Mangiferin Prevents Guinea Pig Tracheal Contraction via Activation of the Nitric Oxide-Cyclic GMP Pathway

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Abstract
Previous studies have described the antispasmodic effect of mangiferin, a natural glucoside xanthone (2-C-β-Dglucopyranosyl-1,3,6,7-tetrahydroxoyxanthone) that is present in mango trees and other plants, but its mechanism of action remains unknown. The aim of this study was to examine the potential contribution of the nitric oxide-cyclic GMP pathway to the antispasmodic effect of mangiferin on isolated tracheal rings preparations. The functional effect of mangiferin on allergic and non-allergic contraction of guinea pig tracheal rings was assessed in conventional organ baths. Cultured tracheal rings were exposed to mangiferin or vehicle, and nitric oxide synthase (NOS) 3 and cyclic GMP (cGMP) levels were quantified using western blotting and enzyme immunoassays, respectively. Mangiferin (0.1–10 μM) inhibited tracheal contractions induced by distinct stimuli, such as allergen, histamine, 5-hydroxytryptamine or carbachol, in a concentration-dependent manner. Mangiferin also caused marked relaxation of tracheal rings that were precontracted by carbachol, suggesting that it has both anti-contraction and relaxant properties that are prevented by removing the epithelium. The effect of mangiferin was inhibited by the nitric oxide synthase inhibitor, N-nitro-L-arginine methyl ester (L-NAME) (100 μM), and the soluble guanylate cyclase inhibitor, 1H-[1,2,4]oxadiazolo[4,3-a]quinoxalin-1-one (ODQ) (10 μM), but not the adenylate cyclase inhibitor, 9-(tetrahydro-2-furyl)adenine (SQ22536) (100 μM). The antispasmodic effect of mangiferin was also sensitive to K⁺ channel blockers, such as tetraethylammonium (TEA), glibenclamide and apamin. Furthermore, mangiferin inhibited Ca²⁺-induced contractions in K⁺ (60 mM)-depolarised tracheal rings preparations. In addition, mangiferin increased NOS3 protein levels and cGMP intracellular levels in cultured tracheal rings. Finally, mangiferin-induced increase in cGMP levels was abrogated by co-incubation with either ODQ or L-NAME. These data suggest that the antispasmodic effect of mangiferin is mediated by epithelium-nitric oxide- and cGMP-dependent mechanisms.

Introduction

The xanthone mangiferin is an active phytochemical compound with therapeutic potential that is primarily found in mango tree leaves and stem bark (Mangifera indica) [1,2]. This substance usually occurs as a glucoside and is also found in a variety of other plant families, including Anemarrhena asphodeloides [3], Bersama abyssinica [4], Cyclopia genistoides [5], Cyclopia subternata [6], Gentiana lutea [7], Guida involucrata [8], Rhizophora belamcand [9] and Salacia oblonga [10], among others. Notably, M. indica and several of the above listed plants have been traditionally used to treat important human diseases, such as diabetes [10], obesity [10], cancer [11], and asthma [12].

An extract obtained via the decoction and drying of mango stem bark was developed at industrial scale in Cuba for use as a nutritional supplement and phytomedicine [13]. Vimang® is the brand name of this commercial preparation, and it contains a standardised mixture of terpenoids, steroids, fatty acids and polyphenols, including phenolic acids, phenolic esters and the predominant component mangiferin [13]. Similar to other polyphenol compounds, such as anthocyanins, curcumin and resveratrol, mangiferin has a broad spectrum of pharmacological effects. The most prominent and best-studied property of this class of phytochemicals is their antioxidant activity [1,14]. The ability to scavenge and decrease the formation of reactive oxygen species, as well as to activate enzymatic antioxidant systems, seems to be crucial for the outstanding antioxidant activity of mangiferin [1,14,15]. Apart from its capacity to interfere with oxidative stress, mangiferin exhibits a number of other properties, including immune-modulatory [16–18], anti-inflammatory [19–21] and anti-cancer [11,22,23] activities, suggesting that this substance could be used as a molecular template for innovative therapeutic applications.

