l Revieu

The Potential Health Benefits of Noni Juice:

3 A Review of Human Intervention Studies

- 4 Brett J. West 1,*, Shixin Deng 1, Fumiyuki Isami 2, Akemi Uwaya 2 and C. Jarakae Jensen 1
 - ¹ Research and Development, Morinda, Inc. 737 East 1180 South, American Fork, Utah 84003, USA
- 6 Research and Development, Morinda, Inc. 3-2-2 Nishishinjuku, Shinjuku-ku, Tokyo 160–0023, Japan.
 - * Correspondence: brett_west@morinda.com

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

5

2

Abstract: Noni juice is a globally popular health beverage that originates from the tropics. Traditional Tahitian healers believe the noni plant to be useful for a wide range of maladies, and noni juice consumers throughout the world have similar perceptions. Nevertheless, human clinical trials are necessary for an understanding of what the health benefits of noni juice truly are. A review of published human intervention studies suggest that noni juice may provide protection against tobacco smoke-induced DNA damage, blood lipid and homocysteine elevation, and systemic inflammation. Human interventions studies also suggest that noni juice may improve joint health, increase physical endurance, increase immune activity, inhibit glycation of proteins, aid weight management, help maintain bone health in women, help maintain normal blood pressure, and improve gum health. Further, these studies point to noni juice possessing notable antioxidant activity, more so than the other fruit juices that served as placebos. It is this antioxidant effect, and its interaction with the immune system and inflammation pathways, that may account for many of the observed health benefits of noni juice. However, the existing evidence does have some limitations in its application to noni juice products in general as all the peer-reviewed human interventions studies to-date have involved only one source of French Polynesian noni juice. Geographical factors and variations in processing methods are known to produce commercial noni juice products with divergent phytochemical and nutrient compositions. Therefore, other sources of noni products may have different toxicological and pharmacological profiles.

Keywords: Noni juice; Morinda citrifolia; clinical trial; antioxidant; immune system; inflammation

2728

29

30

31

32

33

34

35

36

37

38

39

40

41

42

43

44

45

1. Introduction

Noni juice has become one of the most popular health supplements. In the first dozen years of its commercial marketing, more than 106 million liters of noni juice were consumed in more than 80 nations [1]. Noni juice was one of the first whole foods approved under the European Union's 1997 novel food regulations [2]. The Chinese government has also approved one source of noni juice as a safe new resource and approved it as a functional food that can enhance immunity [3].

Noni is the common name for *Morinda citrifolia* and is a small to medium sized tree (3–10m high) that is widely distributed across the tropics [4]. Noni fruit and leaves have a history of food use among Pacific Islanders and in Southern and Southeast Asia. In 1769 Sydney Parkinson, a naturalist sailing with Captain James Cook, recorded that Tahitians ate noni fruit [5]. This was likely the first written description of its use as a food. Since then, others have recorded similar observations. Rarotongans ate the fruit often, and the Burmese used it to prepare curries [6, 7]. Australian Aborigines ate noni fruit during the cool-dry season from May to August in the Northern Territory of Australia [8, 9]. Noni leaves were eaten either raw or cooked in Java and Thailand [10]. In Tahiti, fish were wrapped in the leaves for baking to impart an appealing flavor to the cooked fish [11].

More recently, the fruit is used to produce a juice that is often consumed to improve overall health. A major source of this juice is French Polynesia where noni fruit puree constitutes one of the

largest agricultural exports [12]. In fact, more than 21,000 metric tons of this puree was exported in the past decade [13]. The noni plant was the most important and widely used Polynesian medicinal plant prior to the arrival of Europeans, with Tahitian healers using it in many remedies [14]. Some traditional uses of the fruit include the treatment of inflammation, abscesses, angina, diabetes, ranula, abdominal fibromas and scorpionfish stings [15, 16]. In other parts of French Polynesia, noni fruit is reportedly used to treat osteoarthritis, rheumatism, backache, joint problems, hemorrhoids, skin allergies, burns, and warts [17]. The global popularity of the juice is due to its perceived health value. Indeed, European consumers of a Tahitian sourced noni juice beverage reported that increased energy, improved well-being, reduction of pain, fewer infections, improved sleep, improved digestion, reduction in allergy and asthma symptoms were the most frequently experienced health benefits [18].

There is a large body of published research on the potential health effects of the noni plant. But much of this involves in vitro and animal studies [19, 20]. While the findings of these types of studies are useful in explaining possible mechanisms of action, they may be of limited application to humans due to interspecies differences in absorption, distribution, metabolism, and excretion. Therefore, human studies are necessary to determine what effects noni juice may really have on human health. These studies should also provide information on phytochemical composition since there are differences in noni fruit from various regions, as well as disparities in the ingredients and processing methods used to produce various commercial noni juice products [21, 22]. Accordingly, this review summarizes published human intervention studies involving noni fruit juice and discusses mechanisms of action that may explain the health benefits described in ethnobotanical studies. Also considered are the limitations on the applicability of these human studies to noni juice products in general.

2. Clinical safety study

A 28-day clinical trial with 96 healthy adult volunteers was conducted to evaluate the safety of daily ingestion of noni juice from French Polynesia [23]. The volunteers were divided equally into four dose groups; 0 mL (placebo), 30 mL, 300 mL, or 750 mL. Comprehensive hematology, biochemistry, and urinalysis tests were completed with each volunteer at the start of the trial, as well as at week 2, week 4 (end of noni juice ingestion) and during a two-week follow up (week 6). Electrocardiography tests were performed with each subject during a pre-study screening and at week 6. Hematological measurements included hemoglobin, hematocrit, mean cell volume, red cell count, prothrombin time, activated partial thrombin time, total and differential white cell count (basophils, eosinophils, lymphocytes, monocytes, and neutrophils), and platelet count. Biochemistry analysis included alkaline phosphatase, alanine aminotransferase, aspartate aminotransferase, total bilirubin, lipids (LDL, HDL, cholesterol, triglycerides), creatine kinase, creatinine, gamma-glutamyl transferase, glucose, total protein, and uric acid. Urinalysis involved semi-quantitative analysis for leucocytes, nitrite, urobilinogen, protein, blood, ketones, bilirubin, glucose, pH and specific gravity. Blood pressure, heart rate, weight, and adverse events were also recorded also at baseline and weeks 2, 4, and 6.

Test results revealed that there were no adverse dose-related effects from noni juice ingestion. During the entire study, no clinically significant differences were observed in any of the measurements between the placebo and noni juice groups, apart from there being 20 to 50% fewer total adverse events (i.e., headache and other body aches) in the noni juice groups than in the placebo group. The lower adverse event rates in the noni juice groups suggest possible health benefits from noni. The results of this study indicate that drinking up to 750 mL noni juice each day is safe.

3. Antioxidant and DNA protection

A series of double-blind, placebo-controlled, 30-day intervention studies involving cigarette smokers has demonstrated the substantial antioxidant activity of noni juice. The first of these involved 285 adult volunteers who smoked more than 20 cigarettes per day [24]. The participants were assigned to either a placebo, 29.5 mL noni juice/day, or a 118 mL noni juice/day group. The noni

juice evaluated in these studies was a mixture of noni fruit juice from puree (from French Polynesia), grape, and blueberry juice. The placebo was composed of grape and blueberry juices, with a cheese flavor added to mimic the noni flavor. Blood plasma levels of superoxide anion radicals (SAR) and lipid hydroperoxide (LOOH) levels were measured at the beginning and end of the trial, with participants continuing to smoke throughout the trial. Noni juice ingestion reduced average plasma SAR by 26.9 % and 30.8 % in the 29.5 and 118 mL groups, respectively. Average LOOH levels in the 29.5 mL group were reduced by 24.5 %, while the 118 mL group experienced a 27.3 % reduction. No significant reductions in SAR or LOOH levels occurred in the placebo group. Despite the placebo containing high antioxidant fruit juices, the inhalation of large quantities of tobacco smoke resulted in more oxidative stress than could be overcome with grape and blueberry juices alone. Therefore, the SAR and LOOH reduction was due to the antioxidant properties, whether direct or indirect, of noni juice.

