Spices in a High-Saturated-Fat, High-Carbohydrate Meal Reduce Postprandial Proinflammatory Cytokine Secretion in Men with Overweight or Obesity: A 3-Period, Crossover, Randomized Controlled Trial

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ABSTRACT

Background: Postprandial inflammation that occurs concurrently with hyperglycemia and hyperlipidemia after ingestion of a high-saturated-fat, high-carbohydrate meal (HFCM) is a risk factor for cardiovascular disease (CVD). Numerous preclinical and clinical studies demonstrate anti-inflammatory effects of individual spices. However, the effect of consumption of a spice blend on inflammatory mediators has not been examined in a randomized controlled trial.

Objectives: The objective of this study was to investigate the postprandial effect of a blend of spices in a HFCM on inflammatory cytokine responses.

Methods: Nonsmoking men (40–65 y old) with overweight/obesity (25 ≤ BMI ≤ 35 kg/m²), elevated waist circumference (≥ 94 cm), and ≥ 1 CVD risk factor were recruited for a 3-period crossover study (n = 12). In random order, participants consumed the following: a HFCM (~1000 kcal, 33% kcal from saturated fat and 36% kcal from carbohydrate), a HFCM containing 2 g spice blend, or an HFCM containing 6 g spice blend. The spice blend consisted of basil, bay leaf, black pepper, cinnamon, coriander, cumin, ginger, oregano, parsley, red pepper, rosemary, thyme, and turmeric. Blood was collected before, and hourly for 4 h after the HFCM. Peripheral blood mononuclear cells (PBMCs) were isolated, and the percentage of CD14+Human Leukocyte Antigen-DR isotype + (HLA-DR +) monocytes and proinflammatory cytokine concentrations in plasma and LPS-stimulated PBMCs were quantified as secondary outcomes.

Results: There was a significant spice-by-time interaction on IL-1β (P < 0.001), IL-8 (P = 0.020), and TNF-α (P = 0.009) secretion from LPS-stimulated PBMCs. IL-1β secretion from LPS-stimulated PBMCs was significantly reduced (1314%) at 240 min after HFCM consumption containing 6 g, but not 2 g, of spice blend compared with 0 g spice blend.

Conclusions: A HFCM containing 6 g spice blend attenuated HFCM-induced postprandial IL-1β secretion in men with overweight/obesity. This trial was registered at clinicaltrials.gov as NCT03064958.

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Keywords: monocytes, inflammatory cytokines, obesity, nutritional intervention, randomized controlled trial

Introduction

Overweight or obesity affects nearly 72% of adults in the United States, and the prevalence of obesity has nearly tripled since 1962 (1). There is a strong association between overweight/obesity and the incidence of cardiovascular disease (CVD) and other metabolic diseases (2), which may be mediated, at least in part, by chronic low-grade inflammation (3). In US adults, the risk of cardiometabolic disease is strongly associated with unhealthy dietary habits (e.g., high intake of saturated fats, refined carbohydrates, and sugar-sweetened beverages, and low intake of whole grains, fruits, and vegetables) (4), which may contribute to elevated postprandial inflammation (5). Postprandial hyperglycemia and hyperlipidemia, which are identified as independent CVD risk factors, induce postprandial inflammation (6). Furthermore, postprandial inflammation is associated with increased risk of CVD (5). Thus, gaining a better understanding of the relation between postprandial hyperglycemia, hyperlipidemia, and inflammation has important clinical implications.

In the postprandial state, increased glucose and free fatty acids enter the tricarboxylic acid (TCA) cycle, overwhelm the oxidative phosphorylation capacity of mitochondria, and lead...
to oxidative stress that further activates inflammatory pathways (7, 8). Repetitive and prolonged inflammation increases the risk of CVD, which is mediated by several cytokines including IL-1β, IL-6, IL-8, monocyte chemoattractant protein (MCP)-1, and TNF-α. IL-1β, MCP-1, and TNF-α increase the expression of adhesion molecules on vascular endothelial cells and serve as chemoattractants, inducing the recruitment of inflammatory cells to the intima of blood vessels (9, 10). IL-6 and IL-8 are involved in leukocyte trafficking across the vascular endothelium (11, 12), contributing to vascular inflammation and atherosclerotic plaque formation (13, 14). Postprandial inflammation and oxidative damage are augmented in a state of metabolic dysfunction (e.g., obesity and type 2 diabetes) (15–18). Furthermore, chronic low-grade inflammation may be exacerbated by postprandial inflammation (19). Thus, the identification of novel dietary strategies to reduce postprandial inflammation is of public health importance given the prevalence of chronic low-grade inflammation and cardiometabolic diseases in the population.

Spices not only add flavor to food, but also may confer various health benefits likely due to the presence of bioactive compounds (20). Numerous preclinical and clinical studies demonstrate anti-inflammatory effects of individual spices and their bioactive compounds such as turmeric (curcumin), ginger (gingerol), and cinnamon (cinnamaldehyde). However, the effect of the consumption of a spice blend, as typically occurs in a meal, on inflammatory mediators has not been examined in humans in a randomized controlled trial.

Therefore, the purpose of this study was to examine the effect of consuming a spice blend in a standardized high-saturated-fat, high-carbohydrate test meal compared with the same meal without the spice blend, on postprandial inflammation in middle-aged men with overweight/obesity. Similar to previous research in this area (21–23), we chose to recruit a small number of homogeneous subjects, and used a crossover design to minimize interpersonal variability in order to ascertain whether spice consumption induced an anti-inflammatory effect in the postprandial setting.

Methods

Subjects

We recruited 12 nonsmoking men (40–65 y old) with overweight or obesity (25 ≤ BMI ≤ 35 kg/m²), elevated waist circumference (≥ 94 cm), and 1 of the following CVD risk factors: altered lipid profile [LDL cholesterol > 130 mg/dL, triglyceride (TG) ≥ 150 mg/dL, or HDL cholesterol < 40 mg/dL], elevated C-reactive protein (CRP) (> 1 mg/dL), elevated blood pressure [systolic blood pressure (SBP) ≥ 130 mm Hg and DBP ≥ 85 mm Hg], or elevated fasting glucose (≥ 100 mg/dL). Exclusion criteria included having diabetes (fasting glucose > 126 mg/dL) or hypertension (SBP > 160 mm Hg or DBP > 100 mm Hg); receiving any antihypertensive or glucose-lowering drugs; having established CVD, stroke, diabetes, liver, kidney, or autoimmune disease; using cholesterol/lipid-lowering medication or supplements (psyllium, fish oil, lecithin, and phytoestrogens) and botanicals; and having weight loss of ≥ 10% body weight within the 6 mo before enrolling in the study. We originally planned to enroll 6 participants; however, a preliminary analysis revealed a trend toward a different postprandial inflammatory response based on fasting blood glucose concentration (normal fasting glucose range: <100 mg/dL, compared with impaired fasting glucose range: ≥100 mg/dL). Thus, we enrolled 7 additional participants with the goal of having half of the participants with normal fasting glucose and the remainder with impaired fasting glucose. One participant was excluded from the analysis because we failed to collect peripheral blood mononuclear cells (PBMCs) from the participant.