The free radical nitric oxide is a neurotransmitter of the inhibitory nonadrenergic noncholinergic respiratory system [24,25]. It is produced by neural fibres that innervate airway smooth muscle cells, epithelial ciliated cells, type II alveolar cells and macrophages, and nitric oxide has been described as an effective antispasmodyc mediator in the airway [26]. The molecular mechanism underlying the antispasmodic effect of
nitric oxide is the direct activation of soluble guanylate cyclase and subsequent elevation of intracellular cGMP levels [27].

The aim of the present study was to assess the potential protective effect of mangiferin on the contractile response presented by the rat tracheal smooth muscle, following exposure to distinct pro-spasmodic agents, such as histamine, 5-hydroxytryptamine (5-HT), carbachol and allergen in vitro. All these spasmogens are supposed to play important role in the pathogenesis of airway obstruction noted in atopics asthma. Indeed, earlier investigations have shown that M. indica stem bark aqueous extract is an effective inhibitor of rat tracheal contraction caused by acetylcholine [28,29] and histamine [29]. However, exactly how the extract is acting to induce anti-contraction effects and whether or not this effect is accounted for by mangiferin has not been studied. Furthermore, our intention with this study was to test the hypothesis that mangiferin might be acting as an antispasmodic agent via activation of the nitric oxide-cGMP pathway. The results show that mangiferin can indeed inhibit smooth muscle spasms triggered by immunological and non-immunological stimuli. Such an effect is associated with nitric oxide production by epithelial cells, up-regulation of intracellular cGMP and the opening of K\textsuperscript{+}ATP and small-conductance Ca\textsuperscript{2+}-activated K\textsuperscript{+} channels in airway smooth muscle cells.

Methods

Ethics Statement
Experimental conditions and procedures involving animals were performed with direct approval of the Committee on Use of Laboratory Animals of the Oswaldo Cruz Foundation under license no. CEUA-FIOCRUZ 00085-01.

Animals
Male guinea pigs (300–400 g) were obtained from the Oswaldo Cruz Foundation breeding unit (Rio de Janeiro, Brazil). They were housed under conditions of constant temperature and controlled illumination, and food and water were available ad libitum.

Drugs and Chemical Reagents
Sodium chloride (NaCl), potassium chloride (KCl), potassium dihydrogen phosphate (KH\textsubscript{2}PO\textsubscript{4}), sodium hydrogen carbonate (NaHCO\textsubscript{3}), magnesium sulphate heptahydrate (MgSO\textsubscript{4} \cdot 7H\textsubscript{2}O), calcium chloride dihydrate (CaCl\textsubscript{2} \cdot 2H\textsubscript{2}O) and dimethyl sulfoxide (DMSO) were purchased from Merck (Darmstadt, Germany). Mangiferin, Glucose, EGTA, histamine, 5-hydroxytryptamine (5-HT), ovalbumin, carbachol, N-nitro-L-arginine methyl ester (L-NAME), tetraethylammonium (TEA), glibenclamide, apamin and SQ22536, an inhibitor of adenylate cyclase; 10\textsuperscript{-5} M L-NAME, an inhibitor of NOS; 100 \textmu M SQ22536, an inhibitor of adenylate cyclase; 10 \textmu M TEA, a nonselective K\textsuperscript{+}-dependent K\textsuperscript{+} channel blocker; 1 \textmu M glibenclamide, a K\textsuperscript{+}ATP channel blocker or 1 \textmu M apamin, Ca\textsuperscript{2+}-dependent K\textsuperscript{+} channel blocker of small conductance. All responses were expressed as the percentage of the maximal carbachol-induced contractile responses.

Protocols for Measurement of Tension Development

The experimental protocols were previously described [30]. Briefly, the tracheal rings were allowed to stabilise for 60 min, whereas the bathing solution was exchanged at 10 min intervals. At the end of the equilibration period, isolated tracheal rings, in absence or presence of epithelium, were contracted with carbachol (2.5 \textmu M), and once the contractions had reached a plateau, various concentrations of vehicle (DMSO) or mangiferin (0.1–1000 \textmu M) were added. All relaxations are expressed as the percentage of the maximal carbachol-induced contractile responses.

