Inhibition of Heme Oxygenase-1 by Zinc Protoporphyrin IX Improves Adverse Pregnancy Outcomes in Malaria During Early Gestation

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The enzyme heme oxygenase-1 (HO-1) has cytoprotective effects by catalyzing the degradation of heme to produce carbon monoxide, iron and biliverdin. Furthermore, HO-1 activity has been associated with successful pregnancy. On the other hand, in the context of certain inflammatory conditions, HO-1 can induce iron overload and cell death.

To investigate the role of HO-1 in gestational malaria, pregnant BALB/c mice were infected with Plasmodium berghei ANKA in early, mid and late gestation. We found that malaria affected the pregnancy outcome in the three periods evaluated. However, only poor pregnancy outcomes in early pregnancy were related to HO-1 upregulation, iron overload, lipid peroxidation and necrosis of the decidua, which were prevented by HO-1 inhibition.

In conclusion, HO-1 expression must be finely tuned in gestational malaria to avoid the deleterious effect of increased enzyme activity.

Keywords: HO-1, malaria, pregnancy, uterine natural killer cells, iron overload, ferroptosis, ZnPPIX.

INTRODUCTION

Malaria in pregnancy is a serious public health problem. According to the World Health Organization, the prevalence of exposure to malaria during pregnancy was 29% in areas of moderate to high transmission in sub-Saharan Africa in 2018, where more than 11 million pregnant women acquired malaria in that year, and approximately 872,000 children were born with low birthweight due to the infection. These children are at high risk of stunted growth, poor cognitive development and death (1). Malaria in pregnancy can induce different outcomes for the mother and her offspring, potentially causing maternal anemia, severe malaria and maternal mortality. In addition, if infection occurs between the first and second trimester of pregnancy this may result in early abortion, intrauterine growth restriction and low birthweight (2).

Malaria is a hemolytic disease caused by protozoan parasites of the genus Plasmodium, in which heme released during the rupture of red blood cells leads to oxidative damage due to the production of reactive oxygen species that induce inflammation and tissue damage (3, 4). Because heme is also
toxic to parasites, they turn heme into a non-toxic product, the malarial pigment known as hemozoin (5). However, the main mechanism of catalysis of heme is through the activity of the enzyme heme oxygenase 1 (HO-1), which is a rate-limiting enzyme that catalyzes the degradation of heme to produce carbon monoxide (CO), iron and biliverdin (BV). The enzyme heme oxygenase 1 (HO-1) is a key enzyme that is involved in the metabolism of heme and plays a critical role in the regulation of cell death. HO-1 exerts cytoprotective effects through the activity of the lipid repair enzyme glutathione peroxidase 4 (GPX4) (18). GPX4 is the only enzyme capable of reducing hydroperoxides within membranes, making it a ferroptosis regulator molecule (19).

In mice, embryo implantation takes place when the blastocyst attaches to the cells of the luminal epithelium of the endometrium, inducing the process of decidualization, through which uterine stromal cells proliferate and differentiate into decidual cells (20). Simultaneously with decidualization, a highly organized lymphoid structure is formed adjacent to the decidua, the mesometrial lymphoid aggregate of pregnancy (MLAp) (21). Uterine natural killer cells (uNK) are abundant in decidual tissue and MLAp, constituting approximately 70% of the leukocytes infiltrating the decidua, where they fulfill important functions for the maintenance of pregnancy, among them the promotion of spiral artery (SA) remodeling (22, 23). Decidual cells supply nutritional support to the developing embryo until the establishment of the definitive placenta, which has a maternal layer (the decidua) and a fetal part formed by the junctional zone (which contains several trophoblast cells types), below which is the labyrinth layer, where gas and nutrient exchanges between mother and fetus occur (20, 24). HO-1 expression has been shown to be important for fetal tolerance, maintenance of placental architecture, uNK cell differentiation, SA remodeling, and preservation of normal blood pressure during pregnancy (25–28).

Previous studies have shown that experimental infection by abortive-causing bacteria such as *Listeria monocytogenes* and *Brucella abortus* leads to the reduced expression of HO-1 in trophoblast giant cells of the placenta, and the pharmacological induction of the enzyme prevented infectious abortion (29, 30). On the other hand, it has been reported that *Plasmodium berghei* ANKA infection induces an increase in protein and mRNA expression of HO-1 in the brain of BALB/c mice and in the lungs of DBA mice, conferring protection against the onset of experimental cerebral malaria and acute lung injury, respectively (31, 32). Interestingly, a previous report showed that iron overload in trophoblast cells in placentas from mice infected with *Plasmodium berghei* was associated with fetal death (33). However, the exact role of HO-1 and the iron status at the fetomaternal interface throughout pregnancy remains unclear in the context of malaria infection.

Despite the beneficial aspects frequently associated with HO-1 activity, excessive formation of degradation products as a result of increased expression can also lead to adverse effects. Iron overload and cell death have been associated with increased HO-1 activity in some inflammatory contexts such as *Mycobacterium tuberculosis* infection (13, 14), neuroinflammation during aging (15) and septic shock (16). The detrimental effects of the elevated expression of HO-1 are commonly attributed to iron overload due to its pro-oxidant nature, suggesting that the expression of this enzyme requires delicate modulation to preserve its beneficial effects (8). Labile iron is one of the main causes of lipid peroxidation, modifying physical properties of cell membranes, leading to loss of membrane permeability and cell death (17). In this sense, iron overload and the resulting lipid peroxidation may cause ferroptosis, a regulated non-apoptotic form of cell death marked by loss of activity of the lipid repair enzyme glutathione peroxidase 4 (GPX4) (18). GPX4 is the only enzyme capable of reducing hydroperoxides within membranes, making it a ferroptosis regulator molecule (19).

In order to verify the decidualization and embryo development in early pregnancy and to describe some of the alterations induced by infection during this stage, we inoculated mice with $5 \times 10^5$ infected red blood cells by intraperitoneal route in the first day of pregnancy (dP) and euthanized them 7 days later (Figure 1A). We observed that pregnant infected mice presented parasitemia levels similar to those of non-pregnant infected mice (NP/7dI) (Figure 1B). Since anemia, hypoglycemia and lower serum levels of TGF-β are common alterations during malaria [reviewed by (34–36)] and have been observed in pregnant women with malaria (1, 37–40), these parameters were measured in the animals in this experimental model. We noted that mice infected during early pregnancy displayed lower levels of hemoglobin (p<0.0068), glyceramine (p<0.0101) and serum TGF-β (p<0.0011) than non-pregnant uninfected (NP/NI) and pregnant uninfected mice (8dP/NI) (Figures 1C–E).

