Human recombinant glutamate oxaloacetate transaminase 1 (GOT1) supplemented with oxaloacetate induces a protective effect after cerebral ischemia

M Pérez-Mato, P Ramos-Cabrera, T Sobrino, M Blanco, A Ruban, D Mirelman, P Menendez, J Castillo and F Campos

Blood glutamate scavenging is a novel and attractive protecting strategy to reduce the excitotoxic effect of extracellular glutamate released during ischemic brain injury. Glutamate oxaloacetate transaminase 1 (GOT1) activation by means of oxaloacetate administration has been used to reduce the glutamate concentration in the blood. However, the protective effect of the administration of the recombinant GOT1 (rGOT1) enzyme has not been yet addressed in cerebral ischemia. The aim of this study was to analyze the protective effect of an effective dose of oxaloacetate and the human rGOT1 alone and in combination with a non-effective dose of oxaloacetate in an animal model of ischemic stroke. Sixty rats were subjected to a transient middle cerebral artery occlusion (MCAO). Infarct volumes were assessed by magnetic resonance imaging (MRI) before treatment administration, and 24 h and 7 days after MCAO. Brain glutamate levels were determined by in vivo MR spectroscopy (MRS) during artery occlusion (80 min) and reperfusion (180 min). GOT activity and serum glutamate concentration were analyzed during the occlusion and reperfusion period. Somatosensory test was performed at baseline and 7 days after MCAO. The three treatments tested induced a reduction in serum and brain glutamate levels, resulting in a reduction in infarct volume and sensorimotor deficit. Protective effect of rGOT1 supplemented with oxaloacetate at 7 days persists even when treatment was delayed until at least 2 h after onset of ischemia. In conclusion, our findings indicate that the combination of human rGOT1 with low doses of oxaloacetate seems to be a successful approach for stroke treatment.

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Stroke is a leading cause of mortality and morbidity in developed countries, with increasing incidence because of the progressive aging of the population. Pharmacological or mechanical reperfusion therapy is the most effective treatment during the acute phase of ischemic stroke and it is associated with good outcome in 50–70% of cases. However, these treatments are only applicable to <10% of patients because of the short therapeutic window. Owing to these therapeutic limitations, the development of new and effective therapies to be used during the acute phase of ischemic stroke is in high demand.

After ischemic stroke, there is a rapid but transient elevation of glutamate into the extracellular space followed by a marked increase in intracellular calcium, which provokes a neuronal death through an excitotoxicity mechanism. Consequently, calcium and glutamate antagonists have been widely studied as protective agents in experimental models of cerebral ischemia with encouraging results. Unfortunately, they failed or displayed severe adverse effects when they were tested in clinical trials. Nevertheless, because of the central role of glutamate in the ischemic cascade, the mitigation of glutamate excitotoxicity remains one of the most promising strategies for the development of effective treatments to minimize neurological damage following an acute ischemic stroke. In this sense, novel therapeutic approaches should aim at reducing the increased glutamate levels produced early after the acute phase of ischemia.

Previous studies have shown that a decrease in blood glutamate concentration leads to a larger natural glutamate gradient between the brain and blood, facilitating the efflux of glutamate metabolites into the extracellular space.
excess extracellular brain glutamate into the blood. Reduction in glutamate concentration in blood is induced through the activity of the blood-resident enzyme glutamate oxaloacetate transaminase 1 (GOT1), which catalyzes the reversible transformation of oxaloacetate and glutamate to aspartate and α-ketoglutarate. Thus, the increase in the co-substrate (oxaloacetate) shifts the equilibrium of the reaction to the right side, thereby decreasing the blood glutamate concentration and lowering the brain glutamate levels (see Teichberg et al.4 and Campos et al.5 for review).

