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Viral Infection of the Placenta Leads to Fetal Inflammation and Sensitization to Bacterial Products Predisposing to Preterm Labor

Ingrid Cardenas,*† Robert E. Means,†,‡ Paulomi Aldo,* Kaori Koga,* Sabine M. Lang,† Carmen Booth,‡ Alejandro Manzur,§ Enrique Oyarzun,§ Roberto Romero,§ and Gil Mor*†

Pandemics pose a more significant threat to pregnant women than to the nonpregnant population and may have a detrimental effect on the well being of the fetus. We have developed an animal model to evaluate the consequences of a viral infection characterized by lack of fetal transmission. The experiments described in this work show that viral infection of the placenta can elicit a fetal inflammatory response that, in turn, can cause organ damage and potentially downstream developmental deficiencies. Furthermore, we demonstrate that viral infection of the placenta may sensitize the pregnant mother to bacterial products and promote preterm labor. It is critical to take into consideration the fact that during pregnancy it is not only the maternal immune system responding, but also the fetal/placental unit. Our results further support the immunological role of the placenta and the fetus affecting the global response of the mother to microbial infections. This is relevant for making decisions associated with treatment and prevention during pandemics. The Journal of Immunology, 2010, 185: 1248–1257.

Pregnant women are more susceptible to the effects of microbial products (i.e., endotoxins) and were the most vulnerable subjects during the 1918 pandemic (influenza A subtype H1N1), with a mortality rate that ranged between 50 and 75% (1). Exposure to the virus during pregnancy may also have overt or subclinical effects that become apparent only over time. Although substantial progress has been made in the understanding of the immunology of pregnancy, many unanswered questions remain, especially those associated with the susceptibility and severity of infectious agents of mothers and unborn children (2), (3).

Epidemiological studies have demonstrated an association between viral infections and preterm labor (4, 5) and fetal congenital anomalies of the CNS and the cardiovascular system (6–8). Although some viral infections during pregnancy may be asymptomatic (9), approximately one-half of all preterm deliveries are associated with histological evidence of inflammation of the placenta, termed acute chorioamnionitis (10), or chronic chorioamnionitis (10). Despite the high incidence of acute chorioamnionitis, only a fraction of fetuses have demonstrable infection (11). Most viral infections affecting the mother do not cause congenital fetal infection, and only in a small number of cases is the virus found in the fetuses (12–17), attesting to the unique ability of the placenta to act as a potent barrier with an immune-regulatory function that protects the fetus from systemic infection (10, 12, 18, 19).

Recent observations indicate that rather than acting as a mechanical barrier, the placenta functions as a regulator of the trafficking between the fetus and the mother (20–22). Fetal and maternal cells move in two directions (23, 24); similarly, some viruses and bacteria can reach the fetus by transplacental passage with adverse consequences (25). Although viral infections are common during pregnancy (26), transplacental passage and fetal infection appear to be the exception rather than the rule (27, 28; reviewed in Ref. 29 and subsequent references).

There is a paucity of evidence that viral infections lead to preterm labor (10, 19, 22–24); however, there are several areas of controversy and open questions. For example, what effects do subclinical viral infections of the decidua and/or placenta during early pregnancy have in response to other microorganisms, such as bacteria; and what is the effect of a subclinical viral infection of the placenta on the fetus?

The trophoblast is an important component of the placenta, and it is able to recognize and respond to microorganisms and their products through the expression of TLRs (30–32). TLRs are a family of innate immune receptors that have an essential role in the recognition of pathogen-associated molecular patterns (33–35). Trophoblasts are able to produce cytokines/chemokines and antiviral factors following TLR-3 ligation in vitro, suggesting the potentially active role of these cells in the control of viral infections (20, 36). Some of these receptors (chemokine and TLRs) may also function as viral receptors mediating viral recognition and entry into the...
trophoblast. Signaling through TLRs has been shown to induce murine γ-herpesvirus 68 (MHV-68) reactivation in vivo (37).