Recruitment, screening, and random assignment

Participants were recruited through advertisements using flyers on campus and in the community. Individuals who were on the university email lists and who had previously participated in other studies were contacted. They were given further information about the study and screened for eligibility by telephone if interested. Potential participants were scheduled for a screening visit at the Clinical Research Center on the Pennsylvania State University-University Park campus. Participants fasted for 12 h and abstained from alcohol for 48 h before the screening visit. Height, weight, blood pressure, and waist circumference were measured. A blood sample was collected and sent to Quest Diagnostics (Pittsburgh, PA) for biochemical measurement of glucose, lipid profile, and CRP. Eligible participants (n = 12) were randomly assigned to treatment sequences using a computer-generated scheme (www.randomization.com). The randomization code was kept by the kitchen staff preparing the meals and the code was broken at the end of the study.

Study design

The study was a 3-period, crossover, randomized controlled trial (NCT03064938) that was designed to assess the effect of spice consumption on serum TGs as the primary outcome. The study was powered to detect a change in TGs between the treatment groups, and 13 subjects were estimated to yield a TG difference of 15 ± 32 mg/dL (mean ± SD) with 80% power (α = 0.05). These results are summarized in a separate manuscript (under review). We quantified inflammatory responses in 12 subjects because we could not collect enough blood from 1 subject to perform the assays needed to assess inflammatory cytokine secretion in cultured PBMCs. In random order, participants consumed the following meals: 1) a ~1000-kcal high-saturated-fat, high-carbohydrate meal (HFCM) containing 33% kcal from saturated fat and 36% kcal from carbohydrate with no additional spices; 2) a HFCM containing 2 g spice blend; or 3) a HFCM containing 6 g spice blend, with a ≥3-d washout period between intervention days. The nutrient profile of the HFCM and composition of the spice blend in the HFCM are presented in Tables 1 and 2, respectively. The doses of spices used in the current study were chosen to incorporate doses on the low and high ends of daily spice consumption per capita in the United States in 2015 (mean 4.5 g/d) (24). The spices chosen were based on previous studies that reported benefits on CVD and inflammatory outcomes (25–28). In addition, the spices chosen are among the most widely consumed spices in the US diet (29). The test meal was coconut chicken curry, a corn muffin, and a cinnamon biscuit. All the experiments in this study were performed with the approval of the Institutional Review Board of the Pennsylvania State University-University Park campus.
TABLE 1 Nutrient profile of the high-saturated-fat, high-carbohydrate meal

| Nutrient profile                  | Coconut chicken curry | Corn muffin | Cinnamon biscuit | Total   |
|----------------------------------|-----------------------|-------------|------------------|---------|
| Energy, kcal                     | 617                   | 354         | 105              | 1076    |
| Total fat, g                     | 39.7                  | 15.2        | 3.9              | 59.5    |
| Total fat, % kcal                | 3.2                   | 2.3         | 1.1              | 3.3     |
| Saturated fat, g                 | 32.5                  | 5.7         | 1.1              | 39.3    |
| Saturated fat, % kcal            | 7.2                   | 8.9         | 2.9              | 22.0    |
| Carbohydrates, g                 | 40.4                  | 41.3        | 16.1             | 97.7    |
| Carbohydrates, % kcal            | 15.0                  | 15.4        | 6.0              | 36.3    |
| Protein, g                       | 26.6                  | 14.9        | 1.4              | 42.9    |
| Protein, % kcal                  | 8.3                   | 5.5         | 0.5              | 15.9    |
| Dietary fiber, g                 | 0.4                   | 1.3         | 1.3              | 2.9     |

1Nutrient values were determined using the Nutrient Data System for Research (Minneapolis, MN).

Blood sample collection
A baseline blood sample was collected in sterile EDTA (K2)-coated blood tubes (BD Biosciences), and participants were asked to consume the test meal within 15 min. Blood samples were collected at timed intervals (60, 120, 180, and 240 min) after meal consumption. No other foods or drinks (other than water) were allowed for the remainder of the 240-min testing period. Blood samples were centrifuged at 1800 × g for 15 min at room temperature. Plasma was dispensed into microcentrifuge tubes and frozen at −80°C until analysis.

Proinflammatory cytokine secretion assay
PBMCs were isolated from blood as previously described (30). PBMCs (2 × 10^6/mL) were stimulated with 0.625 μg/mL LPS (Sigma-Aldrich) in round-bottomed 96-well plates, and supernatants were harvested after 24-h incubation and frozen at −80°C until analysis.

Measurement of proinflammatory cytokine concentrations
Cytokines and chemokines (IFN-γ, IL-1β, IL-2, IL-4, IL-6, IL-8, IL-10, IL-12p70, IL-13, TNF-α, and MCP-1) in plasma and supernatants were measured using the V-PLEX Proinflammatory Panel 1 Human Kit and V-PLEX Human MCP-1 kit (Meso Scale Diagnostics) as per the manufacturers’ instructions. For the data points that were below the detection range, half of the lower limit of detection was used as a value in the analyses.

TABLE 2 Composition of spice blend in the high-saturated-fat, high-carbohydrate meal

| Spice          | 2 g spice blend | 6 g spice blend |
|----------------|-----------------|-----------------|
|                | Coconut chicken curry | Corn muffin | Cinnamon biscuit | Total |
|                | 0.35            | 0.13            | 0.08             | 0.35  |
| Turmeric, g    | —               | —               | —                | 0.35  |
| Ginger, g      | 0.13            | —               | 0.15             | 0.26  |
| Cinnamon, g    | 0.08            | —               | 0.13             | 0.23  |
| Oregano, g     | —               | 0.19            | —                | 0.19  |
| Parsley, g     | —               | 0.14            | —                | 0.14  |
| Basil, g       | —               | 0.13            | —                | 0.13  |
| Coriander, g   | 0.13            | —               | —                | 0.13  |
| Cumin, g       | 0.13            | —               | —                | 0.13  |
| Red pepper, g  | 0.13            | —               | —                | 0.13  |
| Rosemary, g    | —               | 0.10            | —                | 0.10  |
| Black pepper, g| 0.08            | —               | —                | 0.08  |
| Bay leaf, g    | —               | 0.06            | —                | 0.06  |
| Thyme, g       | 0.06            | —               | —                | 0.06  |
| Total, g       | —               | —               | —                | 2.00  |

Flow cytometric analysis
PBMCs were stained with fluorescently labeled antibodies as previously described (30). Antibodies for immune cell markers included CD3, CD4, CD8, CD14, CD19, CD56, and Human Leukocyte Antigen-DR isotype (HLA-DR). Antibody isotype controls included mouse IgG2b, mouse IgG2a, and mouse IgM. CD56 was purchased from BD Biosciences and all remaining antibodies were purchased from BioLegend. A total of 50,000 events were acquired with BD LSR-Fortessa (BD Biosciences). Data were analyzed and plotted using FlowJo 10 (FlowJo, LLC). The monocyte population within the total PBMC population was gated based on forward scatter and side scatter (Figure 1A). The percentage of CD14+ /HLA-DR+ cells in the monocyte gate was quantified per sample (Figure 1B) and the percentage of CD14+/HLA-DR+ monocytes within PBMCs was calculated. We could not collect enough blood in 5 subjects to run flow cytometric analyses; thus, we assessed the percentage of CD14+/HLA-DR+ in only 7 subjects.