The second smoker study was completed with 203 subjects [25]. The study intervention was the same as that mentioned above. In this study, however, peripheral blood lymphocytes (PBLs) were isolated from pre-and post-trial whole blood samples and evaluated for the degree of aromatic DNA adduct formation using a 32 P-postlabeling assay. By the end of the 30-day study period, average aromatic DNA adduct levels dropped by 44.9 % (P < 0.001) among those drinking either 1 fluid ounce (29.5 mL) or 4 fluid ounces (118 mL) of noni juice per day. There were no significant differences in gender-specific responses. But at the lower dose, males experienced a greater decrease in aromatic DNA adducts than females (56.1 % vs. 43.1% reduction, respectively). No adverse effects were observed in this trial.

Another randomized, double-blind, placebo-controlled study was completed with 245 heavy cigarette smokers [26]. This trial measured changes to lipid peroxidation—derived DNA adducts after 30 days of noni juice ingestion. Participants in this study were assigned to the same dose groups, with the same placebo and noni juice product, that were used in the antioxidant trial. Again, a ³²P-postlabeling assay was used, following isolation of DNA from PBLs. The previous DNA protection study measured aromatic DNA adducts, which are more likely to be formed by direct reactions with chemicals in cigarette smoke or their immediate metabolites. In this second study, measurements were made specifically of DNA adducts resulting from cigarette smoke-induced oxidative stress and consequent lipid peroxidation. At the completion of the trial period, the placebo group did not experience any reduction in DNA damage. However, the lipid peroxidation—derived DNA adduct levels in those who drank noni juice declined significantly by 46.9% to 57.4%. This effect is consistent with both previous studies, where reactive oxygen species and aromatic DNA-adduct concentrations were significantly reduced.

4. Blood lipid normalization, high sensitivity C-reactive protein (hs-CRP) and homocysteine reduction

As with the antioxidant and DNA adduct studies, adult smokers provided further insight into the effect of 30 days of noni juice ingestion on blood lipids and hs- CRP. In a randomized, double blind, placebo-controlled clinical trial with 132 volunteers [27]. Heavy smokers (≥ 20 cigarettes/day) were chosen as subjects because smoking is reported to increase blood lipids, systemic inflammation and serum homocysteine [28-30]. Pre- and post-study serum cholesterol, triglyceride, low density lipoprotein cholesterol (LDL), high density lipoprotein cholesterol (HDL), hs-CRP, and homocysteine were measured for all participants, who continued smoking during the trial. In this study, the noni juice product and dose groups were the same as in the previously discussed smoker studies.

After drinking noni juice for 30 days, the placebo group had no significant changes in any measurements. But among those who drank noni juice, there were significant declines in average hs-CRP (15.2%) and homocysteine (23.9 %), with an increase in HDL (from 49 to 57 mg/dL). Mean total cholesterol, LDL, and triglycerides were also decreased among those in the noni juice groups. However, the degree of change was dependent on initial (pre-trial) values. Larger declines in mean values were associated with greater initial total cholesterol, LDL or triglycerides levels. For example, decreases in average total cholesterol in the low (190–219 mg/dL), middle (220–299 mg/dL), and high

(>300 mg/dL) baseline strata of the 29.5mL group were 12.1, 17.3, and 36.4%, respectively. No participants in either noni juice group experienced any changes that resulted in individual blood lipid values being under normal reference values. As such, the changes experienced were towards, or remained within, normal healthy ranges. These results indicate that noni juice helped normalized lipid levels in heavy smokers.

A second, although smaller, study examined the cholesterol lowering activity of noni juice in nonsmokers [31]. This open-label pilot study involved older adults (> 40 years) who had normal or borderline high cholesterol levels and were not taking cholesterol-lowering medication nor drinking alcohol. Each participant drank two fluid ounces (59 mL) of noni juice twice per day for 30 days. No other changes to lifestyle habits were made during the trial period. There was no significant difference between pre- and post-trial total cholesterol, HDL, and LDL levels, as well as TC/HDL ratio. These results suggest that the cholesterol modulating effect of noni juice in heavy smokers is likely from its ability to protect against cigarette smoke induced dyslipidemia, via antioxidant activity.

5. Improvement of joint pain and mobility

Treatment of pain and inflammation is one of the most common uses of the noni plant in tropical folk medicine, an effect that appears to be substantiated by numerous in vivo and in vitro studies [32-34]. In fact, one of the traditional names for noni in the Caribbean was "pain killer" [35]. Among the published anti-nociceptive and anti-inflammatory studies of noni juice are two open-label clinical trials that demonstrate potential joint health benefits. The earlier of these two trials reported pain reduction and improved range of motion (flexion, extension, lateral flexion, and rotation) in patients suffering from cervical spondylosis after four weeks of ingesting 15 mL noni juice every morning and evening [36]. This trial enrolled 90 patients who were assigned to one of three treatment groups: standard physiotherapy alone, noni juice alone, and combined treatment (physiotherapy plus noni juice). Pre-and post-treatment measurements of pain intensity and neck flexibility (cervical range of motion) were compared within and among the treatment groups. At the start of the trial, all subjects in the noni juice alone group fell within the 5 – 7 (moderate to severe) pain intensity range. By the end of the study, the pain intensity range of this group decreased to 0-4 (none to very moderate), with complete relief of neck pain in 60 percent of patients. The physiotherapy alone and combined treatment groups also experienced significant reduction in pain symptoms, with the combined treatment group experiencing the greatest reduction in pain intensity. Range of motion improved among all three treatment groups by the end of the trial period. For example, mean lateral flexion and rotation approximately doubled in the noni alone and physiotherapy alone groups. Although improvements in the noni alone group were no different than those of the standard physiotherapy alone group, a significantly greater improvement occurred in the combined treatment group.

A second joint health trial of noni juice involved osteoarthritis patients [37]. In this open label intervention study, 82 volunteers drank 3 fluid ounces (88.5 mL) of noni juice (1 ounce before breakfast, 1 ounce before lunch, and 1 ounce before bedtime) every day. Those enrolled in the study were adults (40 - 75 years), had an X-ray diagnosis of osteoarthritis of the hip or knee, were not taking prescription medication for arthritis, and were willing to consume noni juice for 90 days. Blood samples were collected from the participants for clinical laboratory analysis at enrollment and after the 90-day intervention period. The Arthritis Impact Measurement Scales (AIMS2) were used to measure pre- and post-study pain/discomfort levels. The Short Form-36, version 2 (SF-36 V2) was used to measure pre- and post-study patient quality of life.

By the end of the intervention period, significant improvements in mean quality of life measurements occurred. These included a reduction in the duration of arthritis pain, including a 23.7% decrease in the frequency (in days) of severe pain, and 16.4% decrease in pain severity. Patients also experienced an improved psychological state and mood, and improved mobility. Patient satisfaction with personal health also increased by approximately 19%. As with previous studies, noni juice was well tolerated and appeared to be safe. No significant changes to liver or kidney

functions occurred after three months of noni juice ingestion, nor were there changes to blood glucose, total cholesterol, or triglyceride levels.

