Beneficial Effect of a Multistrain Synbiotic Prodefen® Plus on the Systemic and Vascular Alterations Associated with Metabolic Syndrome in Rats: The Role of the Neuronal Nitric Oxide Synthase and Protein Kinase A

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Abstract: A high fat diet (HFD) intake is crucial for the development and progression of metabolic syndrome (MtS). Increasing evidence links gut dysbiosis with the metabolic and vascular alterations associated with MtS. Here we studied the use of a combination of various probiotic strains together with a prebiotic (synbiotic) in a commercially available Prodefen® Plus. MtS was induced by HFD (45%) in male Wistar rats. Half of the MtS animals received Prodefen® Plus for 4 weeks. At 12 weeks, we observed an increase in body weight, together with the presence of insulin resistance, liver steatosis, hypertriglyceridemia and hypertension in MtS rats. Prodefen® Plus supplementation did not affect the body weight gain but ameliorated all the MtS-related symptoms. Moreover, the hypertension induced by HFD is caused by a diminished both nitric oxide (NO) functional role and release probably due to a diminished neuronal nitric oxide synthase (nNOS) activation by protein kinase A (PKA) pathway. Prodefen® Plus supplementation for 4 weeks recovered the NO function and release and the systolic blood pressure was returned to normotensive values as a result. Overall, supplementation with Prodefen® Plus could be considered an interesting non-pharmacological approach in MtS.

Keywords: metabolic syndrome; synbiotic; hypertension; superior mesenteric artery; perivascular nitrergic innervation; nitric oxide; neuronal nitric oxide synthase; protein kinase a

1. Introduction

Obesity affected two billion people worldwide in 2015 with annual costs reaching two trillion USD [1]. The global prevalence has increased dramatically in the last four decades, reaching 10.8% in
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adult men and 14.9% in adult women [2]. Chronic consumption of a high-fat diet (HFD) induces overweight and is the main trigger for the development of metabolic syndrome (MtS) characterized by dyslipidemia, hypertension and impaired glucose homeostasis [3,4]. MtS is associated with high mortality and morbidity due to an increase in the prevalence of cardiovascular diseases, diabetes, chronic kidney diseases, cancer and musculoskeletal diseases [2].

The accumulation of adipose tissue promotes a pro-inflammatory and pro-oxidative microenvironment, leading to a chronic low-grade inflammation state [5]. Moreover, the secretion of adipokines from the adipose tissue alters the function of multiple vasoactive factors, increases peripheral vascular resistance and leads to hypertension. Evidence indicates that the blood flow and tissue perfusion required for cardiovascular system homeostasis is affected by nitric oxide (NO) synthesis [6,7]. Other factor implicated in the regulation of vascular resistance is perivascular innervation, specifically in certain vessels such as the mesenteric vascular bed, where blood flow is approximately 20–30% of the total cardiac output [8]. Previously we have reported that the neural control of mesenteric vasomotor tone is altered in rats fed on HFD for a short term, partly due to a decrease in neuronal NO release from perivascular nitrergic innervation [9,10].

Increasing evidence links the development of MtS and chronic low-grade inflammation to dysbiosis, an imbalance in the intestinal flora [11–13]. Some studies demonstrated that dysbiosis promotes the development of low-grade inflammation and consequently MtS, and others that dysbiosis is the result of low-grade inflammation during obesity and MtS. Either way, the modulation of dysbiosis by dietary strategies based on alterations of gut microbiome improves the health outcome of MtS [14]. The supplementation of probiotics, live bacterial strains, have been shown to modulate the alteration of gut microbiome [15] and to improve MtS symptoms, such as insulin resistance, lipid profile and high blood pressure in animal models [16,17].

It is considered that multistrain and/or multi-species probiotics have been shown in animal models to be more effective than monostrain probiotics [18]. Additionally, the use of prebiotics, non-digestible oligosaccharides that stimulates the growth of desired bacteria, together with a probiotic is more effective than probiotics alone by improving survival and implantation of live microbes in the gastrointestinal tract [19]. The commercial symbiotic formulation Prodefen® Plus combines various probiotic strains (Lactobacillus rhamnosus, Lactobacillus casei, Lactobacillus acidophilus, Lactobacillus bulgaricus, Streptococcus thermophilus, Bifidobacterium breveis and Bifidobacterium infantis) together with a prebiotic, fructooligosaccharides. Both Prodefen® Plus, and a similar symbiotic commercial formula Prodefen® exert a beneficial effect in antibiotic-associated diarrhoea and acute gastroenteritis in children, respectively [20,21]. Regarding MtS, Prodefen® has shown an antihypertensive effect in spontaneously hypertensive rats [22]. Nevertheless, the possible beneficial effect of Prodefen® Plus in MtS remains unknown.

Given the above, we hypothesized that a supplementation with the commercially available symbiotic formula Prodefen® Plus might improve the cardiometabolic alterations related to MtS, leading to a reduction in the administration of any pharmacological treatment. Hence, in the present study we aim to assess the effect of Prodefen® Plus in rats fed on HFD that develop MtS, with a special focus on the understanding of the vascular mechanisms implicated in the development of hypertension.

2. Materials and Methods

2.1. Animals and Diet

All experimental procedures were approved by the Ethical Committee of the Universidad Autónoma de Madrid, and the Comunidad de Madrid (PROEX 322/16), are in compliance with NIH guidelines and follow the European Parliament Directive 2010/63/EU guidelines. Twenty-four male Wistar rats (initial weight 209.5 ± 3.1 g) were purchased from Harlan Ibérica SL, Barcelona, Spain and housed in the Animal Facility of the Universidad Autónoma de Madrid (Registration number EX-021U). Rats were held in groups of 2 in appropriate cages in controlled environmental conditions (20–
24 °C, 55% relative humidity and 12 h light-dark cycle). All rats had access to drinking water and specific rat chow ad libitum.

Animals were randomly divided into three groups: (I) Rats fed a commercial standard diet (4.0% fat; 2.9 Kcal/g; Tekland global rodent diet 2014, ENVIGO, Huntingdon, UK) for 12 weeks as a control group (CT; \( n = 8 \)); (II) rats with metabolic syndrome (MtS, \( n = 8 \)), fed a HFD (45% fat; 4.6 Kcal/g; 58V8, TestDiet, Columbia, USA) for 12 weeks; and (III) rats with MtS, fed a HFD (45% fat, 4.6 Kcal/g) for 12 weeks, supplemented with the multistrain symbiotic Prodefen® Plus (adjusted doses to \( 10^7 \) colony forming units (c.f.u.)/day) for the last 4 weeks (MtS-SYNB, \( n = 8 \)).

Prodefen® Plus (formulation which combines 990 mg of fructooligosaccharides with various probiotic strains −10^{10} c.f.u. Lactobacillus rhamnosus LGG and 10^9 c.f.u. of a mixture of: Lactobacillus casei, Streptococcus thermophilus, Bifidobacterium brevis, Lactobacillus acidophilus, Bifidobacterium infantis and Lactobacillus bulgaricus), was generously provided by Italfarmaco S.A. This symbiotic formula was administered to the rats dissolved in drinking water, once the obesity had been established [9,10]. The dose (\( 10^7 \) c.f.u./day) and administration time (4 weeks) of Prodefen® Plus was chosen based on preliminary pilot studies, choosing the lowest dose/time in which we found a systemic effect. The body weight was measured every two weeks. During the last 4 weeks, the water intake was measured every two days to assure the intake of appropriate dose of the symbiotic. The summary of the experimental procedure is represented in Figure 1.

**Figure 1.** Diagrammatic representation of the 12-weeks experimental procedure. Abbreviations: body weight (BW), glucose tolerance test (GTT), systolic blood pressure (SBP).