Based on this mechanism, we6 and others4 have demonstrated that the intravenous (i.v.) administration of oxaloacetate in a rat model of ischemia induced by transient occlusion of the middle cerebral artery occlusion (MCAO) decreases the glutamate concentration in blood and induces a lowering of infarct volume and edema after ischemia. These effects were also associated with a significant reduction in the sensorimotor deficit. To confirm that the protective effect was mediated by a decrease in the brain glutamate levels, we performed magnetic resonance spectroscopy (MRS) in the infarct region. Spectroscopic analysis revealed that the increase in brain glutamate seen in control animals after MCAO was significantly reduced in animals treated with oxaloacetate.4,6 These experimental results showed that systemic reduction in glutamate concentration induced by means of exogenous oxaloacetate administration could be used as a novel and efficient protective strategy in ischemic stroke. Importantly, however, the i.v. administration of humans with oxaloacetate has potential serious limitations as the required effective dosage of oxaloacetate in humans, corresponding to that used in rats, might be toxic.7 In a previous retrospective study with >400 human patients, we have found that ischemic patients with good outcome at 3 months after stroke, showed lower glutamate concentration and higher GOT activity (>18 U/l) in serum.8,9 Therefore, it is tentative to postulate that treatments based on the administration of recombinant GOT1 (rGOT1) alone or supplemented with low doses of oxaloacetate could improve the protective effects of oxaloacetate administered on its own.

The purpose of this study was to analyze the protective effect of oxaloacetate and human rGOT1 alone and in combination with a low dose (non-effective dose) of oxaloacetate in the rat model of ischemia induced by transient MCAO, and to compare their effects with a control group treated with saline.

Results

Animals included in the study. A total of 119 animals were used in this study (Table I). Thirty-six animals were used in the rGOT1 dose–response study, six animals per group (six groups of treatments). In this study, no animal deaths were observed within the first 24 h after treatments administration. In the protective study, 60 animals were included, six animals/group (four groups in the spectroscopic study and other six groups in the non-spectroscopic study). Eighteen animals were excluded, 16 of them were not included because of unsuccessful MCA occlusion or reperfusion and two animals died 24 h after surgery (one treated with saline and one treated with rGOT1). Finally, five animals were used as sham-operated control rats (sham) without MCAO.

Serum glutamate concentration is reduced by oxaloacetate and rGOT1 treatments in healthy animals. Administration of saline (control group) did not affect (P = 0.373) the basal concentration of glutamate during 24 h after the i.v. administration (Figure 1a), demonstrating that the surgical procedure did not interfere with the serum glutamate concentration. Injection of glutamate (15 mM) induced a significant (P < 0.05) increase in serum glutamate concentration 1 h after administration (Figure 1b). Basal concentration was normalized 3 h after treatment administration. This effect shows that the administration of glutamate in healthy animals could be used to simulate the increase in serum glutamate, a hallmark biochemical parameter after ischemia.

On the basis of previous reports where rGOT1 has been used to study other pathologies,6,10,11 we started with a dose of 6.44 μg per 100 g to analyze the capacity of the enzyme to metabolize serum glutamate. Administration of 6.44 μg per 100 g rGOT1 induced a reduction (P < 0.01) (Figure 1c) in serum glutamate concentration with respect to the maximum increase observed after glutamate injection. The maximum effect appeared between 2 and 4 h after administration of treatment. A higher dose of 12.88 μg per 100 g rGOT1 induced a robust reduction (P < 0.01) in serum glutamate concentration (Figure 1d). Similar to the previous dose, the maximum lowering effect appeared between 2 and 4 h after administration of treatment; however, in this case, the capacity to reduce serum glutamate concentration was much higher. Administration of rGOT1 25.76 μg per 100 g induced a significant (P = 0.007) reduction in serum glutamate concentration (Figure 1e). The maximum effect appeared between 2 and 4 h after administration of the treatment; however, the capacity of this dose to reduce serum glutamate concentration was similar to the 12.88 μg per 100 g dose. These findings showed that 12.88 μg per 100 g seems to be the most appropriate dose of rGOT1 to achieve a significant reduction in the serum glutamate concentration in the animals. Of note, serum glutamate concentrations were normalized 4–6 h after rGOT1 administration, demonstrating that the effect of the enzymatic treatment is transient.

In order to determine whether the effect of the enzyme could be potentiated by oxaloacetate (co-substrate of the enzymatic reaction), the dose of rGOT1 (12.88 μg per 100 g) was supplemented with a non-effective dose of oxaloacetate (1.5 mg per 100 g). In our previous study (already published6), we had described that a dose of 1.5 mg per 100 g of oxaloacetate did not affect the serum glutamate levels. Similar results were observed again in this study (Figure 1f). Administration of rGOT1 12.88 μg per 100 g with oxaloacetate (1.5 mg per 100 g) induced an immediate and significant (P < 0.01) lowering of serum glutamate concentration, which is the gold standard approach for reducing increased glutamate levels after brain ischemia. The decrease in glutamate...
appeared 30 min (with respect to the basal values) after treatment administration and remained during at least 3 h (Figure 1g); however, no significant differences with respect to rGOT1 12.88 μg pet 100 alone (Figure 1d) were observed.

