Congenital Zika Syndrome: A Nationwide Cohort Study in Brazil



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Introduction



Zika Virus (ZIKV)

- 1. Flavivirus (DENV, Yellow Fever)
- 2. Transmission through arthropods (Aedes), sexual contact, blood transfusion, or **vertical transmission**

Congenital Zika syndrome

- Spectrum of structural anomalies (e.g., cortical atrophy with microcephaly);
- 2. Functional impairments (e.g., dysphagia);
- 3. Clinical sequelae (e.g., epilepsy).

What is the prognosis for live births with CZS?

Objectives



- (i) investigate the overall and cause-specific mortality rates among live births with CZS compared with those without CZS,
- (ii) examine the overall hospital admission rates among live births with CZS compared with those without CZS in a smaller subset of our cohort.



Methods



- Live Birth Information System (SINASC)
- Mortality Information System (SIM)
- Public Health Event Record (RESP)
- Hospitalization Information System (SIH)
- Unified Register for Social Programs (CADU)



Interested on administrative data come to our session on Friday 16:30

Methods



Exclusion: (i) other congenital abnormalities registered in SINASC,

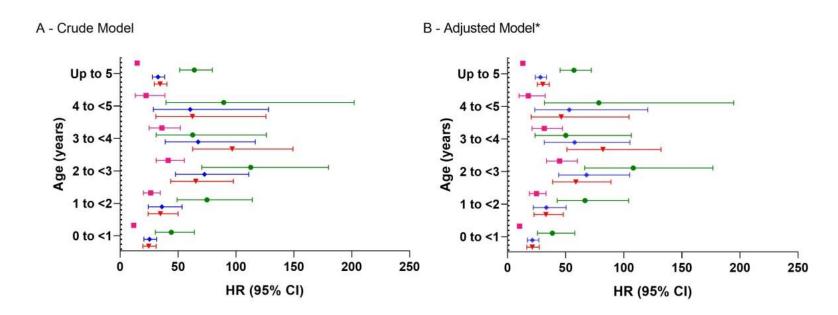
(ii) other congenital STORCH infections registered in RESP, and/or (iii) an inconclusive, under investigation or ruled out classification of C7S in RESP

Estimated rates and ratios with 95% (CIs);

Adjusted: region of birth, year of birth, maternal age, maternal education, maternal race/ethnicity, maternal marital status and sex of the newborn

Results 1 (Mortality)





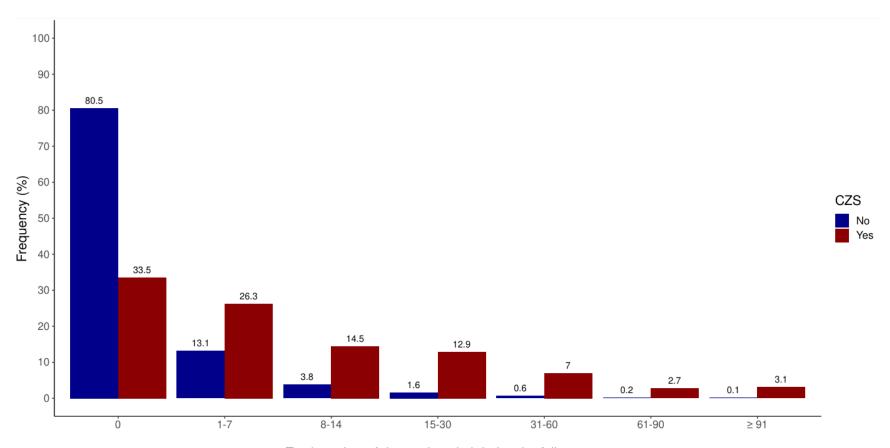
HR: Hazard Ratio CI: Confidence Interval

*Adusted by region, year of birth, maternal age, maternal education, maternal race/ethnicity, marital status and sex of the newborn

- Respiratory System Diseases
- Infectious and Parasitic Diseases
- Nervous System Diseases
- General

Results 2 (Hospital admission)



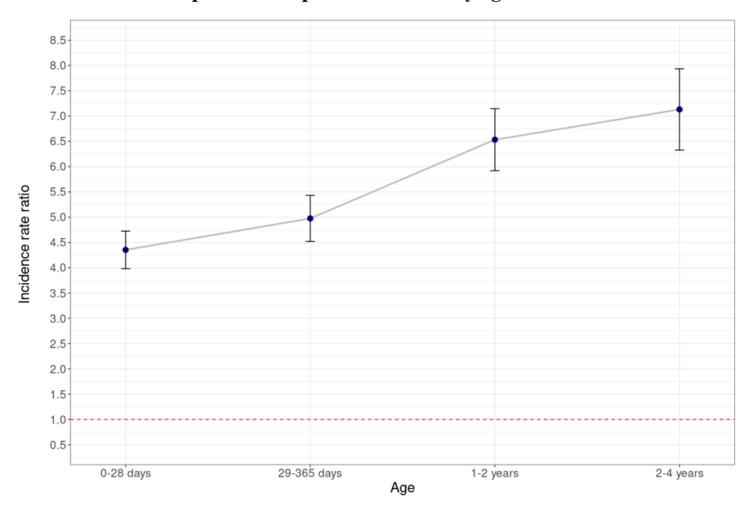


Total number of days at hospital during the follow-up

Results 3 (Hospital admission)



Figure 2 – Adjusted incidence rate ratios and 95% CI for all-cause hospital admissions for CZS patients compared to controls by age at admission.



Results 4 (Hospital admission)



CZS

No microcephaly (N = 1098)

| Age categories | Person- years | Rate* |
|----------------|------------------|-------|
| 0-28 days | 80.8 | 582.9 |
| 29 – 365 days | 913.9 | 46.8 |
| 1-2 years | 903.1 | 38.5 |
| 2-4 years | 772.9 | 28.5 |

Microcephaly (N = 796)

| Person- | Rate* | |
|---------|-------|--|
| years | | |
| 59.7 | 524.4 | |
| 691.4 | 36.3 | |
| 697.0 | 41.6 | |
| 574.4 | 28.6 | |

^{*}Rate represents number of hospital admissions per 100 person-years.

Discussion



Strengths

- Large sample size (Confirmed and probable CZS)
- Included a populationrepresentative comparison group
- Control for confounding.
- Including only confirmed cases showed the robustness of our findings.

Limitations

- Lack of relevant clinical data
- Potential underreporting in the RESP, mainly among those fetuses prenatally exposed to ZIKV during pregnancy, but without detectable malformations at birth.
- Linkage error.

Conclusion



Protocols

There is a higher mortality and morbidity risk in live births with CZS than live births without CZS that persists throughout the first five years. Highlights a need for instituting well-established postnatal protocols, including early intervention, which may help to lessen adverse medical and developmental sequelae and improve survival of these children.





Thank you!

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