### 2.1.1. Glucose Tolerance Test

The oral glucose tolerance test (GTT) was performed at 0, 8 and 12 weeks according to a standard protocol [23,24]. After an overnight fasting, a single oral dose (2 g/kg of body weight) of glucose was delivered. Blood glucose was then measured from the tail vein just before, and at 15, 30, 60, 90 and 120 min after glucose intake, using test strips and reader (FreeStyle Optium®, Abbot Laboratories S.A., Madrid, Spain) (Figure 1).

### 2.1.2. Systolic Blood Pressure

The systolic blood pressure was measured at 0, 8 and 12 weeks in awake rats by a tail-cuff method, using a caudal artery plethysmograph (NIPREM 645, Cibertec S.A., Madrid, Spain) as published previously [9]. (Figure 1)

### 2.2. Animal Euthanasia, Sample Collection and Sample Analysis

After an overnight fasting, rats were anesthetized (100 mg/kg ketamine hydrochloride and 12 mg/kg Xylazine; i.m.) and euthanized by exsanguination by the infrahepatic inferior cava vein puncture. Visceral and epididymal white adipose pads were collected for posterior dissection. Left tibia length was measured as a parameter to evaluate body weight.
2.2.1. Liver Histology

Liver was extracted, rinsed in a saline solution and weighed. The left lateral liver lobe was cryoprotected (30% w/v sucrose in phosphate saline buffer) and frozen at −70 °C until use. Liver samples were embedded in optimum cutting temperature compound (OCT Tissue Tek, Sigma-Aldrich, Madrid, Spain). 5 μm cryostat sections were stained with lipid dye Sudan III and random images were taken at 20 × magnification. Quantification was performed using Image-Pro Plus 7 software (Media Cybernetics, Maryland, USA).

2.2.2. Serum Biochemical Parameters

Blood samples were kept at room temperature for 2 h and were centrifuged afterwards (2000×g 15 min, 4 °C). The serum was then collected and kept at −70 °C until use. Serum concentration of total cholesterol (TC), triglycerides (TG) and high density lipoproteins (HDL) were measured by biochemical analyser RAL LC5 (Barcelona, Spain). Low density lipoprotein (LDL) levels were indirectly calculated using the Friedewald formula: LDL = TC−((TG/5) + HDL) as published previously [25].

2.2.3. Vascular Reactivity

The superior mesenteric artery was carefully dissected, cleaned of connective tissue and maintained at 4 °C in Krebs-Henseleit solution (in mmol/L: 115 NaCl, 25 NaHCO3, 4.7 KCl, 1.2 MgSO4.7H2O, 2.5 CaCl2, 1.2 KH2PO4, 11.1 glucose, and 0.01 Na2EDTA) bubbled with a 95% O2/5% CO2 mixture until use. The segments were endothelium-denuded by gently rubbing the luminal surface of the segments with a thin wooden stick to eliminate the main source of vasoactive substances. Some segments were quickly frozen in liquid nitrogen and maintained at −70 °C.

Vascular reactivity experiments were performed in endothelium denuded segments of 2 mm length according to previously published method [26]. Briefly, two parallel stainless-steel pins were introduced through the lumen of the vascular segment: one was fixed to the bath wall and the other connected to a force transducer (Grass FTO3C; Quincy, MA, USA) that was connected to a model 7D Grass polygraph. For electric field stimulation (EFS) experiments, segments were mounted between two platinum electrodes 0.5 cm apart and connected to a stimulator (Grass, model S44) modified to supply the appropriate current strength. Segments were suspended in an organ bath containing 5 mL of KHS at 37 °C continuously bubbled with a 95% O2/5% CO2 mixture (pH 7.4). The segments were subjected to a tension of 4.9 mN, which was readjusted every 15 min during a 90 min equilibration period before drug administration. After this, the vessels were exposed to 75 mmol/L KCl to confirm their functional integrity. The absence of vascular endothelium after the washout period was tested by the inability of 10 μmol/L acetylcholine (ACh) to relax segments precontracted with noradrenaline. Endothelium removal did not alter the contractions elicited by KCl [10,23].

Frequency-response curves to EFS (1, 2, 4, 8 and 16 Hz) were performed. The parameters used for EFS were 200 mA, 0.3 ms, 1-16 Hz, for 30 s with an interval of 1 min between each stimulus, the time required to recover the basal tone. A washout period of at least 1 h was necessary to avoid desensitisation between consecutive curves. Two successive frequency-response curves separated by 1 h intervals produced similar contractile responses [26].

To analyse the participation of NO in the EFS-induced response in our experimental procedure, 0.1 mmol/L Nω-nitro-L-arginine methyl ester (L-NAME), a non-specific inhibitor of nitric oxide synthase, was added to the bath 30 min before performing the second frequency–response curve. The vasodilator response to the NO donor, diethylamine NONOate, (DEA-NO, 0.1 mmol/L–0.1 mmol/L was determined in noradrenaline-precontracted segments from all experimental groups.

2.2.4. Nitric Oxide Release

Nitric oxide (NO) release experiments were performed in endothelium-denuded mesenteric segments using the fluorescence probe 4,5-diaminofluorescein (DAF-2) as previously described [9,10,27]. Briefly, segments from all the experimental groups were subjected to an equilibration period
of 60 min in HEPES buffer (in mmol/L: NaCl 119; HEPES 20; CaCl₂ 1.2; KCl 4.6; MgSO₄ 1; KH₂PO₄ 0.4; NaHCO₃ 5; glucose 5.5; NaH₂PO₄ 0.15; pH 7.4) at 37 °C. Afterwards, arteries were transferred to a HEPES-filled 400 μL chamber and incubated with 2 μmol/L DAF-2 for 30 min. Then the medium was collected to measure basal NO release. Once the organ bath was refilled, cumulative EFS pulses of 30 s (1, 2, 4, 8 and 16 Hz) were applied at 1 min intervals. The fluorescence of the medium was measured at room temperature using a spectrofluorometer (Jenway 6280 Fluorimeter) with excitation wavelength set at 492 nm and emission wavelength at 515 nm. Some segments were incubated with 1 μmol/L H89 (a PKA (protein kinase A) inhibitor), 0.1 μmol/L calfostin C (a PKC (protein kinase C) inhibitor), or 10 μmol/L LY 294002 (a phosphatidylinositol 3-kinase -PI3K- inhibitor). The EFS-induced NO release was calculated by subtracting basal NO release from that evoked by EFS. Furthermore, blank samples were collected in the same way from segment-free medium in order to subtract background emission. The amount of NO released was expressed as arbitrary fluorescence units (A.F.U.)/mg tissue.

2.2.5. PKA and PKC Activity Assays

PKA and PKC activities were determined using a PKA and PKC kinase activity assay kit (Abcam, Cambridge, UK), respectively following the manufacturers’ protocols. Briefly, frozen arteries from all experimental groups were homogenised in a lysis buffer (0.01 mmol/L Tris-HCl, 1 mmol/L sodium vanadate, 1% SDS, pH 7.4), and centrifuged at 12,000 g for 10 min at 4 °C. The supernatant was then collected and used for the assay. Protein content was measured using a DC protein assay kit (BioRad, Madrid, Spain). Results were expressed as optical density (O.D.) units/μg protein.

2.2.6. Western Blot

Western blot analysis was performed as previously described [8,9]. Briefly, arteries were homogenized, and 30 μg protein was loaded in each lane. A mouse monoclonal antibody against neuronal nitric oxide synthase (nNOS, 1:2000, BD Biosciences), a rabbit polyclonal anti-pan-AKT antibody (1:1000, Abcam), or a rabbit polyclonal anti-pan-AKT (phospho T308, 1:500, Abcam) were used. The development and quantification of the images were performed using Quantity One software (Windows v4.6.6, Bio-Rad, Madrid, Spain). The same membrane was used to correct protein expression in each sample, by means of a monoclonal anti-β-actin–peroxidase antibody (1:50,000; Sigma-Aldrich, Madrid, Spain).