We next evaluated the impact of infection on pregnancy outcome. It was shown that infection did not affect the number of embryos implanted in the uterine horns of each female (Figure 1F), even the parasites were inoculated before implantation, the latter estimated to occur 4–5 days after vaginal plug detection (41), which coincides with the rising of parasitemia levels on day 5 after infection (Figures 1A, B). Despite this observation, we detected a significantly reduced proportion of normal implantation sites (33%) in infected pregnant females when compared with non-infected pregnant females.
Plasmodium berghei ANKA infection causes poor pregnancy outcome during early pregnancy. (A) An experimental design for the study of malaria in early gestation using BALB/c mice infected with P. berghei ANKA was performed. (B) Parasitemia levels were estimated by flow cytometry (n = 5 mice/group). (C–E) Hemoglobin (NP/NI, n=7; 8dP/NI, n=11; 8dP/7dl, n=7), blood glucose (NP/NI, n=8; 8dP/NI, n=6; 8dP/7dl, n=6) and serum TGF-β levels were measured (NP/NI, n=4; 8dP/NI, n=6; 8dP/7dl, n=6). (F–H) The number of implantations sites per female was counted (8dP/NI, n=7; 8dP/7dl, n=10) (F) and classified as normal or abnormal (G) according to macroscopic aspect (H) (8dP/NI, n=7; 8dP/7dl, n=10). (I–N) Uterine tissue was histologically processed and tissue sections were stained by H&E. Decidual area was delimited (I) and implantation sites of uninfected (J) and infected mice (K) were evaluated. e, embryo; d, decidua; n, necrosis; h, hemorrhage. Most of the infected mice presented intrauterine hemorrhages, which are indicated by red arrows (L), decidual edema (M) and decidual hemorrhage, where infected red blood cells were frequently observed (N). uMCs were detected and quantified in Toluidine Blue stained implantation sites (8dP/NI, n=7; 8dP/7dl, n=8). (P) uNK cells were detected and quantified in DBA lectin-stained implantation sites (8dP/NI, n=6; 8dP/7dl, n=7). Results represent mean ± SEM. Data was analyzed by Two-way ANOVA followed by Bonferroni’s post-test (B), One-way ANOVA with Bonferroni’s post-hoc comparisons tests (C–E), unpaired t-test (F, I, O, P) or Mann-Whitney test (G). NP, non-pregnant; NI, non-infected; dP, days of pregnancy; dl, days of infection. *, p <0.05; **, p <0.01; ***, p <0.001; ****, p <0.0001.
counterparts (100%) (p=0.0278) in which no abnormal implantation sites were detected (Figures 1G, H). We then performed histological evaluations of the implantation sites and found that the infected females had implantation sites with a marked reduction in the size of the decidual tissue (p<0.0001) (Figures 11–K).

Decidual cell morphology on the mesometrial side of the uterus of pregnant uninfected females was consistent with a previous description (42), these cells being large, with oval nuclei, several punctate nucleoli and pale eosinophilic cytoplasm (Figure 1J). In contrast, we observed that mesometrial decidua in the uterus of pregnant infected females showed edema (Figure 1M), extensive hemorrhagic areas full of parasitized red blood cells (Figure 1N) and decidual cells with pyknotic nuclei (Figure 1K). At the same time, while uninfected females displayed normal embryonic development for the gestational age (Figure 1I), infected females presented necrosis of the entire decidua including the embryo (Figure 1K). Moreover, while uninfected females displayed normal embryonic development for the gestational age (Figure 1I), infected females presented necrosis of the entire decidua including the embryo (Figure 1K). At the same time, while uninfected females displayed normal embryonic development for the gestational age (Figure 1I), infected females presented necrosis of the entire decidua including the embryo (Figure 1K). Moreover, vaginal bleeding and intrauterine hemorrhage (Figures 1H, L) between implantation sites of infected females was often observed at the time of euthanasia.

Chymases secreted by uterine mast cells (uMCs) and uNK cells are essential for the vascular changes required to support pregnancy (43). It has been suggested that the effects induced by a deficiency in one of these cell types are counterbalanced by the other, promoting spiral artery remodeling and placentation (44). In order to verify whether infection with Plasmodium affects these cells in early pregnancy, we quantified these cell phenotypes using histochemical techniques. We observed that although uMC numbers were not affected (Figure 1O), uNK cells were significantly reduced in infected pregnant animals in relation to non-infected pregnant females (p=0.0303) (Figure 1P). Thus, Plasmodium infection leads to significant alterations in both mother and developing embryo, leading to pregnancy loss associated with a reduction in uNK cells numbers when mothers are infected in early pregnancy.

Plasmodium Infection Impairs the Pregnancy Outcome During Mid-Gestation Without Altering uNK Cell Numbers

In order to determine the impact of Plasmodium infection in mid-pregnancy, mice were infected on day 5 of pregnancy and euthanized 7 days later (Figure 2A). As in early pregnancy (Figure 1B), parasitemia levels were similar between pregnant and non-pregnant animals (Figure 2B). Additionally, infected pregnant mice displayed lower levels of hemoglobin (p=0.0233), blood glucose (p=0.0004) and serum TGF-β (p=0.0003) in relation to uninfected pregnant controls (Figures 2C–E). When evaluating the pregnancy outcome, it was observed that non-infected mice presented a small number of abnormal implantation sites, whereas on the other hand, infected mice exhibited twice as many abnormal implantation sites as uninfected females (p=0.0430) (Figures 2F, G). Additionally, it was observed that vascular spaces in the placenta labyrinth layer of infected mice were reduced due to the thickening and disorganization of the trophoblast barrier. Simultaneously, infected red blood cells were detected in maternal blood vessels of placentas (Figure 2G).

Crucial changes take place in uterine vasculature during mid-gestation. Artery remodeling is a process that renders the spiral arteries into low-resistance, high-capacity vessels, allowing the correct flow of blood to reach the developing fetus (45, 46). Meanwhile, each placental layer fulfills important functions for the maintenance of pregnancy, with the suggestion that inappropriate proportions of the placental layers and poor remodeling of the spiral arteries can lead to intrauterine growth restriction and even embryo loss (47). Although we did not detect significant changes in the size of the MLAp, decidua and trophoblast layers in response to infection (Figures 2H–J), poor remodeling of the spiral arteries in infected mice during mid-gestation was observed, as indicated by an increased wall-to-lumen ratio (p=0.0378) (Figure 2K), which means that the arteries had a thicker wall and smaller lumen. In accordance, the α-smooth muscle actin (α-SMA) staining was increased (p=0.0177) in deciduas of infected animals, revealing a higher proportion of smooth muscle cells in spiral arteries (Figure 2L), a well-known feature of unremodeled arteries. Unexpectedly, the numbers of uNK cells and uMCs, two cell populations associated with the promotion of SA remodeling, were unchanged after infection (Figures 2M, N). We speculate that even in the presence of a similar number of uNK and uMCs cells, they could be differentially expressing molecules that assist in arterial remodeling, as observed in previous studies (48, 49).

