Transient receptor potential vanilloid 1 (TRPV1) is a Ca\(^{2+}\)-permeable channel expressed on neuronal and nonneuronal cells, known as an oxidative stress sensor. It plays a protective role in bacterial infection, and recent findings indicate that this receptor modulates monocyte populations in mice with malaria; however, its role in cerebral malaria progression and outcome is unclear. By using TRPV1 wild-type (WT) and knockout (KO) mice, the importance of TRPV1 to this cerebral syndrome was investigated. Infection with *Plasmodium berghei* ANKA decreased TRPV1 expression in the brain. Mice lacking TRPV1 were protected against *Plasmodium*-induced mortality and morbidity, a response that was associated with less cerebral swelling, modulation of the brain expression of endothelial tight-junction markers (junctional adhesion molecule A and claudin-5), increased oxidative stress (via inhibition of catalase activity and increased levels of H\(_2\)O\(_2\), nitrotyrosine, and carbonyl residues), and diminished production of cytokines. *Plasmodium* load was not significantly affected by TRPV1 ablation. Repeated subcutaneous administration of the selective TRPV1 antagonist SB366791 after malaria induction increased TRPV1 expression in the brain tissue and enhanced mouse survival. These data indicate that TRPV1 channels contribute to the development and outcome of cerebral malaria.

1. Introduction

Malaria is an infectious disease of great morbidity and mortality, which claimed the lives of more than 400 thousand people worldwide in 2015 [1]. Cerebral malaria is a clinical syndrome of the severe form of the disease and is characterized by neurological complications (coma and convulsions) associated with brain inflammation (for review, see [2]) which can be lethal or cause irreversible neurological and/or cognitive sequelae in surviving patients (for review, see [3]).

Several mechanisms were found to contribute to cerebral malaria including alterations in nitric oxide availability, unbalanced oxidative stress responses, changes in the pattern of expression of inflammatory molecules, vascular leakage, and blood brain barrier disruption, amongst others [4–10]. However, its treatment has proven to be difficult and of low efficacy depending on timing and parasite resistance [3, 11],...
with nearly 50% of the infected patients presenting this syndrome [3]. Importantly, 10-40% of the children with cerebral malaria die and a significant percentage develop sequelae [3, 12, 13]. In this context, the host response to infection plays a decisive role in the clinical evolution of malaria and therefore influences disease outcome.

The transient receptor potential vanilloid 1 (TRPV1) is a Ca²⁺-permeable channel expressed on neuronal and non-neuronal cells such as brain endothelial and immune cells [14–18], which plays a role in the inflammatory response of different pathologies (for review, see [16, 19]) and an emerging role in neuroinflammation (for review, see [20]). It was found that TRPV1 is protective against bacterial infection [21–24] and modulates the innate immune response to malaria [25]. These studies also indicate that TRPV1 is detrimental to macrophage/monocyte-mediated responses, including their ability to produce inflammatory mediators, especially those related to oxidative stress [23, 26–30], in addition to regulating body temperature [21, 23]. However, the relevance of TRPV1 to the brain inflammation and symptoms of cerebral malaria has never been investigated.

Here, we used TRPV1 wild-type (WT) and knockout (KO) mice to evaluate the role of TRPV1 in cerebral malaria. Disease progression and brain inflammation were assessed in mice infected with Plasmodium berghei ANKA. It was found that TRPV1 contributes to disease severity and mortality, by mediating brain inflammation.

2. Methods

2.1. Mice. Nonfasted male C57BL/6 wild-type (WT) and TRPV1 knockout (TRPV1KO) mice (2–3 months of age; 22–28 g) were used. Animals were obtained from the animal’s facility of the Faculdade de Medicina de Ribeirão Preto, Universidade de São Paulo (USP). Mice (n = 3–4/cage) were housed in a climatically controlled environment (room temperature of 22 ± 2°C) and humidity of around 60%, on a 12-12 h light/dark cycle (lights on at 07:00), with free access to water and food. All experiments were conducted in accordance with the Brazilian Society for Animal Welfare (SBCAL), following approval by the Ethics Committee of USP. Animals were randomly assigned into groups, and the experimenter was blinded towards the genetic background of animals during the experiment. In some cases, C57BL/6 mice received the selective TRPV1 antagonist SB366791 (0.5 mg/kg, twice a day; Sigma-Aldrich, Brazil) or vehicle (10% DMSO in saline) for up to 14 days, starting at 24 h postmalaria induction. All assays were conducted in a blinded manner.

