CONGENITAL TRANSMISSION OF *TRYPANOSOMA CRUZI* IN EUROPE (SPAIN):
A CASE REPORT

CRISTINA RIERA,* ANNA GUARRO, HOUSSEIN EL KASSAB, JOSÉ MARIA JORBA, MONTserrat CASTRO, ROSER ANGRILL, MONTserrat GALLEGó, ROSER FISA, CARMEN MARTIN, ALEJANDRO LOBATó, AND MONTserrat PORTús

Laboratori de Parasitologia, Facultat de Farmàcia, Universitat de Barcelona, Barcelona, Spain; Servei de Pediatria, Hospital Residència Sant Camil, Sant Pere de Ribes, Barcelona, Spain; Servei d’Anatomia Patològica, Hospital Residència Sant Camil, Sant Pere de Ribes, Barcelona, Spain; Servei de Microbiologia, Hospital Residència Sant Camil, Sant Pere de Ribes, Barcelona, Spain

Abstract. Here we report a documented case of congenital transmission of *Trypanosoma cruzi* from a Bolivian mother with chronic Chagas disease living in Spain. The serology and blood nested polymerase chain reaction (PCR) were positive for the mother, and amastigote forms were observed in histopathological study of the placenta and umbilical cord. Direct examination, culture, and nested PCR were positive in the blood of the neonate. At the age of 8 days, the neonate began treatment with 5-7.5 mg/kg/day of benznidazol, which was continued for 60 days. Direct examination, blood culture, and nested PCR were negative to *T. cruzi* 20 days after the start of treatment and remained negative 4 and 7 months thereafter. Serological tests were negative at 4 months. To detect congenital infection and initiate early treatment of infected newborns, protocols are required to detect Chagas disease in pregnant women who migrate from endemic to non-endemic areas.

INTRODUCTION

Chagas disease or American trypanosomiasis is a widely distributed endemic zoonosis in Central and South America, where ~11 million people are infected and an estimated 100 million people are at risk of infection. The disease is caused by the protozoan *Trypanosoma cruzi* and is usually considered as one affecting poor people in rural communities. In endemic areas, *T. cruzi* is transmitted mainly by triatomin insect vectors, which release excreta infected with the parasite into lacerated skin or mucosa. In addition to insect transmission, the infection may also occur through blood transfusion of whole blood or blood derivatives, congenital transmission from infected mothers, organ transplantation (kidney, heart, bone marrow, and others), and accidental contamination in the laboratory. Infection occurs to a lesser extent by oral transmission through food, such as meat, sugar cane juice, and fruit juice, contaminated with infected triatomines or their dejections.1,2

At present, Chagas disease is a potential public health problem in Spain as a result of the increasing number of migrants from countries in which it is endemic. To our knowledge, a case of asymptomatic congenital transmission has been reported in Europe in a 5-year-old Latin American child living in Romania who had never traveled to an endemic area.3 Another congenital case has recently been diagnosed in a 2-year-old child in Barcelona (J. Muñoz and others, unpublished data). Here we report the first case of congenital Chagas disease transmission diagnosed in a neonate in Europe (Spain). We studied *T. cruzi* infection in the mother and the neonate, and a parasitological and serological longitudinal follow-up was performed on the child to monitor response to treatment.

CASE REPORT

A 28-year-old expectant Bolivian mother with chronic Chagas disease informed physicians of her condition at delivery. The patient was asymptomatic, and anamnesis established that she had been infected at the age of 6 and diagnosed and treated at the age of 22 in Bolivia. The woman had two other children, 10 and 12 years of age, who lived in Bolivia and whose clinical condition was unknown.

The neonate was delivered at 37 weeks of gestation by caesarean section. Vaginal and rectal cultures for the mother were positive for *Streptococcus agalactiae*, and she was treated with endovenous ampicillin during dilation. Caesarean delivery was required because attempts to induce birth failed. The newborn was a boy weighing 2.905 kg. Preliminary examination of the infant detected no complications. The mother nursed the neonate, who showed a satisfactory evolution. Splenomegaly, hyperbilirubinemia, and encephalitis were not detected, and additional hematological and biochemical parameters were normal.