When we investigated the spasmytic effect of mangiferin on isolated tracheal rings. At the end of the equilibration period, the response to carbachol (2.5 \textmu M) was recorded. After carbachol was washed out and a stable baseline tone was re-established, the tissues were exposed to carbachol (0.01–100 \textmu M), histamine (0.1–1000 \textmu M), 5-HT (0.01–30 \textmu M), or antigen (ovalbumin; 0.001–100 \mu g/ml) in the presence or absence of mangiferin (0.1–10 \mu M). The preparations were pre-incubated with mangiferin for 15 min before the addition of each spasmogen. All responses were expressed as a percentage of the initial response to 2.5 \mu M carbachol. In some experiments, the epithelial cells were removed mechanically by rubbing the internal tracheal surface with a fine silver wire (200 \mu m in diameter), as described previously [31]. During the experiment, the contractile response to carbachol (0.01–100 \mu M) was measured before and after exposing intact or denuded epithelium tracheal rings to 10 \mu M mangiferin for 15 min.

To evaluate the putative interference of mangiferin with calcium influx, Ca\textsuperscript{2+}-concentration-response curves were established. Briefly, the responses of tracheal ring segments from naive guinea pig to 2.5 \mu M carbachol were recorded. After the carbachol was washed out and a stable baseline tone was re-established, the tissues were exposed to successive cycles of 60 mM KCl stimulations/washouts in Ca\textsuperscript{2+}-free Krebs' solution containing 2 mM EGTA until complete desensitisation to the 60 mM KCl-evoked contractile response was achieved. Next, the tracheal rings were immersed in Ca\textsuperscript{2+}-free Krebs' solution containing 60 mM KCl, and the extracellular Ca\textsuperscript{2+} concentration was increased stepwise by the cumulative addition of CaCl\textsubscript{2} (0.01–30 mM), in the presence or absence of mangiferin (0.1–10 \mu M) or vehicle (0.1% DMSO). All responses were expressed as a percentage of response to 2.5 \mu M carbachol.

To further investigate the mechanisms of action of mangiferin, the tracheal rings were pretreated 10 min before mangiferin application with 10 \mu M ODQ, an inhibitor of guanylate cyclase; 100 \mu M L-NAME, an inhibitor of NOS; 100 \mu M SQ22536, an inhibitor of adenylate cyclase; 10 \mu M TEA, a nonselective K\textsuperscript{+} channel blocker; 1 \mu M glibenclamide, a K\textsuperscript{+}ATP channel blocker or 1 \mu M apamin, Ca\textsuperscript{2+}-dependent K\textsuperscript{+} channel blocker of small conductance. All responses were expressed as a percentage of the response to 2.5 \mu M carbachol.

Western Blotting for NOS3
Three pools of three 3 rat tracheas were incubated with 10 \mu M mangiferin or vehicle (0.1% DMSO) for 15 min., and then
homogenized in ice cold lysis buffer containing the protease inhibitor cocktail Complete (F. Hoffmann-La Roche Ltd., Basel, Switzerland) and 0.1% Triton X-100 in PBS. The lysate was centrifuged at 13,000 g for 10 min at 4 °C. Supernatant was recovered and protein concentration was determined using the BCA assay (Sigma-Aldrich Corp., St Louis, USA). Equal amounts of sample protein (100 µg/lane) were separated by SDS-PAGE using polyacrylamide gels and proteins were transferred to nitrocellulose membranes (GE Healthcare, Little Chalfont, UK). Nonspecific binding was blocked with 5% (w/v) skimmed milk.
powder in TTBS for 1 h followed by incubation with rabbit polyclonal antibody NOS3 (1: 500; Santa Cruz Biotechnology, CA, USA) or mouse monoclonal antibody β-actin (1: 1000; Santa Cruz Biotechnology, CA, USA) overnight at 4°C. Membranes were incubated with HRP-conjugated secondary antibody (1: 10,000, R&D Systems, MN, USA) for 1 h at room temperature. The membranes were washed in TTBS and protein expression was detected using enhanced chemiluminescence (SuperSignal West Dura, Thermo Fisher Scientific Inc., Rockford, USA). Bands intensity was quantified by densitometry (Image-Pro Plus Media Cybernetics, Bethesda, MD).