A total of 16 infected TRPV1 WT and 14 KO mice were used for analysis of survival rates, disease stage, and severity score; these data were obtained from two independent experiments. For performing the different biochemical, qPCR, and cytokine measurement experiments, samples were collected from 12 TRPV1 WT (5 noninfected and 7 infected) and 16 TRPV1 KO (5 noninfected and 11 infected) mice with stage III/IV malaria, in three independent experiments.

In two separate experiments, 20 WTs were used for assessment of mortality rates with the TRPV1 antagonist (10 vehicle-treated and 10 SB366791-treated). Disease stage and severity score experiments included 17 WT mice (8 vehicle-treated and 9 SB366791-treated), in two independent experiments. For experiments in animals with stage III/IV malaria, 11 TRPV1 WTs were used (5 vehicle-treated and 6 SB366791-treated) in two independent experiments.

2.2. Induction of Cerebral Malaria. Malaria was induced by a single intraperitoneal (i.p.) injection of 10⁷ red blood cells infected with P. berghei ANKA as previously described [31, 32]. Parasitaemia and disease progression were evaluated from day 1 postinfection, by daily recording of parasitaemia and clinical neurological signs of cerebral malaria.

2.3. Blood Parasitaemia. The percentage of parasitaemia was determined by flow cytometry. For this, a drop of blood from the tail was collected directly into 2 ml of PBS for flow cytometry analysis. Each sample was run on a FACSCalibur (Becton Dickinson, San Jose, CA, USA) flow cytometer with a 488 nm argon laser and BD CellQuest™ Pro software version 6.0.1 (Becton Dickinson, San Jose, CA, USA). Erythrocytes were identified on the basis of their specific forward (FSC) and side (SSC) light-scattering properties, and a total of 100,000 events were counted for each sample.

2.4. Analysis of the Clinical Neurological Signs of Cerebral Malaria and Mortality Rates. The neurological signs were evaluated as described by Linares et al. [33], in order to determine disease progression (stages I-IV) as follows: stage I—presence of parasitaemia and absence of neurological symptoms; stage II—presence of head deviation or hemiparesis, in the absence or presence of piloerection, altered gait or ambulation, muscle weakness, tremor, rollover response, and/or anaemia; stage III—presence of significant neurological symptoms, including head deviation, paraplegia/hemiparaplegia, immobility, muscle weakness, pilierection, anaemia, pelvic elevation, lack of responses to external stimuli, tremor, and swollen eyes; and stage IV—presence of exacerbated neurological symptoms in comparison to those observed at stage III.

Disease severity was analysed by using the Rapid Murine Coma and Behavior Scale as previously described by Carroll et al. [34], with minor modifications. Briefly, a score from 0 (normal) to 2 (severe alteration) was attributed to each one of the following parameters as follows: (i) coordination (gait and balance), (ii) exploratory behavior (motor performance), (iii) strength and tone (body position and limb strength), (iv) reflexes and self-preservation (touch escape, pinna reflex, toe pinch, and aggression), and (v) hygiene-related behavior (grooming). The summation of the scores attributed to each of the parameters for each animal was taken as severity score index, with the highest scores corresponding to the worst outcome of disease.

The animals were observed for up to 14 days postinfection and were culled by anaesthetic overdose (90 mg/kg ketamine + 2 mg/kg xylazine; i.p.) as soon as they reached stage III/IV (premortality end point). Blood samples were collected, and the plasma was obtained. Brain samples were also
collected and weighted. Collected plasma and brain tissue samples were immediately frozen and stored at -80°C until further processing for analysis of different parameters, except those used for qRT-PCR to which RNAlater was added according with the manufacturer’s instructions (Sigma-Aldrich, Brazil).

Also, body weight and temperature were registered before (baseline) and at stage III/IV postinfection. All those which did not reach stage III/IV during the observation period were culled, and their measurements and samples were collected at the 14th day postinfection. Noninfected mice were used as controls.

In a separate series of experiments, mortality rates were evaluated over 14 days following induction of cerebral malaria, in independent groups of mice.

2.5. Brain Parasite Load. Tissue parasite load was evaluated in brain samples (left hemisphere) collected from infected TRPV1 WT and KO mice, as previously described [33], and modified. Tissue parasite loads were determined by quantitative PCR and expressed as copy numbers of *P. berghei* ANKA 18S DNA per milligram of host tissue. For this, RNA was extracted in RNeasy Microarray Tissue Mini Kit, according to the manufacturer’s instructions (Qiagen, Brazil). Then, the cDNA was prepared by reverse transcription of 2 μg of RNA with ImProm-II Reverse Transcriptase (Promega, USA). The cDNA was assayed by qRT-PCR using the Taq-Man® system (Applied Biosystems, USA) with *P. berghei* probes (AI 38261, PN 4332079). GAPDH levels were assessed by TaqMan Mouse GAPDH System (TaqMan®, Applied Biosystems, USA) and were used as housekeeping gene controls.