A double-blind placebo-controlled trial, involving female university students suffering from dysmenorrhea, was completed to evaluate the efficacy of a capsule containing milled "noni herb powder" [38]. Over the course of three menstrual cycles, noni capsule ingestion did not improve menstrual pain when compared to the placebo. However, there was some evidence of a significant decline in mean erythrocyte sedimentation rate (ESR) among those taking the noni capsule. This indicated some degree of anti-inflammatory activity, although not enough to affect pain symptoms. While this study did not involve noni fruit juice, it is discussed because it highlights issues regarding noni product identity and variation in potential efficacy. The product used in this study was identified as a "noni herb powder" with no indication as to the geographic origin, plant part used, or harvesting and processing conditions. As will be discussed in more detail later, there is wide variability among nutrient and phytochemical compositions of commercial noni products [39]. These differences preclude assumptions that results from clinical trials of a specific noni product are applicable to other noni products. The two previously discussed trials involved the French Polynesian derived noni juice blend used in the smoker studies. Whereas, this latter trial evaluated a noni herb powder of relatively unknown identity. Therefore, the lack of significant pain reduction in the latter trial does not necessarily refute the pain reduction observations of the cervical spondylosis and osteoarthritis trials. It is interesting, however, that all three trials did provide some evidence of anti-inflammatory activity.

6. Increased physical endurance

Some Pacific Islanders believed that ingesting ripe noni fruit invigorated the body during long fishing trips and ocean voyages [40]. Three clinical trials have been completed which lend some credence to this traditional use. The first of these was a 21-day, placebo controlled intervention study involving 40 highly trained middle and long-distance runners [41]. For three weeks, these athletes drank 100 mL noni juice, twice per day, or a blackberry juice placebo. The athletes participated in pre- and post-intervention time-to-fatigue treadmill tests, during which the workload increased every minute. Pre-and post-intervention medical examinations (including electrocardiogram, heart rate, and blood pressure) and blood tests were also performed (including total protein, urea, glucose, hemoglobin, and lactate). The noni juice group had a 21% increase in average time-to-fatigue, with an accompanying 25% decrease in average blood chemiluminescence (a marker of lipid oxidation), whereas no changes occurred in the placebo group. An accompanying study did not reveal the presence of any substances banned for athletic competition (including beta-blockers, diuretics, narcotics, anabolic steroids, stimulants or masking agents). Therefore, it appears that the increase in physical endurance is associated with the antioxidant activity of noni juice.

A second similar trial was completed but with some modifications [42]. This trial involved 46 university athletes equally divided, along with 14 non-athlete controls, into a noni juice group and a blackberry juice group. For 30 days, these athletes drank 100 mL noni juice, twice per day, or blackberry juice. This trial also included pre- and post-intervention time-to-fatigue treadmill tests, with accompanying blood analyses. The noni juice group experienced a significant decrease in mean serum creatine kinase (CK) concentration, while no such decline occurred in the blackberry juice group. This finding suggests that the improvement in endurance is due to the ability of noni juice to mitigate exercise-induced muscle tissue damage. This protective effect likely involves the antioxidant properties of noni, as CK concentration increased along with oxidative stress markers in athletes undergoing intensive training [43]. Indeed, intense exercise causes increases in free radical production and inflammation [44], both of which are inhibited by noni juice.

Noni juice may also have an influence on oxygen uptake during physical exertion. Semiprofessional male cyclists were enrolled in a double blind, placebo-controlled, randomized study to evaluate the effect of noni fruit juice on aerobic fitness [45]. Volunteers drank either 120 mL of noni juice or a placebo every day for 14 days. At the beginning of the trial and after two weeks, volunteers completed exercise tests on a cycle ergometer (with 30 watts/min workload increase) to

- 248 the point of exhaustion, electrocardiographic alterations, or to the theoretical maximal heart rate.
- Oxygen uptake was measured by spirometry. Increases in the average maximal oxygen uptake, VO2
- 250 max, and in oxygen uptake at 50 Watt workload occurred in the noni juice group. The noni juice
- evaluated in this trial was the same source as those evaluated in the other anti-fatigue, antioxidant,
- and joint pain studies and did not contain any performance enhancing substances.

7. Increased immune activity

Smoking suppresses immune function [46]. Exposure to tobacco smoke is a risk factor for various bacterial and viral infections [47, 48]. A smoking habit can have a significant negative impact of peripheral blood lymphocyte subsets and their function [49, 50]. The ability of noni juice to protect lymphocyte DNA damage, as discussed above, may have a positive effect on the immune function of smokers. But noni juice may have an influence on immune function that goes beyond the protection of cells involved in the adaptive immune system. For instance, the feeding of French Polynesian noni fruit puree to newborn cattle (Holstein bull calves) for the first two weeks of life significantly increased whole blood phagocytic activity against E. coli and Staphylococcus epidermis [51]. A follow up study in Holstein bull calves revealed that noni puree supplementation for the first three weeks of life also reduced all required medical treatments by 54%, with a 61% reduction in respiratory treatments and a 52% reduction in gastrointestinal treatments [52]. The immune modulating activity of noni juice was also observed in mice, where the feeding of the French Polynesian derived noni juice blend for 15 days resulted in inhibition of mitogen-stimulated interleukin-4 production in splenocytes and peritoneal exudate cells [53]. This treatment also significantly increased interferongamma (IFN- γ) production. As IFN- γ is involved in macrophage activation, these results agree with the increase in phagocytic activity seen in newborn calves.

To verify the immune modulating potential of noni juice in humans, an intervention study was conducted with 12 healthy adult volunteers for 8 weeks, during which each consumed 330 mL of Tahitian Noni® Juice every day [54]. Vital signs, serum malondialdehyde (MDA) concentration, interleukin 2 (IL-2) concentration, and ex vivo natural killer-cell (NK) activity were measured before and after the intervention period. During the study, there was no change in vital signs. But there was a significant decrease in average MDA levels over the course of the trial, due to reduced oxidative stress. On the other hand, mean IL-2 concentration and NK activity both increased by approximately 30%. The findings of this pilot study reveal noni juice's potential for supporting immune function, along with concurrent antioxidant activity, in a healthy population.

It is important to note that the antioxidant properties of noni juice likely contribute to its immunomodulation activity, as there is documented connection between oxidative processes and the immune system. Reactive oxygen species (ROS) are involved in the modulation of immune function and serve as secondary messengers in cell signal transduction [55]. Multiple studies have demonstrated the ability of antioxidants, including those from fruits, to modulate immune responses [56-59]. Epidemiology studies have revealed associations between lower cancer incidence and diets rich in antioxidant nutrients, with antioxidant-induced improvement in immune function having been suggested as an important factor [60]. Antioxidant supplementation also improved immune function in the elderly and in other groups [61, 62].

8. Control of advanced glycation end products (AGEs) and glycosylated hemoglobin

Several in vitro studies indicate that noni fruit may be helpful in controlling the formation of advanced glycation end products (AGEs) and, subsequently, reduce their adverse effects. An extract from noni fruit collected in French Polynesia prevented AGE-induced reactive oxygen species generation in human umbilical vein endothelial cells [63]. This extract also inhibited the in vitro formation of glucose-human serum albumin, glucose-collagen, and glucose-keratin AGEs, as well exhibited AGE-crosslink breaking activity [64]. Another noni fruit extract inhibited the development of glucose-induced bovine eye lens opacity [65].

The antiglycation potential of noni fruit juice was further demonstrated in an 8-week open-label intervention study and in a cross-sectional population study [66]. Both studies were conducted within

the context of iridoid content, as iridoids are major phytochemicals in noni fruit and are well known for their anti-AGE biological activities [67, 68]. Both studies utilized skin autofluorescence as a marker for AGE accumulation in the body [69]. The 8-week intervention study measured changes in the skin autofluorescence of 34 adults who consumed, every day, a beverage composed of noni fruit puree (from French Polynesia), cornelian cherry juice, and olive leaf extract. Overweight or obese, prehypertensive or grade 1 hypertensive males and females, with impaired fasting glucose and who were not using prescription medication, were included in the trial. Previously published population reference values revealed that the average initial skin autofluorescence of this group was typical of healthy 44-year-old adults, even though their average actual age was 40. As such, their AGE associated age (ASA) was 4 years greater than their average actual age. But 8 weeks of noni juice supplementation reduced the ASA of this group to 39 years, demonstrating a significant antiglycative effect.

