

# Background

- The rate of marijuana (MJ) use among pregnant patients has risen over the past decade.<sup>1</sup>
- There is growing evidence to support that today's marijuana is more potent and consumed in higher quantities than the previous decade.<sup>2</sup>
- Studies have drawn mixed conclusions on whether maternal MJ use during pregnancy correlates to adverse fetal outcomes
- Fetal biometrics or measurements of fetal femur (FL), humerus (HL), biparietal diameter (BPD), head and abdominal circumferences (HC, AC) via ultrasound (US) can be used to calculate estimated fetal weight and monitor fetal growth.<sup>6</sup>

## Objectives

To identify if there is an increased risk for abnormal fetal growth as determined by fetal biometrics on 2<sup>nd</sup> trimester US in fetuses exposed to MJ in-utero.

## Methods

## Study Design and Data Abstraction

- This is a retrospective cohort study of pregnant patients (pts) with urine drug screens (UDS) from Jan. 2012 to Dec. 2018.
- Cases included pts with positive MJ UDS while controls were identified by negative MJ UDS.
- Controls were matched to cases 1:1 on maternal age at delivery (18-25, 26-35, or >35), parity at entry into care, fetal sex, insurance status (private, income-dependent, and uninsured), and year of delivery (within 4 years of case delivery year).
- Excluded were pts <18 yo. at time of conception, multifetal pregnancies, deliveries outside the UCHealth system, those missing a 2<sup>nd</sup> trimester US (defined as 16-27 weeks in this study), or US with missing FL, HL, BPD, HC, and or AC.

## Statistical Methods

- The MJ exposed infants were compared to the control infants on all growth parameters (BPD, HC, AC, FL, EFW, and growth percentile) via Mann-Whitney U tests for continuous parameters, and Chi square or Fisher's exact tests for categorical parameters.
- In order to see a 5% difference in parameters we estimated a sample size of 200 cases and 200 controls (N=400) would be needed at a power of 90% and a p of <0.01.