2.6. TRPV1, Junctional Adhesion Molecule-A (JAM-A), and Claudin-5 Gene Expression by Real-Time qPCR. qRT-PCR was performed using GoTaq qPCR Master Mix (Promega, USA) and a Rotor-Gene 6000 real-time PCR machine (Corbett Life Science, Australia) in a final volume of 12 μl (hold: 2 min at 95°C; cycling: 40 cycles: 15 s at 95°C and 30 s at 60°C; melt: 68-90°C). The following primers were used: TRPV1 (forward 5′-GGTCAGCATCCACCTCACTGT-3′, reverse 5′-CTTGCGATGGCTGAAGTACA-3′; 94 bp; accession number NM_001001445.2), JAM-A (forward 5′-GCGACCATCCCTCAAGAGT-3′, reverse 5′-AGGTCTTGGCAGTGGCTGAAGTACA-3′; 109 bp; accession number NM_0001001445.2), JAM-A (forward 5′-GGTCAGCATCCCTCAAGAGT-3′, reverse 5′-AGGTCTTGGCAGTGGCTGAAGTACA-3′; 109 bp; accession number NM_0001001445.2), Claudin-5 (forward 5′-GCGACCATCCCTCAAGAGT-3′, reverse 5′-AGGTCTTGGCAGTGGCTGAAGTACA-3′; 109 bp; accession number NM_0001001445.2), and GAPDH primers, respectively.

2.7. Cytokine Measurements. Brain samples (right hemisphere) were prepared, and the supernatant was obtained as previously described [35] and used in the assays. The tissue and plasma levels of TNFα, IFNγ, and IL-6 were evaluated by using mouse cytometric bead array (CBA) cytokine kits according to the manufacturer’s instructions (BD Biosciences, Brazil). Data analysis was performed on a FACSCalibur flow cytometer (BD Biosciences Immunocytometry Systems). Results were calculated in FCAP Array Software version 3.0.1 (BD Biosciences, Brazil) and expressed as picograms of cytokine per mg of tissue protein (pg/mg of protein) or picograms per milliliter of plasma (pg/ml).

2.8. Tissue Sample Preparation for Biochemical Analysis of Oxidative Stress Pathways. Brain samples (100 mg; right hemisphere) were homogenized in 1000 μl of 0.05 M NaPO₄ (pH 7.4) containing ethylenediaminetetraacetic acid (EDTA, 1 mM) and centrifuged at 10,000 g, for 10 min, at 4°C, and then the supernatant was collected and stored at -80°C for analysis of enzyme activities.

2.9. Superoxide Dismutase (SOD). SOD activity was measured as described by Abreu et al. [36]. Briefly, 10 μl of each sample was incubated with 260 μl of sodium carbonate buffer (50 mM; pH 9.4 containing 3 mM EDTA), 10 μl of 3 mM xanthine, 10 μl of 153 μM/ml of 2,3-bis-(2-methoxy-4-nitro-5-sulfophenyl)-2H-tetrazolium-5-carboxanilide (XTT), and 10 μl of 1.87 μM/ml xanthine oxidase. Then, 200 μl of the mixture was added per well in 96-well plates and the absorbance was read at 470 nm for 20 min. Blank reactions were prepared for each sample by boiling them for 5 min in order to inactivate SOD. Results are expressed as milliunits (mU) of SOD/mg of protein. Enzyme activity was defined as the ability of one unit of SOD to dismutate 1 μmol of O₂⁻/min.

2.10. Catalase. Catalase activity was measured as previously described [36], by incubating 30 μl of brain homogenates or plasma samples with 500 μl of hydrogen peroxide (H₂O₂, 10 nmol/l) for 20 min, at 25°C. Reactions were stopped with 500 μl of sodium azide (1 mmol/l), and the concentration of the remaining H₂O₂ was determined by the oxidation of o-dianisidine. For this, 20 μl of each reaction was incubated with 200 μl of phosphate buffer (5 mM; pH 6.0) containing 0.167 mg/ml o-dianisidine and 0.095 mg/ml horseradish peroxidase (HRP). The absorbance was immediately read at 460 nm (SpectraMax Plus 384, Molecular Devices Inc., Sunnyvale, EUA) for 10 min. The remaining reactions were incubated at 60°C for 2 h, in order to inactivate catalase, and used as controls. A standard curve of H₂O₂ (11.3-8820 μM) was used for comparison. Results are expressed as international units (IU) of catalase per milligram (mg) of protein. One IU of catalase was defined as the amount of H₂O₂ (in μmol) degraded per minute.