Plasmodium Infection in Late Gestation Impairs Fetal Growth

Most of the pregnancy-associated malaria complications in both human and mice have been reported to take place in late pregnancy (2, 50–52). To verify whether the maternal and fetal parameters were altered after infection in our experimental model in late pregnancy, mice were infected on 12dP and euthanized 7 days later on 19dP (Figure 3A). It was observed that parasitemia levels in pregnant infected were similar to those in infected non-pregnant mice (Figure 3B). On the other hand, infected pregnant mice exhibited lower hemoglobin (p=0.0181) and glycemia levels (p=0.0022) than uninfected pregnant females (Figures 3C, D), but unlike the previous gestational periods, no change in TGF-β serum levels was observed in response to infection (Figure 3E).

When examining the ratio of normal and abnormal implantation sites, no differences were found between infected and uninfected animals (Figures 3F, G). Nevertheless, infection caused a significant decrease in fetal weight (p<0.0001) (Figures 3H, I) and size (Table 1), without affecting placental weight (Figure 3J), the size of placental layers (Figures 3K–N) or the vascular space in the labyrinth layer of the placenta (Figure 3O).

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HO-1 is upregulated during malaria in early pregnancy inducing iron overload at implantation sites.

HO-1 activity has protective effects during pregnancy (11, 12, 25, 53). In parallel, Plasmodium infection is characterized by the release of large amounts of heme (54). However, there is a lack of studies about the role of HO-1 in the pregnancy outcome during malaria. Thus, we measured the gene and protein expression of this enzyme in the uterus/placenta of infected mice during early, mid and late gestation. In early pregnancy, HO-1 mRNA expression levels were increased in the uterine tissue of pregnant infected mice in relation to uninfected non-pregnant (p=0.0018) and uninfected pregnant mice (p=0.0129) (Figure 4A) and the quantification of HO-1+ cells by immunohistochemistry showed that infected pregnant mice had a

FIGURE 2 | Malaria interferes in the remodeling of spiral arteries without changes in uNK cells numbers in mid-pregnancy. (A) An experimental design for the study of malaria in mid-gestation using BALB/c mice infected with P. berghei ANKA was performed. (B) Parasitemia levels were estimated by flow cytometry (n = 6–7 mice/group). (C–E) Hemoglobin (12dP/NI, n=6; 12dP/7dI, n=5), blood glucose (12dP/NI, n=7; 12dP/7dI, n=5) and serum TGF-β levels were measured (12dP/NI, n=7; 12dP/7dI, n=6). The dashed lines indicate the mean results of non-pregnant uninfected mice for each parameter evaluated. (F, G) The implantations sites were classified as normal or abnormal (F) according to macroscopic aspect (12dP/NI, n=7; 12dP/7dI, n=6), histologically processed and stained by H&E (G), scale bar 50 µm. (H–J) Decidua (H), trophoblast (I) and M/LAp (J) areas were delimited and quantified (12dP/NI, n=6; 12dP/7dI, n=6). (K) Wall-to-lumen area ratio was evaluated in the decidua of implantation sites stained by H&E (12dP/NI, n=6; 12dP/7dI, n=5), scale bar 100 µm. (L) α-SMA staining was quantified in decidual tissue for assessment of spiral arteries remodeling (12dP/NI, n=6; 12dP/7dI, n=5), scale bar 100 µm. (M) uMCs were detected and quantified in Toluidine Blue-stained implantation sites (12dP/NI, n=5; 12dP/7dI, n=5), scale bar 50 µm. (N) uNK cells were detected and quantified in the decidua of DBA lectin-stained implantation sites (12dP/NI, n=6; 12dP/7dI, n=5), scale bar 50 µm. Results represent mean ± SEM. Data was analyzed by Two-way ANOVA followed by Bonferroni’s post-hoc comparisons tests (B) and unpaired t-test (F–J, H–N). NP, non-pregnant; NI, non-infected; dP, days of pregnancy; dl, days of infection. *: p <0.05; ***: p <0.001.
greater number of positive cells in relation to non-pregnant uninfected mice (p=0.0375) (**Figure 4B**). Moreover, the upregulation of HO-1 was associated with an increased iron deposition at the implantation sites of infected mice when compared with uninfected pregnant females (p=0.0137) (**Figures 4C, R, S**). HO-1 protein detection by immunohistochemistry revealed that staining was mainly found throughout the placenta (**Figures 4J, K, N, O**), in scattered cells in the decidua (**Figures 4D–G, I, M**), in stromal cells in the endometrium (**Figures 4H, I**) and in the chorionic plate (**Figures 4P, Q**).

Unlike in early pregnancy, during mid-pregnancy infection did not affect the gene (**Figure 4A**) or protein expression of HO-1. **![Diagram](image-url)**

**FIGURE 3** | Malaria impairs maternal and fetal outcomes in late pregnancy. (A) An experimental design for the study of malaria in late gestation using BALB/c mice infected with *P. berghei* ANKA was performed. (B) Parasitemia levels were estimated by flow cytometry (n = 5 mice/group). (C–E) Hemogoblin (19dP/NI, n=5; 19dP/7dI, n=6), blood glucose (19dP/NI, n=6; 19dP/7dI, n=11) and serum TGF-β levels (19dP/NI, n=5; 19dP/7dI, n=6) were measured. The dashed lines indicate the mean results of non-pregnant uninfected mice for each parameter evaluated. (F, G) The implantation sites were classified as normal or abnormal (F) according to macroscopic aspect (19dP/NI, n=7; 19dP/7dI, n=11) (G). (H–J) Fetal and placental weight were recorded. Each point represents the measurement of a placenta or fetus (19dP/NI, n=42 fetuses/placentas derived from 7 litters; 19dP/7dI, n=77 fetuses/placentas derived from 11 litters). (K–N) Placental sections were stained by H&E (K) for delimitation of decidua (L), junctional zone (M) and labyrinth (N) layers (19dP/NI, n=7; 19dP/7dI, n=9), scale bar 1 mm. (O) Vascular space area in labyrinth layer was estimated in placenta sections stained with H&E (19dP/NI, n=16 placenta derived from 7 litters; 19dP/7dI, n=29 placenta derived from 11 litters), scale bar 50 μm. Results represent mean ± SEM. Data was analyzed by Two-way ANOVA followed by Bonferroni’s post-hoc comparisons tests (B), unpaired t-test (C–F, J, M–N) and Mann-Whitney test (H, L, O). NP, non-pregnant; NI, non-infected; dP, days of pregnancy; dI, days of infection; *: p <0.05; ****: p <0.0001.
TABLE 1 | Fetal biometry measurements in fetuses of BALB/c mice infected or not with *P. berghei* ANKA.

| Parameter   | 19dP/NI   | 19dP/7dI | p-value |
|-------------|-----------|----------|---------|
| CRL (cm)    | 2.234 ± 0.0134 | 2.053 ± 0.0285 | <0.0001 |
| Minimum     | 1.943     | 1.488    |         |
| Maximum     | 2.481     | 2.437    |         |
| SCD (cm)    | 1.104 ± 0.0069 | 1.048 ± 0.0096 | <0.0001 |
| Minimum     | 0.950     | 0.885    |         |
| Maximum     | 1.201     | 1.196    |         |
| APD (cm)    | 0.927 ± 0.0128 | 0.876 ± 0.0112 | 0.0080  |
| Minimum     | 0.848     | 0.690    |         |
| Maximum     | 1.031     | 1.053    |         |
| BPD (cm)    | 0.760 ± 0.0136 | 0.733 ± 0.0091 | 0.1204  |
| Minimum     | 0.687     | 0.589    |         |
| Maximum     | 0.878     | 0.853    |         |

CRL, Crown-rump length, SOD, snout-occipital distance, APD, abdominal anteroposterior diameter, BPD, biparietal diameter, SEM, standard error of the mean. 19dP/NI group (n= 8 litters, 62 pups); 19dP/7dI group (n=5 litters, 46 pups). The differences between the groups were determined by unpaired t-test. p-values <0.05 were considered significant.