Herpesviruses are the most common cause of viral-related perinatal neurologic injury in the United States (38). However, among the eight known human herpesviruses, most reported adverse pregnancy and neonatal outcomes are the result of the HSVs (HSV-1 and HSV-2) and CMV (39) and usually occur due to a primary infection of the mother during the first trimester or infection of the infant during delivery. MHV-68 (murid herpesvirus 4 [NC_001826.2]) is a γ-herpesvirus of rodents that shares significant genomic colinearity with two human pathogens, EBV and Kaposi’s sarcoma-associated herpesvirus (40). As in these two viruses, the effect of MHV-68 in pregnancy is unknown.

We developed a novel murine model to evaluate the role of viral infection in pregnancy and fetal development. Our data suggest that even in the absence of placental passage of the virus, the fetus could be adversely affected by an inflammatory response mounted in response to viral invasion of the placenta. Furthermore, we demonstrate that a viral infection in early pregnancy sensitizes the pregnant mother to the effects of bacterial products later on in gestation, and specifically, to premature labor. These data suggest that exposure to early viral infections may program the immune response of mother and fetus. Such observations have important consequences for understanding the potential risk of viral infections during pregnancy and the importance of adequate surveillance to prevent maternal mortality and subclinical fetal injury, leading to long-term consequences.

Materials and Methods

Virus culture

MHV-68 expressing GFP (provided by R. Sun, University of California, Los Angeles, CA) was passaged in NIH 3T3 cells with DMEM plus 10% FCS. After lysis, supernatant was harvested, filtered (0.45-μm pore), and titered by 2-fold serial dilutions. To determine virus load in infected mice, frozen homogenized tissues were minced and subjected to 10-fold serial dilutions, and endpoint titers were determined in NIH 3T3 cells by GFP (41, 42). A single virion or DNA copy was sufficient to show a positive result by plaque assay or PCR.

Animal procedures

C57BL/6 mice were obtained from The Jackson Laboratory (Bar Harbor, ME), and TLR-3 knockout (KO) was provided by R. A. Flavell (Yale University, New Haven, CT). Adult mice (8–12 wk of age) with vaginal plugs were infected i.p. at embryonic day (E) 8.5 postconception with either 1 × 106 PFU MHV-68 expressing GFP (in 200 μl vol) or DMEM (vehicle). Three or 9 d postinfection (dpi), animals were sacrified, and organs were removed, fixed in 4% paraformaldehyde, and/or stored at −80°C. All animals were maintained in the Yale University School of Medicine Animal Facility under specific pathogen-free conditions. All experiments were approved by the Yale Animal Research Committee.

Reagents and Abs

LPS (Escherichia coli O111:B4) was purchased from Sigma-Aldrich (St. Louis, MO). Lymphocyte separation media was purchased from MP Biomedicals (Solon, OH).

For NK and macrophage detection, biotinylated lectin Dolichos biflorus agglutinin from Sigma-Aldrich and rat anti-mouse F4/80 Ab (eBioscience, San Diego, CA) were used, respectively. Anti-NF-κB p65 mAb was purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Dominant-negative (DN) TLR1 Toll/IL-1 receptor (TIR) and TLR2TIR, incapable of transducing a signal after ligand binding, were purchased from InvivoGen (St. Louis, MO). Lymphocyte separation media was purchased from MP Biomedicals (Solon, OH).

Human first trimester trophoblast HTR-8 cells were gifted from C. Graham (Queen’s University, Kingston, Ontario, Canada). Human first trimester trophoblast 3A cells were stably transfected with DN TLR2 and TLR1 genes, as previously described (22). Following transfection, cells expressing the TLR1/TLR2 (TLR1-DN) or the TLR2/TLR3 (TLR2-DN) were selected with puromycin. Vector alone-transfected cells served as negative control.

Human trophoblast isolation

Three or 9 d postinfection (dpi), animals were sacrificed, and organs were removed, fixed in 4% paraformaldehyde, and/or stored at −80°C. After three washes, 50 μl streptavidin-PE was added to each well and incubated for 10 min at 37°C in the dark at room temperature. After three washes, 50 μl streptavidin-PE was added to each well and incubated for 30 min in the dark at room temperature. After a final wash, the beads were resuspended in 125 μl assay buffer for measurement with the LUMINEX 200 (LUMINEX, Austin, TX). The cytokines included in the Multiplex assay were as follows: IL-1β, IL-10, GM-CSF, IFN-γ, TNF-α, IL-1α, IL-6, IL-12p40, IL-12p70, G-CSF, KC, MIP-1α, RANTES, MCP-1, and MIP-1β.

