



1a. Introduction

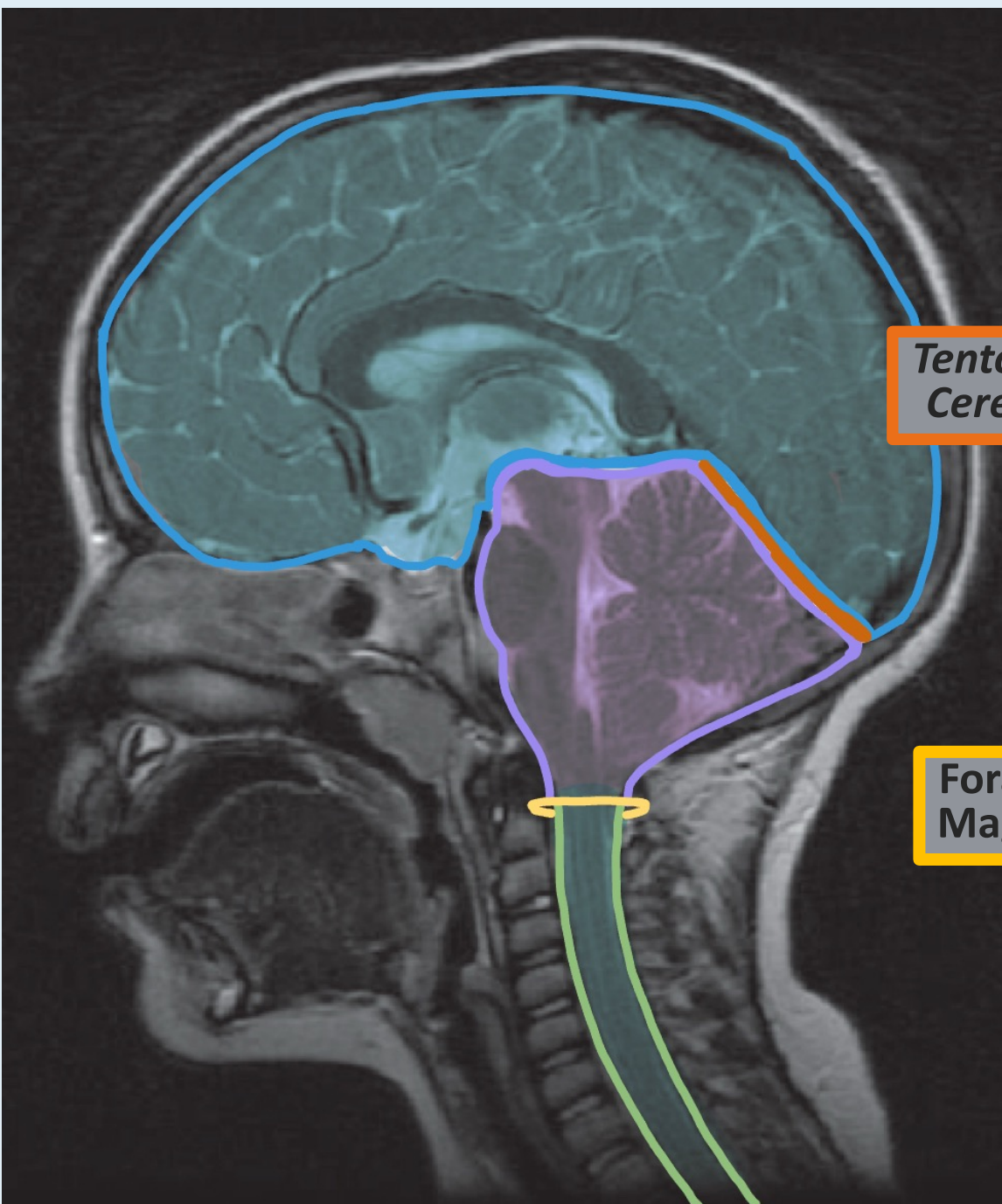
Learning that your baby has a *central nervous system (CNS) anomaly* during pregnancy is among the most distressing experiences a family can face. Despite advances in ultrasound and fetal MRI, accurately **characterizing these anomalies** and **distinguishing them** from insignificant findings remains challenging.¹

Understanding how *clinical factors and location of the anomaly* contribute to these challenges is essential for improving prenatal counseling for fetal CNS anomalies.