(FIGURES 4B, J, K), nor did it lead to iron accumulation at the implantation sites (FIGURES 4C, T, U). In late pregnancy, despite a significant increase in the mRNA expression of HO-1 in the infected group (p<0.0004) (FIGURE 4A), there was no increase in the number of HO-1+ cells or in iron deposition in the placentas (FIGURES 4B, C, V, W). Therefore, our results suggest that HO-1 plays a role in the pathogenesis of malaria during early pregnancy.

Inhibition of HO-1 Ameliorates the Deleterious Effects of Malaria During Early Pregnancy

Since HO-1 is upregulated at the implantation sites during malaria in early pregnancy, we hypothesized that the adverse effects observed in these animals were induced by the increased activity of the enzyme. To investigate this, mice were infected on the day of vaginal plug detection, and treated with an inhibitor of HO-1 (ZnPPIX) or vehicle on gestational days 5, 6 and 7, as expected, we found that ZnPPIX treatment prevented iron overload at implantation sites (p=0.0426) (FIGURE 5N).

Additional experiments were also conducted to modulate the expression of HO-1 in mid and late pregnancy. Interestingly, ZnPPIX treatment was able to prevent infection-induced fetal growth restriction in late pregnancy (Supplementary Figure S2, Table S1). Despite of this, the overall results indicated that the protective effect of HO-1 modulation decreased as pregnancy progressed and that the main adverse effects of infection, associated with the activity of the enzyme, were restricted to early pregnancy (Supplementary Figures S1–S3).

Malaria During Early Pregnancy Induces Oxidative Stress and Pro-Inflammatory Cytokine Production at the Implantation Sites, Both of Which Are Prevented by HO-1 Inhibition

We found that infection led to an increase in serum heme concentrations (p=0.0018) and that the levels of this molecule appeared to be modulated in mice treated with ZnPPIX (Figure 6A). Similarly, while the infected animals presented high serum levels of bilirubin (p=0.0283) indicating an increase...
in HO-1 activity, mice treated with ZnPPIX presented lower bilirubin levels ($p=0.0103$), confirming inhibition of the enzyme’s activity (Figure 6B). Additionally, there was iron overload ($p=0.0021$) (Figure 6C) and induction of lipid peroxidation ($p=0.0011$) (Figure 6D) at the implantation sites during infection ($p=0.0034$), both of which were prevented by HO-1 inhibition treatment ($p=0.0307$ and 0.0042, respectively).

On the other hand, infection caused an increase in serum heme concentration in late pregnancy, but without causing iron accumulation or lipid peroxidation in this period; and in mid-
FIGURE 5 | Inhibition of HO-1 ameliorates the deleterious effects of malaria during early pregnancy. (A) An experimental design for the pharmacological inhibition of HO-1 by ZnPPIX during malaria in early gestation was performed. (B) Parasitemia levels were estimated by flow cytometry. Each symbol represents the mean of all animals within the indicated group ± SEM. (C) Hemozoin was detected in decidual tissue using polarized light microscopy, scale bar 50 μm. (D–F) Hemoglobin, blood glucose and serum TGF-β levels were measured. (G, H) The number of implantation sites per female was counted (G) and classified as normal or abnormal according to macroscopic aspect (H) (n = 7 mice/group). (I–K) Uterine tissue was histologically processed and tissue sections were stained by H&E (I), scale bar 1 mm. Decidual area was delimited (J) and implantation sites alterations were evaluated (K), scale bar 50 μm. (L) uMCs were detected and quantified in Toluidine Blue stained implantation sites, scale bar 50 μm. (M) uNK cells were detected and quantified in DBA lectin-stained implantation sites, scale bar 50 μm. (N) Quantification of iron deposition and representative images of iron-stained tissue, scale bar 50 μm. Unless otherwise stated, each symbol represents the measure of one mouse and results represent mean ± SEM (n = 6-7 mice/group). The dashed lines indicate the mean results of pregnant uninfected and untreated mice for each parameter evaluated. Data was analyzed by Two-way ANOVA followed by Bonferroni’s post-hoc comparisons tests (B), unpaired t-test (D, E, G, J, L–N) or Mann-Whitney test (C, F, H). NP, non-pregnant; NI, non-infected; dP, days of pregnancy; dl, days of infection. *: p <0.05; **: p <0.01.
pregnancy there were no changes in heme or bilirubin serum levels or tissue iron accumulation or lipid peroxidation. These data suggest that the pathogenetic mechanisms of the disease in these periods are not dependent on HO-1 activity, in contrast to early pregnancy (Supplementary Figure S3).

Since ferroptosis is a cell death process that involves the accumulation of lipid peroxides induced by iron (55, 56), we investigated whether this type of cell death occurs at the implantation sites of animals infected by Plasmodium, through the evaluation of the mRNA expression of PTGS2, which is increased during ferroptosis (51) and GPX4 protein and mRNA expression, which has anti-ferroptotic activity (17, 57). PTGS2 mRNA expression was increased in implantation sites in infected mice in comparison to uninfected pregnant (p=0.0043) and uninfected non-pregnant mice (p=0.0010) (Figure 6E). Furthermore, western blot analysis revealed a significant reduction in the protein levels of this enzyme at implantation sites in infected mice (p=0.0136) (Figures 6F, G). Therefore, these data indicate that cell death by ferroptosis could be taking place at implantation sites in infected mice. Accordingly, mRNA analysis showed that the transcript levels of GPX4 were lower at implantation sites in pregnant infected mice in relation to uninfected pregnant mice (p=0.0444) and that GPX4 mRNA is upregulated at implantation sites in infected mice treated with ZnPPIX when compared to uninfected non-pregnant mice (0.0051) and infected pregnant mice treated with vehicle (p=0.0013) (Figure 6H), suggesting that upregulation of GPX4 occurs at the transcriptional level, preventing ferroptosis.
T helper 1 (Th1) cytokines are produced during *Plasmodium* infection to induce mechanisms to control the proliferation of the parasite, but when they are produced in an exacerbated manner, they induce tissue damage (58). In order to evaluate local cytokine production in infected mice we measured the levels of IL-6, IFN-γ, and TNF in tissue homogenates of implantation sites. The results demonstrated that infection was able to induce an increase in the tissue levels of IFN-γ (p<0.0036) (Figure 6I) and TNF (p<0.0258) (Figure 6K) in infected mice treated with vehicle in relation to pregnant non-infected and non-pregnant uninfected mice as well as the levels of IL-6 in relation to non-pregnant uninfected mice (p<0.0006) (Figure 6J). Despite not controlling the increase in IL-6 and IFN-γ, inhibition of HO-1 significantly reduced TNF levels when compared with pregnant infected mice treated with vehicle (p=0.0392) (Figure 6K), suggesting that ZnPPIX is able to control, at least in part, infection-induced inflammation at implantation sites. In fact, we observed a positive correlation between the levels of IL-6 and TNF (Figures 6M, N), but not IFN-γ (Figure 6L), and the levels of iron at the implantation sites. In mid and late gestation, inflammation was driven by an increase in IFN-γ, but not TNF or IL-6 levels (Supplementary Figure S3).