2.11. Glutathione Peroxidase (GPx) and Reductase (GR). GR activity was assessed by measuring the consumption of nicotinamide adenine dinucleotide phosphate (NADPH) as a cofactor in the reduction of oxidized glutathione (GSSG) to reduced GSH [36]. For this, 10 μl of the sample was
incubated with 190 μl of a solution containing 2 mg/ml GSGG and 0.4 mg/ml NADPH, at 37°C. Absorbances were then recorded for 30 min (incubation period), at 340 nm. The results are expressed as μmol of NADPH per min normalized per mg of protein (μmol of NADPH/min/mg of protein).

GPx activity was determined as previously [36]. For this, 30 μl of sample per well (diluted 1:3) was incubated for 5 min at 37°C, with 145 μl per well of 0.05 M phosphate buffer (pH 7.4) containing 0.1 M EDTA, 5 μl of glutathione (GSH, 80 mM), and 5 μl glutathione reductase (0.0096 U/μl). After incubation, 5 μl of 0.46 % tert-butyl hydroperoxide solution and 10 μl of 1.2 mM NADPH were added to each well. Absorbances were monitored at 340 nm for 10 min. The results are expressed as μmol of GSH/min/mg of protein.

2.12. Thioredoxin Reductase (TrxR). TrxR activity was determined by incubating 20 μl of the sample with 140 μl of assay buffer (0.05 M phosphate buffer (pH 7.4) containing 0.1 M EDTA, 50 mM potassium chloride, and 0.2 mg/ml bovine serum albumin), 20 μl of 2 mM NADPH, and 20 μl of 5 mM 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB), in the presence and absence of a TrxR inhibitor (sodium aurothiomalate; 20 μM) [37]. Absorbances were read at 412 nm for 5 min. The results are expressed in IU of TrxR per mg of protein (IU/mg of protein). Enzyme activity was defined as the NADPH-dependent production of 2 μmol of 2-nitro-5-thiobenzoate per min at 22°C.

2.13. Protein Nitrotyrosine and Carbonyl Levels. For analysis of protein nitrotyrosine and carbonyl levels, 2.5 μg of each sample was assayed by slot blotting. The presence of proteins containing 3-nitrotyrosine residues was analysed in the samples as previously described [38]. After sample derivatization by addition of Laemmlí buffer (0.125 M Trizma, pH 6.8; 4% SDS and 20% glycerol; 20 min at room temperature and boiling for 2 min), the membrane was incubated with mouse monoclonal anti-nitrotyrosine primary antibody (1:2,000; Merck Millipore Co., Germany) overnight at 18°C.

Carbonylated proteins were determined according to the method described by Robinson et al. [39]. After the derivatization reaction by addition of 2,4-dinitrophenylhydrazine (DNPH) solution (0.1 mg/ml in 2N HCl, 5 min), the membrane was incubated with anti-DNPH primary antibody (1:25,000 in blocking buffer, Abcam, UK) overnight at 18°C.

Immunoactive bands were detected by chemiluminescence, and their intensities were estimated by densitometric analysis (ChemiDoc Image Systems, Bio-Rad, USA). Results were normalized by the band intensity values obtained after staining with Ponceau red.

2.14. Plasma and Tissue Hydrogen Peroxide Measurements. The levels of H₂O₂ were measured in brain homogenates and plasma samples by using a H₂O₂/peroxidase assay kit (Amplex Red H₂O₂/peroxidase assay kit; Molecular Probes, Invitrogen, Brazil) according to the manufacturer’s instructions. Results were obtained by comparison of each sample with a H₂O₂ (0–10 μM) standard curve and are expressed as H₂O₂ levels in μM (plasma) and in picomoles of H₂O₂ per mg of protein (brain samples).

2.15. Statistical Analysis. The results are presented as mean ± standard error (SE). The percentage of inhibition is reported as the mean for each individual experiment. For multiple statistical comparisons between groups, data were analysed by repeated-measures analysis of variance (ANOVA) or one-way ANOVA followed by the Bonferroni test with FDR correction. Paired and unpaired t tests were used when appropriate. Survival curves were analysed by the nonparametric Mantel-Cox test. All data were analysed in GraphPad Prism 5.0. \( p < 0.05 \) was considered significant. All n numbers are indicated on the graphs.