These results led us to examine whether the protective efficacy of rGOT1 (12.88 μg per 100 g) or rGOT1 (12.88 μg per 100 g) supplemented with oxaloacetate (1.5 mg per 100 g) in ischemic animals was higher than that observed previously with oxaloacetate 3.5 mg per 100 g.
rGOT1 with or without supplementation of oxaloacetate induces a protective effect on ischemic animals. As it was previously shown,6 MCAO induced an increase of 30% of serum glutamate concentration 1 h after reperfusion (Figure 2), returning to normal levels 3 h later. In sham-operated rats in which MCA was not occluded, no changes in plasma glutamate concentration were detected (data not shown). The increase in serum glutamate concentration after cerebral ischemia was inhibited ($P<0.05$) by each one of the three treatments; rGOT1 (12.88 µg per 100 g), oxaloacetate (3.5 mg per 100 g) and rGOT1 plus oxaloacetate (12.88 µg per 100 g + 1.5 mg per 100 g, respectively) (Figure 2). The small differences observed between the treatments were not statistically significant.

To confirm that treatment with rGOT1 induced an increase in systemic GOT1 activity, their levels were measured at different time points (Figure 3). GOT1 activities in the control and oxaloacetate groups were not altered after treatment, whereas rGOT1 treatments induced a significant ($P<0.01$) increase in serum GOT activity 1 h after administration, which returned close to normal levels 24 h after treatment administration.

Quantitative analysis of MRS revealed a persistent increase in brain glutamate levels after the occlusion of MCA in the control group. Glutamate levels in the brain parenchyma were significantly decreased ($P<0.05$ relative to the control animals), with each of the three treatments tested (Figures 4a and b). To demonstrate that the reduction in serum and brain glutamate levels observed after treatment administration leads to a protective effect in our animal model of cerebral ischemia, infarct volumes were measured at 24 h and 7 days after ischemia and were compared with the control group. Figure 5a shows that, although all treatments were able to induce a significant ($P<0.05$) protection against the ischemic damage both 24 h and 7 days after the onset of ischemia, rGOT1 + oxaloacetate revealed to be the more effective treatment ($P<0.01$, with respect to the control group). Analysis of infarct volumes adjusted to the ipsilateral hemisphere showed a similar protective profile (data not shown). Infarct volumes measured in those animals subjected to the spectroscopy protocol (Figure 5b) showed the same protective results for the three treatments. Diffusion images revealed that all animals subjected to spectroscopy presented the same ischemic damage before the treatment administration.

To determine the therapeutic window of rGOT1 treatments, two groups of animals were treated 1 h after reperfusion (140 min after occlusion). Analysis of serum glutamate concentration in animals treated with rGOT1 (12.88 µg per 100 g) or rGOT1 plus oxaloacetate (12.88 µg per 100 g + 1.5 mg per 100 g, respectively) showed that both treatments caused a significant ($P<0.05$) inhibition of the increase in glutamate observed in the control group (Figure 6). Analysis of infarct volume determined that both treatments displayed similar infarct volume reduction at 24 h when administered immediately after reperfusion (Figures 7a and b), however, while the protective effect induced for rGOT1 disappears when this treatment is given 1 h after reperfusion on day 7; when rGOT1 is administered in combination with oxaloacetate, the effect persists beyond 1 h after reperfusion ($P<0.05$, with respect to the control).

Somatosensory test, an important end point assay of drug screening in stroke, confirmed that reduction in infarct volume observed with each of the three treatments was associated with a better neurological outcome measured 7 days after ischemia, showing the treatment rGOT1 + oxaloacetate to be a superior effect ($P<0.01$, with respect to the control) (Figure 8).