Immunohistochemistry

For Ag retrieval with Retrieval Agent (pH 6.0; BD Biosciences), macrophages were detected in paraffin-embedded murine uterus with rat anti-mouse F4/80 Ab at 1:20. NK cell detection was performed, as previously reported (31, 51).

For the localization of the p65 subunit of NF-κB, MHV-68-infected human trophoblast cells were fixed and incubated with the mouse anti-NF-κB p65 Ab. Slides were then incubated with Alexa Fluor546 anti-mouse IgG and counterstained with Hoechst 33342 dye (Molecular Probes, Eugene, OR).

Total RNA isolation

Total RNA was extracted using TRIzol, and cDNA was prepared using a Verso cDNA kit per manufacturer’s protocol (Thermo Scientific, Waltham, MA).

Real-time PCR

Real-time PCR was performed in duplicate using SYBR Green (Invitrogen, Carlsbad, CA) in an ABI Prism 7500 (Applied Biosystems, Foster City, CA). cDNA sample (1 μl) was amplified with gene-specific primers using optimized PCR cycles. GAPDH was used as an endogenous control for relative comparison of human TLR-2, TLR-3, and TLR-4. GAPDH expression did not vary with treatments. TLR-2 forward, 5′-ATGCCTACT-GGGTGCGAAGC-3′; TLR-2 reverse, 5′-AGTGCACCACCTCACA-3′; TLR-3 forward, 5′-GGTGGGTTGCTTGGGCA-3′, TLR-3 reverse, 5′-AGTGGTCTCGATGTCTGGTG-3′; TLR-4 forward, 5′-CAGCTCCTTGGT- GGAAGTGTGA-3′; TLR-4 reverse, 5′-CACAGAAGACCATCAGGG-3′, GAPDH forward, 5′-TCTGACACGCGGTTGGTTG3′, GAPDH reverse, 5′-GACAAGCTCCGTTTCAAGC-3′.

The TLR-2, TLR-3, and TLR-4 cycle threshold value was analyzed using the ΔΔCT cycle threshold Livak method (25).

ELISA

Serum from wild-type (wt) and TLR3 KO mice was analyzed for the presence of anti-viral IgM or IgG Abs. For coating of ELISA plates, MHV-68 GFP stocks were filtered through a 0.45-μm-pore membrane, and virions were centrifuged through a 5% sucrose cushion (SW28, 20,000 rpm for...
75 min at 4°C. Virion pellets were resuspended in PBS containing 0.5% Triton X-100 and 0.5% FBS to achieve 5000× concentration. Maxisorb plates (Nunc-immuno plates) (Corning Life Sciences, Corning, NY) were coated with 91 ng virus protein/well, washed, blocked with 0.3 mg BSA, and incubated with serum from either wt or TLR3 KO with or without MHV-68 infection. After washing, goat anti-mouse IgG or IgM, HRP-conjugated Ab (Southern Biotechnology Associates, Birmingham, AL) was added at 1/4000 dilution in PBS/1% BSA for 2 h at room temperature, washed, and detected with tetramethylbenzidine substrate at 405 nm.

Statistical analysis
Data are expressed as mean ± SE for in vitro study and median ± first or third quartiles for in vivo study. Statistical significance (p < 0.05) was determined using either two-tailed unpaired Student t tests or Mann-Whitney U test for nonparametric data. Unless stated otherwise, all experiments were performed in duplicate.

Results
Maternal infection with MHV-68 does not induce preterm labor
Systemic administration of polyinosinic:polycytidylic acid [poly (I:C)] to pregnant mice induced preterm labor and delivery, and production of proinflammatory cytokines (53, 54). The inflammatory response to the TLR-3 ligand was found in the placenta at E17.5 (local response) as well as in the spleen (systemic response) (30, 55) characterized by upregulation of IL-6, IL-12p40, MCP-1, MIP-1β, growth-related oncogene-α, and RANTES. These observations indicated that the placenta was able to recognize and respond to viral products. To understand the effects of viral infection in pregnancy, we used MHV-68. C57BL/6 pregnant mice received 1×10⁶ PFU MHV-68 i.p. on E8.5 of pregnancy, and were followed up to E17.5. Control animals received media as a placebo. This viral dose has previously been shown to infect mice organs and produce a systemic viral response (37).