After delivery, the mother and the newborn were immediately examined for Chagas disease using parasitological and serological methods. The serological diagnosis of *T. cruzi* infection in sera was performed using a commercial ELISA with recombinant antigens (BioELISA Chagas; Biokit S.A., Lliça d’Amunt, Barcelona, Spain) and a non-commercial conventional ELISA with a whole lysate antigen obtained from epimastigotes of *T. cruzi* I strain. The bands recognized by the sera from mother and child were studied by Western blot (WB) analysis with the same antigen used in the conventional ELISA. A nested polymerase chain reaction (PCR) was performed following the standard technique (external primers TCZ1 and TCZ2 and internal primers TCZ3 and TCZ4, which amplify a DNA fragment of 149 bp of a repetitive sequence of nuclear DNA of 195 bp).4

A blood sample from the mother was taken for laboratory diagnosis of Chagas disease. Microscopic observation of blood and culture in NNN medium were negative, and nested PCR was positive for *T. cruzi* DNA. Maternal seroreactivity to *T. cruzi* antigens was positive by ELISA and WB analyses (Table 1; Figure 1). Histopathological study of the placenta and umbilical cord was performed, and *T. cruzi* amastigote forms were observed in both samples (Figures 2 and 3).

Trypomastigote forms in the blood stream were detected using the microhematocrit concentration technique. The

---

* Address correspondence to Cristina Riera, Laboratori de Parasitologia, Facultat de Farmàcia, Universitat de Barcelona, Avda. Joan XXIII s.n., E-08028 Barcelona, Spain. E-mail: mcriera@ub.edu
parasite was isolated by blood culture in NNN and *T. cruzi* DNA was detected by nested PCR. The strain was identified as *T. cruzi* I. Serological study by ELISA showed high levels of specific antibodies (Table 1), and the WB pattern of bands was identical in the newborn and mother (Figure 1).

Once congenital infection was confirmed, the neonate was treated with benznidazol (5–7.5 mg/kg/day) every 12 hours for 60 days. Parasitological and serological follow-up was performed at 20 days and 4 and 7 months after beginning treatment. Parasitemia, monitored by nested PCR and blood culture, was negative at day 20 after the start of treatment and remained negative at 4 and 7 months. At 4 months, ELISA tests were negative (Table 1), and the WB pattern showed a reduction in the number and intensity of bands (Figure 1). Clinical, hematological, and biochemical follow-up of the infant was performed for 12 months, during which time values were normal, and the child remained clinically asymptomatic. The infant showed good tolerance to treatment and normal development.

### DISCUSSION

Chagas disease enters the chronic phase 2–4 months after the acute clinical symptoms disappear and generally develops

---

**TABLE 1**

Serological, molecular, and parasitological blood analysis of the mother and the newborn

<table>
<thead>
<tr>
<th>Methods</th>
<th>Mother at delivery</th>
<th>At birth</th>
<th>20 days</th>
<th>4 months</th>
<th>7 months</th>
</tr>
</thead>
<tbody>
<tr>
<td>Microscopy</td>
<td>Negative</td>
<td>Positive</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Culture</td>
<td>Negative</td>
<td>Positive</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Nested PCR</td>
<td>Positive</td>
<td>Positive</td>
<td>Negative</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>ELISA R*</td>
<td>6.6</td>
<td>6.2</td>
<td>5.8</td>
<td>0.5</td>
<td>0.2</td>
</tr>
<tr>
<td>ELISA W†</td>
<td>61</td>
<td>51</td>
<td>36</td>
<td>10</td>
<td>7</td>
</tr>
</tbody>
</table>

* ELISA R (recombinant antigen, BioELISA Chagas, Biokit); cut-off = 1.  
† ELISA W (whole antigen, in house); cut-off = 20.

---

**Figure 1.** Trypanosoma cruzi antigen polypeptide recognition of sera from mother at delivery (m) and newborn at 1 day (a), 20 days (b), and 4 months (c) after initiating anti-parasitic treatment. Whole antigen from *T. cruzi* epimastigotes was electrophoretically separated in sodium dodecyl sulphate-polyacrylamide gels (12%) and transferred to nitrocellulose sheets. Immunoreaction was performed with sera at 1:50 dilution, and bound immunoglobulins were detected with protein A peroxidase conjugate (Sigma) at 1:1,000 dilution.