Statistical analyses
Statistical analyses were performed using SAS version 9.4 (SAS Institute). The percentage of CD14+/HLA-DR+ monocytes and inflammatory cytokine concentrations were secondary outcomes in the study design. Normality of the data was confirmed by a Q–Q plot assessment. Equal variance was confirmed by residual-versus-predicted value plots. Plasma cytokine concentration and cytokine secretion from PBMCs are reported as the difference between baseline and individual time points after meal consumption. Data points that were >3 SDs from the mean were considered as outliers and removed. A mixed-effects model for repeated measures was used to test the effects of time and/or treatment (spice) on the distribution of immune cell populations (monocytes, CD4+ T cells, CD8+ T cells, B cells, and NK cells) and postprandial proinflammatory cytokine concentrations in the plasma and in culture supernatants (Supplemental Table 1). Outcomes were modeled as repeated measures with a first-order autoregressive matrix. For postprandial cytokine secretion after the HFCM challenge, plasma glucose and its interaction with treatment (spice) were included as covariates because the change in plasma glucose was associated with changes in cytokine secretion. Baseline outcomes were not included as covariates because the models that included baseline outcome as a covariate were not statistically different from models that did not. Participant was designated as a random factor, and the treatment (spice), time, and covariates were fixed factors. In cases where significant treatment (spice) effects or time-by-treatment (spice) interactions were detected, separate analyses were performed to determine between-treatment (spice) effects at each time point, and time effects for each treatment (spice) group, using mixed-effects analyses for repeated measures followed by Tukey’s post hoc test. For all outcomes, α was
set as 0.05 and statistical significance was accepted at \( P < 0.05 \). Graphs were plotted using Prism 7 (GraphPad). Values are reported as mean ± SEM.

**Results**

**Baseline characteristics**

Table 3 presents anthropometric measurements, blood pressure, and biochemical measurements at screening. The participants were middle-aged nonsmoking men (51.8 ± 2.7 y) with overweight/obesity (BMI 29.4 ± 0.7 kg/m²) and elevated waist circumferences (100.1 ± 1.3 cm). There was a wide range in blood pressure and biochemical measures among participants because only 1 additional risk factor for CVD was required as per the inclusion criteria.

**HFCM challenge increased the percentage of CD14⁺/HLA-DR⁺ monocytes**

There was a significant postprandial increase in the percentage of CD14⁺/HLA-DR⁺ monocytes (main effect of time, \( P = 0.001 \)) (Figure 1C) after the HFCM challenge. However, there was no effect of spice and no time-by-spice interaction on the percentage of CD14⁺/HLA-DR⁺ monocytes in circulation.

**Table 3** Baseline characteristics of participants with overweight or obesity at risk of cardiovascular disease

| Characteristic | Participants |
|----------------|--------------|
| Age, y         | 51.8 ± 2.7(40–64) |
| BMI, kg/m²     | 29.4 ± 0.7(25.9–33.8) |
| Waist circumference, cm | 100.1 ± 1.3(94.2–101) |
| Blood pressure, mm Hg | 122 ± 3.3(106–138) |
| Systolic       | 78.2 ± 1.6(70–87) |
| Diastolic      | 97.1 ± 3.5(75.0–112) |
| Glucose, mg/dL | 200 ± 10.4(149–269) |
| Total cholesterol, mg/dL | 45.0 ± 2.6(34.0–64.0) |
| HDL cholesterol, mg/dL | 131 ± 8.5(88.0–195) |
| Triglycerides, mg/dL | 121 ± 16.2(51.0–202) |
| C-reactive protein, mg/dL | 1.3 ± 0.3(0.3–3.5) |

\( n = 12 \). Values are mean ± SEM (range).

(Figure 1C). The percentage of monocytes in circulation was significantly higher at 240 min than at 0, 60, 120, and 180 min after the HFCM challenge (Tukey’s post hoc test, \( P < 0.05 \)). The percentages of CD3⁺/CD4⁺ T cells, CD3⁺/CD8⁺ T cells, and CD3⁺/CD56⁺ NK cells were not affected by time or spice (Supplemental Figure 1A, B, C). There was a significant postprandial decrease in the percentage of B cells (main effect of time, \( P = 0.016 \)), but no effect of spice or a time-by-spice interaction on the percentage of CD19⁺ B cells. The percentage of B cells was significantly lower at 180 min than at 0, 60, 120, or 240 min after the HFCM challenge (Tukey’s post hoc test, \( P < 0.05 \)) (Supplemental Figure 1D).

**Consumption of a HFCM containing a spice blend attenuated postprandial inflammation**

The HFCM challenge alone induced postprandial inflammation, as evidenced by a significant increase in plasma IL-6 concentration and IL-1β secretion from LPS-stimulated PBMCs (data not shown).

There was no significant effect of time on plasma IL-1β, IL-8, and MCP-1 (Figure 2A, C, D) after HFCM consumption. There was a significant effect of time on plasma concentrations of IL-6 (\( P < 0.001 \)) and TNF-α (\( P = 0.044 \)) (Figure 2B, E). There was no significant spice effect or time-by-spice interaction on plasma IL-1β, IL-6, IL-8, MCP-1, and TNF-α concentrations after HFCM consumption (Figure 2A–E).

There was no significant overall effect of time on IL-1β, IL-6, IL-8, MCP-1, and TNF-α secretion from LPS-stimulated PBMCs (Figure 3A–E). There was a significant spice effect on IL-1β (\( P < 0.001 \)), IL-6 (\( P = 0.049 \)), and IL-8 secretion (\( P = 0.007 \)) after the meal challenge (Figure 3A–C), but no spice effect on MCP-1 and TNF-α secretion (Figure 3D, E). A significant time-by-spice interaction was observed for IL-1β (\( P < 0.001 \)), IL-8 (\( P = 0.020 \)), and TNF-α secretion (\( P = 0.009 \)) after the meal consumption (Figure 3A, C, E), resulting in spice-induced reduction in these cytokines. There was no time-by-spice interaction for IL-6 and MCP-1 secretion (Figure 3B, D).