Maternal infection with MHV-68 had no effect on pregnancy outcome, including litter size, weight, or gestational age at delivery (Fig. 1). We then evaluated whether the absence of viral infection to the placenta and decidua was responsible for the lack of effect in pregnancy outcome. To test this hypothesis, replicative viral loads (PFU/ml) were determined using a limiting dilution plaque assay on frozen tissue taken from the placenta and decidua (local), and spleen and lymph node (systemic and classical target organ for MHV-68) (56) from pregnant mice who had received MHV-68 on E8.5 of pregnancy and sacrificed 3 or 9 d after viral administration (dpi).

Three days after viral administration, we observed a high viral load in the spleen and lymph nodes. Interestingly, viral titers in the decidua were significantly higher than those in the spleen (Fig. 2A). The placenta was also infected, but overall viral titers were lower than those in the spleen.

Nine days after viral administration, we observed a substantial increase in splenic viral titers (higher than 3 logs), a slight decrease in the decidua, and increasing viral titers in the placenta (Fig. 2B). Most notably, no viruses were detected in the fetus of infected mothers using the plaque assay or by PCR, even 9 d after viral administration (Fig. 2A and data not shown).

These results suggest that the viral load administered to the mice is able to infect all the organs, including placenta and decidua; however, in contrast to the effect observed with poly(I:C), the viral infection in the placenta and decidua did not seem to have an effect on pregnancy outcome. To better understand the differences between the two responses [virus versus poly(I:C)], we evaluated the cytokine/chemokine profile in the placenta and spleen of mice infected with MHV-68. MHV-68 infection did not induce the production of chemokines and inflammatory cytokines seen with poly(I:C) administration (Fig. 2C, 2D). Moreover, we observed inhibition on the production levels of IL-6, MIP-1β, and RANTES in the placenta of MHV-68–infected mice. These data suggest that the change in the cytokine profile may play an important role in the induction of preterm labor/delivery.

TLR-3 is necessary for the control of early viral infection
The effects of poly(I:C) in pregnancy outcome (and, in particular, preterm labor) are mediated through TLR-3 (30, 35), and this pattern recognition receptor plays an important role in the immune response to herpesviruses (57). Moreover, some viruses such as influenza A and Kaposi’s sarcoma-associated herpesvirus have been associated with activation of the TLR-3 pathway in humans (58, 59). Thus, we evaluated the role of TLR-3 on MHV-68 infection during pregnancy by using TLR3 KO mice. Similarly, as in the wt mice, administration of MHV-68 had no effect on the duration of pregnancy (i.e., there was no premature delivery).
FIGURE 2. Effect of MHV-68 viral infection in pregnant mice. Viral titers as PFU/ml were determined in wt pregnant mice infected with MHV-68 (1 \times 10^6 PFU) 3 d (E11.5; A) and 9 d (E17.5; B) postinfection. Viral titers were observed in lymph nodes, placenta, decidua, and spleen, but were absent in the fetuses. *p < 0.05, decidua versus spleen. Placenta (C) and spleen (D) cytokine profile was determined in wt pregnant mice treated with poly(I:C) or MHV-68 4 and 9 d postinfection, respectively. *p < 0.05, MHV-68 versus control; #p < 0.05, poly(I:C) versus MHV-68. E and F, Viral titers as PFU/ml were determined in TLR-3 KO pregnant mice infected with MHV-68: E, 3 d (E11.5), and F, 9 d (E17.5) postinfection. Note the high levels of viral titers in lymph nodes, placenta, decidua, and spleen, but absent in the fetuses. Bars show median ± SEM. n = 6 mice per group.
labor). However, we observed higher viral titers in all tissues, including the decidua and placenta of TLR-3 KO mice, both at 3 and 9 dpi (Fig. 2E, 2F). The fetuses of either wt or TLR-3 KO-infected mothers were not infected, suggesting that TLR-3 is not required for the protection of fetuses against viral invasion.