3. Results

3.1. P. berghei Infection Reduces TRPV1 mRNA Expression in the Mouse Brain, a Response Attenuated by TRPV1 Antagonism. We initially investigated whether infection with \( P. berghei \) ANKA, a plasmodium strain known to cause cerebral malaria in mice, influences TRPV1 expression in the mouse brain. Infected WT mice expressed lower TRPV1 mRNA levels (56%) in their brain tissue than non-infected controls did (Figure 1(a)). On the other hand, the systemic administration of SB366791 in C57BL/6 mice with malaria increased TRPV1 expression (2.1-fold increase) in comparison with vehicle controls (Supplementary Material Figure S1a).

3.2. Loss of TRPV1 Signaling Protects against Cerebral Malaria. We next assessed whether the ablation of TRPV1 influences cerebral malaria progression and mortality. Data depicted in Figure 1(b) show that infected TRPV1KO mice exhibit attenuated disease in comparison with WT controls. Of note, TRPV1KOs only presented parasitaemia without any other sign or symptom of cerebral malaria, suggesting they do not develop this syndrome. Accordingly, malaria was less severe and remained at stage I in these mice whilst it progressed into stages III and IV in the majority of the WT animals over the 14-day observation period (Figures 1(b) and 1(c)). Mortality was markedly prevented by TRPV1 ablation as 90% of the TRPV1KO mice survived infection in contrast with WT animals (19% survival; Figure 1(d)). Mice treated with SB366791 presented similar disease severity and course to those receiving vehicle until day 6 postinfection, improving their condition over the 14-day observation period (Supplementary Material Figures S1b and S1c). Twenty percent of those receiving the TRPV1 antagonist survived (Supplementary Material Figure S1d). As lack of TRPV1 was previously shown to exacerbate hypothermia in mice with bacterial infection [23], mouse body temperatures were registered. At baseline conditions, both genotypes exhibited similar body temperatures; however, hypothermia was only observed in stage III/IV WT but not TRPV1KO mice (Figure 1(e)). A similar response was registered in those receiving SB366791 (Supplementary Material Figure S1e).

Blood parasitaemia was similar in both genotypes, although WT mice exhibited higher parasitaemia than TRPV1KOs did at days 6 and 7 postinfection (Figure 2(a)). On the other hand, \( P. berghei \) ANKA 18S levels were elevated
Figure 1: Brain TRPV1 mRNA expression and cerebral malaria progression. (a) TRPV1 mRNA expression in brain samples of noninfected and infected (at stage III/IV) TRPV1 wild-type (WT) mice. Disease progression (b) and stage (c); survival rates (d) and body temperature (e) recordings from TRPV1 WT and knockout (KO) mice infected with P. berghei ANKA. Disease progression, stage, and survival rates were registered over 14 days postinfection. Mouse body temperatures were evaluated at baseline and postmalaria induction (at stage III/IV or at day 14 for those that survived the observation period). Results represent the mean ± SEM of all mice per group, obtained from two-three independent experiments. n is indicated on each graph. Data were analysed by repeated-measures analysis of variance (ANOVA) followed by the Bonferroni test with FDR correction (panels b and c). Paired and unpaired t tests were used when appropriate (panels a and e). Survival curves were analysed by the nonparametric Mantel-Cox test (panel d). *p < 0.05 differs from noninfected WTs or baseline readings; #p < 0.05 differs from infected WT mouse.
in the brain samples of TRPV1KO (2.8-fold) in comparison with those obtained from WT mice (Figure 2(b)).

3.3. Lack of TRPV1 Increases the mRNA Expression of Blood Brain Barrier Integrity Markers and Attenuates Oedema Formation in the Brains of Infected Mice. Loss of integrity of the blood brain barrier is a hallmark of cerebral malaria, contributing to increased oedema formation and neuronal damage as disease progresses [40, 41]. Possible effects of TRPV1 ablation in brain oedema formation and in the gene expression of the tight junction components claudin-5 and JAM-A [41, 42] were then, investigated. Data depicted in Figure 3(a) demonstrates that infection with \textit{P. berghei} ANKA promotes brain swelling in WT (1.7-fold) and TRPV1KO (1.2-fold) mice in comparison with their respective noninfected controls; however, this response was reduced by 25% in those lacking TRPV1. Additionally, analysis of claudin-5 and JAM-A mRNA levels revealed that infected WT mice express diminished levels of both genes (49% and 80%, respectively), in comparison with noninfected controls, a response that was attenuated in infected TRPV1KO mice (Figures 3(b) and 3(c)). Genotype did not affect brain weight/body weight ratios or claudin-5 mRNA expression in noninfected mice (Figures 3(a) and 3(b)). However, noninfected TRPV1KOs presented with lower expression of JAM-A (47%) in comparison with their WT counterparts (Figure 3(c)).