2.2.7. Detection of Superoxide Anions

Superoxide anion levels were measured using lucigenin chemiluminescence, as previously described [10,27]. Briefly, endothelium-denuded segments from all experimental groups were equilibrated for 30 min in HEPES buffer at 37 °C, transferred to test tubes that contained 1 mL HEPES buffer (pH 7.4) with lucigenin (5 μmol/L) and then kept at 37 °C. The luminometer (Optocom I, GEM Biomedical Inc., Hamden, CT, USA) was set to report arbitrary units of emitted light; repeated measurements were collected (10 s intervals for 5 min) and averaged. 4,5-Dihydroxy-1,3-benzene-disulphonic acid “Tiron” (10 mmol/L), a cell-permeant, non-enzymatic superoxide anion scavenger, was added to quench the superoxide anion-dependent chemiluminescence. Blank samples (HEPES without arterial segment) were collected in the same way from culture medium without mesenteric segments to subtract background emission.

2.3. Drugs Used

L-Noradrenaline hydrochloride, ACh chloride, diethylamine NONOate diethylammonium salt, Nω-nitro-L-arginine methyl ester (L-NAME) hydrochloride, lucigenin, tiron, DAF-2, calfostin C, H89, and LY294002 were used. All drugs were purchased from Sigma-Aldrich (Spain) except for LY294002 and H89, which were obtained from Tocris (Spain). Stock solutions (10 mmol/L) of drugs were made
in distilled water, except for NA, which was dissolved in a NaCl (0.9%)-ascorbic acid (0.01% w/v) solution, or DAF, H89 and calfofin C and LY294002, which were dissolved in dimethyl sulfoxide.

2.4. Data Analysis

Graph representation and statistical analysis were performed using GraphPad Prism 6.0 software (California, USA). The adipose tissue pads and liver weight were normalised using tibia length. The responses induced by EFS were expressed as a % of the initial contraction elicited by 75 mmol/L KCl. To compare the effect of preincubation with L-NAME in EFS-induced contraction experiments, differences between areas under the curve (dAUC) were analysed. The relaxation induced by DEA-NO was expressed as a % of the initial contraction elicited by NA. Results were expressed as mean ± S.E.M. The body weight evolution, the glucose tolerance test and the vasomotor responses to DEA-NO or EFS were compared by means of an unpaired two-way analysis of variance (ANOVA). When comparing the effect of L-NAME on the EFS-induced contraction, a paired two-way ANOVA was used. For the body weight gain, food and water intake, lipid profile, blood pressure, KCl, dAUC, NO, superoxide anion, PKA activity, PKC activity and Western blot densitometry analyses, a Shapiro-Wilk test was applied to confirm the normality of the population data, followed by a one-way ANOVA with a Newman–Keuls post-hoc test. \( p < 0.05 \) was considered significant.

3. Results

3.1. The Effect of HFD and Synbiotic Supplementation on the Body Weight Gain and Lipid Profile

In the present study we induced obesity in Wistar rats by the administration of a HFD (45% fat). The obesity was established after 8 weeks. At week eight, eight rats out of 16 fed on a HFD were administered a commercial synbiotic supplement Prodefen® Plus. After 12 weeks, the body weight increase was significantly higher in MtS rats compared to CT. The supplementation of Prodefen® Plus did not affect the body weight in MtS-SYNB rats (MtS-SYNB) (Figure 2).

![Figure 2](image)

**Figure 2.** Body weight evolution in CT (control group), MtS (metabolic syndrome) and MtS-SYNB (MtS rats supplemented with Prodefen® Plus) groups over 12 weeks. Body weight is expressed as mean ± S.E.M. Statistical analysis: Unpaired two-way ANOVA. *\( p < 0.05 \) CT vs. MtS; +\( p < 0.05 \) CT vs. MtS-SYNB. \( n = \) eight animals per group.

The body weight increase in MtS and MtS-SYNB group was accompanied by an increase in epididymal and visceral adipose pads. The food intake in MtS and MtS-SYNB group was lower and conversely the calories intake was higher. Water intake was comparable among groups (Table 1).
Total cholesterol, LDL and HDL levels did not differ between groups, while the levels of plasma TG were significantly increased in the MtS group as compared to CT group and were returned to control levels in MtS-SYNB group (Table 1). Given the fact that increased TG levels leads to a development of liver steatosis, we next analysed the presence of lipids in liver sections by Sudan III lipid dye. Even though liver weight was comparable among groups, the HFD led to a development of steatosis as demonstrated by the presence of lipid vacuoles throughout the hepatic left lateral lobe. Interestingly, supplementation of Prodefen® Plus decreased the level of steatosis, even though not to the control state (Figure 3).

Table 1. Body weight gain, food and water intake and lipid profile.

| Parameters                        | CT                | MtS               | MtS-SYNB          |
|-----------------------------------|-------------------|-------------------|-------------------|
| Body weight increase 1 (%)        | 87.8 ± 2.1        | 111.3 ± 7.1 *     | 106.1 ± 9.1 *     |
| Epididymal adipose pad 2 (g/cm)   | 1.66 ± 0.08       | 3.30 ± 0.34 *     | 3.44 ± 0.33 *     |
| Visceral adipose pad 2 (g/cm)     | 1.39 ± 0.06       | 3.05 ± 0.35 *     | 3.54 ± 0.30 *     |
| Liver weight 2 (g/cm)             | 1.91 ± 0.07       | 2.01 ± 0.09       | 1.86 ± 0.09       |
| Food intake 3 (g/day)             | 20.96 ± 0.47      | 16.48 ± 0.47 *    | 16.67 ± 0.86 *    |
| Calorie intake 3 (Kcal/day)       | 60.75 ± 1.34      | 75.80 ± 2.14 *    | 76.69 ± 3.93 *    |
| Water intake 3 (mL/day)           | 36.59 ± 2.08      | 35.91 ± 1.02      | 38.16 ± 0.76      |
| TC 4 (mg/dL)                      | 88.88 ± 5.07      | 81.8 ± 5.37       | 90.38 ± 5.93      |
| TG 4 (mg/dL)                      | 112.20 ± 55.80    | 156.00 ± 16.37 *  | 115.40 ± 6.89 *   |
| HDL 4 (mg/dL)                     | 37.29 ± 2.32      | 30.37 ± 2.19      | 35.41 ± 2.57      |
| LDL 4 (mg/dL)                     | 20.61 ± 2.47      | 25.31 ± 3.53      | 31.89 ± 3.08      |

Notes: Data are expressed as mean ± S.E.M. Statistical analysis: One-way ANOVA followed by a Newman–Keuls post-hoc test. * p < 0.05 CT vs. MtS; # p < 0.05 MtS vs. MtS-SYNB; † p < 0.05 CT vs. MtS-SYNB. n = 8 animals per group. 1 Body weight increase is expressed as % of the initial body weight. 2 Respective weights were normalized to tibia length (in cm; CT: 5.25 ± 0.03; MtS: 5.26 ± 0.09; MtS-SYNB: 5.19 ± 0.09). 3 Food and water intake were measured every two days in the last 4 weeks of the experimental procedure coinciding with the administration of the synbiotic. 4 Serum total cholesterol (TC), triglycerides (TG), high density lipoproteins (HDL) and low density lipoproteins (LDL) were analysed at the end of the experimental period of 12 weeks. CT (control group), MtS (metabolic syndrome), MtS rats supplemented with Prodefen® Plus (MtS-SYNB)
Figure 3. Liver sections were stained with Sudan III lipid dye. Representative images (20 × magnification) from CT (a), MtS (b) and MtS-SYNB (c) rats are shown. The scale bar represents 100 μm. (d) Quantitative analysis of Sudan III lipid dye staining. Results are represented as % of stained area (mean ± S.E.M). Statistical analysis: One-way ANOVA followed by a Newman–Keuls post-hoc test. * p < 0.05 CT vs. MtS; # p < 0.05 MtS vs. MtS-SYNB.