We then evaluated whether the viral dose used in this study is able to mount an adaptive immune response by assessing seroconversion. IgG anti–MHV-68 were significantly higher in both wt and TLR-3 KO-infected pregnant mice than in noninfected animals. However, when we compared the response between wt and TLR-3 KO, we observed that IgG anti MHV-68 levels were significantly lower in the TLR-3 KO (Fig. 3). These results confirm that MHV-68 infections during pregnancy are able to mount a specific adaptive immune response characterized by the presence of anti–MHV-68 IgG. The presence of higher viral titers and low levels of anti–MHV-68 IgG in the TLR-3 KO mice indicated that TLR-3 expression is required to elicit a potent antiviral Ab response against this dsDNA virus.

Effect of MHV-68 infection on the placenta

Because we observed viral infection of the placenta and decidua in mothers, the next objective was to determine the effect of MHV-68 infection on the placental and decidual pathology. Thus, uteroplacental units were collected at 9 dpi, and H&E staining was performed. All histological samples were analyzed in a blinded manner by an independent animal pathologist (C.B.). Sites of edema were observed only in the decidua of infected mice; necrosis and inflammation foci were observed in the labyrinth of infected mice (Fig. 4A). Significant pathologic changes present within the labyrinth of TLR3 KO-treated mice included an overall tissue hyperesinophilia, nuclear pyknosis, cellular fragmentation, and multifocal loss of tissue of architecture (necrosis) in the labyrinth (Fig. 4B).

We then evaluated changes of the number and distribution of NK (lectin-positive) cells and macrophages (F4/80 positive). NK cells were observed in decidua of control as well as infected mice. No change in the location of these cells was observed as a result of the infection (data not shown). Macrophages are mainly localized in the myometrium and decidua of the pregnant mice (Fig. 4C, arrows). Few macrophages are also found in the placenta. No differences in the distribution and number of macrophages were found in the decidua and placenta from infected and control groups.

In addition, we observed an increase in collagen deposition in the perivascular spaces of infected animals, predominantly in the labyrinth layer, as compared with control mice (Fig. 4D). The presence of collagen in the perivascular areas suggests that an active repair process was taking place in the placenta of infected mice. Similar changes were observed in TLR-3 KO mice (Fig. 4D).

Fetal response to placental infection

Although we observed high viral titers in the placenta, no virus was detected in any of the fetuses, as determined by the limiting dilution plaque assay and confirmed by PCR. To determine whether the lack of fetal infection was due to inability of the virus to infect fetal cells, mouse embryonic fibroblast cells were isolated and infected with MHV-68 in vitro with a similar dose as that used for trophoblast cells (see below). Eighty percent of embryonic fibroblasts were infected by MHV-68 in less than 12 h, as shown by GFP-positive signal; however, the viral infection induced a lytic effect (data not shown). These results suggest that the placenta is functioning as an immunological barrier, capturing the virus and preventing it from reaching the fetus. To determine whether the infection of the placenta could have an effect on the developing fetus, we next assessed fetal morphology from mice infected with MHV-68 during pregnancy versus those receiving a placebo.

Analysis of the fetuses revealed that viral infection of the mother has a transient effect on development. Three dpi, fetuses of infected mothers were smaller and had a lower weight (in both wt and TLR-3 KO mice), although this effect was more evident in the TLR-3 KO group (Supplemental Fig. 1A). Furthermore, we observed a delay in the process of differentiation of the eye, tails, and limbs (Supplemental Fig. 1B). However, after 9 d, the differences between the fetuses from infected and noninfected mice from both wt and TLR-3 KO were no longer detectable (Supplemental Fig. 1C). These important observations are evidence of the remarkable plasticity of the developing fetus.

Because we observed an early effect on fetal development, we then evaluated the integrity of fetal organs and tissues using microscopic sections. Despite the absence of viruses in the fetuses, we noted severe pathological changes in the fetal tissues of infected mothers from both wt and TLR-3 KO. We observed hydrocephalus, defined as an increase in the subarachnoid space, in the brains of all fetuses from infected mothers (Fig. 4E). We did not see any changes in the lateral ventricles, nor did we detect abnormal immune infiltration or white matter damage.