In vitro tests for neurotoxicity of oxaloacetate. To determine the potential toxicity of oxaloacetate treatment used in the study, a neuronal culture was exposed to a range of concentrations of oxaloacetate (100 nm to 1 mM) during 48 h. Our analysis showed that, under these experimental conditions, this compound did not affect the viability of the neurons (Supplementary Figure I).
Discussion

Nowadays, the concept of blood/brain glutamate scavenging is well recognized as a novel and attractive protective strategy to reduce the excitotoxic effect of excess extracellular glutamate that accumulates in the brain following an ischemic stroke. The efficacy of this strategy has been demonstrated in different types of ischemic animal models, and it has also been tested in other pathologies associated with an increase in brain glutamate levels, such as traumatic brain injury (TBI) or glioma with successful results. GOT activation by means of exogenous administration of oxaloacetate has been used as the most common approach to reduce the serum glutamate concentration. In this study, we found that the i.v. administration of rGOT1 was also able to induce a reduction in serum and brain glutamate levels, which also resulted in a significant reduction in the infarct volume and improvement in the sensorimotor deficit after ischemia. These effects were similar to those observed with the higher doses of oxaloacetate. Importantly, the protective effect of rGOT1 was more evident when it was administered in combination with a low concentration of oxaloacetate that had no effect by itself, indicating that the combined administration of rGOT1 and oxaloacetate could be an efficient serum glutamate-scavenging approach. The i.v. administration of an endogenous serum enzyme such as GOT1 as a novel protective treatment against ischemia is an interesting strategy because it is unlikely that the administration of rGOT1 can induce toxic effects in humans as the levels of this enzyme varies among healthy human subjects (7–45 U/l), and has been shown to increase >10-fold in patients with liver damage. On the other hand, the increase in the enzyme activity after the treatment is not much longer than 24 h, and serum glutamate concentrations come back to normal within the first 6 h, which reduces long-term potential adverse effects. However, the effective dose of GOT (12.88 μg per 100 g) used in the animals cannot be extrapolated to humans, as GOT levels in rats are almost 10-fold higher than those in humans (10 U/l in humans versus 95 U/l in rats). In this regard, further (pre)-clinical studies based on the administration of GOT should be conducted in humans to determine the effective therapeutic dose of GOT able to induce a significant reduction in serum glutamate in humans. In line with this, we have previously observed that stroke patients who had high GOT activity (average of 17 U/l) at admission had reduced serum glutamate concentrations and a significantly better neurological outcome than patients with lower average activity of GOT of (average of 17 U/l). Therefore, it can be hypothesized that the administration of rGOT1 doses able to increase the systemic GOT activity, at least two or three-fold, are necessary for clinical studies in stroke patients.

Regarding the use of oxaloacetate as a serum glutamate-scavenging treatment, such a treatment could present some limitations associated with the high dose likely to be necessary in patients to achieve the same effects than those observed in experimental animals. It should be noted that there are no consistent data in humans confirming this toxic effect, a study published in the 60’s, in which oxaloacetate was used to treat diabetic patients (from 200 to 1000 mg per day in three divided doses given orally) did not report any effect on liver function tests and blood acetone and cholesterol levels at the doses administered. Oral administration is not comparable to i.v. administration used in this study; nevertheless, this clinical study represents evidence for the relative safety of the administration of this molecule in humans. In this line, we also observed that oxaloacetate does not affect the viability of the neuronal culture, which is used as a current assay for neurotoxicity. The protective effects of oxaloacetate have also been discussed. As this molecule participates in the energy metabolism and provides an antioxidant protection of cells subjected to stress, such as hydrogen peroxide, it has been suggested that the protective role of oxaloacetate could be related to other mechanisms different from serum glutamate reduction. Studies in a TBI model showed that the protective effect of oxaloacetate was abolished in the presence of excess glutamate in serum or when it was administered in combination with maleate (a GOT inhibitor), which effectively demonstrates its serum glutamate-scavenging action.
Interestingly, the protective effect of rGOT1 (12.88 μg per 100 g) was potentiated when it was administered in combination with a non-effective low dose of oxaloacetate (1.5 mg per 100 g) but not with higher doses of the enzyme (25.76 μg per 100 g). These findings are in agreement with a previous glioblastoma study reporting that the combination of rGOT1 with low amounts of oxaloacetate together with temozolodine significantly extends the lifespan of tumor-bearing animals.12 The therapeutic synergistic effect observed between oxaloacetate and GOT can be attributed to the fact that the endogenous oxaloacetate concentration can become a limiting factor of the enzymatic reaction when GOT activity is increased after treatment. Therefore, attending to the rapid and maintained reduction in serum glutamate concentration, treatments based on the combined administration of rGOT1 supplemented with a low concentration of oxaloacetate could be optimal to reach the maximum protective effect in ischemia. Of note, the low dose of oxaloacetate is sufficient to increase the glutamate-scavenging activity of rGOT1, which reduces the potential complications associated with a high dose of this molecule.