When the effect of time was analyzed in each treatment group, IL-1β secretion from cultured PBMCs was significantly increased 60 min after HFCM consumption containing 0 g spice blend compared with 0, 120, 180, and 240 min after the meal (Tukey’s post hoc test, \( P < 0.05 \)) (Supplemental Table 2). IL-1β secretion was significantly reduced at 240 min after
HFCM consumption containing 2 g spice blend compared with 0, 60, 120, and 180 min after the meal (Tukey’s post hoc test, \( P < 0.05 \)) (Supplemental Table 2). There was no effect of time on IL-\( \beta \) secretion after HFCM consumption containing 6 g spice blend. However, all cytokine change scores were negative after the meal, indicating a decrease from baseline after the meal containing 6 g spice blend (Supplemental Table 2).

When the effect of spice was compared at each time point, IL-1\( \beta \) and IL-8 secretion from cultured PBMCs was significantly lower at 180 min after HFCM consumption containing 6 g spice blend, than for 2 g spice blend, by 171% and 155%, respectively (Figure 3A, C). At 240 min after HFCM consumption containing 6 g spice blend, IL-1\( \beta \) secretion was significantly reduced (1314%) compared with 0 g spice blend (Figure 3A). In addition, at 240 min after HFCM consumption containing 6 g spice blend, IL-6 and IL-8 secretion were significantly lower than for 2 g spice blend, by 445% and 829%, respectively (Figure 3B, C).

There was an effect of spice consumption on plasma IFN-\( \gamma \) (\( P < 0.001 \)), IL-2 (\( P < 0.001 \)), IL-4 (\( P = 0.043 \)), and IL-10 concentrations (\( P = 0.034 \)) (Supplemental Table 3) and on IL-2 secretion from LPS-stimulated PBMCs (\( P = 0.040 \)) (Supplemental Table 4).

**Discussion**

In the current study, we demonstrated that the percentage of monocytes significantly increased in the circulation at 240 min after the HFCM challenge. Furthermore, we demonstrated a significant spice effect on IL-6, and a significant spice-by-time interaction on IL-1\( \beta \), IL-8, and TNF-\( \alpha \) secretion from LPS-stimulated PBMCs. Consumption of 6 g spice blend significantly reduced IL-1\( \beta \) secretion at 240 min after HFCM consumption compared with 0 g spice blend. Furthermore, the change in IL-6, IL-8, MCP-1, and TNF-\( \alpha \) secretion from LPS-stimulated PBMCs at 240 min after HFCM consumption containing 6 g spice blend was below baseline values for each cytokine. These results suggest that spice consumption may reduce HFCM-induced postprandial inflammation.

Anti-inflammatory effects of spices and their bioactive compounds are well documented in preclinical and clinical studies. Turmeric is the most commonly used spice in the world. Curcumin, the main bioactive compound in turmeric, alleviates oxidative stress in LPS-injected BALB/c mice and in mouse peritoneal macrophages (31), and reduces IL-1\( \beta \) and TNF-\( \alpha \) secretion from LPS-stimulated human monocyte cell lines (32). Curcumin reduces serum inflammatory cytokine concentrations in adults with obesity (33) and metabolic syndrome (34). Ginger has a long history of medical use, some of which is based on its broad anti-inflammatory function. Supplementation with ginger reduces circulating inflammatory markers in subjects with type 2 diabetes (35, 36) and nonalcoholic fatty liver disease (37). Cinnamon exhibits similar anti-inflammatory and immunomodulatory properties in both *in vitro* and *in vivo* animal studies (38, 39), and this is likely due to the action of the bioactive compound cinnamaldehyde (40). In addition, other spices included in our test meal from the current study, such as basil, bay leaf, black pepper, coriander, cumin, red pepper, rosemary, and thyme, may have anti-inflammatory effects.
FIGURE 3  Change in IL-1β (A), IL-6 (B), IL-8 (C), MCP-1 (D), and TNF-α (E) secretion from LPS-stimulated peripheral blood mononuclear cells after a HFCM challenge containing 0 g, 2 g, or 6 g spice blend in men with overweight or obesity at risk of cardiovascular disease. Labeled means at each time point without a common letter differ, \( P < 0.05 \). Data are mean ± SEM. \( n = 12 \). HFCM, high-saturated-fat, high-carbohydrate meal; MCP, monocyte chemoattractant protein.

Thus, the use of a combination of spices may reduce inflammatory mediators to a greater degree than any individual spice owing to possible differences in bioactivity, duration of effect, and mechanism of action of the individual spices in the blend. Although numerous individual spices show anti-inflammatory properties in vitro and in vivo in animal models and humans, few studies have examined the effect of the consumption of a spice blend in humans in a randomized controlled trial. Also, few studies have investigated the anti-inflammatory effect of spices in the context of daily meal consumption, which is the typical vehicle of spice consumption. Thus, we developed a test meal containing a spice blend in order to investigate the anti-inflammatory effect of spice in the context of meal consumption. In addition, the spices were selected from the most widely consumed spices in the United States at doses that may be consumed (29), thus representing a blend of spices that may be consumed in the average American diet.

Postprandial inflammation is the response of immune cells, such as monocytes, dendritic cells, and lymphocytes, to the acute postprandial overload of macronutrients that culminates in oxidative stress and inflammation. Metabolism of high amounts of glucose and free fatty acids results in excessive acetyl-CoA production, mitochondrial metabolism of acetyl-CoA via the TCA cycle, and an increase in reactive oxygen species (ROS) within immune cells. This change in the redox status can activate redox-sensitive transcription factors, including NF-κB, which triggers an inflammatory cascade resulting in cytokine secretion (7, 8). There are nutrient-independent and nutrient-dependent factors that influence postprandial inflammation. Nutrient-independent factors include obesity, type 2 diabetes, and a sedentary lifestyle, which are associated with chronic low-grade inflammation. Individuals with metabolic disorders such as obesity and diabetes have an augmented postprandial inflammatory response in addition to the elevated inflammatory mediators in the circulation in the fasting state when compared with healthy controls (15). Patel et al. (17) demonstrate prolonged ROS generation and greater intranuclear NF-κB binding activity in PBMCs after high-fat, high-carbohydrate meal consumption in subjects with obesity compared with healthy subjects. Nutrient-dependent factors of postprandial inflammation include caloric content, glycemic index, and lipid profile of the meal (41). Thus, high-fat and/or high-carbohydrate challenge tests are often used to study postprandial inflammation and the effectiveness of nutritional interventions in reducing postprandial inflammation. An acute homeostatic perturbation, such as an HFCM challenge, can be used to capture small changes in inflammatory mediators in the circulation that are not easily observed by static homeostatic measures (42). In the current study, we induced postprandial inflammation by administrating a high-saturated-fat, high-carbohydrate test meal (~1000 kcal, 33% kcal from saturated fat and 36% kcal from carbohydrate) that included coconut chicken curry, a corn muffin, and a cinnamon biscuit, and assessed plasma inflammatory cytokine concentration and inflammatory cytokine secretion from LPS-stimulated PBMCs after consumption of the test meal containing 0 g, 2 g, and 6 g spice blend in a crossover design.
In optimal physiologic (healthy) conditions, postprandial inflammation is adequately controlled by a variety of nutrient-sensing regulatory mechanisms, making it challenging to capture the therapeutic effect of nutritional interventions on postprandial inflammation. In contrast, subjects at risk of metabolic syndrome have a higher systemic stress response to high-fat challenges than do healthy subjects (43). Given the exaggerated postprandial inflammatory response in this population with elevated systemic inflammatory tone, it is often easier to detect the effectiveness of therapeutic nutritional interventions. The participants recruited for the current study were men with overweight/obesity who also had an elevated waist circumference and had ≥ 1 other CV risk factor (elevated LDL cholesterol and TGs, reduced HDL cholesterol, elevated CRP, elevated blood pressure, or elevated fasting glucose). Therefore, these participants were expected to have chronic low-grade inflammation with prolonged, increased oxidative and inflammatory stress in response to the HFCM challenge. We hypothesized that spice consumption in conjunction with the HFCM would reduce postprandial inflammation in this population with elevated inflammatory mediators.