3.2. The Effect of Synbiotic Supplementation on MtS Induced Alterations

MtS is related to several metabolic alterations, therefore next we analysed the effect of HFD and synbiotic supplementation on glucose homeostasis. Although basal glycemia was comparable among groups, insulin resistance was developed already at week eight of HFD and was maintained until week 12. The insulin resistance was ameliorated by Prodefen® Plus supplementation, even though not to the control state (Figure 4).

Figure 4. Evolution of insulin resistance measured by glucose tolerance test in CT, MtS and MtS-SYNB rats at the beginning of procedure (a), after 8 weeks of diet intervention (b) and after 12 weeks once the synbiotic supplementation terminated (c). Results are expressed as mg glucose/dL (mean ± S.E. M). Statistical analysis: Unpaired two-way ANOVA. * p < 0.05 CT vs. MtS; † p < 0.05 MtS vs. MtS-SYNB; ‡ p < 0.05 MT vs. MtS-SYNB.
Hypertension is also related to MtS, therefore next we measured the systolic blood pressure using tail cuff method. At week 0 all the animals showed a normotensive phenotype. At week eight, animals fed on a HFD showed an increased systolic blood pressure reaching levels considered as hypertension. The supplementation of Prodefen® Plus for 4 weeks normalized the systolic blood pressure reaching levels comparable to the control group (Table 2).

Table 2. Systolic blood pressure.

|          | CT       | MtS      | MtS-SYNB |
|----------|----------|----------|----------|
| Week 0   | 118.5 ± 1.8 | 110.9 ± 3.0 * | 110.1 ± 2.7 * |
| Week 8   | 119.5 ± 2.4 | 141.1 ± 1.6 * | 137.4 ± 1.0 * |
| Week 12  | 120.3 ± 3.7 | 145.3 ± 2.1 * | 120.6 ± 1.6 * |

Notes: Results are expressed as mmHg (mean ± S.E. M). Statistical analysis: One-way ANOVA followed by a Newman–Keuls post-hoc test. *p < 0.05 CT vs. MtS; #p < 0.05 MtS vs. MtS-SYNB; +p < 0.05 CT vs. MtS-SYNB. n = eight animals per group. 1 Beginning of procedure. 2 Eight weeks of dietary intervention. 3 Twelve weeks of dietary intervention (4 weeks of synbiotic supplementation in MtS + SYNB group).

3.3. Mechanism of HFD-Induced Hypertension

3.3.1. Nitric Oxide Release

The alterations in blood pressure are partially linked to modifications in peripheral vascular resistance. Among the multiple vasoactive factors which regulate vascular tone in superior mesenteric artery, NO has a relevant role, acting as a potent vasodilator. Given the great relevance of nitrergic innervation in the regulation of mesenteric resistance, we analysed the release of neuronal NO release in segments from all experimental groups. The application of an EFS pattern induced NO release in mesenteric segments without endothelium from all groups. As expected, NO release was significantly decreased in MtS group as compared to CT group. Interestingly, the supplementation with Prodefen® Plus for 4 weeks recovered the NO release levels comparable to the control group (Table 3).

Table 3. Nitric oxide and superoxide anion release in endothelium-denuded mesenteric rings.

|                  | CT       | MtS      | MtS-SYNB |
|------------------|----------|----------|----------|
| NO release       | 2518 ± 147 | 1065 ± 10 * | 2498 ± 279 * |
| Superoxide anion | 55.26 ± 3.01 | 55.78 ± 8.13 | 55.74 ± 6.54 |

Notes: 1 Results are expressed as arbitrary fluorescence units (A.F.U.)/mg tissue. Mean ± S.E.M. Statistical analysis: One-way ANOVA followed by a Newman–Keuls post-hoc test. *p < 0.05 CT vs. MtS; #p < 0.05 MtS vs. MtS-SYNB. n = five to eight animals per group. 2 Results are expressed as chemiluminescence units (C.U.)/minute/mg tissue. Mean ± S.E.M. n = five to eight animals per group.

3.3.2. Functional Role for Neuronal Nitric Oxide: Vascular Function

Alteration in blood pressure is linked to modifications in smooth muscle sensitivity to nitric oxide (NO). However, in our model we did not find any differences in the vasodilator response to NO donor DEA-NO among groups (Figure 5a).
Next, we aimed to determine whether the observed alteration in NO release have a relevant functional role. We applied EFS to induce a frequency-dependent contractile response in endothelium-denuded mesenteric segments. An enhanced vasoconstriction response was observed in MtS group, and the supplementation with Prodefen® Plus for 4 weeks diminished this vasoconstriction response (Figure 5b). We next inhibited the NO synthesis with an unspecific NOS inhibitor L-NAME (0.1 mmol/L) and observed a potentiation in the vasoconstriction response in CT and MtS-SYNB groups, while it exerted no effect in arteries from animals from MtS group (Figure 6a–d).
3.3.3. Mechanisms Implicated in Neuronal Nitric Oxide Release

Alterations in nNOS expression and/or activity can be the responsible for the differences in NO release observed. We found that the expression of nNOS was comparable among groups (Figure 6e).

PKA, PKC and PI3K/AKT signalling pathways play a crucial role in the activation of nNOS. To analyze the involvement of the PKA pathway, we preincubated the endothelium-denuded mesenteric segments with a PKA inhibitor H89 (1 μmol/L) before EFS stimulation and measurement of NO release. We observed that H89 diminished EFS-induced NO release similarly in the CT group (60.69 ± 3.45% of inhibition) and in MtS-SYNB group (50.19 ± 13.95% of inhibition) while it produced a lower effect in MtS group (20.48 ± 11.79% of inhibition). We next analysed the involvement of the PKC pathway by using a PKC inhibitor Calphostin C (0.1 μmol/L), finding a similar inhibition by in EFS-induced NO release among the experimental groups (% of inhibition: CT: 76.41 ± 6.53; MtS: 57.33 ± 22.64; MtS-SYNB: 57.54 ± 12.71). Similar results were found after preincubation with LY394002 (10 μmol/L), a PI3K inhibitor (% of inhibition: CT: 83.96 ± 8.61; MtS: 60.51 ± 10.06; MtS-SYNB: 64.22 ± 8.46) (Figure 7a–c).
Mesenteric segments from CT (a), MtS (b) and MtS-SYNB (c) rats were preincubated with a protein kinase A –PKA- inhibitor H89 (1 μmol/L), a protein kinase C –PKC- inhibitor Calphostin C (0.1 μmol/L) or a phosphatidylinositol 3-kinase -PI3K- inhibitor LY294002 (10 μmol/L). NO release was induced by EFS. Data are expressed as arbitrary fluorescence units (A.F.U.)/mg tissue. Statistical analysis: One-way ANOVA followed by a Newman–Keuls post-hoc test. $p < 0.05$ conditions without inhibitor vs. conditions with respective inhibitor.

This finding coincides with the PKA activity, which was significantly reduced in the MtS group as compared to the other two groups (Figure 8a). Similarly to previous findings, PKC activity and AKT expression and phosphorylation were comparable among groups (Figure 8b,c).

3.3.4. Neuronal Nitric Oxide Bioavailability: Oxidative Stress

Although oxidative stress can modulate NO function by diminishing its bioavailability, we observed that superoxide anion release was similar among groups (Table 3).
4. Discussion

Dietary factors, especially hypercaloric diets, are the main contributors to the aetiology of MtS [28]. Chronic consumption of HFD induces overweight along with metabolic and cardiovascular alterations, such as insulin resistance, dyslipidemia or hypertension, among others, being all of these symptoms of the MtS [3,4]. Interestingly, studies in animal models demonstrated that HFD intake is related to gut dysbiosis [29,30]. Gut dysbiosis can develop as early as 2 weeks on HFD, long before the development of MtS-related symptoms [31].

