Results. Seventy-two non-CZS infants had neurodevelopmental tests: 40 were at a mean (SD) of 5.7 (0.9) months and 66 were at 13.5 (3.2) months of age. Thirty-four had two assessments. The total WIDEA, social cognition, and mobility domain scores became more abnormal with postnatal age (figure). The AIMS scores were similar to the normative sample. Three infants had an AIMS score < 2 SDs below the norm. On cranial US, 19 infants (26%) had a nonspecific finding (lenticulostriate vasculopathy, choroid plexus cysts, subependymal cysts, and/or calcification). Infants with a US finding had a lower WIDEA mobility score than infants with normal US (P = .054). There was a trend toward lower AIMS scores in infants with US findings compared with infants with normal US (P = .06). AIMS Interrater agreement on video-based scoring was good (ICC = 0.73, 95% CI 0.42, 0.87).

Conclusion. ZIKV-exposed infants without CZS are at risk for neurodevelopmental delay. Nonspecific cranial US findings may represent mild ZIKV-related injury. Long-term neurodevelopmental follow-up is important for all ZIKV-exposed infants.

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1873. Pregnancy and Birth Outcomes Among Colombian Women with Zika Virus Disease in 3 Surveillance Sites, Proyecto Vigilancia de Embarazadas con Zika
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Background. Proyecto Vigilancia de Embarazadas con Zika (VEZ) was an intensified surveillance system built upon existing national surveillance of pregnant women with symptoms of Zika virus (ZIKV) disease and conducted in three Colombian cities with a high prevalence of Zika. This analysis of data from VEZ estimates the risk of Zika-associated birth defects among pregnant women with symptoms of ZIKV disease, and among a subset with laboratory evidence of possible ZIKV infection during pregnancy.

Methods. During April–November 2016, pregnant women were enrolled if they were reported to the surveillance system (Sigivila) or visited participating clinics with symptoms of ZIKV disease. Maternal and pediatric data were abstracted from prenatal care, ultrasound, and delivery records, as well as from pediatric or specialist visits.

Results. Of 1,223 women enrolled, 47.8% and 34.3% reported first or second trimester symptom onset, respectively. Of 381 pregnancies with maternal and/or infant specimens tested, 108 (29%) had laboratory evidence of possible ZIKV infection during pregnancy; half of these (53.3%) were positive for ZIKV RNA only, 37.4% for IgM antibodies only, and 9.3% for both. Of 1,190 of pregnancies with known outcome, 63 (5%) had Zika-associated brain or eye defects; among the subset with any laboratory evidence, 12 (11%) had Zika-associated brain or eye defects. The prevalence of Zika-associated brain or eye defect was 5% (95% CI 0.6–4.5) among pregnant women with normal US and 9.3% for both in the first and second trimester, respectively.

Conclusion. Among pregnant women with symptoms of ZIKV disease enrolled during the height of the ZIKV epidemic in Colombia, prevalence of any Zika-associated brain or eye defect was 5%, with a higher prevalence among those with laboratory evidence of possible ZIKV infection. Rapid enhancements to Colombia’s national surveillance enabled the estimation of the risk of birth defects associated with ZIKV disease in pregnancy.

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1874. Comparison of the Risk of Birth Defects in Live Births From Pregnant Women Infected and Not Infected by Zika Virus in Guadeloupe, 2016–2017
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Background. In the French Territories in the Americas (FTA), the risk of birth defects possibly associated with Zika virus (ZIKV) infection was estimated at 7% among fetuses/infants in a cohort of 546 women who developed a presumptive RT-PCR confirmed ZIKV infection during pregnancy (NEJM 2018;378:985–94). There was no concomitant prospective cohort of pregnant women without ZIKV infection to use as a control group.

Methods. In Guadeloupe, one of the 3 FTA that participated in the FTA cohort study, pregnant women were recruited at the time of delivery and tested for ZIKV infection. Women who had a confirmed negative IgG serology test for ZIKV at delivery and no other positive ZIKV test during pregnancy were considered to be ZIKV non-infected. Information on the course of the pregnancy was collected retrospectively and outcomes of live born infants of ZIKV noninfected women were analyzed, using the same definition criteria as those used for the FTA cohort study. Pregnancy outcomes were compared with those of the 241 ZIKV-exposed live born infants in Guadeloupe, extracted from the FTA cohort.

Results. Of the 490 live born infants without in-utero exposure to ZIKV, 42 infants (8.6%) had neurological abnormalities that were described as “potentially linked to ZIKV infection”; all but one of these were microcephaly without any other brain or clinical abnormalities. The proportion of such abnormalities was not statistically different from that observed in the 241 live born infants of ZIKV infected women exposed, using a ~2 SD cut-off with international growth curves, may lead to an overestimation of the rate of neurological complications of ZIKV infection during pregnancy.

Conclusion. Isolated anthropometric and other mild neurological abnormalities had the same prevalence among live born infants with and without in utero ZIKV exposure. The high prevalence of isolated microcephaly among ZIKV noninfected women in our study population suggests that the sensitivity definition for microcephaly, using a ~2 SD cut-off with international growth curves, may lead to an overestimation of the rate of neurological complications of ZIKV infection during pregnancy.

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1875. La Crosse Virus Neuroinvasive Disease in Children: A Contemporary Review and Evaluation for Predictors of Disease Severity
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Background. La Crosse Virus (LACV) is the most common neuroinvasive arboviral disease in children. Contemporary data on clinical presentation, management, outcomes, and predictors of disease severity are lacking.

Methods. A retrospective analysis was performed of children (0–18 years) admitted to Nationwide Children’s Hospital from January 2009 to December 2018 diagnosed with LACV neuroinvasive disease (LACV-ND). LACV-ND diagnosis was defined as a compatible clinical illness and serum serologic detection of LACV in the absence of other infectious etiologies. Demographic, clinical, laboratory, electroencephalography (EEG), radiologic, and outcome data were recorded. Severe disease was defined as the presence of clinical or electroencephalographic status epilepticus, SEADH, PICU