3.4. \( \text{H}_2\text{O}_2 \), Protein Nitrotyrosine and Carbonyl Residues Are Raised in Infected TRPV1KO Animals. Oxidative stress normally occurs as part of the host response to malaria [5, 43]. TRPV1 is an oxidative stress sensor [28], which not only does modulate oxidative stress [23, 27] but also can have its expression regulated by endogenous oxidant molecules [26]. Therefore, the impact of TRPV1 ablation in malaria-associated oxidative stress was investigated. Higher levels of \( \text{H}_2\text{O}_2 \) and protein nitrotyrosine residues (indicative of NO-dependent oxidative stress; [44]) were detected in infected mice of both genotypes in comparison with their noninfected controls (Figures 4(a) and 4(b)). WT mice presented 4.8-fold and 3.7-fold increases and TRPV1KOs 6.0-fold and 2.7-fold increases for tissue \( \text{H}_2\text{O}_2 \) and protein nitrotyrosine residue levels, respectively. Protein carbonyl residues (indicative of lipid peroxidation-dependent oxidative stress; [45]) were only increased (1.9-fold) in brain samples of infected mice lacking TRPV1 (Figure 4(c)). Analysis of plasma \( \text{H}_2\text{O}_2 \) levels, and protein nitrotyrosine and carbonyl levels indicated these were raised in TRPV1KO but not WT mice infected with \textit{P. berghei} ANKA (Figures 4(d)–4(f)). TRPV1KOs presented greater levels of plasma \( \text{H}_2\text{O}_2 \) (13.9-fold increase), protein nitrotyrosine (1.5-fold increase), and carbonyl (1.4-fold increase) residues in comparison with those observed for WT animals with cerebral malaria (Figures 4(d)–4(f)).

As TRPV1KO mice presented with an exacerbated production of oxidants, the activity of antioxidant enzymes was then, investigated. The tissue activity levels of SOD, GPx, and GR were attenuated (by \(~35\%\), \(~20\%\), and \(~34\%\) of reduction, respectively) in infected mice irrespective of genotype when compared to noninfected controls (Figures 5(a), 5(d), and 5(e)). Also, TrxR activity was enhanced in both infected genotypes (1.5-fold increase; Figure 5(f)). On the other hand, brain catalase activity was markedly diminished (49%) in infected TRPV1KO but not WT mice (Figure 5(b)). Infected TRPV1KO mice also displayed lower levels of catalase activity (70% less) in their plasma in comparison with WT controls (Figure 5(c)).

3.5. Diminished Cytokine Production Is Detected in Infected TRPV1KO Mice. Cytokines are involved in neuronal survival [46, 47] and therefore may affect cerebral malaria

\[ \text{H}_2\text{O}_2 \]

\[ \text{protein nitrotyrosine} \]

\[ \text{carbonyl residues} \]
progression. Thus, the levels of both tissue and plasma IFNγ, TNFα, and IL-6 were assessed in WT and TRPV1KO mice with malaria. Tissue and plasma TNFα production was markedly reduced (52% and 64%, respectively; Figures 6(a) and 6(d)) in TRPV1KO in comparison with WT controls. A similar profile was observed for IL-6 as mice lacking TRPV1 exhibited significant lower levels of this cytokine at both tissue (65% reduction) and plasma (86% reduction) levels (Figures 6(b) and 6(d)). Genotype did not affect IFNγ levels in a significant manner (Figures 6(c) and 6(f)).

4. Discussion

Since its discovery, the TRPV1 channel has been pointed out as an essential receptor in a variety of physiological and pathological responses. This is due to its wide expression and ability to transduce signals in both neuronal and nonneuronal cells, therefore participating in responses that range from cell differentiation to death [16, 23, 48–50]. Novel and recent findings on its role indicate that the endogenous activation of TRPV1 protects mammals from bacterial infections [21–24]. More recently, a nonselective TRPV1 antagonist (capsazepine) was found to modulate the peripheral immune response to malaria [25], but no studies have reported to date, a role for TRPV1 in cerebral malaria development and outcome. Here, we show for the first time that in the absence of TRPV1, P. berghei ANKA infection does not progress into cerebral malaria in the majority of the infected mice, protecting them from death and from the development of any disease symptoms and signals apart from blood parasitaemia. Protection was also observed in mice receiving the TRPV1 antagonist SB366791 repeatedly after malaria was induced. Of note, this effect was more pronounced in TRPV1KOs than in mice treated with SB366791. Although these results suggest that an intervention with a TRPV1 antagonist may be an alternative to avoid malaria progression, its use should be carefully considered as it may increase mortality upon bacterial infection [23]. Interestingly, although TRPV1 ablation exacerbates hypothermia in bacterial infection [21, 23], it was found herein that TRPV1KO mice and WTs treated with the selective TRPV1 antagonist SB366791 are protected from this condition in comparison with infected WTs.