**Figure 2.** Histologic section of the extra-placental membranes, stained with hematoxylin-eosin, showing amastigotes of *T. cruzi* (H&E, ×60).
Maria infection was diagnosed in the neonate. The data obtained in the immunologic profile of the mother and child, detected by WB analysis, allows differentiation between passively transmitted maternal antibodies and those newly synthesized in the infant, which are used in early postnatal diagnosis of congenital toxoplasmosis. In our case, no differences in the band pattern of the sera from the mother and newborn were detected after delivery, and no new bands were recognized by child sera during the follow-up period. Therefore, neonate production of antibodies could not be appreciated.

Microscopic observation of bloodstream trypomastigotes and/or culture of theuffy coat fraction of blood samples using the microhematocrit concentration technique are commonly used to diagnose congenital infection and are highly sensitive when parasitemia is high, as in the case reported here. In contrast, when parasitemia is low, conventional parasitological and serological techniques are not efficient enough to provide early diagnosis of congenital Chagas disease. However, PCR is an easy, rapid, and sensitive diagnostic method that requires only a small amount of blood, although unusual false-positive results have been described. In addition, this technique can be used to monitor parasitemia in infants after treatment.

Early diagnosis of T. cruzi infection is essential for the rapid administration of anti-parasitoc therapy. Studies on nitrofurazom or benzimidazol treatments show that the cure rate depends on the speed with which treatment is initiated. Most infants treated early test negative when examined 6 months to 2 years after starting treatment. The data obtained in our study are consistent with these results and show that treatment is effective when started during the first months of life.

Given the increasing number of women of fertile age migrating to Spain from regions with endemic Chagas disease, new cases of congenital transmission may occur. To increase the early detection of congenitally infected babies and thus facilitate their early treatment, we recommend screening for Chagas disease in pregnant women from endemic areas and the establishment of diagnostic protocols to detect the infection in the neonate.

Received January 5, 2006. Accepted for publication July 5, 2006.

Acknowledgments: The authors thank Dr. P. Bonay, Universidad Autónoma de Madrid, for genotyping T. cruzi strains. We also thank P. Lopez-Chajade, M. Vérges, and S. Tobar (Laboratorio de Parasitología, Facultat de Farmàcia, Universitat de Barcelona) for providing technical assistance.

Authors’ addresses: Cristina Riera, Montserrat Gallego, Roser Fisa, and Montserrat Portús, Laboratorio de Parasitología, Facultat de Farmàcia, Universitat de Barcelona, Avda. Joan XXIII s.n., E-08028 Barcelona, Spain, Telephone: 34 93 402 45 04, E-mails: mcricia@ub.edu, mgallego@ub.edu, rfisa@ub.edu, mportus@ub.edu. Anna Guaro, Houssemon El Kassab, José María Jorba, Carmen Martin, and Alejandro Lobo, Servei de Pediatría, Hospital Residencia Sant Camil, Sant Pere de Ribes, Carretera Puigmoltó Km 0.8, Sant Pere de Ribes, 08810 Barcelona, Spain, Telephone: 34 93 896 00 25, Fax: 34 93 896 12 87, E-mails: santcamil@hrsantcamil.es, 34578hk@comb.es, C.MEDIC@terra.es, santcamil@hrsantcamil.es. Montserrat Castro, Servei de Anatomia

ELISA detected high levels of IgG-specific antibodies in the newborn and in the mother. Nevertheless, high levels of IgG in the former has low predictive value of congenital transmission because of the passive transfer of maternal IgG antibodies, which could be present for up to 7 months in the non-infected child. Comparison of the immunologic profile of the mother and child, detected by WB analysis, allows differentiation between passively transmitted maternal antibodies and those newly synthesized in the infant, which are used in early postnatal diagnosis of congenital toxoplasmosis. In our case, no differences in the band pattern of the sera from the mother and newborn were detected after delivery, and no new bands were recognized by child sera during the follow-up period. Therefore, neonate production of antibodies could not be appreciated.
CONGENITAL TRANSMISSION OF TRYPANOSOMA CRUZI IN EUROPE

REFERENCES