Monocytes are the main inflammatory cell type that invade the arterial wall during the development of atherosclerosis, and their subsequent differentiation into macrophages and the formation of foam cells are implicated in all stages of atherosclerotic lesion formation (44). Postprandial hyperlipidemia after a high-fat challenge is accompanied by a transient increase in monocyte number with concomitant production of proinflammatory cytokines, which may contribute to endothelial dysfunction (45–47). Consistent with previous studies, we demonstrated a significant increase in the percentage of monocytes (CD14+/HLA-DR+) after HFCM consumption. Schildberger et al. (48) report that PBMCs and monocytes secrete comparable amounts of IL-1β, IL-6, IL-8, and TNF-α after LPS stimulation, suggesting that the monocyte subset in PBMCs is primarily responsible for the production of inflammatory cytokines. We, therefore, adjusted cytokine secretion from LPS-stimulated PBMCs by CD14+/HLA-DR+ monocyte number to determine cytokine secretion per monocyte.

Glucose was included as a covariate in the model based on data collected in previous studies, and the association observed between change in plasma glucose and cytokine secretion in our model. In a cross-sectional study including subjects aged 35–75 y (n = 5176) living in Lausanne, Switzerland, a positive correlation between plasma cytokines (IL-6 and TNF-α) and fasting plasma glucose was observed (49). In addition, proinflammatory cytokine secretion from LPS-stimulated PBMCs is significantly elevated after the pretreatment of PBMCs with glucose compared with no pretreatment (50), suggesting that glucose modulates cytokine secretion from LPS-stimulated PBMCs. In the current study, adding change in plasma glucose to the mixed-effect model significantly changed plasma cytokine concentration and cytokine secretion from LPS-stimulated PBMCs after consumption of the HFCM without spices, demonstrating that the variability in plasma glucose after consumption of the HFCM may be influencing cytokine response of the meal.

Our data on cytokine secretion from LPS-stimulated PBMCs after consumption of the HFCM suggest that the pattern of secretion in response to this stimulus may be altered by consumption of the HFCM containing the 6 g spice blend. IL-1β and IL-8 secretion at 180 min, and IL-6 and IL-8 secretion at 240 min after HFCM consumption containing the 6 g spice blend were significantly lower than the 2 g spice blend. Consumption of the 6 g spice blend significantly reduced IL-β secretion at 240 min after HFCM consumption compared with the 0 g spice blend. Lastly, the change in IL-6, IL-8, MCP-1, and TNF-α secretion from LPS-stimulated PBMCs at 240 min after HFCM consumption containing the 6 g spice blend, was below baseline values for each cytokine, suggesting that the HFCM-induced increase in inflammatory cytokine secretion may be blunted by the consumption of an HFCM containing 6 g spice blend.

Although a dose–response relation between phytochemical consumption and chronic disease risk has been reported (51), we only observed a dose-response effect of the spice blend on IL-1β secretion. It is possible that there is a dose threshold between 2 and 6 g spice blend below which the anti-inflammatory effect of spice consumption does not occur. However, additional studies are needed to test this hypothesis. In contrast to the change in cytokine secretion from LPS-stimulated PBMCs after spice consumption with the HFCM, we observed no effect of spice consumption on plasma cytokine concentrations. Herieka and Erridge (32) demonstrate that whereas elevated postprandial inflammatory markers are reliably detected in leukocytes (either extracellularly or intracellularly), they are not consistently raised in plasma. Our data support these findings, and suggest that LPS stimulation may be a more sensitive way to assess monocyte function because we are directly quantifying the cytokine response to LPS stimulation by monocytes in the PBMC compartment rather than measuring the accumulation of cytokines secreted from various cell sources in the circulation.

Obesity-induced, chronic, low-grade inflammation results from activation of several inflammatory signaling cascades, which includes the NF-κB pathway. NF-κB is a key mediator of inflammation because it regulates a large array of genes encoding proinflammatory cytokines (53). Accumulating evidence suggests that the bioactive metabolites of spices can prevent chronic inflammation by targeting the NF-κB signaling pathway (20). Turmeric, ginger, and cinnamon were the spices that comprised the highest proportion of the spice blend in our study. Numerous in vitro and in vivo animal and human studies have demonstrated anti-inflammatory effects of these spices. In randomized controlled trials, curcumin supplementation reduced serum IL-1β and IL-4 in adults with obesity (1 g/d for 8 wk) (33) and reduced serum IL-6, MCP-1, and TNF-α concentrations in subjects with metabolic syndrome (1 g/d for 4 wk) (34). In diabetic rats, curcumin reduced circulating IL-6, MCP-1, and TNF-α concentrations (100 mg · kg body weight−1 · d−1 for 7 wk) (54). In vitro studies have demonstrated that curcumin treatment suppresses IL-6 and TNF-α secretion after the inhibition of the NF-κB activity from LPS-stimulated RAW264.7 cells (55, 56) and LPS-stimulated THP-1 cells (57). Ginger supplementation reduced CRP in subjects with type 2 diabetes (1.6–3 g/d for 12 wk) (35, 36) and nonalcoholic fatty liver disease (2 g/d for 12 wk) (37). In mice with LPS-induced acute systemic inflammation, supplementation of ginger reduced serum IFN-γ and IL-6 (100–1000 mg · kg body weight−1 · d−1 for 3 d) (58). The mechanism underlying the anti-inflammatory effect of ginger is likely the suppression of NF-κB activity by the bioactive 6-gingerol. 6-Gingerol inhibited the LPS-stimulated NF-κB activity in murine peritoneal macrophages, resulting in decreased IL-1β secretion from the macrophages (59). Similarly, oral administration of cinnamon extract significantly reduced serum TNF-α after LPS stimulation in BALB/c mice (20–500 mg · kg body weight−1 · d−1 for 6 d) (60). Preincubation of THP-1 human monocytes with cinnamon extract significantly
reduced LPS-stimulated IL-8 secretion (61). In vitro studies have reported that cinnamaldehyde inhibits NF-κB activity in RAW264.7 murine macrophages (40, 62, 63), suggesting the anti-inflammatory property of cinnamon may be due to cinnamaldehyde, a major bioactive compound from cinnamon bark. Moreover, other spices and their bioactive compounds, such as basil (e Estragole) (64, 65), black pepper (piperine) (66), coriander (linalool) (67), cumin (cuminaldehyde) (68), red pepper (capsaicin) (69, 70), rosemary (rosmarinic acid) (71), and thyme (thymol) (72), may also have anti-inflammatory effects. Overall, based on previous studies, the reduction in IL-1β, IL-6, and IL-8 secretion from LPS-stimulated PBMCs after the meal challenge in the current study may be mediated via spice-induced inhibition of NF-κB activity, which should be explored in future studies.

