Case report of Zika virus during controlled ovarian hyperstimulation: results from follicular fluid, cumulus cells and oocytes

Edilberto Araújo Filho¹, Cássio Leão Fáció¹, Ligiane Alves Machado-Paula¹, Mariana Angelozzi de Oliveira², Ciro Dresch Martinhago², Leonardo Previato Araújo², Lígia Fernanda P. Araújo¹

¹Center of Human Reproduction of São José do Rio Preto, São José do Rio Preto - SP, Brazil
²Chromosome Genomic Medicine, São Paulo - SP, Brazil

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ABSTRACT

We describe a case of a 37-year-old female, indicated for in vitro fertilisation. She developed skin rash on her trunk and limbs, during the treatment. RT-PCR results were positive in her blood and negative in her husband's blood and semen. Oocyte aspiration was performed, retrieving 7 oocytes, follicular fluid, and cumulus cells. RT-PCR results for the follicular fluid and cumulus cells were negative for ZIKV, and positive for only 2 oocytes. This is the first report in the literature analysing ZIKV in the follicular fluid, cumulus cells, and oocytes, and will contribute to the understanding of ZIKV infection and transmission.

Keywords: Zika virus, in vitro fertilisation, follicular fluid, oocytes, RT-PCR

INTRODUCTION

Zika virus (ZIKV) is a member of the Flaviviridae virus family and is primarily transmitted by infected domestic mosquitoes such as Aedes aegypti and Aedes albopictus to a lesser extent. It was first isolated in 1947 from a Rhesus monkey in Zika forest in Uganda and was recognised in humans in 1952 in Nigeria (Dick, 1952). ZIKV infection was subsequently observed in numerous countries in Africa and Southeast Asia (Hayes, 2009; Faye et al., 2014). In 2007 there was the major outbreak on Yap Island in the Federated States of Micronesia, followed by French Polynesia in 2013 and 2014 (Cao-Lormeau et al., 2014; Lessler et al., 2016). In 2015 and 2016 there was an epidemic in the Americas, starting in 2015 in Brazil, and spreading throughout South America, Central America, North America and the Caribbean (Henssey et al., 2016).

During this outbreak of ZIKV in Brazil, an increased number of cases of newborns with microcephaly were reported in northeastern Brazil (Ministério da Saúde, 2016). In 2016, the causal association between ZIKV during pregnancy and foetal microcephaly and other brain anomalies was acknowledged by the World Health Organization (WHO) and the U.S. Center for Disease Control and Prevention (CDC) (WHO, 2016; Center for Disease Control and Prevention, 2016; Rasmussen et al., 2016).

Intrauterine and perinatal transmission of ZIKV has also been found (Besnard et al., 2014), and there are evidences of sexual transmission (Hills et al., 2016; McCarthy, 2016). These facts must be considered in reproductive-assisted treatments. Here we present a case report of a patient who contracted ZIKV during in vitro fertilisation (IVF). The objective of this observational study is to discover if ZIKV is present in the follicular fluid, cumulus cells and oocytes during an IVF cycle.

CASE DESCRIPTION

Patient

A 37-year-old female, indicated for IVF due to bilateral tubal factors, initiated the ovulation induction on day 2 of her menstrual cycle. One day later, she travelled to São Paulo on business with her mother and sister, taking the medications. She returned and underwent an ultrasound on day 5 of the treatment, showing a skin rash on her trunk, spreading to limbs. She reported that her mother and sister had the same symptoms. The patient’s temperature was 37°C, and she had mild low back pain and insignificant joint pain. A complete blood count (CBC) did not show changes in the platelets and leukocytes. The patient and her husband had a blood test for ZIKV using a reverse transcriptase polymerase chain reaction (RT-PCR). The patient offered to follow the treatment until oocyte aspiration at the clinic’s expense. Oocyte aspiration was performed and seven MII stage oocytes were retrieved. The patient’s follicular fluid and cumulus cells were donated for study in addition to her husband’s semen.

This is a case report of a single-patient at our private IVF center, thus no Institutional Review Board (IRB) approval was obtained, because our IRB designates a single-patient case report as not subject to IRB review because it does not meet the definition of human subjects research. Nevertheless, informed written consent was taken from the couple.

METHODOLOGY

Ovarian stimulation protocol

A basal ultrasound was performed on the second day of the patient’s cycle. Recombinant follicle-stimulating hormone (r-FSH; Gonad Sereno, Rio de Janeiro, Brazil) at a dosage of 150 IU was administered from days 2 to 4 of her menstrual cycle. Human menopausal gonadotropin (hMG; Menopur, Ferring, São Paulo, Brazil) at a dosage of 150 IU was administered from days 2 to 6. Ultrasound was performed on day 7 of the cycle and daily afterward until the administration of recombinant human chorionic gonadotropin (r-hCG; Ovidrel, Merck Serono, Rio de Janeiro, Brazil).