The cross-sectional population study included 3913 people from ten locations throughout Japan. During health education and promotion events, a questionnaire was used to collect demographic data and information on daily ingestion rates of noni juice beverages blended with either grape and blueberry juices, or with cornelian cherry juice and olive extract. Daily iridoid intake was calculated from the ingestion rates by applying the data obtained from chemical analyses of the beverages. As with the 8-week intervention study, skin autofluorescence was used as a marker for AGE burden in 2,790 noni juice consumers and 1,123 controls (those who did not drink noni juice beverages). Regression analysis revealed that noni juice intake is associated with lower AGE levels, with noni consumers having an average ASA that was 2.07 years less than the general population. In fact, for every mg of iridoids consumed, ASA decreased by 0.017 years. Among those who never smoked, the average ASA of noni juice consumers was 3.52 years less than the general population. The significant anti-AGE activity of noni juice is likely due, at least in part, to its ability to induce antioxidant enzyme activity, specifically superoxide dismutase and catalase [70].

Pacific Islanders reported that noni fruit was helpful in controlling blood sugar levels [71]. In vitro and animal studies have provided some supporting evidence for this traditional use [72-75]. Even so, data from human clinical trials is limited. A small, 8-week, human pilot study had been conducted with type-2 diabetes patients [76]. In this study, ingestion of 2 mL of noni juice per kg body weight, twice daily, was associated with improved blood glucose profiles and a significant reduction in glycosylated hemoglobin (HbA1c). It should be noted, however, that noni juice did not influence blood glucose levels in those without diabetes [23]. Therefore, the anti-diabetic property of noni juice appears to only involve the regulation of normal blood glucose level and is limited by feedback mechanisms, rather than involving the pharmacological pathways that are targets of conventional diabetes medications.

9. Weight management and mitigation of osteoporosis, hypertension, and gingivitis

Noni plant preparations were used in folk medicine to promote the loss of excess body weight [77]. Multiple animal studies have demonstrated the anti-obesity potential of noni juice [78]. Two clinical trials have further investigated this potential. The first of these was a 12-week, open-label trial of a weight-loss program that included noni juice, daily calorie restriction, and exercise interventions [79]. By the end of the trial, all participants experienced weight loss, with significant reductions in fat mass, percent body fat and body mass index. Greater weight loss occurred in individuals with an initial weight in the 75-111 kg range, with \geq 9 kg lost by each. Five of 22 participants moved from overweight to the normal weight category while another five switched from obese to overweight.

The second weight loss study included 90 grade 3 overweight (morbidly obese) adults who were divided into three treatment groups [80]. For 6 weeks, all participants followed a low calorie and low sodium diet. Also for the duration of the study, participants in two separate noni juice groups were assigned to drink one of two noni juice beverage blends (either with grape and blueberry juices or with cornelian cherry juice and olive leaf extract). Those in the control group did not consume any noni juice beverages. Muscle mass loss was significantly lower in those drinking noni juice beverages, as compared to the control group. Maintenance of weight loss throughout the study was greater in

those drinking noni juice than in the control group, as was reduction of waist circumference and body mass index. The preservation of active muscle cell mass seems to be a likely mechanism by which the improved weight loss was achieved by those in the noni juice groups. Protection of muscle tissue is also one mechanism responsible for improved endurance, as previously discussed.

Perhaps a somewhat expected property of noni juice is its potential influence on osteoporosis and conductive hearing loss. A three-month placebo-controlled pilot study was conducted with post-menopausal women wherein participants drank either 2 fluid ounces (59 mL) of noni juice or placebo twice per day [81]. Before and after the study, the women completed the Short Form 36 (SF-36) Quality of Life Survey and provided urine samples for deoxypyridinoline analysis (a marker for bone turnover), as well as received hearing tests. Those who drank noni juice had slightly greater average deoxypyridinoline crosslinks, indicating increased bone resorption. This finding is consistent with the observed attenuation of hearing loss that also occurred in the noni juice group, as the transmission of sound vibrations by the bones of the middle ear is influenced by the severity of osteoporosis. These positive effects on bone health also accompanied improvements in several quality of life scores, including mental health scores.

Two additional small human pilot studies suggest that noni juice may also have a positive impact on hypertension and gingivitis. In a one-month open label trial, hypertensive adults drank two fluid ounces (59 mL) of noni juice twice per day [82]. Pre-and post-trial diastolic and systolic blood pressure measurements were made in triplicate and compared. By the end of the study, all participants experienced reductions in systolic blood pressure, with average diastolic and systolic readings decreasing from 83 to 76 mm Hg and from 144 to 132 mm Hg, respectively. Accompanying in vitro testing suggested that noni juice may have some effect on angiotensin converting enzyme (ACE) and angiotensin receptors (AR). However, this trial included a small number of participants (n=10), and there was no effect on the blood pressure of normotensive participants in other clinical trials of noni juice. Therefore, ACE inhibiting and AR blocking activities are not likely to occur in real-life conditions. If noni juice truly does have an impact on high blood pressure it may involve other mechanisms, such reduction of aortic SAR and the release of nitric oxide by macrophages [83]. These same antioxidant and anti-inflammatory mechanisms are reportedly responsible for the hypotensive activity of olive leaf extract [84, 85].

To evaluate the influence of noni juice on oral health, eleven patients with moderate to severe gingivitis or periodontitis were enrolled in a four-week pilot study [86]. These volunteers were divided into a control group (n=5) and a noni juice group. Those in the noni juice group rinsed their mouths for two minutes with 30 mL noni juice plus 30 mL water, followed by swallowing. This was done twice per day for four weeks. Bacterial samples were isolated from the oral cavity and gingival pouches. Papillae bleeding (PBI), plaque and approximal plaque indices were also scored for each volunteer before and after the treatment period. Those in the noni juice group experienced a significant decline in PBI, especially when compared to the control group. Interestingly, noni juice had only weak bacteriostatic activity in vitro and bacterial composition of patient oral cavities did not change significantly during the treatment phase. Therefore, the study authors concluded that the improvement in clinical outcomes in the noni juice group was likely due to the anti-inflammatory properties of noni juice.

10. Noni extract studies

There have been several clinical trials with various noni extracts. The phytochemical compositions of these extracts are certainly different than that of noni fruit juice. Therefore, the conclusions drawn from these studies may not be applicable to noni juice. However, they are briefly described in this review as they may provide some additional insight into the general health-promoting properties of the noni plant.

A phase 1 clinical trial of freeze-dried noni fruit capsules found no evidence of toxicity, but did reveal dose-dependent improvements in physical functioning, pain and fatigue in patients with progressing advanced cancer for which no standard treatment was available [87]. A randomized, placebo-controlled clinical trial with 100 adult patients revealed that an aqueous extract of dried,

whole noni fruit prevented postoperative nausea when ingested 1 hour before surgery [88]. Ingestion of an aqueous noni fruit extract increased ranitidine absorption in healthy adult volunteers, indicating a possible effect on gastrokinetic activity that might make it useful as a carminative and appetite-stimulating agent and in the relief of heartburn [89]. A noni fruit extract exhibited analgesic properties, comparable to or better than ibuprofen, in a randomized, parallel clinical trial involving 51 patients who had undergone simple tooth extraction procedures [90]. The results of a three-month open-label longitudinal study demonstrated the potential anti-stress (adaptogenic) effect of an herbal mixture that included noni [91]. Human trials involving the topical application of noni extracts, as well as noni fruit juice, indicate anti-inflammatory, anti-acne, and anti-aging activities within the skin [92-96].

11. Discussion

There is a preponderance of evidence among the results of these human studies which demonstrates that noni juice has notable antioxidant activity, even more so than the other more common fruit juices that were used in the placebos. It seems that the antioxidant effects of noni juice from French Polynesia are universal, as it has been observed in heavy smokers, athletes, and in ordinary healthy people. This antioxidant activity may involve direct chemical reactions between phytochemicals and reactive oxygen species or may involve the induction of antioxidant enzyme systems. Examples of these two mechanisms are found in both in vitro and in vivo studies of noni juice.