Measurement of cGMP

Intracellular cGMP concentrations in guinea pig tracheal rings were assayed as described previously [30]. Isolated tracheas were cut into rings, quickly immersed in Krebs’ nutritional solution, and incubated with mangiferin (0.1–10 μM), L-NAME (100 μM) or ODQ (10 μM) in the presence of 100 μM 3-isobutyl-1-methylxanthine (IBMX) for 20 min. Some tracheal rings were pretreated with 100 μM L-NAME or 10 μM ODQ for 10 min before the addiction of 10 μM mangiferin. Tissue sections were rapidly frozen in liquid nitrogen, and the frozen tissue rings were homogenised in ice-cold 6% trichloroacetic acid (TCA). The homogenate was centrifuged at 2000×g for 15 min at 4°C. To remove TCA, the supernatants were washed 4 times with 5 volumes of water-saturated diethyl ether. The top ether layer was removed after TCA removal. The supernatants were lyophilised, and the cGMP of each sample was determined using commercially available enzyme immunoassay kits (GE Healthcare, Chalfont St. Giles, UK).

Statistical Analysis

The results were expressed as the mean ± S.E.M. EC50 values were calculated by fitting the log (agonist) vs. normalised response using GraphPad Software, and the results are displayed as the negative logarithm (pEC50). Significant differences were determined using one-way analysis of variance (ANOVA), followed by the Student-Newman-Keuls test. P values of 0.05 or less were considered to be statistically significant.

### Results

**Mangiferin Prevents Allergen-, Histamine-, 5-HT- and Carbachol-induced Tracheal Contraction**

It was previously reported that the aqueous extract of *M. indica*, of which mangiferin is the major active constituent [1], inhibited acetylcholine- or histamine-induced rat tracheal contractions [28,29]. To better characterise the putative antispasmodic property of mangiferin, we first investigated its ability to inhibit the anaphylactic contraction of tracheal rings obtained from ovalbumin-sensitised guinea pigs. The cumulative addition of ovalbumin (0.001–100 μg/ml) led to concentration-dependent contractile responses with a pEC50 of 6.08±0.20 and a maximal effect (Emax) of 97.1±8.5% (mean ± S.E.M) [n = 6]. Pretreatment with mangiferin (0.1–10 μM) for 15 min prior to the addition of cumulative concentrations of ovalbumin inhibited anaphylactic contractions in a concentration-dependent manner (Figure 1A). Furthermore, mangiferin (0.1 μM, 1 μM and 10 μM) affected both the potency and maximal response to ovalbumin, causing rightward shifts of 2-, 22- and 562-fold, respectively (Table 1). Next, tracheal rings from naive guinea pigs were used to demonstrate that the contraction elicited by cumulative concentrations of distinct spasmogens, such as histamine (Figures 1B), 5-HT (Figure 1C) or carbachol (Figure 1D), were also inhibited by mangiferin in a concentration-dependent manner. As shown in Table 1, 1 μM and 10 μM mangiferin caused rightward shifts of 3.8- and 67.6-fold for histamine, 12.3- and 51.3-fold for 5-HT and 1.4- and 4-fold for carbachol, respectively. At these concentrations, mangiferin also reduced the maximal response to histamine (from 111±4% (n = 7) to 90±2% (p<0.01, n = 6) and 65±4% (p<0.001, n = 5)), 5-HT (from 83±2% (n = 5) to 46±3% (p<0.01, n = 6) and 30±6% (p<0.001, n = 8)) and carbachol (from 137±2% (n = 6) to 111±4% (n = 7) and 79±5% (p<0.001, n = 7)).