**DISCUSSION**

This study found that *P. berghei* ANKA infection affected embryo viability in early and mid-pregnancy, while in late pregnancy infection compromised the success of the pregnancy by reducing fetal weight, as previously observed in other studies using various models of murine malaria in pregnancy, and reviewed by Barateiro et al. (34). In addition, it was demonstrated that poor pregnancy outcome and inflammation caused by malaria during early pregnancy are associated with increased expression and activity of HO-1 at the implantation sites, which lead to iron overload, lipid peroxidation and ferroptosis. HO-1 inhibition prevented most of the alterations provoked by the infection leading to a reduction in parasitemia levels as well as hemozoin accumulation, TNF levels, iron deposition and lipid peroxidation at implantation sites.

HO-1 has been shown to be elevated in plasma samples from infected pregnant women (59), and its mRNA expression was increased in the liver of pregnant BALB/c mice infected with *P. berghei* ANKA and conceptuses from C57BL/6 mice infected with *P. chabaudi* AS (60, 61). Furthermore, a previous study found that conceptuses of Swiss mice infected with *P. chabaudi* AS displayed an increase in HO-1 mRNA that was correlated with hemozoin density in placental tissues, showing that higher tissue hemozoin levels are related to higher HO-1 expression (62), which is consistent with our results.

The antimalarial effect exerted by ZnPPIX observed in this study could be explained in two ways: the first, could be due to competitive inhibition of HO-1, preventing the formation of heme catabolism products, including iron, which are necessary for the growth of the parasite (63, 64); the second could be attributed to its ability to prevent the process of hemozoin formation, similarly to the mechanism of action of many quinolones, such as chloroquine (65). It is possible that these mechanisms are restricted to the initial pregnancy stages due to the higher rate of parasitemia observed in the latter period in relation to the previous ones, exhausting the possible mechanisms of antiparasitic action exerted by ZnPPIX.

The reduction in uNK cells numbers during early pregnancy was associated with greater loss of embryonic viability in this study. The role of uNK cells in gestational malaria has not been well elucidated, since there is a lack of studies on these cells in murine models of malaria in pregnancy, and previous studies of placentas from women with malaria have focused on detecting NK cells in peripheral blood and blood from placental intervillous spaces (66, 67) or even placental tissue samples, although without quantifying NK cells in the decidua (68), which is the main tissue in which these cells are found (69). At delivery, the NK cell numbers were similar in placental blood from mothers with placental *Plasmodium falciparum* infection (PM+) and ones without placental malaria (PM-); however, IFN-γ production by NK cells in placental blood was higher in PM- individuals than in PM+ mothers, suggesting a protective role of placental IFN-γ-producing NK cells (67). Another study demonstrated that more NK cells were seen in *P. falciparum*-infected placentas from the third trimester than uninfected placentas, suggesting that NK cells in placentas may participate in mechanisms involved in the poor birth outcome associated with parasite infection (62). These studies were conducted using samples collected at parturition, at which time the number of these cells had declined (70, 71). In the present study it was shown for the first time that *P. berghei* infection induced a decrease in uNK cells in early gestation that was accompanied by poor pregnancy outcome, suggesting a protective role of this cell phenotype.

In rodent models, reduced numbers of uNK cells have been associated with decidual abnormalities and poor spiral artery remodeling, compromising the nutrient supply to the developing fetus (43, 72). HO-1 is required to support placentation and fetal development, an effect that can be mimicked by CO supplementation. In addition, it was previously reported by Salifu et al. (59) that iron accumulation in trophoblast cells during experimental malaria was associated...
with fetal death. Accordingly, in the present investigation, an increase in serum heme, upregulation of HO-1 and iron overload at the feto–maternal interface were observed, but not in late gestation as observed by Penha-Gonçalves et al. (33). The different results observed in our experimental investigation could be related to the parasite load utilized and period of gestation and infection, since Penha-Gonçalves et al. (33) examined animals inoculated with a higher parasite load than that used in the present investigation and only in late gestation. Together, these results highlight the importance of studying iron status at the feto–maternal interface to prevent infection-induced adverse pregnancy outcomes.

Our data suggest that iron accumulation, consequent lipid peroxidation and the adverse outcomes caused by infection during early pregnancy are mediated by HO-1, since its inhibition led to an improvement in the gestational outcome associated with reduced iron levels, lipid peroxidation and TNF levels at the implantation sites in parallel with upregulation of GPX4 mRNA expression levels. One hypothesis is that once iron-induced oxidative stress is established, positive feedback occurs as HO-1 increases, which could also increase the accumulation of iron at the implantation sites. In this sense, Fisher et al. (74) demonstrated that iron overload causes increased expression of HO-1 in placental endothelial cells, and exposure to iron overload and inflammation have a synergistic effect on the induction of oxidative stress and cell death in a TNF-α-dependent manner. In fact, increased levels of circulating TNF have been associated with fetal loss in the first trimester of human pregnancy (75). TNF signaling induces the production of reactive oxygen species (ROS) capable of overwhelming endogenous cellular antioxidant mechanisms and decreasing GPX4 expression, leading to lipid peroxidation (76). Conversely, treatment with a GPX4 activator inhibited the TNF-induced activation of the NF-κB pathway promoting the resolution of inflammation and prevention of ferroptosis (77). Thus, in the present investigation embryonic loss could be due to a combination of iron accumulation and TNF production at implantation sites in infected mice during early gestation.

In the brain, aging was able to upregulate HO-1 expression, which in association with an inflammatory stimulus induced iron-mediated cell death and ferroptosis, while ZnPPIX treatment was able to decrease inflammatory markers (15). Moreover, HO-1 has been described as important for lipid peroxidation during ferroptosis. Treatment of fibrosarcoma cells with erastin, an ferroptosis inducer, increase mRNA and protein levels of HO-1 while treatment with ZnPPIX prevented erastin-induced cell death (78). Our results indicate that the adverse effects of *P. berghei* ANKA infection during early pregnancy are related to iron overload, lipid peroxidation and necrosis of the decidua provoked by the increased expression and activity of HO-1 at the implantation sites, suggesting ferroptotic cell death. Accordingly, higher levels of lipid peroxidation have been observed in placentas of BALB/c mice infected with *P. berghei* ANKA (79) and in serum samples of pregnant women with malaria when compared with non-infected pregnant women and infected non-pregnant women (80).