Tahitian Noni Juice was also evaluated for its in vitro antioxidant activity in the LOOH and tetrazolium nitroblue (TNB) assays [97]. The SAR scavenging activity of noni juice, as measured in the TNB assay, followed a linear positive dose-response. The SAR scavenging activity of 7 μ L/mL Tahitian Noni Juice was also compared to those of 13.3 μ g/mL vitamin C, 13.3 μ g/mL Pycnogenol®, and 22.2 μ g/mL of grape seed powder, the latter three being well known for their antioxidant activity. The SAR scavenging activity of TNJ was 2.8 times greater than vitamin C, 1.4 times greater than Pycnogenol®, and 1.1 times greater than grape seed powder. Very relevant to this comparison is the fact that the total solid material in 7 μ L TNJ is approximately 0.7 μ g. Therefore, a much smaller amount of noni juice solids, 3 to 5% of the total weight of the other antioxidants, exhibited greater SAR scavenging activity. As with SAR scavenging activity, in vitro LOOH quenching activity followed a linear positive dose-response, revealing a consistent antioxidant action.

Ingestion of noni juice protected the livers of Sprague-Dawley (SD) rats exposed to carbon tetrachloride and resulted in decreased hepatotoxic lesions and significant reductions in serum alanine aminotransferase (ALT) and aspartate aminotransferase (AST), when compared to a placebo group [98]. Further, SAR and LOOH levels were significantly lower in the liver tissue of noni fed rats [99]. Carbon tetrachloride (CCl4) causes extensive liver damage from lipid oxidation processes, due to the formation of the trichloromethyl and trichloromethyl peroxide radicals during metabolism. These trichloromethyl radicals initiate lipid peroxidation of unsaturated fatty acids resulting in cytotoxic and genotoxic LOOH and other decomposition products, as well as increased proinflammatory SAR production [100, 101]. As such, the liver protective results in these in vivo studies suggest that noni juice may contain compounds that are capable of scavenging trichloromethyl radicals or subsequent peroxidation products.

Ingestion of Tahitian Noni Juice by ICR mice for 30 days resulted in a dose-dependent decrease in blood MDA, as well as dose-dependent increases in glutathione peroxidase and superoxide dismutase activities in the liver [102]. Feeding of deacetylasperulosidic acid (DAA), a major phytochemical constituent of noni juice, for 7 days to Wistar rats resulted in significant increase in enzymatic antioxidant activity. Further, a dose-dependent reduction in serum MDA concentration, a downstream oxidation product of SAR action, accompanied an increase in superoxide dismutase activity, suggesting that DAA ingestion increases catalase activity [70].

As previously discussed, the antioxidant properties of noni juice contribute to its immune modulating activity. The attributes may also be at least partially responsible for noni's anti-inflammatory activity. The relationship between inflammation, the immune system, and reactive

oxygen species (ROS) is well established in the scientific literature. Leukocytes release proinflammatory cytokines and ROS, which in turn cause a "respiratory," or oxidative, burst by NADPH oxidase. This leads to further inflammation, as well as recruits and activates other leukocytes. This inflammatory process leads to tissue damage, such as that occurring in certain types of muscle injury [103]. Antioxidants have been shown to suppress neutrophil chemotaxis (recruitment) and reactive oxygen intermediates [104, 105]. Antioxidants also have an influence on cyclooxygenase-2 (COX-2) expression and activity. Reactive oxygen intermediates are involved in signaling pathways that lead to COX-2 expression in cells. Inhibition of COX-2 expression has been observed in vitro, in vivo, and ex vivo for a variety of radical scavengers [106-108]. As the rate of prostaglandin synthesis by cyclooxygenases is dependent on peroxide, antioxidants and enzymes that limit peroxide availability may inhibit cyclooxygenase activity [109, 110]. Similarly, lipoxygenase is activated by ROS [111]. Therefore, antioxidants may exhibit anti-inflammatory activity by way of multiple mechanisms.

The antioxidant mechanisms induced by noni juice appear to be involved, to some degree, in all of the effects reported in human interventions studies. It is very likely that this property of noni juice is central to its observed effects on physical fatigue, weight control, osteoporosis, hypertension, and gingivitis. Certainly, antioxidant action is a major mechanism by which noni juice controls AGE formation [68].

There are limitations in applying the results of the human studies discussed in this review to noni products in general. First, these noni juice studies were conducted on one commercial source. This source is noni fruit puree from French Polynesia, and the safety and efficacy of other sources may not be the same. As briefly mentioned above, there is wide variability among the nutrient and phytochemical compositions of commercial noni products. An analysis of 177 commercial noni juice products revealed large differences among the mineral content of these products [39]. One reason for this is the inclusion of different ingredients in these juice products. Another reason is the use of varying amounts of different types of noni materials. From a safety perspective, this variability is significant. An example of this is provided in a liver injury case involving a 14-year old boy who had been ingesting what was purportedly a noni juice dietary supplement [112]. But subsequent chemical analysis of this supplement revealed that it contained less than 1% noni juice, as well as included other ingredients that were not declared on the product label. Ingredients other than noni were responsible for the liver injury [113]. Another example is a product that was marketed as a noni fruit extract, but was devoid of any of the well-known chemical markers of noni fruit. This counterfeit noni extract was reported to be potentially less safe, when evaluated in a reproductive toxicity test, than authentic noni fruit that had been evaluated in several other reproductive toxicity tests [114, 115]. A third example is a case of liver injury from a noni product of unknown identity that was purchased from a local market in Ecuador and brought to Europe for personal use [116]. This product was not produced with the same quality controls that ensure an authentic and safe noni juice product and, certainly, was not a product approved for use by the European Union.

These particular instances of product adulteration are a warning that a label declaration of "noni juice" does not guarantee that a product is indeed safe or authentic. Authentic noni juice made from noni fruit puree from French Polynesia (Tahitian Noni Juice) has been subject to thorough safety assessments by European Union officials. Consumption of this source of noni fruit juice has been determined to be safe, as well as ingredients derived from the same source [117, 118]. Nutrient and chemical specifications for these ingredients were set in place to provide an identity for what has been demonstrated to be safe [119]. When noni fruit is processed into puree, the seeds and skin are removed. There are a number of consequences to this, in terms of phytochemical content. Among these, is an increase the safety profile due to the prevention of possible anthraquinone contamination [120]. This is one reason why processing of noni fruit into puree is a recommended and approved method.

Differences in noni fruit harvesting and processing methods will also have an impact on potential efficacy. For example, the degree of ripeness at harvest, and subsequent post-harvest aging, may influence antioxidant activity, as well as vitamin C and phenolic compound content [121, 122]. Also, changes that occur during long periods of fermentation include significant losses in antioxidant

activity [123]. On the other hand, production of noni fruit puree does not seem to significantly affect deacetylasperulosidic acid, scopoletin or vitamin C content [67, 124].

Geographical, or environmental, factors (soil, sunlight, temperature, precipitation, etc.) may significantly influence noni fruit composition. Scopoletin, rutin, quercetin, and 5,15-dimethylmorindol were detected in greatly varying concentrations in noni fruits and noni fruit juice products sourced from 13 different nations located in the Caribbean, Central America, the Central Pacific, the South Pacific, and Asia [125]. Iridoid content was also substantially different among samples of noni fruit collected from French Polynesia, Tonga, the Dominican Republic, Okinawa, Thailand and Hawaii, with French Polynesia containing the highest concentration and the Dominican Republic containing the lowest [67]. Principle component analysis of commercial noni juice products has also revealed regionally distinct phytochemical profiles [126]. Further, iridoid glucoside, scopoletin, rutin, and fatty acid glucoside concentrations were considerably different among various samples of noni fruits, powdered fruit capsules, noni juices, and mixed noni fruit juices. Noniosides B and C, as well as scopoletin were even absent in some samples of commercial noni juice [127].