Diets with a 45–60% of the energy derived from fats have been reported to promote a great weight gain in animal models [32,33]. Here we have induced obesity by a diet containing 45% fat and consistently with previous studies, we observed an increase in caloric intake together with body weight gain coinciding with an increased weight of both visceral and epididymal adipose pads [9,10]. Furthermore, in our study, the supplementation of Prodefen® Plus for 4 weeks did not alter the body weight nor the adiposity. Inconsistent results regarding the effect of a synbiotic supplementation on body weight or adiposity index have been reported, showing either improvements [34,35] or no alterations in the weight gain [36,37]. The discrepancies observed are probably caused by different experimental procedures including the use of various probiotic strains. Clinical trials have also reported inconsistent results in body weight changes after a probiotic or synbiotic supplementation [38], moreover clinical trials also usually included changes in life or nutritional habits [39]. Therefore, more studies are needed to evaluate the role of synbiotic supplementation on body weight change.

In our study, the level of TG was increased in rats fed with HFD as compared to standard diet, while TC, HLD and LDL were not, similarly to previously published studies [9], but contrasting with others, in which the dietary intervention was different from ours [24,40]. Liver steatosis is linked to an increase in either TG and/or cholesterol levels, therefore in our study the liver steatosis observed in rats fed with HFD was caused by an increase in TG levels, which is comparable to other studies [40,41]. The supplementation of a synbiotic in our study decreased the TG levels to those achieved with a standard diet together with the diminishment of hepatic lipid deposition as previously described [41,42]. Even though basal glycemia was not altered in the MtS group, an insulin resistance was established already at week eight and was improved by synbiotic administration. A number of studies demonstrated that synbiotic supplementation leads to amelioration or a complete recovery of glucose homeostasis [34,36,37], that can be linked to a normalization in adipokine levels [41,43]. Furthermore, the lowering of plasma TGs levels can also lead to the insulin resistance improvement [44].

The adipokines secreted by the adipose tissue alter the function of multiple vasoactive factors participating in the increase in the peripheral vascular resistance and consequently in the development of MtS-linked hypertension [5]. The HFD in rodents is associated with an increase or no modification of blood pressure depending on the diet composition and duration [9,36,45]. Here we observed that HFD induced an increase in systolic blood pressure, reaching levels that are considered as hypertension. Synbiotic supplementation for 4 weeks diminished systolic blood pressure to values considered as normotensive. The release of antihypertensive factors, such as propionate or butyrate, by the probiotic strains, as well as the normalization of adipokine levels might explain the decrease in blood pressure [41,43,46].

Previous works from our group have demonstrated that the mesenteric vasculature, especially superior mesenteric artery, plays a pivotal role in the maintenance of peripheral resistance through the release of vasodilators, such as NO [8]. Although alterations in blood pressure have been linked with modifications in smooth muscle sensitivity to NO [6,47], we found no differences in the vasodilator response to NO donor DEA-NO among our experimental groups, allowing us to rule out this possibility.

NO in the vascular tissue is predominantly released from the endothelium, and it has been demonstrated that the decrease in endothelial NO release in obesity and MtS is the main factor for endothelial dysfunction [3,48]. Aside from endothelium, NO is also released from perivascular nitrergic innervation and this release is altered in situations in which vascular resistance is decreased [47,49]. Previously we have demonstrated that HFD intake diminishes NO release in superior
mesenteric artery [9,10]. Similarly to observed with endothelial NO [16,36], here we have shown that the supplementation of MtS rats with synbiotics restore neuronal NO release, reaching levels comparable in the CT group. NO released from nitrergic nerve terminals is able to decrease in a great extent the maximal arterial tone elicited by the sympathetic neurotransmitter noradrenaline [23,50]. In situations where the vascular resistance is increased, we have previously observed either a potentiation in nitrergic function, which aims to compensate the greater vascular resistance [23,51], or a blunted nitrergic role, thereby participating in the enhancement of peripheral vascular resistance consequently, in the development of hypertension [47,49]. Here we observed an enhanced EFS-induced vasoconstrictor response in MtS rats, as expected [9,10], and that the supplementation with Prodefen® Plus diminished this vasoconstriction. When inhibiting NO synthesis with the unspecific NOS inhibitor L-NAME, we observed a potentiation in the vasoconstriction in segments from CT and MtS-SYNB rats, while it exerted no effect in arteries from MTS animals, as expected [9,10]. Consequently, we can confirm that the supplementation with Prodefen® Plus restored the nitrergic function lost by MtS, therefore improving the vascular resistance in this pathology.

Next, we wanted to find out whether the expression or the activity of nNOS, the enzyme responsible for the synthesis of NO, is affected by MtS and synbiotic supplementation. We and others have reported that the decrease in the different NOS isozymes, in different tissues is related to obesity and MtS [9,10,52–54] and that probiotic or synbiotic supplementation led to an increase or no modification in the expression of these isoforms [16,55,56]. In the present study we did not observe any alteration in the nNOS expression among groups, a surprising finding given the fact that previously we observed a diminishment of nNOS expression in superior mesenteric artery in rats on a HFD [9,10]. Therefore, we assumed that the increased fat content of the diet and prolonged administration activated compensatory mechanisms, which might explain this difference. Consequently, we assumed that the differential NO released among groups might be due to an alteration in nNOS activation rather than nNOS expression, as we found in spontaneously hypertensive rats supplemented with the similar formula Prodefen® [22].

Different kinases, such as PKA, PKC or PI3K/AKT pathways are essential for multiple physiological responses. Alterations in these pathways have been reported in different MtS models, including increases, decreases and no modifications in their activities [53,57–59]. Even though only few inconsistent results were reported regarding the effect of probiotics and synbiotics on those pathways [60–66], their role in the phosphorylation and subsequent activation of nNOS is well demonstrated [27,51,67,68]. We found that the inhibition of either PKC or PI3K with calfostin C or LY294002 respectively reduced EFS-induced NO release in a similar extent in the three experimental groups. These results suggest that neither MTS nor supplementation with Prodefen® Plus altered the participation of PKC or PI3K/AKT signalling pathways in our experimental conditions. This result agrees with the fact that both the PKC activity, and the expression/activation of AKT were similar in the three experimental groups. Concerning PKA, the preincubation with its specific inhibitor H89 blunted EFS-induced NO release similarly in both CT and MtS-SYNB rats, while it exerted no effect in arteries from MtS animals. This result correlates with the diminished PKA activity observed in segments from MtS rats, which was restored to levels similar to those reached in CT after the administration of Prodefen® Plus. Overall, the observed reduction of PKA activity in mesenteric arteries from MtS rats presumably induced a decrease in nNOS activation and, consequently, a diminished NO release. This inactivation of the PKA-nNOS-NO pathway was recovered by supplementation of Prodefen® Plus for 4 weeks.

NO function can be also influenced by oxidative stress. The amount and the kind of dietary fatty acids can regulate complex intracellular signalling systems, thereby modulating cellular function and promoting a pro-oxidative microenvironment, which reduces NO bioavailability [5,62]. At the vascular level, an increase in superoxide anion metabolizes NO, forming peroxynitrite [27]. Hence, superoxide anion reduces the availability of NO and participates in the development of the vascular disturbances observed in MtS. Since different probiotic strains have been reported to potentiate several antioxidant routes [16,69], the alterations in NO observed in our experimental conditions might be due to modifications in vascular oxidative stress status. However, we found no differences
among the different groups when measuring superoxide anion formation, allowing us to rule out a possible role of oxidative stress in the nitrergic modifications observed in the present study. We should point out that, although both smooth muscle and adventitial layers can be a source of superoxide anions, the endothelial layer, removed in the present study, also has a relevant role in producing these reactive oxygen species [70–72].

5. Conclusions

Overall, our data describe a beneficial effect of Prodefen® Plus regarding the different metabolic associated to MtS. Additionally, Prodefen® Plus improves different mechanisms implicated in the regulation of vascular tone, therefore ameliorating the hypertension associated to this condition. In conclusion, the commercially available formula Prodefen® Plus could be considered an interesting non-pharmacological approach in MtS.

