age	0.00	0.03	0.90							
Preterm birth	-0.07	0.04	0.07							
Aspartate transaminase (AST)										
Men										
Small-for-gestational-age	0.03	0.02	0.26							<i>b</i>
Large-for-gestational-age	-0.02	0.03	0.36							0.03
Preterm birth	0.08	0.04	0.04							0.96
Women										
Small-for-gestational-age	0.05	0.02	<0.01							
Large-for-gestational-age	0.02	0.02	0.24							
Preterm birth	0.01	0.03	0.65							
Fatty liver index (FLI)	BMI included in measurement									

^aNot log-transformed.^bInteraction model failed to converge.

Table 6

Birth outcomes as predictors of probable fatty liver on ultrasound, the Cardiovascular Risk in Young Finns Study (*n*=2546)

	Unadjusted		Adjusted for age, sex, smoking, alcohol use, social class, physical activity		Adjusted for previous + BMI	
	OR ^a	95% CI	OR	95% CI	OR	95% CI
Small-for-gestational-age	1.82	(1.32, 2.50)	1.63	(1.16, 2.29)	1.75	(1.21, 2.53)
Large-for-gestational-age	0.78	(0.53, 1.14)	0.78	(0.52, 1.16)	0.68	(0.44, 1.05)
Preterm birth	2.46	(1.59, 3.79)	2.75	(1.72, 4.39)	2.84	(1.70, 4.76)
Low birthweight	2.59	(1.58, 4.25)	2.41	(1.42, 4.09)	2.33	(1.31, 4.16)
Macrosomia	0.82	(0.60, 1.14)	0.70	(0.50, 0.99)	0.60	(0.41, 0.88)

OR, odds ratio; CI, confidence interval.

^aReference category for all models is all participants not categorized as 'case'.