The reduction of lipid peroxides is essential for cell survival and, therefore, its iron-induced increase triggers cell death by ferroptosis. GPX4 activity protects cells against ferroptotic cell death and is important for normal mouse embryo development, since deficiency of this enzyme caused embryo death between 7.5- and 8.5-days post coitum (81, 82). We observed that at 8 days of pregnancy and 7 days of infection with *P. berghei* ANKA mice exhibited substantial embryo loss associated with high expression of HO-1 and hallmarks of ferroptotic cell death such as iron-induced lipid peroxidation, upregulation of *Ptgs2* and downregulation of *Gpx4*. On the other hand, HO-1 inhibition induced the opposite effects, improving pregnancy outcome and highlighting the importance of HO-1 regulation for prevention of ferroptosis at implantation sites and maintenance of pregnancy during malaria in early pregnancy.

In view of the well-known protective effects induced by HO-1 in different contexts and processes, as recently reviewed by Campbell et al. (83), including improvement in cerebral malaria (31) and infections by other pathogens during pregnancy (29, 30, 84), the finding that its increase could be related to tissue damage and, on the contrary, that its inhibition would be protective, was unexpected. However, it has been verified that the enzyme has cytoprotective activity when it increases less than five times the initial expression, while enhancement of HO-1 activity of more than 15 times is cytotoxic, inducing high levels of iron and lipid peroxidation. In this scenario, treatment with the HO-1 inhibitor tin mesoporphyrin (SnMP) decreased HO-1 activity to moderate levels (four to five times compared to the control), which decreased the levels of reactive iron and oxidative damage, highlighting that there is a beneficial HO-1 expression threshold related to the accumulation of reactive iron released in heme degradation and that this accumulation makes it unlikely that the excessive expression of HO-1 offers a protective response (85).

Our results showed that HO-1 expression and activity are enhanced in *Plasmodium* infection during early pregnancy and that ZnPPIX treatment seems to decrease the ability of HO-1 to exert its deleterious effect when in excessive amounts. On the other hand, HO-1 expression was unchanged and moderately increased in mid- and late-pregnancy periods, respectively, in agreement with the reduced protective effects of ZnPPIX as pregnancy progresses, which could mean that other mechanisms overlap in the pathogenesis of the disease in these periods, in addition to the effects of HO-1. The lack of conclusive changes in the expression of HO-1 in the mid and late pregnancy may be due to the strong predominance of the Th2 response profile in those periods, especially during mid-gestation (86). In contrast, early pregnancy is characterized by a Th1 response that favors embryo implantation, trophoblastic invasion, and vascular remodeling (86). Proinflammatory cytokines such as IL-1α and TNF are able to induce HO-1 expression (87). Interestingly, in the present work local levels of TNF were increased by infection in early pregnancy, time in which HO-1 overexpression and related pathological markers are observed, and decreased after HO-1 inhibitor treatment.
In summary, the data presented here suggest that infection with *P. berghei* ANKA causes hemolysis, releasing large amounts of heme that lead to upregulation of HO-1 at implantation sites in early pregnancy. This in turn leads to an increase in iron deposition in this tissue, causing oxidative stress via lipid peroxidation, and a decrease in *Gpx4* expression inducing ferroptosis, which together with the pro-inflammatory response mediated mainly by TNF lead to embryonic loss. Treatment with the enzyme inhibitor, ZnPPIX, modulates the enzyme’s activity and it can exert its beneficial effects, thus improving pregnancy outcome during infection (Figure 7). The results of the present investigation highlight the importance of the study of malaria in early gestation and open new perspectives in the therapeutic management of HO-1 in this context, especially in those women who receive iron supplementation.

**MATERIALS AND METHODS**

**Parasite Strain**
The *P. berghei* ANKA strain expressing green fluorescent protein (GFP) was kindly provided by Dr. Ricardo Gazzinelli (Fiocruz, Brazil) and used to infect mice in this work. Parasites were maintained as frozen stocks and defrosted to infect BALB/c mice through inoculation of infected red blood cells (iRBCs) by the intraperitoneal (i.p.) route. To infect experimental animals, tail blood was collected from a previously infected mouse containing mainly parasites in ring or trophozoite stages, adjusting the inoculum to 5×10⁴ iRBCs per animal, which were injected by the i.p. route.

**Mice and Ethical Approval**
Male and female BALB/c mice at 8- to 12-weeks-old were housed in specific-pathogen-free conditions and received water and food ad libitum. Animal experiments were approved by the Animal Experimental Ethics Committee (CEUA) of the Federal University of Uberlândia under protocol number 066/17 and performed according to institutional guidelines for animal ethics at the animal facilities of the Federal University of Uberlândia (Rede de Biotérios de Roedores - REBIR-UFU).

**HO-1 Inhibitor Treatment**
Zinc protoporphyrin IX (ZnPPIX) (Cayman Chemical, Ann Arbor, MI, USA) a known inhibitor of HO-1 activity, was dissolved in NaOH 0.1M, diluted in PBS, and the pH was adjusted to 7.0 using HCl 1M, to be then administered in doses of 10 mg/kg/day by subcutaneous injection.

*FIGURE 7* | ZnPPIX beneficial effects in malaria during early pregnancy. HO-1 inhibition may have several beneficial effects in malaria during early pregnancy. Infection causes an increase in heme levels and induction of HO-1 in implantation sites leading to iron overload, lipid peroxidation and inflammation that ends up in embryo loss during early pregnancy, being these alterations prevented in ZnPPIX-treated mice. Moreover, ZnPPIX treatment preserves the number of uNK cells, that decrease in infected pregnant mice.
Pregnancy Monitoring and Experimental Infection

BALB/c females were mated with males of the same lineage at a ratio of 2:1. Females were monitored daily and the day of vaginal plug detection was considered as the first day of pregnancy (dP). Pregnant mice were infected or not on gestational day (GD) 1, 5 and 12 and euthanized on GD 8, 12 and 19, respectively. Non-pregnant non-infected (NP/NI) and infected non-pregnant (NP/7dI) mice were used as controls when considered appropriate. All infected mice were euthanized after 7 days of infection (dI) (Figures 1A, 2A and 3A). For this purpose, mice were injected with the anesthetics Ketamine/Xylazine (Vetnil, Louveira, SP, Brazil) by the i.p. route, blood samples were collected from the retro-orbital plexus to obtain serum and from the tail for parasitemia and hemoglobin analysis, then euthanasia was performed by cervical dislocation.