**Epithelial Removal Impairs the Anti-Spasmodic Effect of Mangiferin**

Next, we examined whether the epithelium is involved in the effect of mangiferin by mechanically removing the epithelial cells from the internal tracheal surface with a fine silver wire, as previously reported [31]. Histological examination confirmed that the epithelial layer was removed (data not shown). The relaxant effect of mangiferin on carbachol pre-contracted tracheal segments (Figure 2A) was abolished following epithelial removal (Figure 2B).

Similarly, the ability of mangiferin to impair carbachol-induced

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**Table 1.** Potency (pEC50) and maximal response (Emax) values obtained from concentration-response curves of allergen (ovalbumin, 0.001–100 μg/ml), histamine (0.1–3000 μM), 5-HT (0.01–30 μM) or carbachol challenge (0.01–100 μM) in guinea pig tracheal rings, following co-incubation with mangiferin (0.1–10 μM) or vehicle (0.1% DMSO).

| Mangiferin (μM) | Ovalbumin | Histamine | 5-HT | Carbachol |
|----------------|-----------|-----------|------|-----------|
|                | pEC50     | Emax (%)  | pEC50 | Emax (%)  | pEC50 | Emax (%)  | pEC50 | Emax (%)  |
| 0 (vehicle)    | 6.08±0.20 | 97.1±8.5  | 5.53±0.12 | 111.3±4.3 | 6.15±0.14 | 82.9±2.1 | 7.12±0.06 | 137.1±2.1 |
| 0.1            | 5.78±0.13 | 75.1±6.2  | 5.11±0.10 | 101.9±2.6 | 5.93±0.04 | 74.8±2.5 | 7.13±0.09 | 129.7±2.5 |
| 1.0            | 4.73±0.44* | 51.6±7.6* | 4.95±0.10** | 89.7±2.4* | 5.06±0.17** | 46.4±2.8 | 6.97±0.06 | 110.9±3.7 |
| 10.0           | 3.33±0.09*** | 17.4±2.6*** | 3.70±0.34*** | 65.2±3.9*** | 4.44±0.19*** | 30.3±5.7 | 6.52±0.20* | 79.1±5.3*** |

Data represent the mean ± S.E.M. of 4 to 8 tracheal segments.

*p<0.05,

**p<0.01 and,

***p<0.001 compared with the tracheal responses of vehicle-treated tissues.

doi:10.1371/journal.pone.0071759.t001
Figure 2. Involvement of the epithelium in the protective effect of mangiferin. Tracheal relaxations induced by cumulative concentrations of mangiferin in guinea-pig trachea that were precontracted with 2.5 μM carbachol in the presence (A) or absence (B) of epithelium. The antispasmodic effect of 10 μM mangiferin on tracheal contractions induced by carbachol (0.01–100 μM) in the presence (C) or absence (D) of epithelium. Each point represents the mean ± S.E.M. of 5 to 6 segments. All results are expressed as a percentage of the contractile response induced by 2.5 μM carbachol. *p<0.05 compared with the tracheal responses of vehicle-treated tissues (open circles).

doi:10.1371/journal.pone.0071759.g002
Figure 3. Involvement of nitric oxide in the antispasmodic effect of mangiferin. Effects of mangiferin (10 μM) on carbachol-contracted guinea pig trachea, performed in absence or presence of L-NAME (A), ODQ (B), or SQ22536 (C). Each point represents the mean ± S.E.M. of 6 segments. All results are expressed as a percentage of the contractile response induced by 2.5 μM carbachol. *p<0.05 compared with tracheal responses of vehicle-treated tissues.
doi:10.1371/journal.pone.0071759.g003

Figure 4. Mangiferin-induced up-regulation of NOS3 in cultured tracheas. Expression of NOS3 in rat tracheas was determined by Western blot. a, c and e are lanes of trachea pools incubated with vehicle, while b, d and f are lanes of trachea pools incubated with mangiferin for 15 min. Data were normalized to β-actin and represented as the ratio between expression of NOS3:β-actin. Each bar represents the mean ± S.E.M. of 3 pools containing 3 tracheas each. * p<0.05 as compared with vehicle-treated groups.
doi:10.1371/journal.pone.0071759.g004
contraction (Figure 2C) was abolished after epithelial denudation (Figure 2D).