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References

1. Swinburn, B.A.; Kraak, V.I.; Allender, S.; Atkins, V.J.; Baker, P.I.; Bogard, J.R.; Brinsden, H.; Calvillo, A.; De Schutter, O.; Devarajan, R.; et al. The Global Syndemic of Obesity, Undernutrition, and Climate Change: The Lancet Commission report. *Lancet* 2019, 393, 1–56.
2. Collaborators, G.O. Health Effects of Overweight and Obesity in 195 Countries over 25 Years. *N. Engl. J. Med.* 2017, 377, 13–27.
3. Heras Ndl Valero-Muñoz, M.; Ballesteros, S.; Gómez-Hernández, A.; Martín-Fernández, B.; Blanco-Rivero, J.; Cachofeiro, V.; Benito, M.; Balfagón, G.; Lahera, V. Factors involved in rosvastatin induction of insulin sensitization in rats fed a high fat diet. *Nutr. Metab. Cardiovasc. Dis.* 2013, 23, 1107–1114.
4. Williams, T.D.; Chambers, J.B.; Roberts, L.M.; Henderson, R.P.; Overton, J.M. Diet-induced obesity and cardiovascular regulation in C57BL/6 mice. *Clin. Exp. Pharmacol. Physiol.* 2003, 30, 769–778.
5. Torres, S.; Fabersani, E.; Marquez, A.; Gauffin-Cano, P. Adipose tissue inflammation and metabolic syndrome. The proactive role of probiotics. *Eur. J. Nutr.* 2019, 58, 27–43.
6. Caracuel, L.; Sastre, E.; Llevenes, P.; Prieto, I.; Funes, T.; Aller, M.A.; Arias, J.; Balfagón, G.; Blanco-Rivero, J. Acute-on-chronic liver disease enhances phenylephrine-induced endothelial nitric oxide release in rat mesenteric resistance arteries through enhanced PKA, PI3K/AKT and cGMP signalling pathways. *Sci. Rep.* 2019, 9, 6993.
7. Xavier, F.E.; Blanco-Rivero, J.; Avendaño, M.S.; Sastre, E.; Yela R., Velázquez, K.; Salaices M.; Balfagon, G. Aldosterone alters the participation of endothelial factors in noradrenaline vasoconstriction differently in resistance arteries from normotensive and hypertensive rats. *Eur. J. Pharmacol.* 2011, 654, 280–288.
8. Sastre, E.; Márquez-Rodas, I.; Blanco-Rivero, J.; Balfagón, G. Perivascular innervation of the superior mesenteric artery: Pathophysiological implications. *Rev. Neurol.* 2010, 50, 727–737.
9. Blanco-Rivero, J.; Heras, N.D.L.; Martín-Fernández, B.; Cachoeiro, V.; Lahera, V.; Balfagón, G. Rosuvastatin restored adrenergic and nitrergic function in mesenteric arteries from obese rats. *Br. J. Pharmacol.* 2011, 162, 271–285.
10. Sastre, E.; Caracuel, L.; Balfagón, G.; Blanco-Rivero, J. Aerobic exercise training increases nitrergic innervation function and decreases sympathetic innervation function in mesenteric artery from rats fed a high-fat diet. *J. Hypertens.* 2015, 33, 1819–1830.
11. Ley, R.E.; Backhed, F.; Turnbaugh, P.; Lozupone, C.A.; Knight, R.D.; Gordon, J.I. Obesity alters gut microbial ecology. *PNAS* 2005, 102, 11070–11075.

12. Everard, A.; Cani, P.D. Diabetes, obesity and gut microbiota. *Best Pract. Res. Clin. Gastroenterol.* 2013, 27, 73–83.

13. Vallianou, N.; Stratigou, T.; Christodoulatos, G.S.; Dalamaga, M. Understanding the Role of the Gut Microbiome and Microbial Metabolites in Obesity and Obesity-Associated Metabolic Disorders: Current Evidence and Perspectives. *Curr. Obs. Rep.* 2019, 8, 317–332.

14. Rastelli, M.; Knauf, C.; Cani, P.D. Gut Microbes and Health: A Focus on the Mechanisms Linking Microbes, Obesity, and Related Disorders. *Obesity* 2018, 26, 792–800.

15. Jumperz, R.; Le, D.S.; Turnbaugh, P.J.; Trinidad, C.; Bogardus, C.; Gordon, J.I.; Krakoff, J. Energy-balance studies reveal associations between gut microbes, caloric load, and nutrient absorption in humans. *Am. J. Clin. Nutr.* 2011, 94, 58–65.

16. Gómez-Guzmán, M.; Toral, M.; Romero, M.; Jiménez, R.; Galindo, P.; Sánchez, M.; Zarzuelo, M.J.; Olivares, M.; Gálvez, J.; Duarte, J. Antihypertensive effects of probiotics Lactobacillus strains in spontaneously hypertensive rats. *Mol. Nutr. Food Res.* 2015, 59, 2326–2336.

17. Wang, J.; Tang, H.; Zhang, C.; Zhao, Y.; Derrien, M.; Rocher, E.; van-Hylckama, Vlieg, J.E.; Strissel, K.; Zhao, L.; Obin, M.; et al. Modulation of gut microbiota during probiotic-mediated attenuation of metabolic syndrome in high fat diet-fed mice. *ISME J.* 2015, 9, 1–15.

18. Timmerman, H.M.; Koning, C.J.; Mulder, L.; Rombouts, F.M.; Beynen, A.C. Monostrain, multistrain and multispecies probiotics. — A comparison of functionality and efficacy. *Int. J. Food Microbiol.* 2004, 96, 219–233.

19. Diplock, A.T.; Aggett, P.J.; Ashwell, M.; B ornert, F.; Fern, E.B.; Roberfroid, M.B. Scientific Concepts of functional Foods in Europe Consensus Document. *Br. J. Nutr.* 1999, 81, S1–S27.

20. García-Menor, E.; García-Marin, F.; Vecino-López, R.; Horcayo-Martínez, G.; de Ibarrondo Guerricaba, M.J.; Gómez-González, P.; Velasco-Ortega, S.; Suárez-Almarza, J.; Nieto-Magro, C. A Multicenter, Prospective, Randomized Controlled Trial to Evaluate the Additional Benefit of a Multistrain Synbiotic (*Prodefen®*) in the Clinical Management of Acute Viral Diarrhea in Children. *Glob. Pediatr. Health* 2016, 3, doi:10.1177/2333794X16679587.

21. Romeo, A.; Barreiro, C.; Miegimolle, M.; Cadafalch, J.; Millán, B.; Gallego, S.; Suárez-Almarza J.; Nieto C. Efficacy of a 7-strain synbiotic mixture in combination with oral antibiotic treatment in preventing antibiotic-associated diarrhoea (AAD). (PRODEGGIO study. World Congress of Gastroenterology, Istanbul 2019. *Turk. J. Gastroenterol.* 2019, 30 (Suppl. S3), S298.

22. Blanco-Rivero, J.; Llévenes, P.; Balfagón, G. Supplementation with a synbiotic (*Prodefen®*) increases neuronal nitric oxide bioavailability through an enhanced neuronal nitric oxide synthase activation and diminished oxidative stress in superior mesenteric artery from spontaneously hypertensive rats. In Proceedings of the 2nd World Congress on Nutrition and Obesity Prevention Source 2018, Valencia, Spain, 19–21 April 2018; p. 81.

23. Queiroz, D.B.D.; Sastre, E.; Caracuel, L.; Callejo, M.; Xavier, F.E.; Blanco-Rivero, J.; Balfagón, G. Alterations in perivascular innervation function in mesenteric arteries from offspring of diabetic rats. *Br. J. Pharmacol.* 2015, 172, 4699–4713.