The number of total, normal and abnormal implantation sites was documented. Implantation sites were considered abnormal when they presented a pale, hemorrhagic or necrotic appearance or a small size for gestational age, these features suggest that these implantation sites are being resorbed or have a high probability of undergoing embryonic resorption (Supplementary Figure S4). Additionally, placental and fetal weight were recorded when mice were euthanized on GD 19. Biometric measurements such as crown-rump length (CRL), snout-occipital distance (SOD), abdominal anteroposterior diameter (APD) and biparietal diameter (BPD) were also performed in fixed pups, as previously described (88, 89).

Parasitemia Analysis

tail blood samples were collected at 3, 5 and 7 days of infection, fixed in 4% formaldehyde in PBS and stored at 4°C until processing for detection of infected red blood cells, through GFP expression by flow cytometry in FACSflow cytometer (BD Biosciences, San Jose, CA, USA). A total of 50,000 gated events were acquired and recorded for analysis of each sample using FlowJo v10 software (BD Biosciences).

Biochemical Measurements

Serum glucose, serum bilirubin, blood hemoglobin and iron tissue levels were measured using commercial kits (Labtest Diagnostica S.A. Belo Horizonte, MG, Brazil) according to the instructions of the kit for each parameter.

Histological Processing

Whole implantation sites (8 and 12dP) and placentas (19dP) were fixed in 10% buffered formalin and embedded in paraffin. Tissue sections of 4 μm thick were made and stained with Hematoxylin and Eosin (H&E) for histological analysis. Additionally, Toluidine Blue staining was performed for uMC detection. These cells were quantified by examining 35 microscopic fields of the myometrium and endometrium layers of the uterus.

Images from tissues were obtained using an Aperio AT Turbo scanner (Leica Biosystems Inc., Buffalo Grove, IL, USA) that allowed the management of images of whole tissues for morphometric measurements of utero-placental layer areas or a Leica DM500 microscope connected to a Leica ICC50 digital camera (both Leica Microsystems, Wetzlar, Germany). These uterine and placental layers were delimited using QuPath software (90) (Supplementary Figure S4).

Hemozoin Area Measurement

Hemozoin was detected in H&E-stained sections of uterus/placenta via polarized light microscopy. For this, three microscopic fields of decidual tissue per implantation site were randomly acquired in a Nikon Eclipse Ti-5 microscope and the area of hemozoin was estimated using Image J Fiji software (91). The results are expressed as the percentage of hemozoin area in relation to the total area of the image.

Vascular Space Area Assessment

H&E-stained sections of placentas were used for estimation of vascular space areas as previously described (59). For this, five microscopic fields were randomly acquired from lateral and middle regions of the labyrinth layer of the placentas. These images were analyzed with Image J (NIH, Bethesda, MA, USA), in which an automated light analysis procedure was applied to ensure color and image quality standardization across sections. Then, a threshold color was used to cover the area corresponding to the vascular space lumen. The coverage of vascular space area was determined as the ratio between the number of pixels covered by the area defined by the threshold and, the total number of pixels in the image. Results were expressed as the mean of vascular space area of the placentas from each female.

uNK Cells Detection

uNK cells were stained according to a previously described method (92). Tissue sections of implantation sites of mice in early and mid-pregnancy were deparaffinized, re-hydrated and incubated with 3% hydrogen peroxide for 30 min. Sections were washed in PBS and then incubated with 1% bovine serum albumin (BSA) (Sigma-Aldrich, St. Louis, MO, USA) in PBS for 30 min, followed by overnight incubation at 4°C with biotinylated-‘Dolichos biflorus’ (DBA) lectin (Sigma-Aldrich) diluted 1:150 in PBS. After that, sections were washed with PBS and incubated with avidin-biotin-peroxidase complex (ABC) (ABC kit, PK-4000; Vector Laboratories, Inc., Burlingame, CA, USA) at 37°C for 30 minutes and revealed with 3,3’-diaminobenzidine (Sigma-Aldrich) diluted in PBS containing hydrogen peroxide. The sections were counterstained with Harris’ hematoxylin and mounted with Dammar resin. Regions of interest from mesometrial deciduala were delimited and uNK cells counted using Aperio Imagescope software (Leica Biosystems Inc.). Results were expressed as uNK cells/mm².

HO-1 and α-SMA Immunohistochemistry

Deparaffinized tissue sections were re-hydrated and submitted to antigenic retrieval in citrate buffer (pH 6.0) in a microwave oven for 7 min. Then slides were incubated with 3% hydrogen peroxide for 30 min. Sections were washed in PBS and nonspecific ligation sites were blocked using bovine serum...
albumin (BSA) 1% in PBS (α-SMA staining) or skimmed milk (Molico, Nestlé®, São Paulo, SP, Brazil) 2% in PBS (HO-1 staining) for 30 min, followed by overnight incubation at 4°C with mouse anti-α-actin antibody (1:500, Santa Cruz Biotechnology, Inc., Dallas, TX, USA) or rabbit anti-HO-1 antibody (1:300, Sigma-Aldrich) diluted in PBS. Subsequently, tissue sections were washed with PBS and incubated with goat anti-mouse (1:500, Sigma-Aldrich) or goat anti-rabbit (1:400, Sigma-Aldrich) antibodies at 37°C for 1 h and, then incubated with avidin-biotin-peroxidase complex (1:100; ABC kit, PK-4000, Vector Laboratories Inc.) at 37°C for 30 minutes and revealed with 3,3′-diaminobenzidine (Sigma-Aldrich) diluted in PBS containing hydrogen peroxide. The sections were counterstained with Harris’ hematoxylin and mounted with Dammar resin.

For α-SMA staining quantification, images from the decidua were extracted using QuPath software (90) and analyzed with Image J Fiji software using the Colour Deconvolution method. After applying the “H DAB” vector option, α-SMA area was determined in the “Color 2 window” and normalized by the area of decidua analyzed.

HO-1⁺ cell counting was done in QuPath software by selecting the analyze, cell analysis, and positive cell detection buttons. Then, to segment the cells in the image the following parameters were set: Detection image: hematoxylin OD; requested pixel size:0.5 µm; background radius: 8 µm; median filter radius: 0 µm; sigma: 1.5 µm; minimum cell area: 10 µm²; maximum cell area: 400 µm²; threshold: 0.1; maximum background intensity: 2; Split by shape: on; Exclude DAB: off; Cell expansion: 5 µm; Include cell nucleus: on; Smooth boundaries: on; Make measurements: on; Score compartment: Cell DAB OD mean; Threshold 1+: 0.2; single threshold: on. The number of HO-1⁺ cells was normalized according to the total area of tissue analyzed (Supplementary Figure S4).