In the thoracic cavity, the pathological changes were characterized by the presence of hemorrhage inside the lungs and pericardium in all treated animals compared with the controls (Fig. 4F). However, there was no damage in the abdominal cavity or the limbs.

We then evaluated the cytokine profile in fetuses from infected and control mothers. Interestingly, 9 dpi, we observed a significant increase in the levels of fetal proinflammatory cytokines (Fig. 4G), including high levels of IFN-γ and TNF-α. The presence of these two cytokines may explain some of the morphological changes observed in these fetuses.

Collectively, these data suggest that although there is no demonstrable fetal viral infection, the presence of an active inflammatory response in the placenta and decidua can have a direct effect on fetal development.

Trophoblast-viral interaction

To understand the implications of these observations in humans, trophoblast cells were isolated from first trimester human placentas and infected with GFP-MHV-68 for either 24 or 48 h. Infection was monitored by the presence of GFP. Positive GFP-MHV-68–infected trophoblast cells were observed ≈12 h postinfection and remained viable up to 6 dpi (Fig. 5A).

Next, we determined the cytokine response induced by MHV-68 in trophoblast cells in vitro. Contrary to what we observed with poly
The Journal of Immunology 1253

Discussion

We demonstrated that maternal viral infection can lead to productive replication in the placenta and a fetal inflammatory response, even though the virus is not detected in the fetus. The experiments described in this work are intended to show that viral infection of the placenta can elicit a fetal inflammatory response, which in turn can cause organ damage and, potentially, downstream developmental deficiencies. Furthermore, we demonstrated that a viral infection of the placenta may sensitize to bacterial infection and promote preterm labor.

Pregnant women are exposed to many infectious agents that are potentially harmful to the fetus. The risk evaluation has been focused on whether there is a maternal viremia or fetal transmission (62). Viral infections that are able to reach the fetus by crossing the placenta might have a detrimental effect on the pregnancy (63, 64). It is well accepted that in those cases infection can lead to embryonic and fetal death, induce miscarriage, or induce major congenital anomalies (62, 65). However, even in the absence of fetal viral infection, the fetus could be adversely affected by the maternal response to the infection. Examples are infections with HIV, hepatitis B, varicella zoster virus, and parvovirus B19, among others (5, 28, 66, 67). Indeed, viral crossing of the placenta may be the exception rather than the rule.

One of the main questions of this study was how a microorganism, in this case a virus, might initiate a response that may not lead to preterm labor, but would alter the immunologic balance at the maternal fetal interface. Poly(I:C) has been used in several studies as a model for TLR3 activation and shown to be a potent inducer of preterm labor (68). However, the use of MHV-68, which is able to activate TLR-3 (61), did not show the same outcome. Our results indicate that only a condition characterized by the expression of inflammatory cytokines at the maternal-fetal interface will trigger a cascade of events leading to the termination of the pregnancy. In contrast, a viral infection in the placenta that triggers a mild inflammatory response will not terminate the pregnancy, but is able to activate the immune system not only of the mother, but of the fetus as well.

It is critical to take into consideration the fact that during pregnancy it is not only the maternal immune system responding, but also the fetal/placental unit. Our results further support the immunological role of the placenta and the fetus affecting the global response of the mother to microbial infections. This is relevant for making decisions associated with treatment and prevention during pandemics.

An important observation in this study is the fact that even though there is a high viral titer in the placenta and decidua, no virus was detected in the fetus. This result further confirms our and others’ studies suggesting that the placenta is an active barrier, able to control an infection and protect the fetus (49, 69–72). However, the inflammatory response originated on the maternal side has a negative impact on the fetus and triggers a fetal inflammatory condition.

