Comparing Clinical Research Participation Trends Among Pregnant Cannabis-Exposed and Unexposed Participants

Introduction

- The American College of Obstetrics and Gynecology (ACOG) and American Academy of Pediatrics (AAP) advise against use of cannabis and cannabidiol (CBD) products during pregnancy.
 - However, recent studies suggest that cannabis use is increasing for the relief of pregnancy-related symptoms, such as nausea and anxiety.
- While there is currently limited data on the effects of perinatal cannabis use, in-utero exposure has been linked to adverse neonatal neurodevelopmental outcomes. As perinatal cannabis use rates reportedly increase, it is necessary to obtain conclusive, pregnancy-specific safety data through well-designed studies.

Study Objective: To identify clinical research participation trends in a pilot study among cannabis-exposed and unexposed participants in the perinatal period.

Methods

Study Design and Participants

- Participants were recruited from OBGYN settings affiliated with an academic health system in Florida between July 2023 to January 2025.
- Pregnant, English-speaking patients aged 18-50 and self-reporting cannabis use during prenatal visits were recruited, along with pregnant control patients with no self-reported use. Participants were consented using an IRB-approved protocol (IRB#202300712). Participants were enrolled in any trimester and received compensation for each biospecimen/imaging completion.

Data Collection

- Biological samples were collected: maternal urine in each trimester; maternal urine, placenta, umbilical cord, and neonatal meconium at delivery; and maternal urine and breastmilk in the postpartum period.
- Imaging collection included a fetal ultrasound in each trimester and a third trimester fetal magnetic resonance imaging (MRI) scan to obtain views of the fetal brain and cerebral blood flow.

Data Analysis

Participation trends were identified by calculating proportions of missed/completed biospecimens/imaging, study visit reschedules/cancellations, and losses-to-follow-up (LTFUs).

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> Results July 2023-January 2025







Biospecimens

Of 239 possible biospecimen collections, 71% were completed [66% (n=126) cannabis-exposed and 90% (n=44) control]





Results, continued

- controls.

Discussion



Retention Facilitators and Barriers

- appointments.

Limitations



• 40% (n=8) of cannabis-exposed participants were LTFU, with the majority (88%, n=7) occurring postpartum and 13% (n=1) after the third trimester. Notably, no LTFUs were observed in the control group. • 24% of participants (30%, n=6 cannabis-exposed; 0%, n=0 control) did not complete one or more postpartum breastmilk samples due to not breastfeeding or no longer lactating.

• 45% of cannabis-exposed participants either canceled or required **rescheduling of at least one appointment**, compared to o% (n=o) of controls. Similarly, 40% (n=8) canceled, did not attend, or did not **schedule their postpartum appointment**, compared to o% (n=o) of

Findings suggest that participants with self-reported cannabis use were more likely to miss postpartum visits, become lost-to-follow-up, and require rescheduling of study visits compared to controls; however, cannabis-exposed participants still completed 68% of study visits.

Retention facilitators: financial compensation, trust/rapport-building with study staff, and scheduling study visits onto existing obstetrics

 Retention barriers, as described by participants: transportation, finances, and time constraints/childcare.

To improve retention/compliance rates among cannabis-exposed participants, future research could consider implementing stricter follow up, staggered increase in incentive, and home visits for sample collections.

Small sample size relative to large healthcare setting. L&D hospital staff unfamiliar with study protocols, resulting in missed delivery samples.