Inhibition of NOS or Guanylate Cyclase Prevents the Anti-Spasmodic Effect of Mangiferin

Tracheal rings were pretreated with the NOS inhibitor, L-NAME (100 μM), to examine the role of nitric oxide-mediated signalling in the antispasmodic effect of mangiferin. As shown in figure 3A, L-NAME prevented the relaxing effect of mangiferin after carbachol-induced tracheal contraction. This response was also abrogated by pre-incubation with the soluble guanylate cyclase inhibitor, ODQ (10 μM) (Figure 3B), whereas pretreatment with the adenylate cyclase inhibitor, SQ22536 (100 μM), did not alter the protective effect of mangiferin (Figure 3C).

Figure 5. Mangiferin increases cGMP levels in cultured tracheal rings. Effect of mangiferin on tracheal cGMP levels in the presence or absence of L-NAME or ODQ. Each bar represents the mean ± S.E.M. of 4 segments. *p<0.05 compared with vehicle-treated tissues. +p<0.05 compared with 10 μM mangiferin-treated tissues. doi:10.1371/journal.pone.0071759.g005

Figure 6. Involvement of K⁺ channels in the antispasmodic effect of mangiferin. Effects of mangiferin (10 μM) on carbachol-contracted guinea pig trachea, performed in the presence or absence of TEA (A), glibenclamide (B) or apamin (C). Each point represents the mean ± S.E.M. of 6 segments. All results are expressed as a percentage of the contractile response induced by 2.5 μM carbachol. *p<0.05 compared with vehicle-treated tissues. doi:10.1371/journal.pone.0071759.g006
The contractile response evoked by 2.5 mM calcium on tracheal preparations is expressed as a percentage of the 6 segments. The magnitude of the contractile tension induced by K^+ with vehicle-treated tissues.

**Mangiferin Increases cGMP Levels in Cultured Tracheal Rings**

To determine whether NOS3 isoform was up-regulated by mangiferin treatment, western blotting analyses were performed on tracheal tissue homogenates. We found that constitutive NOS3 expression was significantly increased in cultured tracheal rings exposed to 10 μM mangiferin, as compared to vehicle exposed rings (Figure 4).

**Mangiferin Increases cGMP Levels in Cultured Tracheal Rings**

Next, we demonstrated that mangiferin (0.1–10 μM) up-regulated the levels of cGMP in cultured guinea pig tracheal rings in a concentration-dependent manner (Figure 5). The ability of mangiferin to elevate cGMP levels was abolished by pre-incubation with either 100 μM L-NAME or 10 μM ODQ (Figure 5).

**K^+ Channel Blockers Reduce the Antispasmodic Effect of Mangiferin**

Prior studies revealed that the opening of K^+ channels leads to K^+ efflux, hyperpolarisation and relaxation of respiratory smooth muscle [32]. The nonselective K^+ channel blocker TEA (10 μM), the K^+ATP channel blocker glibenclamide (1 μM) and the small-conductance Ca^{2+}-activated K^+ channel blocker apamin (1 μM) were utilised to assess the potential role of K^+ channels in the antispasmodic activity of mangiferin. The ability of individual K^+ channel blockers to interfere with the protective effect of 10 μM mangiferin was evaluated in epithelia-preserved guinea pig tracheas that were stimulated with 2.5 μM carbachol. We found that a 15-min preincubation with TEA (Figure 6A), glibenclamide (Figure 6B) or apamin (Figure 6C), prior to the application of mangiferin, prevented the antispasmodic effect of this compound.