24. Amor, S.; González-Hedström, D.; Martin-Carro, B.; Inarejos-García, A.M.; Almodóvar, P.; Prodanov, M.; García-Villalón, A.L.; Granado, M. Beneficial Effects of an Aged Black Garlic Extract in the Metabolic and Vascular Alterations Induced by a High Fat/Sucrose Diet in Male Rats. *Nutrients*. 2019, 11, 153.

25. Knophholz, J.; Disserol, C.C.D.; Pierin, A.J.; Schirr, F.L.; Streisky, L.; Takito, L.L.; Massucheto Ledesma, P.; Faria-Neto, J.R.; Olandoski, M.; da Cunha, C.L.; Bandeira, A.M. Validation of the Friedewald Formula in Patients with Metabolic Syndrome. *Cholesterol* 2014, 2014, doi:10.1155/2014/261878.

26. Nielsen, K.C.; Owman, C. Contractile response and amine receptor mechanisms in isolated middle cerebral artery of the cat. *Brain Res.* 1971, 27, 33–42.

27. Llévenes, P.; Balfagón, G.; Blanco-Rivero, J. Thyroid hormones affect nitricergic innervation function in rat mesenteric T artery: Role of the PI3K/AKT pathway. *Vascul. Pharmacol.* 2018, 108, 36–45.

28. Unger, A.L.; Torres-Gonzalez, M.; Kraft, J. Dairy Fat Consumption and the Risk of Metabolic Syndrome: An Examination of the Saturated Fatty Acids in Dairy. *Nutrients* 2019, 11, 2200.
29. Velázquez, K.T.; Enos, R.T.; Bader, J.E.; Sougiannis, A.T.; Carson, M.S.; Chatzistamou, I.; Carson, J.A.; Nagarkatti, P.S.; Nagarkatti, M.; Murphy, E.A. Prolonged high-fat-diet feeding promotes non-alcoholic fatty liver disease and alters gut microbiota in mice. *World J. Hepatol.* **2019**, *11*, 619–637.

30. Hosomi, R.; Matsudo, A.; Sugimoto, K.; Shimon, T.; Kanda, S.; Nishiyama, T.; Yoshida, M.; Fukunaga, K. Dietary fat influences the expression of genes related to sterol metabolism and the composition ofecal microbiota and its metabolites in rats. *J. Oleo Sci.* **2019**, *68*, 1133–1147.

31. Saiyasit, N.; Chunchai, T.; Prus, D.; Suparan, K.; Pittayapong, P.; Apaiaji, N.; Pratchayasakul, W.; Sripetchwanandee, J.; Chattipakorn, N.; Chattipakorn, S. Gut dysbiosis develops before metabolic disturbance and cognitive decline in high-fat diet-induced obese mouse. *Nutrition* **2019**, *69*, 110576.

32. Lamont, B.J.; Waters, M.F.; Andrikopoulos, S. A low-carbohydrate high-fat diet increases weight gain and does not improve glucose tolerance, insulin secretion or β-cell mass in NZO mice. *Nutr. Diabetes* **2016**, *6*, e194.

33. Qin, L.; Zhao, Y.; Zhang, B.; Li, Y. Amentoflavone improves cardiovascular dysfunction and metabolic abnormalities in high fructose and fat diet-fed rats. *Food Funct.* **2018**, *9*, 243–252.

34. Bagarolli, R.A.; Tobar, N.; Oliveira, A.G.; Araújo, T.G.; Carvalho, B.M.; Rocha, G.Z.; Vecina, J.F.; Calisto, K.; Guadagnini, D.; Prada, P.O.; et al. Probiotics modulate gut microbiota and improve insulin sensitivity in DIO mice. *J. Nutr. Biochem.* **2017**, *50*, 16–25.

35. Alard, J.; Lehrter, V.R.; Rhimi, M.; Mangin, I.N.; Peucelle, V.R.; Abraham, A.-L.; Mariadassou, M.; Maguin, E.; Waligora-Dupriet, A.J.; Pot, B.; et al. Beneficial metabolic effects of selected probiotics on diet-induced obesity and insulin resistance in mice are associated with improvement of dysbiotic gut microbiota. *Environ. Microbiol.* **2016**, *18*, 1484–1497.

36. Toral, M.; Gómez-Guzmán, M.; Jiménez, R.; Romero, M.; Sánchez, M.; Utrilla, M.P.; Garrido-Mesa, N.; Rodriguez-Cabáezas, M.E.; Olivares, M.; Gálvez, J.; et al. The probiotic Lactobacillus casei subsp. casei CECT5711 reduces the vascular pro-oxidant and pro-inflammatory status in obese mice. *Clin. Sci.* **2014**, *127*, 33–45.

37. Wanchai, K.; Yasom, S.; Tunapong, W.; Chunchai, T.; Eamworaawuthikul, S.; Thiennimitr, P.; Chaissyat, C.; Pongchaidecha, A.; Chatsudthipong, V.; Chattipakorn, S.; et al. Probiotic Lactobacillus paracasei HII01 protects rats against obese-insulin resistance-induced kidney injury and impaired renal organic anion transporter 3 function. *Clin. Sci.* **2018**, *132*, 1545–1563.

38. Barengolts, E. Gut microbiota, prebiotics, probiotics, and synbiotics in management of obesity and prediabetes: Review of randomized controlled trials. *Endocr. Pract.* **2016**, *22*, 1224–1234.

39. Tenorio-Jiménez, C.; Martínez-Ramírez, M.J.; Castillo-Codes, I.D.; Arraiza-Irigoyen, C.; Tercero-Lozano, M.; Camacho, J.; Chueca, N.; García, F.; Olza, J.; Plaza-Díaz, J.; et al. Lactobacillus reuteri V3401 reduces the vascular pro-oxidant and pro-inflammatory status in obese mice. *Clin. Sci.* **2018**, *132*, 1851–1862.

40. Higarza, S.G.; Arboleya, S.; Güeimonde, M.; Gómez-Lázaro, E.; Arias, J.L.; Arias, N. Neurobehavioral dysfunction in non-alcoholic steatohepatitis is associated with hyperammonemia, gut dysbiosis, and metabolic and functional brain regional deficits. *PLoS ONE* **2019**, *14*, e0223019.

41. Yao, F.; Jia, R.; Huang, H.; Yu, Y.; Mei, L.; Bai, L.; Ding, Y.; Zheng, P. Effect of Lactobacillus paracasei N1115 and fructooligosaccharides in nonalcoholic fatty liver disease. *Arch. Med. Sci.* **2019**, *15*, 1336–1344.

42. Choi, B.-R.; Kwon, E.-Y.; Kim, H.-J.; Choi, M.-S. Role of Symbiotics Containing d-Allulose in the Alteration of Body Fat and Hepatic Lipids in Diet-Induced Obese Mice. *Nutrients* **2018**, *10*, 1797.

43. Mingrone, G.; Rosa, G.; Rocco, P.D.; Manco, M.; Capristo, E.; Castagneto, M.; Vettor, R.; Gasbarrini, G.; Greco, A.V. Skeletal muscle triglycerides lowering is associated with net improvement of insulin sensitivity, TNF-α reduction and GLUT4 expression enhancement. *Int. J. Obes.* **2002**, *26*, 1165–1172.

44. Nascimento, A.R.; Machado, M. Natha. Structural and Functional Microvascular Alterations in a Rat Model of Metabolic Syndrome Induced by a High-Fat Diet. *Obesity* **2012**, *21*, 2046–2054.

45. Marques, F.Z.; Mackay, C.R.; Kaye, D.M. Beyond gut feelings: How the gut microbiota regulates blood pressure. *Nat. Rev. Cardiol.* **2018**, *15*, 20–32.