# Table 1:

Maternal Chara Age

Parity Nulliparous a

Multiparous a

Race/ethnicity Non-Non-F Non-

Baby Sex

nsurance

Uninsur

Delivery/Pregna

2012-2014-2 2016-2018-

Sestational age Maternal pre-pr

weight (kg) Maternal delive

Comorbidities

Pre-existing

Gestationa

Medication use Prescript MAT (Suboxone

Substance use Self-repo

MJ use prio

Tobacco use

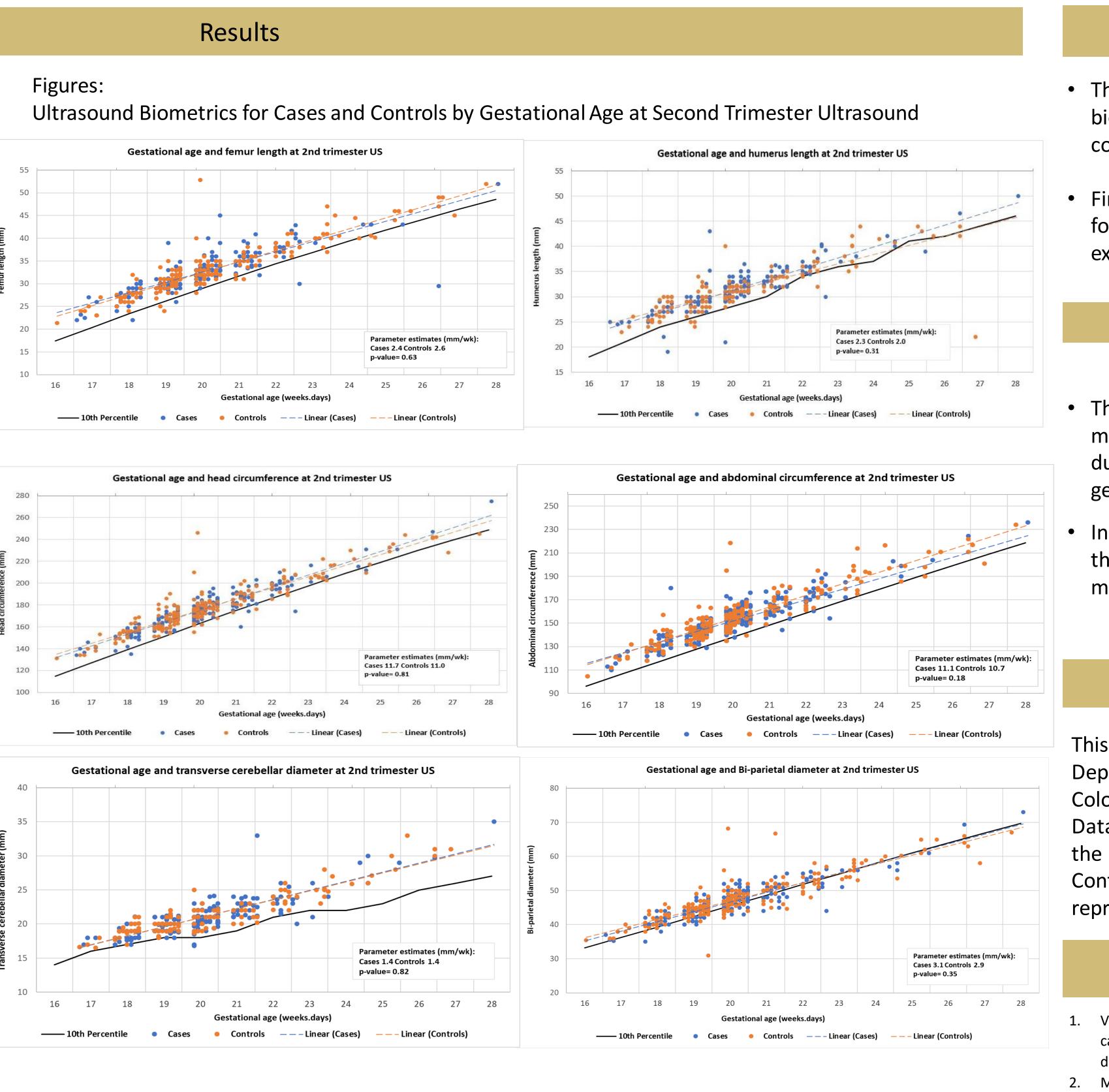
Alcohol us Opioid us

Methamph

Cocaine use in pregnancy

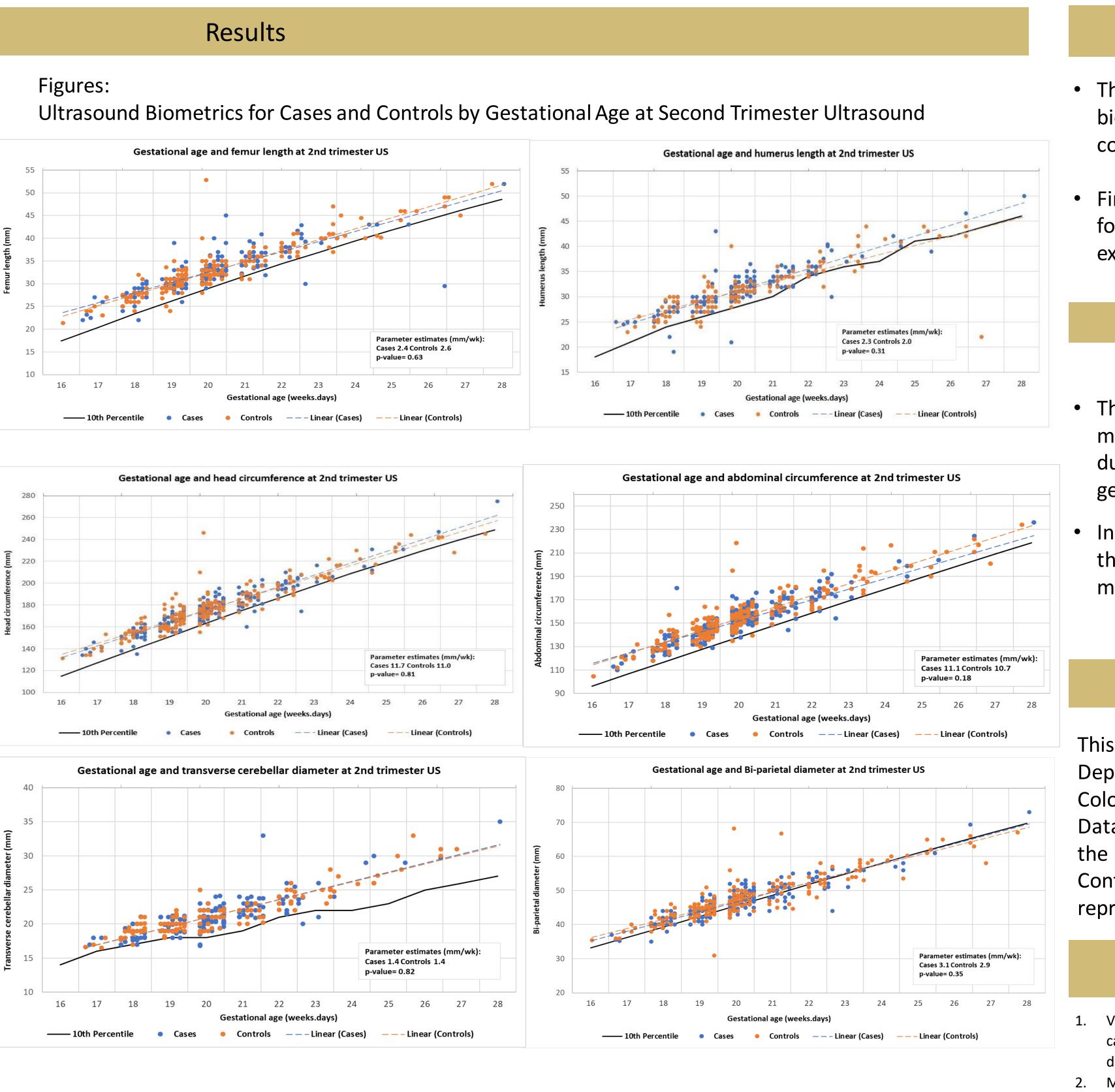
# Does marijuana use in pregnancy increase the risk for abnormal fetal biometrics on prenatal ultrasound?