P. berghei ANKA-infected mice treated with capsazepine were previously demonstrated to present similar blood parasitaemia to those treated with vehicle [25]. Here, we show...
that infected mice lacking TRPV1 present with similar blood parasitaemia to those expressing this receptor. On the other hand, at days 6 and 7 postinfection, infected WTs presented higher parasitaemia than TRPV1KOs did. Despite that, surviving TRPV1KO mice exhibited higher levels of Plasmodium 18S in their brain samples than WTs did with cerebral malaria at stage III/IV. Of note, the techniques used to measure blood parasitaemia and brain parasite load are different as peripheral parasitaemia comprises the detection of live parasites whilst brain 18S expression does not discriminate between live and dead plasmodium. However, it is possible that TRPV1KO mice are able to kill the parasites that reach the brain more efficiently than WTs are, therefore protecting those lacking TRPV1 from death.

Brain oedema formation in patients with cerebral malaria is indicative of a bad disease prognosis, especially in children [3, 51]. In adults, brain oedema is not as usual but affects 25% of these patients [52]. Brain swelling results from increased

**Figure 4:** Levels of H$_2$O$_2$, protein nitrotyrosine, and carbonyl residues. (a) H$_2$O$_2$ concentrations, protein (b) nitrotyrosine and (c) carbonyl residues in brain samples obtained from TRPV1 wild-type (WT) and knockout (KO) mice infected or not with *P. berghei* ANKA. (d) H$_2$O$_2$ concentrations, protein (e) nitrotyrosine, and (f) carbonyl residues in plasma samples obtained from TRPV1 wild-type (WT) and knockout (KO) mice infected or not with *P. berghei* ANKA. Samples were collected at stage III/IV or at day 14 for those that survived the observation period. Results represent the mean ± SEM of all mice per group, obtained from three independent experiments. $n$ is indicated on each graph. Data were analysed by one-way analysis of variance (ANOVA) followed by the Bonferroni test with FDR correction. *$p < 0.05$ differs from noninfected WTs; # $p < 0.05$ differs from infected WT mice.
vascular leakage and disruption of the blood brain barrier [8, 9]. TRPV1 activation promotes vasodilation and oedema formation [53, 54]. Then, the contribution of TRPV1 to brain oedema formation was assessed in infected mice. Infected WT mice exhibited brain swelling and decreased mRNA expression of the markers of blood brain barrier integrity JAM-A and claudin-5 [41, 42]. However, in the absence of TRPV1, there was higher JAM-A and claudin-5 mRNA expression. This response was associated with less brain oedema formation, suggesting that mice lacking TRPV1 are

![Graphs showing activity levels of antioxidant enzymes](image)