A major strength of our study is the study design (3-period, crossover, randomized controlled trial) because each participant served as his own control. Another strength of the study is that we quantified the LPS-stimulated cytokine production from PBMCs, which enabled us to evaluate monocyte function in subjects after a meal with and without the addition of the spice blend. Limitations of this study are the small sample size (n = 12) and homogeneity of the participants. In addition, because this was not a double-blinded study (i.e., participants could differentiate between spiced and nonspiced meals), a potential bias may exist because subjects were aware of their treatment group. However, the outcomes we evaluated in this study are objective markers (i.e., inflammatory cytokine concentrations) measured in blood or in culture supernatants, thus the potential for bias may be small. In addition, the inflammatory mediators evaluated in this study are secondary outcomes; therefore, adjustment for multiple outcomes is not necessary for these data. However, because analysis of multiple outcomes for the same study inflates the false positive rate, the results should be viewed with caution. To ensure the findings from the current study are broadly applicable, the results from the current study need to be replicated in a larger, more diverse population. Moreover, it would be interesting to explore the potential mechanisms underlying the anti-inflammatory effect of spices, such as nuclear translocation and activity of NF-κB in PBMCs, in future studies.

In summary, we demonstrated that the presence of the 6 g spice blend in a HFCM significantly reduced postprandial IL-1β, IL-6, and IL-8 secretion from LPS-stimulated PBMCs in men with overweight/obesity at risk of CVD, suggesting a potential anti-inflammatory effect of spices. These findings are the first to our knowledge to demonstrate that consumption of a HFCM containing a spice blend could modulate postprandial cytokine secretion from culture PBMCs, suggesting the incorporation of spices into daily diet may help reduce postprandial inflammation and concurrently attenuate chronic low-grade inflammation.

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References
1. Fryar CD, Carroll MD, Ogden CL. Prevalence of overweight, obesity, and severe obesity among adults aged 20 and over: United States, 1960–1962 through 2015–2016 [Internet]. NCHS Health E-Stats. Hyattsville (MD): National Center for Health Statistics; 2018. Available from: https://www.cdc.gov/nchs/data/hestat/obesity_adult_15_16/obes_ity_adult_15_16.htm. [Accessed June 2019].
2. Ndueme CE, Matsuhashi K, Lazo M, Bello N, Blumenthal RS, Gerstenblith G, Nambi V, Ballantyne CM, Solomon SD, Selvin E. Obesity and subtype of incident cardiovascular disease. J Am Heart Assoc 2016;5:e003921.
3. León-Pedroza JI, González-Tapia LA, del Olmo-Gil E, Castellanos-Rodriguez D, Escobedo G, González-Chávez A. Low-grade systemic inflammation and the development of metabolic diseases: from the molecular level to the clinical practice. Cirr Cir 2015;83: 543–51.
4. Micha R, Peñalvo JL, Cudhea F, Imamura F, Rahmd CD, Mozaffarian D. Association between dietary factors and mortality from heart disease, stroke, and type 2 diabetes in the United States. JAMA 2017;317: 912–24.
5. O’Keefe JH, Bell DS. Postprandial hyperglycemia/hyperlipidemia (postprandial dysmetabolism) is a cardiovascular risk factor. Am J Cardiol 2007;100:899–904.
6. Bell D, O’Keefe J, Jellinger P. Postprandial dysmetabolism: the missing link between diabetes and cardiovascular events? Endocr Pract 2008;14:112–24.
7. Munoz A, Costa M. Nutritionally mediated oxidative stress and inflammation. Oxid Med Cell Longev 2013;6:10950.
8. Emerson SR, Korti SP, Harms CA, Haub MD, Mgarego T, Logan C, Rosenkranz SK. Magnitude and timing of the postprandial inflammatory response to a high-fat meal in healthy adults: a systematic review. Adv Nutr 2017;8:213–25.
9. Williams JW, Huang IH, Randolph GJ. Cytokine circuits in cardiovascular disease. Immunity 2019;50:941–54.
10. Niu J, Kolatudaty PE. Role of MCP-1 in cardiovascular disease: molecular mechanisms and clinical implications. Clin Sci 2009;117:95–109.
11. Fontes JA, Rose NR, Cháiková D. The varying faces of IL-6: from cardiac protection to cardiac failure. Cytokine 2015;74:62–8.
12. Apostolakis S, Vagiazis K, Amanatidou V, Spandidos DA. Interleukin 8 and cardiovascular disease. Cardiovasc Res 2009;84:333–60.
13. Reiss AB, Siegart NM, De Leon J, Interleukin-6 in atherosclerosis: atherogenic or atheroprotective? Clin Lipidol 2017;12:14–23.
14. Simonetti A, Mosucci M, Muller D, Bates E, Pagani F, Burdick M, Strieter R. IL-8 is an angiogenic factor in human coronary atherectomy tissue. Circulation 2000;101:1519–26.
15. Blackburn P, Despres JP, Lamarche B, Tremblay A, Bergeron J, Lemieux I, Guillard C. Postprandial variations of plasma inflammatory markers in abdominally obese men. Obesity (Silver Spring) 2006;14:1747–54.
16. Esposito K, Ciotola M, Sasso FC, Cozzolino S, Saccornano F, Assaloni R, Cerielo A, Guigliano D. Effect of a single high-fat meal on endothelial function in patients with the metabolic syndrome: role of tumor necrosis factor-α. Nutr Metab Cardiovasc Dis 2007;17:274–9.
17. Patel C, Ghanim H, Rishavshankar S, Sia CL, Viswanathan P, Mohanty P, Dandona P. Prolonged reactive oxygen species generation and nuclear factor-kB activation after a high-fat, high-carbohydrate meal in the obese. J Clin Endocrinol Metab 2007;92:4476–9.
18. Manning PJ, Sutherland WH, McGrath MM, de Jong SA, Walker RJ, Williams MJ. Postprandial cytokine concentrations and meal composition in obese and lean women. Obesity 2008;16:2046–52.
19. Chiu C-H, Li T-L, Chang C-K, Wu C-L. The potential ways how are they linked? J Transl Med 2018;16:14.
20. Kunnumakkara AB, Song L, Banik K, Harsha C, Prasad S, Gupta SC, Chiu C-H, Li T-L, Chang C-K, Wu C-L. The potential ways how are they linked? J Transl Med 2018;16:14.
21. Vors C, Couillard C, Paradis M-E, Gigleux I, Marin J, Vohl M-C, Couture P, Lamarche B. Supplementation with resveratrol and curcumin does not affect the inflammatory response to a high-fat meal in healthy adults: a systematic review. Adv Nutr 2017;8:213–25.
22. Williams JW, Huang IH, Randolph GJ. Cytokine circuits in cardiovascular disease. Immunity 2019;50:941–54.
23. Niu J, Kolatudaty PE. Role of MCP-1 in cardiovascular disease: molecular mechanisms and clinical implications. Clin Sci 2009;117:95–109.
24. Fontes JA, Rose NR, Cháiková D. The varying faces of IL-6: from cardiac protection to cardiac failure. Cytokine 2015;74:62–8.
25. Apostolakis S, Vagiazis K, Amanatidou V, Spandidos DA. Interleukin 8 and cardiovascular disease. Cardiovasc Res 2009;84:333–60.
26. Reiss AB, Siegart NM, De Leon J, Interleukin-6 in atherosclerosis: atherogenic or atheroprotective? Clin Lipidol 2017;12:14–23.
27. Simonetti A, Mosucci M, Muller D, Bates E, Pagani F, Burdick M, Strieter R. IL-8 is an angiogenic factor in human coronary atherectomy tissue. Circulation 2000;101:1519–26.
28. Blackburn P, Despres JP, Lamarche B, Tremblay A, Bergeron J, Lemieux I, Guillard C. Postprandial variations of plasma inflammatory markers in abdominally obese men. Obesity (Silver Spring) 2006;14:1747–54.
29. Esposito K, Ciotola M, Sasso FC, Cozzolino S, Saccornano F, Assaloni R, Cerielo A, Guigliano D. Effect of a single high-fat meal on endothelial function in patients with the metabolic syndrome: role of tumor necrosis factor-α. Nutr Metab Cardiovasc Dis 2007;17:274–9.
30. Patel C, Ghanim H, Rishavshankar S, Sia CL, Viswanathan P, Mohanty P, Dandona P. Prolonged reactive oxygen species generation and nuclear factor-kB activation after a high-fat, high-carbohydrate meal in the obese. J Clin Endocrinol Metab 2007;92:4476–9.
22. Ono-Moore KD, Snodgrass RG, Huang S, Singh S, Freytag TL, Burnett DJ, Bonnel EL, Woodhouse LR, Zunino SJ, Peerson JM. Postprandial inflammatory responses and free fatty acids in plasma of adults who consumed a moderately high-fat breakfast with and without blueberry powder in a randomized placebo-controlled trial. J Nutr 2016;146:1411–9.