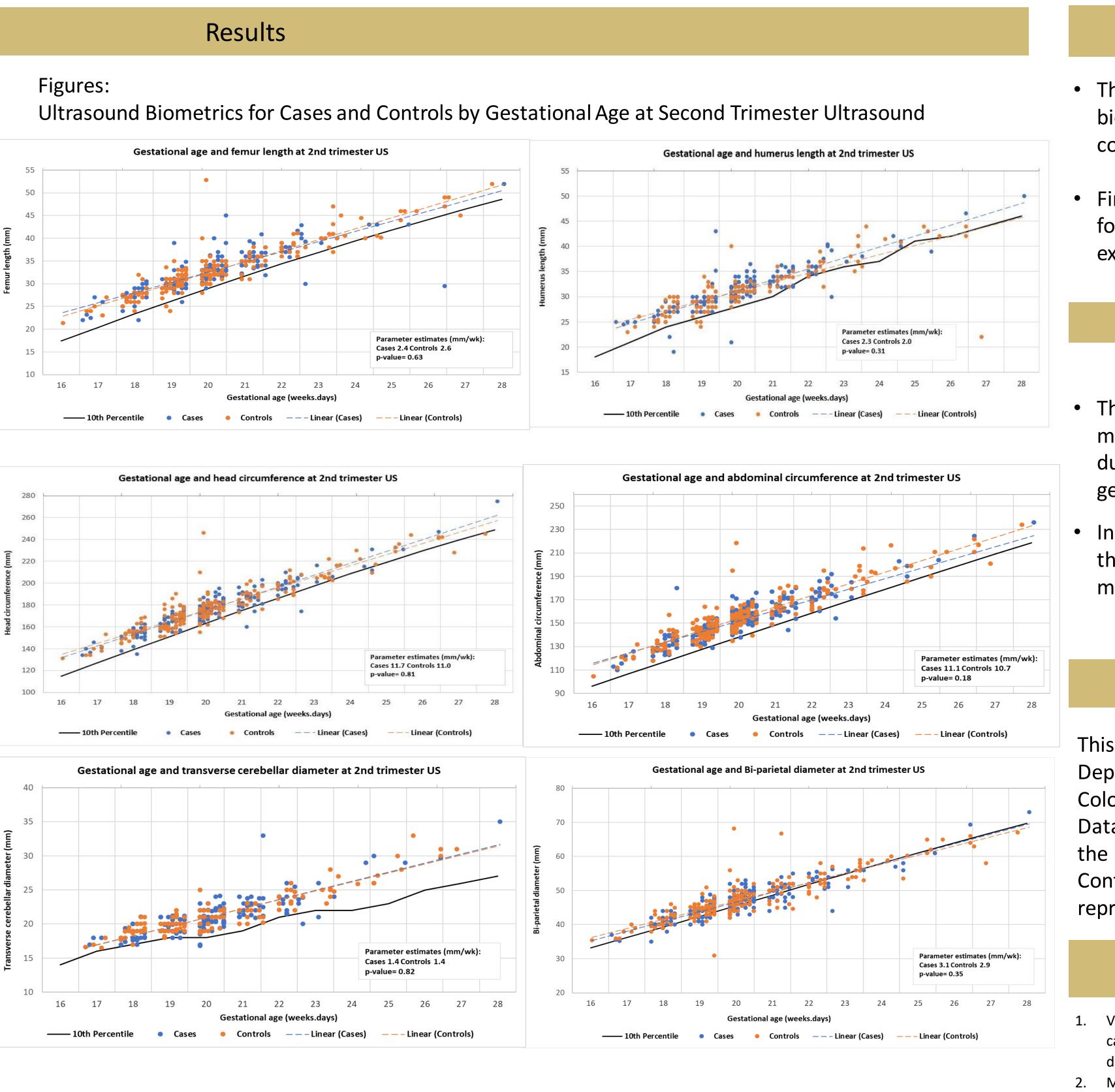
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### Maternal Demographics and Growth Parameter Results

0.98 0.92 0.92 0.92 0.92 1
0.92 0.08 0.92 1
0.92 0.92 0.92 1
0.08
0.08
0.08
0.08
0.92
0.92
1
1
1
1
1
<0.01
<0.01
0.62
0.32
0.56
0.52
0.54
0.02
0.13
1
0.66
0.22
<0.01
0.17
0.67
0.38
1
<0.01
<0.01
<0.01
0.63
0.67
0.38
0.62





- (Table 1).
- 67%, p<0.01) (Table 1).

• There were no significant demographic differences between patients with and without a positive MJ UDS. Patients were generally < 35-years-old (95.5% vs 95.5%, p=0.98), multiparous (62.9% vs 62.4%, p=0.92), Non-Hispanic White (57.4% vs 58.9%, p=0.08) with public insurance (87.6% vs 87.6%, p=1.0)

Among patients with a positive MJ UDS, there was a higher frequency of depression (38.1% vs 24.8%, p<0.01), self-reported tobacco use (46% vs 23%, p<0.01), and self-reported MJ use (58% vs

## Conclusions

• There was no significant difference in neonatal outcomes or fetal biometrics on 2<sup>nd</sup> trimester US in infants exposed to MJ in-utero compared to infants without MJ exposure.

Findings provide support for additional retrospective studies focused on quantifying maternal MJ use and timing of fetal exposure its impact on adverse fetal outcomes.

### Limitations

• The sample was limited to patients who delivered in Colorado, with many of the patients having resided in the state through the duration of their pregnancy. These findings may ultimately not be generalizable to other populations outside of Colorado.

• In the medical charts, there was limited documentation regarding the timing, form (edible, inhaled, etc.) amount, and frequency of marijuana usage.

## Disclosures

This project was supported by the PARITY research team in the Department of Obstetrics and Gynecology at the University of Colorado, the Colorado Center for Personalized Medicine's Health Data Compass Data Warehouse project (healthdatacompass.org), and the NIH/NCATS Colorado CTSA Grant Number UL1 TR002535. Contents are the authors' sole responsibility and do not necessarily represent official NIH views.

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