12. Conclusions

The weight of evidence obtained from human studies point to noni juice possessing notable antioxidant activity, more so than the other fruit juices that served as placebos. It is this activity, and its interaction with the immune system and inflammation pathways, that may account for much of the observed health benefits of noni juice. These health benefits may include protection against tobacco smoke toxicities—including DNA protection, normalization of blood lipids, control of systemic inflammation, and reduction of homocysteine— improvement of joint pain and mobility, increased physical endurance, increased immune activity, control of AGE accumulation, weight management, maintenance of bone health in women, control of blood pressure, and improved gum health.

The existing evidence does have some limitations in its application to noni juice products in general. Geographical factors, along with post-growth factors (harvesting, storage, transportation, processing, and formulation) produce commercial noni juice products with different phytochemical and nutrient profiles. Differences in phytochemical profiles will likely result in variations in biological activity [128]. As such, conclusions drawn from the human clinical trials discussed in this review may be limited to a single source of noni juice. Noni juice products from other sources are likely to have somewhat different toxicological and pharmacological profiles.

- Author Contributions: Brett J. West, Shixin Deng, Fumiyuki Isami, Akemi Uwaya, and C. Jarakae Jensen were
 all involved in the development, review and approval of this manuscript.
- Conflicts of Interest: The authors are employed in the research and development department of Morinda, Inc.,
 a manufacturer of noni juice.

538 References

- 1. European Food Safety Authority. Scientific Opinion of the Panel on Dietetic Products Nutrition and Allergies on a request from the European Commission on the safety of Tahitian Noni® 'Morinda citrifolia (Noni) fruit puree and concentrate' as a novel food ingredient. *EFSA Journal* **2009**, 998, 1-16; DOI:10.2903/j.efsa.2009.998.
- 543 2. European Commission. Commission decision of 5 June 2003 authorising the placing on the market of "noni juice" (juice of the fruit of *Morinda citrifolia* L.) as a novel food ingredient under regulation (EC) No 258/97 of the European parliament and of the council. *Official J. Eur. Union L* 144 2003, 46, 12. Available online: http://data.europa.eu/eli/dec/2003/426/oj (accessed 11 December 2017).
- 547 3. China Food and Drug Administration. *June 27, 2011 Health Food Record Information Release.* Available online: http://www.sda.gov.cn/WS01/CL0613/63379.html (accessed 11 December 2017).
- Morton, J. The ocean-going Noni, or Indian Mulberry (*Morinda citrifolia*, Rubiaceae) and some of its "colorful" relatives. *Econ. Bot.* **1992**, 46, 241-256; DOI:10.1007/BF02866623.

- 551 5. Parkinson S. *A Journal of a Voyage to the South Seas, in His Majesty's Ship, The Endeavor;* South Seas, National Library of Australia, Canberra. Available online: http://southseas.nla.gov.au/journals/parkinson/068.html (accessed 10 January 2018).
- 554 6. Cheesman, T.F. The flora of Raratonga, the chief island of the Cook group. *Trans. Linn. Soc. Lond.*, 2nd Series 1903, 6, 261-313; DOI: 10.1111/j.1095-8339.1903.tb00277.x.
- 556 7. Hedrick, U.P. Sturtevant's Notes on Edible Plants. J.B. Lyon Company, Albany, NY, USA, 1919; pp. 368.
- Maiden, J.H. *Useful Native Plants of Australia (and Tasmania)*. Technological Museum of New South Wales, Sydney, Australia, 1889; pp. 45.
- 559 9. Rae, C.J.; Lamprell, V.L.; Lion, R.J.; Rae, A.M. The role of bush foods in contemporary Aboriginal diets. 560 *Proc. Nutr. Soc. Aust.* **1982**, 7, 45-48.
- 561 10. Ochse J.J.; van den Brink, C.B. Vegetables of the Dutch East Indies (Edible Tubers, Bulbs, Rhizomes and Spices Included): Survey of Indigenous and Foreign Plants Serving as Pot-Plants and Side-Dishes. Archipel Drukkerij, Java, Dutch East Indies, 1931: pp. 630-632.
- 564 11. Henry, T. *Ancient Tahiti: Bernice P. Bishop Museum Bulletin 48.* Bernice P. Bishop Museum, Honolulu, HI, USA, 1928; pp. 59.
- 566 12. West, B.J.; Jensen, C.J.; Westendorf, J. A new vegetable oil from noni (*Morinda citrifolia*) seeds. *Int. J. Food Sci. Tech.* **2008**, 43, 1988-1992; DOI: 10.1111/j.1365-2621.2008.01802.x.
- 568 Bouzerand, E. Points Forts De la Polynésie Française: Bilan Commerce Exterieur 2016. Institut de la Statistique 569 Polynesie Papeete, 2017. online: de la Francaise, French Polynesia, Available 570 http://www.ispf.pf/docs/default-source/publi-pf-bilans-et-etudes/pf-bilan-01-2017-comext-571 2016.pdf?sfvrsn=4 (accessed 12 December 2017).
- 572 14. Whistler, W.A. *Polynesian Herbal Medicine*, Hong Kong: National Botanical Garden, Hong Kong, 1992; pp. 173-174; ISBN: 0-915809-16-8.
- 574 15. Petard, P. *Quelques Plantes Utiles de Polynesie Francaise et Raau Tahiti*. Editions Haere Po No Tahiti, Papeete, French Polynesia, 1986; pp. 280-286; ISBN: 2-904171-06-1.
- 576 16. Brown, F.B.H. Flora of southeastern Polynesia. III. Dicotyledons. Bishop Museum Bulletin 130. Bernice P. Bishop Museum, Honolulu, HI, USA, 1935; pp. 306-308.
- 578 17. Girardi, C.; Butaud, J.F; Ollier, C.; Ingert, N.; Weniger, B.; Raharivelomanana, P.; Moretti, C. Herbal medicine in the Marquesas Islands. *J. Ethnopharm.* **2015**, 161, 200-213; DOI: 10.1016/j.jep.2014.09.045.
- 580 18. Westendorf, J.; Mettlich, C. The benefits of noni juice: an epidemiological evaluation in Europe. *J. Med. Food*581 *Plants* 2009, 1, 64-79.
- 582 19. Assi, R.A.; Darwis, T.; Abdulbaqi, I.M.; Khan, A.A.; Vuanghao, L.; Laghari, M.H. *Morinda citrifolia* (Noni): a comprehensive review on its industrial uses, pharmacological activities, and clinical trials. *Arabian J. Chem.* 2017, 10, 691-707; DOI: 10.1016/j.arabjc.2015.06.018.
- 585 20. Torres, M.A.O.; de Fatima Braga Magalhaes, I.; Mondego-Oliveira, R.; de Sa, J.; Rocha, A.L.; Abreu-Silva, S86 A.L. One plant, many uses: a review of the pharmacological applications of *Morinda citrifolia*. *Phytother*. *Res.* **2017**, 31, 971-979; DOI: 10.1002/ptr.5817.
- 588 21. Brown, A.C. Anticancer activity of Morinda citrifolia (noni) fruit: a review. *Phytother. Res.* 2012; 26, 1427-1440; DOI: 10.1002/ptr.4595.
- 590 22. Gupta, R.K.; Patel, A.K. Do the health claims made for *Morinda citrifolia* (Noni) harmonize with current scientific knowledge and evaluation of its biological effects. *Asian Pac. J. Cancer Prev.* **2013**, 14, 4495-4499; PMID: 24083691.
- West, B.J.; White, L.D.; Jensen, C.J.; Palu, A.K. A double-blind clinical safety study of noni fruit juice. *Pacific Health Dialog* **2009**, 15, 21-32; PMID: 20443518.
- Wang, M.Y.; Lutfiyya, M.N.; Weidenbacher-Hoper, V.; Anderson, G.; Su, C.X.; West, B.J. Antioxidant activity of noni juice in heavy smokers. *Chem. Cent. J.* **2009**, 3, article ID 3:13; DOI: 10.1186/1752-153X-3-13.
- 597 25. Wang, M.Y.; Peng, L.; Lutfiyya, M.N.; Henley, E.; Weidenbacher-Hoper, V.; Anderson, G. *Morinda citrifolia*598 (noni) reduces cancer risk in current smokers by decreasing aromatic DNA adducts. *Nutr. Cancer* 2009, 61,
 634-639; DOI: 10.1080/01635580902825605.
- Wang, M.Y.; Peng, L.; Jensen, C.J.; Deng, S.; West, B.J. Noni juice reduces lipid peroxidation–derived DNA adducts in heavy smokers. *Food Sci. Nutr.* **2013**, 1, 141-149; DOI: 10.1002/fsn3.21.
- Wang, M.Y.; Peng, L.; Weidenbacher-Hoper, V.; Deng, S.; Anderson, G.; West, B.J. Noni juice improves serum lipid profiles and other risk markers in cigarette smokers. *Scientific World Journal* **2012**, 2012, article ID 594657; DOI: 10.1100/2012/594657.