46. Blanco-Rivero, J.; Furieri, L.B.; Vassallo, D.V.; Salaices, M.; Balfagón, G. Chronic HgCl2 treatment increases vasoconstriction induced by electrical field stimulation: Role of adrenergic and nitricergic innervation. *Clin. Sci.* **2011**, *121*, 331–341.
48. Heras, N.D.L.; Martín-Fernández, B.; Miana, M.; Ballesteros, S.; Oubiña, M.P.; López-Farré, AJ.; Cachofeito, V.; Lahera, V. The protective effect of irbesartan in rats fed a high fat diet is associated with modification of leptin–adiponectin imbalance. J. Hypertens. 2009, 27, S37–S41.

49. Campo, L.D.; Ferrer, M.; Balfagón, G. Hypertension alters the function of nitricergic and sensory innervation in mesenteric arteries from female rats. J. Hypertens. 2009, 27, 791–799.

50. Sastre, E.; Blanco-Rivero, J.; Caracuel, L.; Callejo, M.A.; Balfagón, G. Alterations in Perivascular Sympathetic and Nitric Innervation Function Induced by Late Pregnancy in Rat Mesenteric Arteries. PLoS ONE 2015, 10, e0126017.

51. Marin, J.; Ferrer, M.; Balfagón, G. Role of protein kinase C in electrical-stimulation-induced neuronal nitric oxide release in mesenteric arteries from hypertensive rats. Clin. Sci. 2000, 99, 277–283.

52. Li, Z.; Rodriguez-Iturbe, B.; Ni, Z.; Shahkarami, A.; Sepassi, L.; Vaziri, N.D. Effect of hereditary obesity on renal expressions of NO synthase, caveolin-1, AKT, guanylate cyclase, and calmodulin. Kidney Int. 2005, 68, 2766–2772.

53. Marčal, A.C.; Leonelli, M.; Fiamoncini, J.; Deschamps, F.C.; Rodrigues, M.A.M.; Curi, R.; Carpinelli, A.R.; Britto, L.R.; Carvalho, C.R.. Diet-induced obesity impairs AKT signalling in the retina and causes retinal degeneration. Cell Biochem. Funct. 2012, 31, 65–74.

54. Surendran, S.; Konnapaka, S.B. Altered expression of neuronal nitric oxide synthase in the duodenal longitudinal muscle-myenteric plexus of obesity induced diabetes mouse: Implications on enteric neurodegeneration. Biochem. Biophys. Res. Commun. 2005, 338, 919–922.

55. Rashid, S.K.; Khodja, N.I.; Auger, C.; Alhosin, M.; Boehm, N.; Oswald-Mammosser, M.; Schini-Kerth, V.B. Probiotics (VSL#3) Prevent Endothelial Dysfunction in Rats with Portal Hypertension: Role of the Angiotensin System. PLoS ONE 2014, 9, e97458.

56. Suo, H.; Zhao, X.; Qian, Y.; Sun, P.; Zhu, K.; Li, J.; Sun, B. Lactobacillus fermentum Suo Attenuates HCl/Ethanol Induced Gastric Injury in Mice through Its Antioxidant Effects. Nutrients 2016, 8, 155.

57. Engin, A. Human Protein Kinases and Obesity. Adv. Exp. Med. Biol. 2017, 960, 111–134.

58. Hwang, J.H. Effects of obesity on protein kinase C, brain creatine kinase, transcription, and autophagy in cochlea. Metab. Brain Dis. 2017, 32, 735–742.

59. Vagena, E.; Ryu, J.K.; Baeza-Raja, B.; Walsh, N.M.; Syme, C.; Day, J.P.; Houslay, M.D.; Baillie, G.S. A high-fat diet promotes depression-like behavior in mice by suppressing hypothalamic PKA signaling. Transl. Psychiatry. 2019, 9(1): 141.

60. Zhou, Y.-K.; Qin, H.-L.; Zhang, M.; Shen, T.-Y.; Chen, H.-Q.; Ma, Y.-L.; Chu, Z.X.; Zhang, P.; Liu, Z.H... Effects of Lactobacillus plantarum on gut barrier function in experimental obstructive jaundice. World J. Gastroenterol. 2012, 18, 3977–3991.

61. Heuvelin, E.; Lebretcon, C.; Bichara, M.; Cerf-Bensussan, N.; Heyman, M. A Bifidobacterium Probiotic Strain and Its Soluble Factors Alleviate Chloride Secretion by Human Intestinal Epithelial Cells. J. Nutr. 2009, 140, 7–11.

62. Ting, W.-J.; Kuo, W.-W.; Kuo, C.-H.; Yeh, Y.-L.; Shen, C.-Y.; Chen, Y.H.; Ho, T.J.; Viswanadh, V.P.; Chen, Y.H.; Huang, C.Y. Supplementary heat-killed Lactobacillus reuteri GMNL-263 ameliorates hyperlipidaemic and cardiac apoptosis in high-fat diet-fed hamsters to maintain cardiovascular function. Br. J. Nutr. 2015, 114, 706–712.

63. Paveljšek, D.; Juvan, P.; Kosir, R.; Rozman, D.; Hacin, B.; Ivičak-Kocjan, K.; Rogelj, I.. Lactobacillus fermentum L930BB and Bifidobacterium animalis subsp. animalis IM386 initiate signalling pathways involved in intestinal epithelial barrier protection. Benef. Microbes 2018, 9, 515–525.

64. Yang, B.; Luo, Y.; Liu, Z.; Yang, P.; Gui, Y. Probiotics SOD inhibited food allergy via downregulation of STAT6-TIM4 T signaling on DCs. Mol. Immunol. 2018, 103, 71–77.

65. Han, N.; Jia, L.; Su, Y.; Du, J.; Guo, L.; Luo, Z.;Liu, Y.. Lactobacillus reuteri extracts promoted wound healing via PI3K/AKT/NF-κB axis. Stem Cell Res. Ther. 2019, 10, 243.

66. Kumar, R.; Sharma, A.; Gupta, M.; Padwad, Y.; Sharma, R. Cell-Free Culture Supernatant of Probiotic Lactobacillus fermentum Protects Against H2O2-Induced Premature Senescence by Suppressing ROS-Akt-mTOR Axis in Murine Preadipocytes. Probiotics Antimicrob. Proteins 2019, 1–14, doi:10.1007/s12602-019-00957-2.

67. Ferrer, M.; Sánchez, M.; Martín, M.D.C.; Márquez-Rodas, I.; Alonso, M.J.; Salaices, M.; Ballagón, G.et al. Protein kinase A increases electrical stimulation-induced neuronal nitric oxide release in rat mesenteric artery. Eur. J. Pharmacol. 2004, 487, 167–173.
68. Blanco-Rivero, J.; Balfagón, G.; Ferrer, M. Male Castration Increases Neuronal Nitric Oxide Synthase Activity in the Rat Mesenteric Artery through Protein Kinase C Activation. *J. Vasc. Res.* 2005, 42, 526–534.

69. D’Souza, A.; Fordjour, L.; Ahmad, A.; Cai, C.; Kumar, D.; Valencia, G.; Aranda, J.V.; Beharry, K.D. Effects of probiotics, prebiotics, and synbiotics on messenger RNA expression of caveolin-1, NOS, and genes regulating oxidative stress in the terminal ileum of formula-fed neonatal rats. *Pediatr. Res.* 2010, 67, 526–531.

70. Miller, F.J., Jr.; Gutterman, D.D.; Rios, C.D.; Heistad, D.D.; Davidson, B.L. Superoxide production in vascular smooth muscle contributes to oxidative stress and impaired relaxation in atherosclerosis. *Circ. Res.* 1998, 82, 1298–1305.

71. Wang, H.D.; Pagano, P.J.; Du, Y.; Cayatte, A.J.; Quinn, M.T.; Brecher, P.; Cohen, R.A.. Superoxide anion from the adventitia of the rat thoracic aorta inactivates nitric oxide. *Circ. Res.* 1998, 82, 810–818.

72. Li, H.; Horke, S.; Förstermann, U. Vascular oxidative stress, nitric oxide and atherosclerosis. *Atherosclerosis* 2014, 237, 208–219.

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