**Mangiferin Inhibits Ca^{2+}-Induced Contraction in K^+-Depolarised Trachea Rings**

Airway smooth muscle contraction is, in large part, regulated by intracellular Ca^{2+}. Thus, we wanted to assess the effect of mangiferin on extracellular Ca^{2+}-induced tracheal tension using the classical system of isolated organ bath preparations maintained in Ca^{2+}-free medium and depolarised with 60 mM KCl [33]. As expected, when the extracellular Ca^{2+} concentration was increased stepwise by the cumulative addition of CaCl_2 (0.01–30 mM), we observed a concentration-dependent elevation in the tracheal contractile response (Figure 7). As illustrated in Figure 7, preincubation for 15 min with mangiferin (0.1–10 μM) dramatically reduced Ca^{2+}-induced tracheal contraction.

**Discussion**

The current study shows that the xanthone glucoside mangiferin prevents the contraction of guinea pig tracheal rings induced by distinct spasmogens, including carbachol and allergen stimuli. This effect was abrogated by removal of the epithelium and exposure to the NOS inhibitor, L-NAME, or the guanylate cyclase inhibitor, ODQ. Mangiferin up-regulated NOS3 protein levels in the tracheal tissue. It also caused a dose-dependent increase in the intracellular cGMP levels of cultured tracheal rings via a mechanism that was also sensitive to L-NAME and ODQ treatments. These data suggest that mangiferin activates the nitric oxide-cGMP pathway to prevent airway smooth muscle contraction.

A previous study from Alvarez and collaborators [12] showed that asthma patients benefit from oral therapy with Vimang®, the brand name of an aqueous extract of *M. indica* stem bark that is traditionally used in the Caribbean region to treat respiratory disorders [1,34,33]. Accordingly, there are consistent reports demonstrating the efficacy of Vimang® and mangiferin, its prominent active ingredient, in preventing inflammatory changes in murine models of allergy and asthma [21,36]. Interestingly, the *M. indica* extract is also a potent inhibitor of histamine- and acetylcholine-induced rat tracheal contraction in *in vitro* settings [28,29]. Therefore, there is evidence that mangiferin has combined anti-inflammatory and airway-relaxing properties, which greatly increases the likelihood that this compound represents a potential therapeutic agent for the treatment of asthmatic conditions. To the best of our knowledge, this is the first study designed to assess the antispasmodic activity of mangiferin on the airway smooth muscle system and the first to indicate its molecular mechanism of action. Our findings reveal that mangiferin not only reduced the maximal tracheal contraction induced by distinct spasmogenic agents (allergen, histamine, carbachol and 5-HT) but also shifted the spasmogens-induced concentration-response curves to the right, clearly impacting the efficacy and potency of these agents. Moreover, it is clear that the effect of mangiferin on airway smooth muscle contractile responses is not specific for a certain type of receptor.
The observation that tracheal epithelial denudation abrogated the relaxant effect of mangiferin on carbachol pre-contracted or post-contracted segments is of great importance, and it indicates that the integrity of the airway epithelial layer is essential for the antispasmodic activity of mangiferin. Epithelial cells play a paramount role in the modulation of airway tone by working as a physical barrier that protects sensory nerves and smooth muscle cells from inhaled irritants [37]. In addition, the epithelial layer has the ability to release smooth muscle relaxant factors, such as prostaglandin (PG) E2 and nitric oxide, protecting the airway from excessive bronchoconstriction [37].

Nitric oxide activates guanylate cyclase (GC), which increases the level of intracellular cyclic GMP (cGMP). Increased cGMP then activates the protein kinase G (PKG) cascade, enhancing K+ efflux and attenuating Ca2+ influx-associated smooth muscle cell contractility. GTP (guanosine triphosphate); L-NAME (N-nitro-L-arginine methyl ester); ODQ (1H-[1,2,4]oxadiazolo[4,3-a]quinoxalin-1-one); TEA (tetraethylammonium).