It is crucial to develop treatment approaches capable of overcoming the narrow therapeutic window in stroke in order to mitigate the detrimental effects of the excess of brain glutamate after ischemia, which represents a stumbling-block from a clinical point of view. Owing to this limitation, only those treatments (mainly, NMDA antagonists) with protective effects beyond the first 1–2 h after the onset of ischemia have shown clinical interest.3 In our ischemic experimental model, we have observed that brain and serum glutamate levels appear to be increased in the first 2 h after reperfusion. We observed that lowering of glutamate by means of GOT treatments had protective effects even when the treatment was delayed until 1 h after reperfusion. Further experiments need to be carried out to determine the length of the therapeutic window; however, based on our results, it may be speculated that the therapeutic window for the administration of rGOT1 could be up to 2 h after the onset of ischemia. As this treatment does not require a prior computerized tomography scan, it could be given as early as possible, perhaps even as ambulatory treatment suggesting potential clinical application. In comparison with previous glutamate antagonists, another important advantage of
serum-scavenging-based treatments is that their effects are not mediated through the neuronal ionotropic glutamate receptors, thereby avoiding problems of poor blood–brain barrier permeability and potential neurotoxic effects.

In conclusion, in this study we show that oxaloacetate potentiates the protective effect of intravenously administered rGOT1 after ischemia. This effect is mediated through a reduction in serum and brain glutamate levels. The protective effect was observed even when the treatments were initiated as soon as 2 h after the onset of ischemia. The robust protection shown by the combination of rGOT1 and low doses of oxaloacetate proposes that this strategy may represent a valid approach for a successful acute-phase stroke treatment.

Materials and Methods

Animals. Experimental protocols were approved by the local Animal Care Committee according to the European Union (EU) rules (86/609/CEE, 2003/65/CE and 2010/63/EU). Male Sprague–Dawley rats (Harlan Laboratories, Barcelona, Spain) with a weight of 250–300 g were used. Rats were watered and fed ad libitum. Anesthesia was induced by inhalation of 5% sevoflurane in a nitrous oxide/oxygen mixture (70/30). Rectal temperature was maintained at 37±0.5°C using a feedback-controlled heating pad. Glucose levels were analyzed before surgery (ranging from 180 to 220 mg/dl).

Surgical procedures. Transient focal ischemia was induced in rats by using the transient MCAO following the surgical procedures previously described. In brief, under an operating microscope, the left common, external and internal carotid arteries were dissected from connective tissue through a midline neck incision. The left external carotid artery and pterygopalatine artery of the internal carotid artery were separated and ligated by 5–0 silk sutures. A 23-mm segment of 3–0 nylon monofilament suture with the tip rounded by heat was inserted into the stump of the left common carotid artery and advanced into the internal carotid artery to 20 mm from the bifurcation to occlude the origin of the MCA. The suture was removed after 80 min of occlusion. A laser-Doppler flow probe (tip diameter 1 mm) attached to a flowmeter (PeriFlux 5000; Perimed AB, Stockholm, Sweden) was located over the thinned skull in the MCA territory (4 mm lateral to bregma) to obtain a continuous measure of relative cerebral flow during the experiment. Only animals with a cerebral serum flow reduction of over 70% and with reperfusion after occlusion were included in the study. Experimental procedures were performed following five criteria derived from the Stroke Therapy Academic Industry Roundtable (STAIR) group guidelines for preclinical evaluation of stroke therapeutics:11,16 (1) cerebral serum flow was measured to confirm the vascular occlusion as an index of the reliability of the ischemic model; (2) animals were randomly assigned to treatment groups of the study; (3) researchers were blinded to treatment administration; (4) researchers were blinded to treatments during outcome assessment; and (5) temperature was controlled during the ischemic period.