1b. Background

Several factors impact characterization of fetal CNS anomalies including:

A. Region of the Anomaly²



Supratentorial

↑ Structural Variability
Subtle cortical & midline findings
difficult to define prenatally

Posterior Fossa

↓ Visibility
Complex anatomy & shadowing from
bone obscure detail

Spinal Region

Spine Curvature Obscures View
Spine position → Difficulty assessing
open vs. closed neural tube defects

B. Clinical Factors^{3,4}



Imaging Modality: MRI improves detail



Maternal Obesity: ↓ Ultrasound clarity



Advanced Maternal Age (AMA): ↑ Anomaly risk

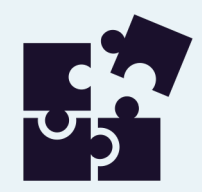


Gestational Age (GA): Development Impacts Visibility
→ Early scans=incomplete anatomy → Late scans=obscured view

1c. Objectives



Assess the accuracy of prenatal CNS anomaly diagnoses made using fetal neuroimaging techniques



Evaluate how anomaly region, imaging modality, maternal factors, and gestational age influence diagnostic consistency.

2a. Cohort Selection

Study type: Retrospective cohort study

Population: Pregnancies diagnosed with fetal CNS anomalies *between 2019 and 2025* under the care of LSUHSC Maternal-Fetal Medicine (MFM) physicians.

105 cases: CNS Anomalies Identified

All pregnancies (105) in the LSUHSC–MFM perinatology-neonatology care (PNC) database with a **documented fetal CNS anomaly**.

Excluding Non-isolated Anomalies → 76 cases

Remaining cases (76) **do not** have **major non-CNS malformations** or multisystem anomalies affecting *other organ systems*.

Excluding Insufficient Records & Fetal Demise → 48 cases

Remaining cases (58) have **sufficient longitudinal records** and cases ending in **live birth**.

Excluding Anomalies Resolved Prenatally → 37 cases

Final cohort (37) characterized by **persistent CNS anomalies at birth** (excluded those with transient or resolved prenatal findings).

Final cohort: 37 pregnancies with **isolated, non-resolving fetal CNS anomalies** and **complete longitudinal records**

2b. Data Collection

For each case, we collected:

- Prenatal data:** Final prenatal CNS diagnosis, imaging modality, **GA**
- Maternal data:** Body-mass index (**BMI**) and age
- Postnatal data:** **Postnatal outcome**, imaging modality

Postnatal outcome categorized as:

- Confirmed:** prenatal diagnosis matched postnatal diagnosis
- Redefined:** prenatal diagnosis modified postnatally
- Insignificant:** prenatal finding deemed clinically insignificant

*Note: “**Reclassified**”=“Redefined” + “Insignificant”

2c. Hypothesis

We hypothesize that accuracy of prenatal CNS anomaly diagnosis is influenced by both the **region of the anomaly** and **clinical or imaging-related factors** that alter fetal visibility.

3a. Results

Legend

- Confirmed (29.73%)
- Reclassified (70.27%)
- Insignificant (37.84%)
- Redefined (32.43%)