**Protein Extraction and Expression Analysis by Western Blot**

Tissue samples of implantation sites of mice at 8dP were weighed and homogenized in RIPA buffer (Tris 0.05M; NaCl 0.150M; Triton X-100 0.2%; sodium deoxycholate 0.2%; SDS 0.1%) with 10 mM EDTA, containing protease inhibitor cocktail (complele™, Roche Diagnostics, Mannheim, Germany), 1 mM Na₂VO₄ and 25 mM NaF. The samples were homogenized in a polycrylamide gels (SDS-PAGE) at 12% and transferred to polyvinylidene fluoride (PVDF) membranes (Immobilon®-FL, Millipore Corporation, Billerica, USA). The membranes were subsequently blocked with skimmed milk (Itambé, Belo Horizonte, MG, Brazil) 5% in blotting buffer (Tris buffer/Tween 20 0.1%) for 1 h, incubated overnight with mouse anti-GPX4 antibody (1:1000, Santa Cruz Biotechnology, Inc.) or mouse anti-β-actin antibody (1:1000; Sigma-Aldrich), both diluted in blotting buffer containing 3% BSA and 0.02% sodium azide. After that, membranes were washed three times of 10 minutes each in blotting buffer and incubated with peroxidase-conjugated goat anti-mouse secondary antibody diluted in 2% skimmed milk (1: 3000; Sigma-Aldrich) for 2 h. Finally, membranes were washed and the reactions were developed using a chemiluminescent substrate (ECL) (SuperSignal® West Pico kit, Thermo Fisher Scientific, USA). Membranes photodocumentation was performed on transilluminator ChemiDoc™ MP Imaging System (Bio-Rad Laboratories, Hercules, CA, USA). Densitometric analysis was performed using Image Lab Software (Bio-Rad Laboratories). Data were expressed as relative expression of the interest protein (GPX4) in relation to endogenous protein control (β-actin).

**Cytokine Analysis**

Tissue samples were homogenized in 5 mM EDTA/PBS buffer containing protease inhibitor cocktail (complele™, Roche) using a Turrax T10 (IKA Works GmbH & Co. KG-Staufen–Germany) homogenizer (200 mg of tissue in 1 mL of buffer). Later, samples were centrifuged at 3000 × g for 10 min at 4°C. After that, supernatants were collected, aliquoted and stored at -80°C until use. IL-6, TNF and IFN-γ concentrations were measured utilizing a mouse Th1/Th2/Th17 CBA kit (BD Biosciences) following manufacturer’s instructions. Data were acquired on a FACSCanto II Flow Cytometer (BD Biosciences) and analyzed with FACSDiva software (BD Biosciences). Pierce™ BCA protein assay kit (Thermo Fisher Scientific) was employed to determine the total protein concentrations that were used for results normalization. The cytokine theoretical limits of detection (pg/mL) are: IL-6, 1.4; TNF, 0.9; IFN-γ, 0.5.

TGF-β levels were measured in serum samples using a sandwich ELISA kit following the manufacturer’s instructions (R&D Systems, Minneapolis, MN, USA). Detection limit for this assay was 15.6 pg/mL.

**Heme Quantification**

Heme measurements in tissue samples were performed as described by (94) using 1-Step™ Turbo TMB (Thermo Fisher Scientific), a mixture of TMB and a stabilized hydrogen peroxide, due to the pseudoperoxidase characteristics of heme. Standard hemin chloride (Sigma-Aldrich) concentrations were used to extrapolate the heme concentrations in experimental samples.

**Iron Deposition Assessment**

Iron was measured in supernatant of tissue homogenates using a commercial kit (Labtest Diagnostica S.A.) following the manufacturer’s instructions and normalized by the total protein concentration. Additionally, iron deposition was evaluated in whole tissue sections stained with Perls’ Prussian Blue. Images of whole tissue were extracted utilizing the QuPath software and blue staining area was quantified making use of the Color Threshold of Image J Fiji. Results were normalized according to the total section area.
**Lipid Peroxidation Test**

Lipid peroxidation was measured according to the thiobarbituric acid reactive substances (TBARS) method. Samples of tissue homogenates were incubated with a mixture of 10% trichloroacetic acid and 0.67% thiobarbituric acid for 2h at 40°C, after that n-butanol was added and mixed. Subsequently, the samples were centrifuged at 3000 ×g for 15 min, upper phase was collected and the fluorescence was measured at 515 nm (excitation) and 553 (emission) (Molecular Devices, Menlo Park, CA, USA). The percentage of lipid peroxidation was calculated as previously described (95).

**RNA Extraction and Quantitative PCR**

Frozen pieces of uterus/placenta were mechanically pulverized in liquid nitrogen. Total RNA was extracted using TRizol® (Ambion®, Life Technologies, Carlsbad, CA, USA) and quantified by measuring the absorbance at 260 nm on GeneQuant spectrophotometer (GE Healthcare). Samples were treated with DNase (Invitrogen, Thermo Fisher Scientific) and transcribed into complementary DNA (cDNA). Reactions were performed using 1 μg of RNA and reverse transcriptase ImProm-II™ (Promega, Madison, WI, USA) on Artik thermocycler (Thermo Scientific). The cDNA was amplified in an ABI PRISM-7500 sequence detection system (Applied Biosystems, USA) using GoTaq® qPCR Master Mix (Promega) and gene-specific primers for HO-1, PTGS2, GPX4 and GAPDH. Cycle threshold data were normalized to the expression of GAPDH and analyzed using 2−ΔΔCt method (96). The sequences for each forward (Fw) and reverse (Rv) primers used in this study were as follows: HO-1 Fw 5’-GCTGGTATGGCTTCTTGTTT-3’, Rv 5’-GTGGGCGATAGACTGGGTCTT-3’; GPX4 Fw 5’-GCAACCAAGTTGGGAGCGAGG-3’, Rv 5’-GAAGGCGTATGGCCATGGAGG-3’; PTGS2 Fw 5’-CTGCGGCTTTTC AAGGATGG-3’, Rv 5’-GGGGATACACCTCTCCACCA-3’; GAPDH Fw 5’-GGGATAACACTCTCACCACA-3’, Rv 5’-CAGTGTAAGCCAAGATGCC-3’.

**Statistical Analysis**

Statistical analysis and graphs were performed using GraphPad Prism version 7.0 (GraphPad Software version 7.0, San Diego, CA, USA). Statistical differences between groups were determined by One-way ANOVA and Two-way ANOVA followed by Bonferroni’s post hoc test, as well as unpaired t-test for normally distributed values. Non-normally distributed values were analyzed by Mann-Whitney test or Kruskal Wallis test followed by Dunn’s post hoc test. Values of p < 0.05 were considered statistically significant.

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**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Animal Experimental Ethics Committee (CEUA) of the Federal University of Uberlândia.

**AUTHOR CONTRIBUTIONS**

NS conceived the idea. YC performed all the experimental procedures, statistical analysis, figure design and wrote the manuscript. MA and YC performed western blot analysis. MP, EA, AD-R and MA provided technical help in the processing of samples for mRNA expression measurement. EA and MA contributed to the cytokine and biochemical measurements. AD-R provided assistance with animal handling and histological analysis. RF and FE contributed with animal handling and histological analysis. YC performed all the experimental procedures, statistical analysis, figure design and wrote the manuscript. All the authors critically reviewed the manuscript.

**FUNDING**

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2022.879158/full#supplementary-material

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