23. Sardo CL, Kitzmiller JP, Apelsof G, Harris RB, Roe DJ, Stoner GD, Jacobs ET. An open-label randomized crossover trial of lyophilized black raspberries on postprandial inflammation in older overweight males: a pilot study. Am J Ther 2016;23:e86–91.

24. Nguyen L, Duong LT, Mentreddy RS. The U.S. import demand for spices and herbs by differentiated sources. J Appl Res Med Aromat Plants 2019;12:13–20.

25. Srinivasan K. Dietary spices as beneficial modulators of lipid profile in conditions of metabolic disorders and diseases. Food Funct 2013;4:503–21.

26. Lal A, Kumar T, Murthy PB, Pilai KS. Hypolipidemic effect of Coriandrum sativum L. in triton-induced hyperlipidemic rats. Indian J Exp Biol 2004;42:909–12.

27. Khan A, Zaman G, Anderson RA. Bay leaves improve glucose and lipid profile of people with type 2 diabetes. J Clin Biochem Nutr 2009;44:52–6.

28. Zare R, Heshmati F, Fallahzadeh H, Nadjarzadeh A. Effect of cumin powder on body composition and lipid profile in overweight and obese women. Complement Ther Clin Pract 2014;20:297–301.

29. Ishit J, Kanjilal S, Khubchandani J. Use of ethnic spices by adults in the United States: an exploratory study. Health Promot Perspect 2018;3:33–40.

30. Meng H, Ba Z, Lee Y, Peng J, Lin J, Fleming JA, Furumoto EJ, Roberts RF, Kris-Etherton PM, Rogers CJ. Consumption of Bifidobacterium animalis subsp. lactis BB-12 in yogurt reduced expression of TLR-2 on peripheral blood-derived monocytes and pro-inflammatory cytokine secretion in young adults. Eur J Nutr 2017;56:649–61.

31. Chan MM-Y, Huang HJ, Fenton MR, Fong D. In vivo inhibition of nitric oxide synthase gene expression by curcumin, a cancer preventive natural product with anti-inflammatory properties. Biochem Pharmacol 1998;55:1955–62.

32. Chan MM-Y. Inhibition of tumor necrosis factor by curcumin, a phytochemical. Biochem Pharmacol 1995;49:1551–6.

33. Ganjali S, Sahebkar A, Mahdipour E, Jamialahmadi K, Torabi M. Antioxidant effects of curcumin on serum cytokines in obese individuals: a randomized controlled trial. ScientificWorldJournal 2014;89:361.

34. Panahi Y, Hosseini MS, Khalili N, Naimi E, Simental-Mendia LE, Maihead M, Sahebkar A. Effects of curcumin on serum cytokine concentrations in subjects with metabolic syndrome: a post-hoc analysis of a randomized controlled trial. Biomed Pharmacother 2016;82:578–82.

35. Abu-Elfoul A, Sayed H, Al-Abdul H, El-Salhy M, El-Etery A, El-Omari A. Effect of curcumin on glycemic status, lipid profile and some inflammatory markers in patients with type 2 diabetes mellitus. Int J Food Sci Nutr 2014;65:515–20.

36. Shidfar F, Rajab A, Rahideh T, Khandouzi N, Hosseini S, Shidfar S. The effect of ginger (Zingiber officinale) on glyceric markers in patients with type 2 diabetes. J Complement Integr Med 2015;12:165–70.

37. Rahimeh M, Yari Z, Hekmatdoost A, Alavian SM, Keshavarz S. Inhibitory effect of curcumin and ginger on monocyte cytokine expression in normolipemic men. Clin Appl Thromb Hemost 2002;8:1477–55.

38. Khan IM, Pokharel YD, Dadu RT, Lewis DE, Hoovenge RC, Wu H, Ballantyne CM. Postprandial monocyte activation in individuals with metabolic syndrome. J Clin Endocrinol Metab 2016;101:4195–204.