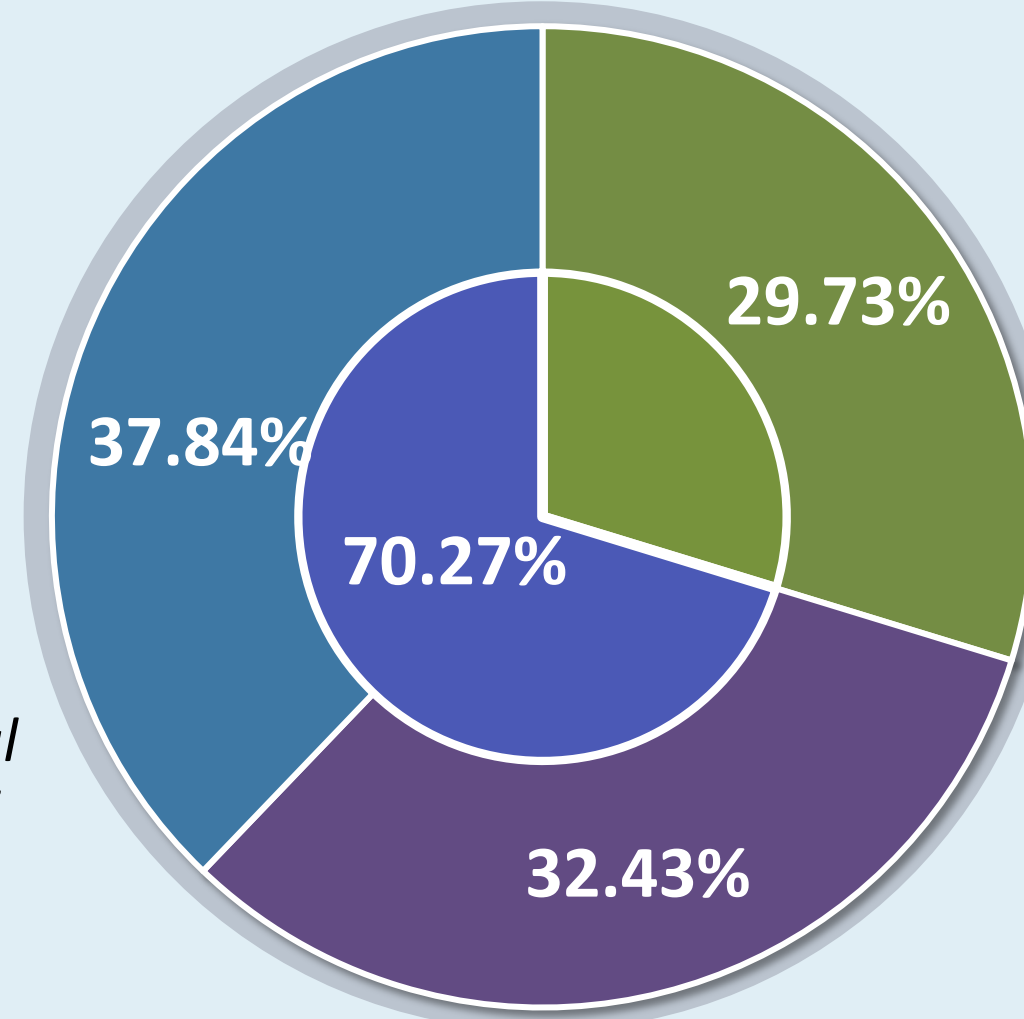


Figure 1. Diagnostic outcomes for 37 fetal CNS anomaly cases describing percent of cases confirmed or reclassified (insignificant + redefined) postnatally.

CNS Region	Confirmed (n)	Reclassified (n)	Total (n)	% Reclassified
Supratentorial	9	18	27	67%
Posterior Fossa	2	2	4	50%
Spine	0	1	1	100%
Multiple	0	5	5	100%

Prenatal MRI Use	Confirmed (n)	Reclassified (n)	Total (n)	% Reclassified
Yes	7	20	27	74%
No	4	6	10	60%

Maternal Obesity	Confirmed (n)	Reclassified (n)	Total (n)	% Reclassified
Yes	6	11	17	65%
No	5	15	20	75%

AMA	Confirmed (n)	Reclassified (n)	Total (n)	% Reclassified
Yes	1	9	10	90%
No	10	17	27	63%

GA at Diagnosis	Confirmed (n)	Reclassified (n)	Total (n)	% Reclassified
<20 weeks	3	2	5	40%
20–24 weeks	2	10	12	83%
25–32 weeks	4	10	14	71%
>32 weeks	2	4	6	67%

Figure 2. Stratification of CNS anomaly cases by region and clinical factors. While minor variations in reclassification were observed among these variables, none reached statistical significance (all $p>0.05$).

3b. Conclusion

Although no significant associations were found between reclassification and anomaly region, MRI use, obesity, age, or timing of diagnosis, the **consistent high reclassification rate** highlights the importance of postnatal confirmation in fetal CNS anomaly counseling.

Recognizing when and why these discrepancies occur may help improve the value of fetal neuroimaging and help clinicians provide more accurate prognoses for families.

3c. Future Steps



A. Expand cohort for greater statistical power.



B. Standardize reclassification terminology.



C. Link prenatal imaging with postnatal neurodevelopmental outcomes.

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