Fetal inflammatory response syndrome (FIRS) is a condition in which, despite an absence of cultivable microorganisms, neonates with placental infections have very high circulating levels of inflammatory cytokines, such as IL-1, IL-6, IL-8, and TNF-α (73–75). We observed a similar outcome in our animal model in which MHV-68 infection of the placenta triggers a fetal inflammatory condition.
FIGURE 4. Effect of MHV-68 viral infection on the maternal/fetal interface. Morphological changes were observed in the placenta and decidua of MHV-68–infected pregnant mice associated with the following: A, edema (++) in the D, but absent in the L and S. Upper and lower panel, scale bar, 200 and 300 μm, respectively. B, Necrosis in placenta, marked loss of cellular detail, fragmentation, hypereosinophilia (boxes) in the labyrinth subjacent to the epithelium (E), and necrosis of scattered giant cells (arrowheads). These changes were more accentuated in the TLR-3 KO compared with wt mice. Scale bars, 200 μm. C, Immunohistochemistry for F4/80-positive macrophages (brown) localized in the myometrium (MYO) and decidua (DEC) at E11.8 and E17.5. Black arrows show the edge between myometrium and decidua. Original magnification ×20. D, Increase in collagen deposition (arrows) in the labyrinth layer of MHV-68–infected mice using Trichromic Mason staining (original magnification ×20). E, Presence of fetal brain hydrocephalus (black arrows) in wt and TLR-3 KO MHV-68–infected mice (middle and right panel) compared with normal controls (left panel). Note the width of LVs. Original
response similar to the one observed in FIRS, even though the virus was not able to reach the fetus. In the case of human FIRS, these cytokines have been shown to affect the CNS and the circulatory system (76). In this study, we showed that fetal morphologic abnormalities may be caused by fetally proinflammatory cytokines, such as IL-1, TNF-α, MCP-1, MPI-β, and IFN-γ. Beyond morphological effects on the fetal brain, the presence of FIRS increases the future risk for schizophrenia, neurosensorial deficits, and psychosis induced in the neonatal period (77–79).

Therefore, we propose that an inflammatory response of the placenta, which alters the cytokine balance in the fetus, may affect the normal development of the fetal immune system, leading to anomalous responses during childhood or later in life (77–79). One example of this is the differential responses in children to vaccination or the development of allergies (11, 80). Antenatal infections can have a significant impact on later vaccine responses. We can observe this type of outcome in other conditions associated with placental infection, such as malaria. A few studies suggest that surviving infants with placental malaria may suffer adverse neurodevelopmental sequelae and may have an abnormal response to a later infection with the parasite (81). In the majority of the cases, the parasite did not reach the fetus, but the inflammatory process in the placenta affected the normal fetal development (82).

The differential cytokine response observed between poly(I:C) and MHV-68 infection questioned the role of TLRs during a viral infection. However, our finding demonstrates a unique interaction between the virus and TLR expression and function at the placenta and decidua. We confirmed that TLR-3 is necessary for the control of viral replication, as demonstrated by the presence of higher titers of MHV-68 found in the TLR-3 KO mice. According to this, clinical data proved that TLR-3 controls herpesvirus infection, because children with a TLR-3 deficiency are very susceptible to HSV-1–induced encephalitis (83). In contrast, the virus requires TLR-2 and TLR-1 expression for its own replication. These findings open the possibility of using TLR-2 or TLR-1 antagonists as potential agents for preventing herpes viral replication.

Viral infection may influence the outcome of a concurrent bacterial infection (84); however, to date there is no evidence indicating whether a viral infection sensitizes to bacterial infection during pregnancy. We showed that MHV-68–infected pregnant mice underwent preterm labor following injection of a low dose of LPS, which has almost no effect on noninfected mice. These results suggest that a viral infection during pregnancy increases the risk of preterm labor.
or maternal death in response to other microorganisms, such as bacterial infection. In the pandemic of 1918, high rates of pregnancy loss and preterm delivery were reported (1), and during the pandemic of 1957–1958, an increase in CNS defects and other adverse outcomes were reported. In the more recent H1N1 influenza virus pandemic of 1957–1958, an increase in CNS defects and other adverse outcomes were reported. In all these cases, a bacterial-associated complication was reported.

In conclusion, we demonstrate that even in the absence of fetal viral infection, the inflammatory response originating in the placenta and decidua induces an inflammatory process with potential damage in fetal organs. It is therefore essential to evaluate the presence of maternal viral infections prenatally to prevent long-term adverse outcomes for the child and the mother. Future studies are needed to develop useful biomarkers for viral infections during pregnancy even in a subclinical stage as a strategy of early detection and prevention of fetal damage and maternal mortality. Furthermore, it is extremely important to take into consideration the possibility of placental infection when determining a response to emerging infectious disease threats.