do:10.1371/journal.pone.0071759.g008

**Figure 8. Proposed mechanism of action of mangiferin on guinea pig tracheal epithelium and smooth muscle cells.** Mangiferin activates nitric oxide synthase 3 (NOS3) isoform that up-regulates the production of nitric oxide (NO) in the epithelial cell. Nitric oxide activates guanylate cyclase (GC), which increases the level of intracellular cyclic GMP (cGMP). Increased cGMP then activates the protein kinase G (PKG) cascade, enhancing K+ efflux and attenuating Ca2+ influx-associated smooth muscle cell contractility. GTP (guanosine triphosphate); L-NAME (N-nitro-L-arginine methyl ester); ODQ (1H-[1,2,4]oxadiazolo[4,3-a]quinoxalin-1-one); TEA (tetraethylammonium).
instance, aminoglutethimide inhibition of NOS2 activity ameliorates cerebral vasospasm after subarachnoid haemorrhage in rabbits via increase of NOS1 mRNA and protein levels [42]. While exploring the fact that respiratory epithelial cells do express NOS3 [43], we demonstrated in the current study that the treatment with mangiferin effectively increased the protein levels of this constitutive form NOS in cultured epitheliun-intact tracheal tissues. These findings add support to the interpretation that mangiferin relaxant properties are closely related with activation of the nitric oxide-formation system.

The relationship between increased tissue levels of cGMP and tracheal smooth muscle relaxation in guinea pigs and other animal species has been reported [44–46]. Most of the nitric oxide- and cGMP-regulated signalling pathways responsible for airway smooth muscle relaxation is mediated by the opening of K+ channels, including small-conductance Ca2+-activated K+ and KATP channels. The activation of K+ channels causes potassium ion efflux, plasma membrane hyperpolarisation, increased closure of voltage-gated calcium channels and, eventually, a decrease in intracellular calcium levels in smooth muscle cells [32,47]. In the current study, we demonstrated that the anti-spasmodic property of mangiferin is lost in the presence of the non-selective K+ channel blocker TEA, suggesting the pivotal involvement of K+ channels in this response. It is also noteworthy that glibenclamide, a KATP channel blocker, and aminap, a blocker of small conductance Ca2+-activated K+ channels, significantly inhibited the anti-contraction effect of mangiferin. We chose these inhibitors because KATP channels and small conductance Ca2+-activated K+ channels have been strongly implicated in the airway smooth muscle relaxant response. These results suggest that both small conductance KATP and KATP channels play a role in the inhibitory effect of mangiferin on tracheal contraction.

There is a significant body of evidence for the existence of voltage-dependent Ca2+ channels in airway smooth muscle [48,49]. In the guinea pig airway, tracheal contraction evoked by KCl is induced by membrane depolarisation and the influx of Ca2+ through voltage-dependent Ca2+ channels [50]. Mangiferin also inhibited Ca2+-induced contractions in K+ depolarised preparations of epitheliun-intact tracheal rings, suggesting that mangiferin could inhibit Ca2+ influx by blocking voltage-dependent Ca2+ channels. Taken together, our results indicate that the effect of mangiferin on tracheal tissue is mediated by activation of the nitric oxide cGMP pathway, leading to enhanced K+ efflux with subsequent attenuation of Ca2+ influx-associated contractility in smooth muscle cells (Figure 8).

Conclusion

Our findings emphasise the ability of mangiferin to inhibit smooth muscle spasms caused by immunological (allergen) and non-immunological (histamine, carbachol and 5-HT) stimuli. These effects seem to be strongly associated with increased NOS3 protein levels and nitric oxide production by epithelial cells, up-regulation of intracellular cGMP and the opening of KATP and small-conductance Ca2+-activated K+ channels in smooth muscle cells. These changes block voltage-dependent Ca2+ channels, resulting in smooth muscle relaxation. Taken together, these findings demonstrate that mangiferin may be beneficial for the treatment of airflow limitation in human lung diseases.

Acknowledgments

We thank Dr. Patricia Jurgilas from the Oswaldo Cruz Foundation for technical assistance with cGMP measurement.

Author Contributions

Conceived and designed the experiments: MAM PMRS. Performed the experiments: ABV LPC DBRI VFC. Analyzed the data: MAM ABV LPC DBRI VFC. Wrote the paper: MAM ABV LPC DBRI.

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