- Craig, W.Y.: Palomaki, G.E.; Haddow, J.E. Cigarette smoking and serum lipid and lipoprotein concentrations: an analysis of published data. *BMJ* **1989**, 298, 784-788; DOI: 10.1136/bmj.298.6676.784.
- 607 29. Jefferis, B.J.; Lowe, G.D.; Welsh, P.; Rumley, A.; Lawlor, D.A.; Ebrahim, S.; Carson, C.; Doig, M.; Feyerabend, C.; McMeekin, L.; Wannamethee, S.G.; Cook, D.G.; Whincup, P.H. Secondhand smoke (SHS) exposure is associated with circulating markers of inflammation and endothelial function in adult men and women. *Atherosclerosis* 2010, 208, 550-556; DOI: 10.1016/j.atherosclerosis.2009.07.044.
- 611 30. Fröhlich, M.; Sund, M.; Löwel, H.; Imhof, A.; Hoffmeister, A.; Koenig, W. Independent association of various smoking characteristics with markers of systemic inflammation in men. Results from a representative sample of the general population (MONICA Augsburg Survey 1994/95). *Eur. Heart J.* 2003, 24, 365-1372: DOI: 10.1016/S0195-668X(03)00260-4.
- 615 31. Palu, A.K.; Brown, A.; Deng, S.; Kaluhiokalani, N.; West, B.J. The effects of noni (*Morinda citrifolia* L.) fruit juice on cholesterol levels: a mechanistic investigation and an open label pilot study. *J. Appl. Pharm. Sci.* 2012, 2, 25-30; DOI: 10.7324/JAPS.2012.2905.
- 618 32. Pawlus, A.D.; Kinghorn, D.A. Review of the ethnobotany, chemistry, biological activity and safety of the botanical dietary supplement *Morinda citrifolia* (noni). *J. Pharm. Pharmacol.* **2007**, 59, 1587-1609; DOI: 10.1211/jpp.59.12.0001.
- 33. Basar, S.; Uhlenhut, K.; Högger, P.; Schöne, F.; Westendorf, J. Analgesic and antiinflammatory activity of *Morinda citrifolia* L. (Noni) fruit. *Phytother. Res.* **2010**, 24, 38-42; DOI: 10.1002/ptr.2863.
- 623 34. Dussossoy, E.; Brat, P.; Bony, E.; Boudard, F.; Poucheret, P.; Mertz, C.; Giaimis, J.; Michel, A. Characterization, anti-oxidative and anti-inflammatory effects of Costa Rican noni juice (*Morinda citrifolia* L.). *J. Ethnopharm.* 2011, 133, 108-115; DOI: 10.1016/j.jep.2010.08.063.
- Little E.L.; Wadsworth, F.H. Common trees of Puerto Rico and the Virgin Islands. Agriculture Handbook No. 249.
 U.S. Department of Agriculture, Forest Service, Washington D.C., USA, 1964.
- 628 36. Akinbo, S.R.A.; Noronha, C.C.; Okanlawon, A.O.; Denesi, M.A. Comparative study of the effect of *Morinda citrifolia* (Noni) with selected physiotherapy modalities in the management of patients with cervical spondylosis. *Nigerian J. Health Biomed. Sci.* **2006**, 5, 6-11; DOI: 10.4314/njhbs.v5i2.11590.
- 631 37. Wang, M.Y.; Lutfiyya, M.N.; Weidenbacher-Hoper, V.; Peng, L.; Lipsky, M.S.; Anderson, G. *Morinda citrifolia* L. (noni) improves the quality of life in adults with osteoarthritis. *Func. Food Health Dis.* 2011, 1, 75-90.
- 634 38. Fletcher, H.M.; Dawkins, J.; Rattray, C.; Wharfe, G.; Reid, M.; Gordon-Strachan, G. *Morinda citrifolia* (Noni) as an anti-Inflammatory treatment in women with primary dysmenorrhoea: a randomised double-blind placebo-controlled trial. *Obstet. Gynecol. Int.* **2013**, 2013, article ID 195454; DOI: 10.1155/2013/195454.
- 637 39. West, B.J.; Tolson, C.B.; Vest, R.G.; Jensen, S.; Lundell, T.G. Mineral variability among 177 commercial noni juices. *Int. J. Food Sci. Nutr.* **2006**, 57, 556-558; DOI: 10.1080/09637480601066794.
- 40. Thaman, R.R. Kiribati Agroforestry: Trees, People and the Atoll Environment. *Smithsonian Institution, Atoll Research Bulletin* **1990**, 333, 1-29; DOI: 10.5479/si.00775630.333.1.
- 641 41. Palu, A.K.; Seifulla, R.D.; West, B.J. Morinda citrifolia L. (noni) improves athlete endurance: Its mechanisms 642 of action. J. Med. Plant Res. 2008, 2, 154-158. Available online 643 www.academicjournals.org/journal/JMPR/article-full-text-pdf/736746415191 (accessed 13 December 2017).
- 42. Anugweje K.C.; Okonko, I.O.; Effect of noni supplementation on the serum creatine kinase (CK) levels of athletes. *World J. Sport Sci.* **2012**, *7*, 41-47; DOI: 10.5829/idosi.wjss.2012.7.1.65170.
- 43. Liu, J.F.; Chang, W.Y.; Chan, K.H.; Tsai, W.Y.; Lin, C.L.; Hsu, M.C. Blood lipid peroxides and muscle damage increased following intensive resistance training of female weightlifters. *Ann. N. Y. Acad. Sci.* **2005**, 1042, 255-261; DOI: 10.1196/annals.1338.029.
- 649 44. Brancaccio, P.; Lippi, G.; Maffulli, N. Biochemical markers of muscular damage. *Clin. Chem. Lab. Med.* **2010**, 650 48, 7575-767; DOI: 10.1515/CCLM.2010.179.
- West, B.J.; Martínez, R.; Cano, M.; White, A.; Palu, A.K. Noni fruit juice may increase oxygen uptake in athletes during exercise. *Curr. Res. J. Biol. Sci.* **2013**, 5, 177-181.
- 653 46. Mehta, H.; Nazzal, K.; Sadikot, R.T. Cigarette smoking and innate immunity. *Inflamm. Res.* **2008**, 57, 497-654 503; DOI: 10.1007/s00011-008-8078-6.
- 47. Huttunen, R.; Heikkinen, T.; Syrjanen, J. Smoking and the outcome of infection. *J. Intern. Med.* **2011**, 269, 258-269; DOI: 10.1111/j.1365-2796.2010.02332.x.
- 48. Zhou, Z.; Chen, P.; Peng, H. Are healthy smokers really healthy? *Tob. Induc. Dis.* **2016**, 14, article ID 35; DOI: 10.1186/s12971-016-0101-z.