Experimental protocol. In our previous study, we demonstrated that an i.v. dose of 3.5 mg per 100 g (weight of animal) of oxaloacetate induced a significant reduction in serum glutamate concentration, whereas a dose 1.5 mg per 100 g...
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In vivo MRS. MRS was acquired as previously described.17,18 Local shimming was performed by manual adjustment of first- and second-order shim coil currents using a proton-stimulated-echo acquisition mode (STEAM)-waterline sequence. The field homogeneity in a 3-mm³ voxel typically resulted in signal line widths of 10–20 Hz for the water signal. Water signal was suppressed by variable power RF pulses with optimized relaxation delays (VAPOR). In vivo 1 H magnetic resonance spectra of the rat brain were acquired by using a STEAM sequence with echo time = 3 ms, mixing time (TM) = 5 ms, repetition time = 2500 ms, 176 averages, cubic voxel = 3 × 3 × 3 mm³ (27 μl). Spectra were processed using MNOVA 7 (Mestrelab Research, Santiago de Compostela, Spain). For the quantitative analysis, glutamate signals were normalized to the creatine peak areas for each single spectrum (Supplementary Figure II). MRS was acquired during the occlusion (80 min) and 180 min after reperfusion.

Image analysis. All images were processed using ImageJ (Rasband WS, ImageJ, NIH, http://rsb.info.nih.gov/ij/). Infarct volumes were determined from (ADC maps) and T2-weighted images by manually selecting areas of reduced ADC values or hyperintense T2 signal by a researcher blinded to the animal protocols.

Sensorimotor test. Sensorimotor deficits after ischemic insult were assessed using the cylinder test as previously described.19 This test consists of evaluation of asymmetry of limbs during the exploratory activity. For this test, the animal was put in a cylinder of transparent base of 20 cm diameter. A video camera is located under this transparent cylinder for recording the vertical exploratory movement of the animal with anterior limbs during 2–10 min, depending on movement grade. For recording analysis, the VirtualDub software was used. Analyzed behaviors were as follows: number of times that the animal touches the cylinder wall and independent use of each limb in contact with the cylinder wall in each upward movement. Laterality index was calculated (the number of times that the animal touches the cylinder with the right leg during the ascendant movement by the number of times that the animals touch with each leg). This index is close to 0.5 for healthy animals and tends to be 0 or 1 for animals that have a preferential use of the left or the right paw, respectively. Somatosensory tests were performed 1 day before MCAO and on day 7 after ischemia during the darkness cycle.

Primary culture of rat cortical neurons. Primary cultures of cortical neurons were performed as described previously.20 Brains were removed from fetal Sprague–Dawley rats (Harlan Laboratories, Barcelona, Spain) on embryonic day 18, and the cortical area was dissected. Neurons were mechanically dissociated in incubation medium consisting of 80% Eagle’s minimum essential medium (MEM) containing 0.6% glucose, 0.029% glutamine, 16 mg/l gentamicin, 10% horse serum and 10% fetal calf serum. The dissociated neurons were plated at a density of 100,000 cells per well in poly-lysine-precoated 24-multwell plates. Medium was replaced twice weekly and studies were performed at in vitro on day 8, when the cultures consisted of ≥95% neurons.

Neuronal toxicity MMT assay. After 8 days in culture, neurons were exposed to different concentrations of oxaloacetate. Cell death was quantified with thiazoyl blue tetrazolium bromide (MTT) (Sigma, St. Louis, MO, USA) 48 h later. Neurons were incubated in 200 μg/ml MTT at 37 °C, and after 2 h, the culture medium was aspirated and cells were lysed in 250 μl DMSO. Color intensity was measured at 570 nm. Results were expressed as the percentage of absorbance of control wells. Vehicle treatment was used in control group and DMSO 2% treatment for control death.

Statistical analysis. Results were expressed as mean ± S.E.M. Statistical analyses were performed using one-way ANOVA followed by a Bonferroni post hoc analysis to determine between-group differences. A P-value < 0.05 was considered as statistically significant. The statistical analysis was conducted using PASW Statistics 18 for Mac (SPSS Inc., Chicago, CA, USA).

Conflict of Interest
The authors declare no conflict of interest.
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Author contributions

MPM, PRC, TM and JA conducted the experiments. MB, DM, JC and FC analyzed the data. MB, JC and FC designed the study. MB and PM helped in the interpretation of the data. All authors participated in the writing and editing of the manuscript. JC and FC were the principal investigators and supervised the project.

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