Gonadotrophin-releasing hormone (GnRH) antagonist (Cetrotide, Merck, Rio de Janeiro, Brazil) was initiated on day 7, when the follicles reached 16 mm in diameter. When two or more follicles reached 20 mm diameter, 250 mcg of recombinant human chorionic gonadotropin (r-hCG; Ovidrel, Merck Serono, Rio de Janeiro, Brazil) was administered and oocyte aspiration was performed 36 hours later using a transvaginal ultrasound-guided needle.

The patient had eight follicles and we collected seven MI stage oocytes. The oocytes, follicular fluid and cumulus corona cells were cryopreserved.
**Genetic analysis**

The blood samples obtained from the patient and her husband, the follicular fluid, cumulus cells and oocytes from the patient, and semen from her husband was tested for ZIKV RNA using a real-time reverse transcriptase polymerase chain reaction (RT-PCR). The genetic analysis was performed at the Chromosome Genomic Medicine laboratory in Sao Paulo, Brazil.

**Purification of viral RNA**

RNA was extracted from 140 μl of the patient and her husband’s samples using a QIAamp Viral RNA Mini Kit (Qiagen, Hilden, Germany), following the manufacturers’ protocol.

Briefly, 140 μl of samples were mixed with AVLCarrier RNA buffer. After 10 min of incubation at room temperature, 560 μl of ethanol (96-100%) was added. The samples were transferred into a column containing silica and centrifuged at 6000 x g for 1 min. The RNA was washed twice with buffer AW1 and AW2, respectively. RNA was eluted in 60 μl of AVE buffer and stored at -20°C until use.

**Real-time RT-PCR**

All assays were performed using the RT-PCR Kit BR ZIKV (LCG Biotechnology, Brazil) with amplification in the StepOnePlus(tm) Real-Time PCR System (Thermo-Fisher Scientific, Waltham, MA, USA), following the manufacturer’s protocol.

For the real time RT-PCR, was used with a 25 µl reaction mixture under the following conditions: 12.5 µl of 2 × RT-PCR buffer, 0.5 µl of One-Step RT-PCR enzyme; 0.5 µl of primers and a BR-ZIKV probe; 0.5 µl of set of primers and a probe of the internal control (IC); 10 µl purified viral RNA (sample), and 1.5 µl nuclease-free water. The negative control consisted of blank reagent and water. Nucleic acid extracted from virus stocks was used for the positive control.

The following thermal profile was used for a single cycle of 30 min at 55°C for reverse transcriptase, a single cycle of 30 sec at 95°C for initial denaturation, followed by 40 amplification cycles of 10 sec at 95°C for denaturation, 15 sec at 55°C for annealing, and 45 sec at 68°C for extension. The data were analysed using StepOne Plus software (SDS software from Thermo-Fisher).

**RESULTS**

The real-time RT-PCR results for ZIKV in the blood was positive for the patient, and negative for her husband, and his semen was also negative. Both the follicular fluid and cumulus cells were negative for ZIKV RNA. ZIKV RNA was present in two of the seven oocytes.

**DISCUSSION**

This is the first report in the literature analysing the presence or absence of ZIKV in the follicular fluid, cumulus cells and oocytes. It will improve the understanding of the transmission and infection of this virus. ZIKV infection has been associated with foetal abnormalities such as congenital microcephaly and can lead to pregnancy loss (Miner et al., 2016; Plourde & Bloch, 2016). ZIKV can be present in the amniotic fluid, suggesting that the virus can cross the placental barrier (Calvet et al., 2016; Plourde & Bloch, 2016). ZIKV has also been isolated from other body fluids, as blood, urine, saliva, breastmilk, and semen (Plourde & Bloch, 2016). Sexual transmission has been described in several publications (D’Ortenzio et al., 2016; Plourde & Bloch, 2016). In this report, the absence of the ZIKV in the husband’s semen indicated that he was not infected sexually. Studies have found that male-to-female transmission is the most common (D’Ortenzio et al., 2016), but there have also been reports of sexual transmission from females to males (Davidson et al., 2016) and from males to males (Deckard et al., 2016).

The presence of ZIKV RNA in the two oocytes shows the importance of testing couples seeking assisted reproduction techniques because the risk of contaminating the embryo is real.

However, only two oocytes, out of seven were infected by the virus. That could be explained, in part, by the fact that RNA quickly degrades, and we may have missed detecting the virus in the other oocytes. Furthermore, we do not know if the virus was in the zona pellucida and in the oolemma, because the PCR did not separate these two areas of the oocyte.

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**CONFLICT OF INTERESTS**

The authors declare no conflict of interests.

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