**Figure 5:** Activity levels of antioxidant enzymes. (a) Superoxide dismutase (SOD), (b) catalase, (d) glutathione peroxidase (GPx), (e) glutathione reductase (GR), and (f) thioredoxin reductase (TrxR) activity levels in brain samples obtained from TRPV1 wild-type (WT) and knockout (KO) mice infected or not with *P. berghei* ANKA. (c) Activity levels of catalase in plasma samples of infected TRPV1 WT and KO mice. Samples were collected at stage III/IV or at day 14 for those that survived the observation period. Results represent the mean ± SEM of all mice per group, obtained from three independent experiments. *n* is indicated on each graph. Data were analysed by one-way analysis of variance (ANOVA) followed by the Bonferroni test with FDR correction (panels a, b, d, e, and f). Unpaired *t* test was used when appropriate (panel c). *p < 0.05 differs from noninfected WTs; *#* *p < 0.05 differs from infected WT mice.
Figure 6: Brain and circulating levels of cytokines. Brain levels of (a) tumor necrosis α (TNFα), (b) interleukin-6 (IL-6), and (c) interferon γ (IFNγ) and plasma concentrations of (d) TNFα, (e) IL-6, and IFNγ (f) in TRPV1 wild-type (WT) and knockout (KO) mice infected with P. berghei ANKA. Samples were collected at stage III/IV or at day 14 for those that survived the observation period. Results represent the mean ± SEM of all mice per group, obtained from three independent experiments. n is indicated on each graph. Data were analysed by unpaired t test. * p < 0.05 differs from infected WT mice.
Intravascular oxidative stress is a common phenomenon in malaria which has been associated with alterations in the endothelium that in turn, facilitate the parasite accumulation into the brain tissue and/or vasculature. As a result of this close interaction between infected RBC, leukocytes, and the endothelium, oxidative stress products (H₂O₂, nitrosylated and carbonylated proteins) and cytokines (TNFα, IL-6, and IFNγ) are detected in the circulation and in the brain tissue; H₂O₂ levels are a lot higher in the brain tissue in comparison with the circulation. Plasma extravasation is increased in the brain and this is associated with reduced mRNA expression of the tight-junction endothelial markers claudin-5 and JAM-A. These alterations may culminate with neuronal death, thus, contributing to the increased morbidity and mortality observed in WT mice following infection with *Plasmodium berghei* ANKA. (b) Infected mice lacking TRPV1 (TRPV1KO) present increased levels of H₂O₂ and nitrosylated and carbonylated proteins than WT animals at both brain tissue and circulation. TRPV1KOs also exhibit lower concentrations of plasma and brain cytokines, especially TNFα and IL-6, and less plasma extravasation than WT mice, a response that is accompanied by higher expression of claudin-5 and JAM-A in their brain vasculature. The inflammatory response profile observed in TRPV1KO mice may reflect in less neuronal damage, as these animals are protected from *P. berghei* ANKA-induced death and symptoms.
outcome. Their clinical use may be limited as they worsen sepsis and may enhance their capacity of killing this parasite. Inflammation is involved in the tissue damage associated with cerebral malaria. Although not significant, a similar profile was observed for IFNy in the same mice. Interestingly, IFNy and TNFa have been linked to cerebral malaria progression by acting on brain endothelial cells, thus promoting their activation and/or apoptosis [67, 68]. Recently, the TRPV1 antagonist AMG9810 was found to confer neuroprotection by attenuating TNFa production in a rodent model of stroke [54]. These evidences and the gathered data allow us to suggest that the diminished cytokine generation by TRPV1KO mice contributes to the diminished brain swelling and damage observed in P. berghei ANKA-infected mice, a response that is associated with a greater ability of these mice to produce higher amounts of oxygen/nitrogen-derived oxidant species which in turn may enhance their capacity of killing this parasite.

Figure 7 summarizes the inflammatory events that occur in the brain of TRPV1 WT and KO mice during cerebral malaria. Overall, the data presented here, indicate that TRPV1 channels contribute to the development and outcome of cerebral malaria. Although antagonists targeting this receptor may be useful to preventing the development of the cerebral syndrome caused by Plasmodium sp., their clinical use may be limited as they worsen sepsis outcome.

Data Availability

The datasets used to support this study will be made available upon request. Requests should be sent to the corresponding author.

Conflicts of Interest

The authors declare no conflicts of interest.

Authors’ Contributions

D.M.S. Pereira, S.A. Teixeira, O. Murillo, E.P.M. Peixoto, M.C. Araújo, and N.C.F. Sousa performed the experiments and data analysis; C.R.F. Marinho, M.N. Muscará, and E.S. Fernandes designed the experiments and supervised the study; E.S. Fernandes secured the funding support and originally drafted the manuscript; V. Monteiro-Neto, J.B. Calixto, T.M. Cunha, C.R.F. Marinho, M.N. Muscará, and E.S. Fernandes critically revised the manuscript.

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Supplementary Materials

Figure S1: Effect of the selective TRPV1 antagonist SB366791 in the brain expression of TRPV1 mRNA and in cerebral malaria progression. (a) TRPV1 mRNA expression in brain samples of infected (at stage III/IV) TRPV1 wild-type (WT) mice. Disease progression (b) and stage (c); survival rates (d) and body temperature (e) recordings from TRPV1 WT mice infected with P. berghei ANKA. Disease progression, stage, and survival rates were registered over 14 days post-infection. Mouse body temperatures were evaluated at baseline and post-malaria induction (at stage III/IV or at day 14 for those that survived the observation period). Mice received the TRPV1 antagonist SB366791 (0.5 mg/kg, s.c., twice a day) or vehicle (10% DMSO in saline), from 24 h postinduction of malaria. Results represent the mean + SEM of all mice per group, obtained from three independent experiments. n is indicated on each graph. *p < 0.05 differs from baseline readings; #p < 0.05 differs from infected WT mice treated with vehicle. (Supplementary Materials)

References


[36] F. F. Abreu, A. C. A. Souza, S. A. Teixeira et al., "Elucidating the role of oxidative stress in the therapeutic effect of..."