Disclosures
The authors have no financial conflicts of interest.

References
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Corrections


The sixth author’s middle initial was omitted. The correct name is Carmen J. Booth.

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Sup Figure 1

Wild Type/TLR3-KO
CONTROL

a

TLR3-KO
MHV 68

b

*
Sup Figure 1c

For E11.5:
- **Wild Type**: CTR, MHV68
- **TLR3 KO**: CTR, MHV68

For E17.5:
- **Wild Type**: CTR, MHV68
- **TLR3 KO**: CTR, MHV68

The graphs compare the expression levels of gm in different conditions at E11.5 and E17.5.
Sup Figure 2

<table>
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<td><img src="image1" alt="WT Phase" /></td>
<td><img src="image2" alt="TLR-2 Δ Phase" /></td>
<td><img src="image3" alt="TLR-1Δ Phase" /></td>
</tr>
<tr>
<td>b</td>
<td><img src="image4" alt="WT Fluorescence" /></td>
<td><img src="image5" alt="TLR-2 Δ Fluorescence" /></td>
<td><img src="image6" alt="TLR-1Δ Fluorescence" /></td>
</tr>
</tbody>
</table>
Sup Figure 3

a  PBS + LPS

b  MHV68 + PBS

c  MHV68 + LPS
Sup Figure 4

a  PBS+LPS  b  MHV68+PBS  c  MHV68+LPS
SUPPLEMENTARY MATERIAL

Supplementary Figure 1
**Effect of MHV-68 infection on fetal development.** Fetal and placental samples from wt and TLR-3KO pregnant mice infected with murine herpes virus 68 (MHV-68) at E8.5 were collected at day E11.5. (a) Control fetus and placenta from WT and TLR3 KO mice treated with PBS. Note the normal development of the fetus* (b) fetus and placenta from TLR3 KO pregnant mice infected with MHV-68. Note the growth retardation of the fetus compared to the control. (* Fetuses). c) Fetal weight of fetuses from WT and TLR3 KO mothers at 11.5 and 17.5 days post conception. Note the significant decrease in weigh of TLR3 KO fetuses from MHV-68 infected mothers. * P<0.05. Bars show median ±SEM. Representative figure of six animals per group and three independent experiments.

Supplementary Figure 2
**Role of TLR-2 in MHV-68 replication in trophoblast cells.** First trimester trophoblast cells or trophoblast cells deficient of TLR-2 (TLR-2 DN) or TLR-1 (TLR-1 DN) were infected with GFP-labeled MHV-68, supernatant from these cultures were collected 48 hours after infection and transferred to new cultures of wild type first trimester trophoblast cells. (a) Wt trophoblast culture with supernatant from trophoblast infected with GFP labeled MHV-68 (b) Wt trophoblast culture with supernatant from TLR-2 DN trophoblast infected with GFP labeled MHV-68. (c) Wt trophoblast culture with supernatant from TLR-1 DN trophoblast infected with GFP labeled MHV-68. Magnification, X20. White arrows indicate the presence of infected cells. n= 3 samples per group. Figures are representative of three independent experiments.

Supplementary Figure 3
MHV-68 viral infection sensitizes to LPS treatment. Wt mice were infected with either MHV-68 (1X10⁹) or PBS at E8.5 followed by a single dose of LPS (20ug/kg) or PBS at E15. (a)Control group where pregnancy mice received PBS followed by LPS. No changes are observed in the vagina. (b) Pregnant mice infected with MHV-68 followed by PBS. No macroscopic changes were observed, similar as the control. (c) Pregnant mice infected with MHV-68 followed by LPS. Note the presence of vaginal bleeding and dilation associated with parturition. Figures are representative of three independent experiments. n=6 mice per group.

Supplementary Figure 4
**MHV-68 infection sensitizes to LPS induced preterm labor.** Groups are similar as in Supp Figure 3. While normal gestational sacs are observed in the control group and in the animals infected with MHV68 followed by PBS treatment; fetal death and preterm delivery is observed in animals infected with MHV68 followed by LPS treatment. Representative figures of at least 6 mice per group.