The emergence of Zika virus (ZIKV) in the Americas has coincided with an abnormal increase in prenatal and neonatal documented cases of microcephaly and other anomalies of the central nervous system (1). These alterations of the brain, along with animal models of vertical transmission of ZIKV, a single-stranded RNA flavivirus, are evidence of the neurotropic nature of the virus (2–4).

Vertical transmission and infection of the fetus during all 3 trimesters of pregnancy with ZIKV has been extensively reported, but little is known about perinatal transmission; only a few cases have been reported (5–8). We report 3 cases of perinatal ZIKV infection during the epidemic in Colombia and provide detailed descriptions of clinical features, diagnosis, and neurodevelopmental outcome at 18 months of age (corrected).

The Study
The Institutional Review Board and Ethics Committee of the Universidad Industrial de Santander approved this study. Formal written consent was obtained from participating women.

Case-patient 1 was a 26-year-old pregnant woman in labor who was admitted to the Hospital Universitario de Santander on August 7, 2015, after she reported fever, exanthema (maculopapular rash on the torso), and osteoid–muscular pain (Figure). Her initial hemogram showed mild thrombocytopenia (112,000 platelets/µL; reference range 150,000–400,000 platelets/µL). Test results for syphilis, toxoplasmosis, rubella, cytomegalovirus, herpes simplex virus, varicella zoster virus, and parvovirus B19 during hospitalization were negative, as was a test result for dengue virus (DENV) IgM.

A boy was born vaginally at 37 weeks of gestation. Apgar scores were normal. Anthropometric measurements yielded a birthweight of 3.03 kg (32nd percentile), a head circumference (HC) of 32 cm (6th percentile), and a length of 54 cm (99th percentile). Results of a physical examination were unremarkable. The infant was admitted for observation, and cord blood samples were sent to the Instituto Nacional de Salud (INS) for additional testing by reverse transcription PCR (RT-PCR) for DENV and chikungunya virus. On the second day of life, the infant had a distal macular–papular rash with hyperalgesia and mild edema of the hands and feet; a test result for DENV IgM was negative. Hyperthermia developed on the sixth day, and generalized exanthema developed on the seventh day.

Because of persistent fever, we initiated a sepsis work-up. Results for initial complete blood count/differential count, erythrocyte sedimentation rate, blood culture, urine culture, and C-reactive protein were within reference ranges. Fever and exanthema
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were present for 4 additional days; a follow-up C-reactive protein level was abnormal (21 mg/dL). Findings for analyses of cerebrospinal fluid and culture were unremarkable. After symptoms improved, the infant was discharged; an RT-PCR result from the INS was negative for chikungunya virus.

Because of the Zika epidemic in Brazil, the INS initiated RT-PCR testing for ZIKV and randomly

Figure. Timelines for 3 case-patients who had perinatal Zika virus infection during a Zika epidemic, Colombia, 2015. A) Case-patient 1; B) case-patient 2; C) case-patient 3. DENV, dengue virus; CHIKV, chikungunya virus; RT-PCR, reverse transcription PCR; ZIKV, Zika virus.
remained asymptomatic, and both hearing and ophthalmologic evaluations were normal. Neurodevelopmental evaluation with the Bayley Scale for Infant and Toddler Development (10) at 7.2, 13.0, and 36.5 months of age for CA was considered within the normal range for age (Table). Neuroimaging was not indicated.

Case-patient 3 was a 25-year-old pregnant woman who was admitted to the Hospital Universitario de Santander hospital on April 5, 2016, at 34 weeks of gestation, with a history of eye irritation, rash, and fever. Test results for syphilis, toxoplasmosis, varicella zoster virus, parvovirus B19, rubella, cytomegalovirus, and herpes simplex virus were negative, but blood and urine RT-PCR results for ZIKV positive. Betamethasone was administered for lung matura-

On the third day of admission, a boy was delivered vaginally. He had normal Apgar scores and a birthweight of 2.24 kg, an HC of 31 cm, and a length of 42 cm. An umbilical cord blood sample tested by RT-PCR for ZIKV was positive. The infant showed signs of respiratory distress syndrome at birth, requiring supplemental oxygen, ventilatory support with continuous positive airway pressure, and surfactant administration. Symptoms improved after 24 hours and he was subsequently discharged.

We conducted neurodevelopmental follow-up at 6, 12, 24, and 36 months of age by using the Bayley Scale for Infant and Toddler Development (10). All evaluations were considered within the normal range for age (corrected). Ophthalmologic and hearing assessments were normal; neuroimaging was not indi-

Conclusions
The symptoms and diagnostic tests for the 3 pregnant women strongly support maternal infection 5 days before delivery. Vertical transmission is shown by suggestive early neonatal symptoms and a positive

| Case-patient | Infant age, mo | Cephalic perimeter | Motor | Scale domain |
|-------------|---------------|-------------------|-------|--------------|
|              | Infant age, mo | Z score           | Score | Z score | Percentile | Language | Percentile | Cognitive | Percentile |
| 1            | 20.1          | −0.50             | 94    | −0.40   | 34         | 94       | −0.40      | 34        | 90         | −0.67      | 25         |
|              | 24.9          | −1.70             | 107   | 0.47    | 68         | 83       | −1.13      | 13        | 100        | 0.00       | 50         |
|              | 40.3          | −0.51             | 88    | −0.80   | 21         | 89       | −0.73      | 23        | 95         | −0.33      | 37         |
| 2            | 7.2           | 0.86              | 100   | 0.00    | 50         | NT       | NT         | NT        | NT         | NT         | NT         |
|              | 13.0          | 0.64              | 97    | −0.20   | 42         | 97       | −0.20      | 42        | 85         | −1.00      | 16         |
|              | 36.5          | 0.21              | 110   | 0.67    | 75         | 91       | −0.60      | 27        | 90         | −0.67      | 25         |
| 3            | 8.4           | 0.72              | 115   | 1.00    | 84         | NT       | NT         | NT        | NT         | NT         | NT         |
|              | 14.5          | 0.89              | 112   | 0.80    | 79         | 94       | −0.40      | 34        | 110        | 0.67       | 75         |
|              | 21.6          | 0.52              | 100   | 0.00    | 50         | 89       | −0.73      | 23        | 110        | 0.67       | 75         |

*NT, not tested.
†Standardized.
RT-PCR result for ZIKV for case-patient 1 and RT-PCR Zika virus–positive cord blood for the other 2 case-patients (Figure). Case-patient 2 had a positive RT-PCR result for ZIKV in blood and urine and a positive result for DENV IgM, suggesting co-infection. Although cross-reactivity in an RT-PCR for ZIKV and other arbovirus infections is possible (11), co-infection with 2 different types of arbovirus is also possible because of the endemic nature of dengue in Colombia where the patients were identified (12). Vertical transmission of DENV is considered a rare event, and there have been no reports of congenital dengue infection in neonates born to mothers infected early during pregnancy (13). Case-patient 3 had a positive result by RT-PCR for ZIKV in blood and urine, and her infant was positive for ZIKV in cord blood, which enabled us to confirm vertical transmission. However, because testing for DENV or chikungunya virus was not available, we cannot speculate on this issue.

Adverse fetal central nervous system anomalies with maternal ZIKV infection have been reported during the third trimester of pregnancy (14,15). Neurodevelopmental follow-up results were uneventful for the 3 case-patients we describe who had perinatal vertical transmission. Two previous cases of perinatal transmission reported showed no evidence of neurodevelopmental impairment, thus supporting our findings (8). In summary, our findings indicate that perinatal infection within the time frame described for these case-patients does not appear to affect neurodevelopmental outcomes of the newborns.

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