39. Schildberger A, Rossmanith E, Eichhorn T, Strassl K, Weber V. Monocytes, peripheral blood mononuclear cells, and THP-1 cells exhibit different cytokine expression patterns following stimulation with lipopolysaccharide. Mediators Inflamm 2013;2013:697792.

40. Marques-Vidal P, Bastardot F, von Känel R, Paccaud F, Preisig M, Waeber G, Vollenweider F. Association between circulating cytokine levels, diabetes and insulin resistance in a population-based sample (CoLaus study). Clin Endocrinol (Oxf) 2013;78:322–41.

41. Wu HP, Chen CH, Hsieh HC, Liu YC. Effects of insulin and glucose on cytokine production from peripheral blood mononuclear cells. Chang Gung Med J 2008;31:253–9.

42. Holst B, Williamson G. Nutrients and phytochemicals: from bioavailability to bioefficacy beyond antioxidants. Curr Opin Biotechnol 2008;19:73–82.

43. Herieka M, Erridge C. High-fat meal induced postprandial inflammatory. Mol Nutr Food Res 2014;58:136–46.

44. Cattyssse L, van Loo G. Inflammation and the metabolic syndrome: the tissue-specific functions of NF-κB. Trends Cell Biol 2017;27:417–29.

45. Jain SK, Rains J, Croad J, Larson B, Jones K. Currinucin suppression lowers TNF-α, IL-6, IL-8, and MCP-1 secretion in high glucose-treated cultured monocytes and blood levels of TNF-α, IL-6, MCP-1, glucose, and glycosylated hemoglobin in diabetic rats. Antioxid Redox Signal 2009;11:241–9.

46. Kim S-J. Curcumin suppresses the production of interleukin-6 in Prevotella intermedia lipopolysaccharide-activated RAW 264.7 cells. J Periodontal Implant Sci 2011;41:157–63.

47. Zhao F, Gong Y, Hu Y, Lu M, Wang J, Dong J, Chen D, Chen L, Fu F, Qiu F. Curcumin and its major metabolites inhibit the inflammatory response induced by lipopolysaccharide: translocation of nuclear factor-κB as potential target. Mol Med Rep 2015;11:3087–93.

48. Zhou Y, Zhang T, Wang X, Wei X, Chen Y, Guo L, Zhang J, Wang C. Currinucin modulates macrophage polarization through the inhibition of the toll-like receptor 4 expression and its signaling pathways. Cell Physiol Biochem 2015;36:631–41.

49. Choi YY, Kim MH, Hong J, Kim S-H, Yang WM. Dried ginger (Zingiber officinale) inhibits inflammation in a lipopolysaccharide-induced mouse model. Evid Based Complement Alternat Med 2013;914563.

50. Tripathi S, Maier KG, Bruch D, Kittur DS. Effect of 6-gingerol on pro-inflammatory cytokine production and costimulatory molecule expression in murine peritoneal macrophages. J Surg Res 2007;138:209–13.

51. Hong J-W, Yang G-E, Kim YB, Eom SH, Lew J-H, Kang H. Anti-inflammatory activity of cinnamon water extract in vivo and in vitro LPS-induced models. BMC Complement Altern Med 2012;12:237.

52. Shink A, Naumoska K, Kitanovski K, Kampf CJ, Frohlich-Nowoisky J, Thines E, Poschl U, Schuppan D, Lucas K. Anti-inflammatory effects of cinnamon extract and identification of active compounds influencing the TLR2 and TLR4 signaling pathways. Food Funct 2018;9:5950–64.

53. Youn HS, Lee JK, Choi YJ, Saitoh SI, Miyake K, Hwang DH, Lee JY. Cinnamaldehyde suppresses toll-like receptor 4 activation mediated through the inhibition of receptor oligomerization. Biochem Pharmacol 2008;75:494–502.
63. Kim BH, Lee YG, Lee J, Lee JY, Cho JY. Regulatory effect of cinnamaldehyde on monocyte/macrophage-mediated inflammatory responses. Mediators Inflamm 2010;529359.
64. Silva-Comar FM, Würzler LAM, Silva-Filho SE, Kummer R, Pedroso RB, Spironello RA, Silva EL, Bersani-Amado CA, Cuman RK. Effect of estragole on leukocyte behavior and phagocytic activity of macrophages. Evid Based Complement Alternat Med 2014;784689.
65. Rodrigues LB, Oliveira Brito Pereira Bezerra Martins A, Cesário FR, Ferreira e Castro F, de Albuquerque TR, Martins Fernandes MN, Fernandes da Silva BA, Quintans Júnior LJ, da Costa JG, Melo Coutinho HD, et al. Anti-inflammatory and antiedematogenic activity of the Ocimum basilicum essential oil and its main compound estragole: in vivo mouse models. Chem Biol Interact 2016;257:14–25.
66. Ying X, Yu K, Chen X, Chen H, Hong J, Cheng S, Peng L. Piperine inhibits LPS induced expression of inflammatory mediators in RAW 264.7 cells. Cell Immunol 2013;285:49–54.
67. Huo M, Cui X, Xue J, Chi G, Gao R, Deng X, Guan S, Wei J, Soromou LW, Feng H, et al. Anti-inflammatory effects of linalool in RAW 264.7 macrophages and lipopolysaccharide-induced lung injury model. J Surg Res 2013;180:e47–54.
68. Wei J, Zhang X, Bi Y, Miao R, Zhang Z, Su H. Anti-inflammatory effects of cumin essential oil by blocking JNK, ERK, and NF-κB signaling pathways in LPS-stimulated RAW 264.7 cells. Evid Based Complement Alternat Med 2015;474509.
69. Vasanthkumar T, Hanumanthappa M, Lakshminarayana R. Curcumin and capsicum modulates LPS induced expression of COX-2, IL-6 and TGF-β in human peripheral blood mononuclear cells. Cytotechnology 2019;71:963–76.
70. Choi SE, Kim TH, Yi SA, Hwang YC, Hwang WS, Choe SJ, Han SJ, Kim HJ, Kim DJ, Kang Y, et al. Capsaicin attenuates palmitate-induced expression of macrophage inflammatory protein 1 and interleukin β by increasing palmitate oxidation and reducing c-Jun activation in THP-1 (human acute monocytic leukemia cell) cells. Nutr Res 2011;31:468–78.
71. Rocha J, Eduardo-Figueira M, Barateiro A, Fernandes A, Brites D, Bronze R, Duarte CM, Serra AT, Pinto R, Freitas M, et al. Anti-inflammatory effect of rosmarinic acid and an extract of Rosmarinus officinalis in rat models of local and systemic inflammation. Basic Clin Pharmacol Toxicol 2015;116:398–413.
72. Gholijani N, Gharagozloo M, Farajdian S, Amirghofran Z. Modulatory effects of thymol and carvacrol on inflammatory transcription factors in lipopolysaccharide-treated macrophages. J Immunotoxicol 2016;13:157–64.