- 49. Fusby, J.S.; Kassmeier, M.D.; Palmer, V.L.; Perry, G.A.; Anderson, D.K.; Hackfort, B.T.; Alvarez, G.K.; D.M.;
 Akhter, M.P.; Swanson, P.C. Cigarette smoke-induced effects on bone marrow B-cell subsets and
 CD4+:CD8+ T-cell ratios are reversed by smoking cessation: influence of bone mass on immune cell
 response to and recovery from smoke exposure. *Inhal. Toxicol.* 2010, 22, 78-796: DOI:
 10.3109/08958378.2010.483258.
- 50. Duvenci Birben, O.; Akcay, S.; Sezer, S.; Sirvan, S.; Haberal, M. Effect of smoking on peripheral blood lymphocyte subsets of patients with chronic renal failure. *Exp. Clin. Transplant.* **2016**, 14, 91-94; DOI: 10.6002/ect.tondtdtd2016.P34.
- 51. Schafer, M.; Sharp, P.; Brooks, V.J.; Xu, J.; Cai, J.; Keuler, N.S.; Peek, S.F.; Godbee, R.G.; Schultz, R.D.; Darien,
 B.J. Enhanced bactericidal activity against *Escherichia coli* in calves fed *Morinda citrifolia* (Noni) puree. *J. Vet.*Intern. Med. 2008, 22, 499-502; DOI: 10.1111/j.1939-1676.2008.0044.x.
- 52. Brooks, V.J.; De Wolfe, T.J.; Paulus, T.J.; Xu, J.; Cai, J.; Keuler, N.S.; Godbee, R.G.; Peek, S.F.; McGuirk, S.M.;
 Darien, B.J. Ethnoveterinary application of *Morinda citrifolia* fruit puree on a commercial heifer rearing facility with endemic salmonellosis. *Afr. J. Tradit. Complement. Altern. Med.* 2013, 10, 1-8; DOI: 10.4314/ajtcam.v10i1.1.
- 674 53. Palu, A.K.; Kim, A.H.; West, B.J.; Deng, S.; Jensen, J.; White, L. The effects of *Morinda citrifolia* L. (noni) on the immune system: Its molecular mechanisms of action," *J. Ethnopharm.* 2008, 115, 502-506; DOI: 10.1016/j.jep.2007.10.023.
- 677 54. Ma, D.L.; Jun, Z.; Jianhua, G.; et al. The effect of Tahitian Noni Juice on antioxidation and immune function. *Chinese Medical Research & Clinical* **2008**, 6, 8-10.
- 55. Droge, W. Free radicals in the physiological control of cell function. *Physiol. Rev.* **2002**, 82, 47-95; DOI: 10.1152/physrev.00018.2001.
- Lee, K.S.; Park, H.S.; Park, S.J.; Kim, A.R.; Min, K.H.; Jin, S.M.; Li, L.; Lee, Y.C. An antioxidant modulates
 expression of receptor activator of NF-κB in asthma. *Exp. Mol. Med.*, 2006, 38, 217-229; DOI: 10.1038/emm.2006.27.
- 684 57. Chen, S.S.; Gong, J.; Liu, F.T.; Mohammed, U. Naturally occurring polyphenolic antioxidants modulate IgE-mediated mast cell activation. *Immunology*, **2000**, 100, 471-480; DOI: 10.1046/j.1365-2567.2000.00045.x.
- 58. Lampe, J.W. Health effects of vegetables and fruit: assessing mechanisms of action in human experimental studies. *Am. J. Clin. Nutr.* **1999**, 70, 475S–490S; DOI: 10.1093/ajcn/70.3.475s.
- 59. Bub, A.; Watzl, B.; Blockhaus, M.; Briviba, K.; Liegibel, U.; Müller, H.; Pool-Zobel, B.L.; Rechkemmer, G. Fruit juice consumption modulates antioxidative status, immune status and DNA damage. *J. Nutr. Biochem.* 2003, 14, 90–98; DOI: 10.1016/S0955-2863(02)00255-3.
- 691 60. Hughes, D.A. Effects of dietary antioxidants on the immune function of middle-aged adults. *Proc. Nutr. Soc.* **1999**, 58, 79-84; DOI: 10.1079/PNS19990012.
- 693 61. De la Fuente, M. Effects of antioxidants on immune system ageing. *Eur. J. Clin. Nutr.* **2002**, 56, S5–S8; DOI: 10.1038/sj.ejcn.1601476.
- 695 62. Carr, A.A.; Maggini, S. Vitamin C and immune function. *Nutrients*, **2017**, 9, article ID 1211; DOI: 10.3390/nu9111211.
- 697 63. Ishibashi, Y.; Matsui, T.; Isami, F.; Abe, Y.; Sakaguchi, T.; Higashimoto, Y.; Yamagishi, S.I. N-butanol extracts of *Morinda citrifolia* suppress advanced glycation end products (AGE)-induced inflammatory reactions in endothelial cells through its anti-oxidative properties. BMC Complement. Altern. Med. **2017**, 17, article ID 137; DOI: 10.1186/s12906-017-1641-3.
- 701 64. Abe, Y.; Yagi, M.; Uwaya, A.; Isami, F.; Yonei, Y. Effect of iridoid (containing plants) on AGE formation and degradation. *Glycative Stress Research* **2016**, 3, 56-64.
- Gacche R.N.; Dhole, N.A. Profile of aldose reductase inhibition, anti-cataract and free radical scavenging activity of selected medicinal plants: an attempt to standardize the botanicals for amelioration of diabetes complications. *Food Chem. Toxicol.* **2011**, vol. 49, 1806-1813; DOI: 10.1016/j.fct.2011.04.032.
- West, B.J.; Uwaya, A.; Isami, F.; S. Deng, S.; Nakajima, S.; Jensen, C.J. Antiglycation activity of iridoids and their food sources. *Int. J. Food Sci.* **2014**, 2014, article ID 276950; DOI: 10.1155/2014/276950.
- 708 67. Deng, S.; West, B.J.; Palu, A.K.; Jensen, C.J. Determination and comparative analysis of major iridoids in different parts and cultivation sources of *Morinda citrifolia*. *Phytochem. Anal.* **2011**, 22, 26-30; DOI: 10.1002/pca.1246.
- 711 68. West, B.J.; Deng, S.; Uwaya, A.; Isami, F.; Abe, Y.; Yamagishi, S.I.; Jensen, C.J. Iridoids are natural glycation inhibitors. *Glycoconj. J.* **2016**, 33, 671–681; DOI: 10.1007/s10719-016-9695-x.

- 713 69. Meerwaldt, R.; Graaff, R.; Oomen, P.H.N.; Links, T.P.; Jager, J.J.; Alderson, N.L.; Thorpe, S.R.; Baynes, J.W.; Gans, R.O.B; Smit, A.J. Simple non-invasive assessment of advanced glycation endproduct accumulation.
 715 Diabetologia 2004, 47, 324-1330; DOI: 10.1007/s00125-004-1451-2.
- 716 70. Ma, D.L.; Chen, M.; Su, C.X.; West, B.J. In vivo antioxidant activity of deacetylasperulosidic Acid in noni. 717 *J. Anal. Methods Chem.* **2013**, 2013, article ID 804504; DOI: 10.1155/2013/804504.
- 71. Pande, M.; Naiker, M.; Mills, G.; Singh, N.; Voro, T. The kura files: qualitative social survey. *Pac. Health Dialog* **2005**, 12, 85-93; PMID: 18181498.
- 72. Horsfal, A.U.; Olabiyi, O.A.; Osinubi, A.A.; Noronha, C.C.; Okanlawon, A.O. Anti diabetic effect of fruit juice of *Morinda citrifolia* (Tahitian Noni Juice®) on experimentally induced diabetic rats. *Nigerian Journal of Health and Biomedical Sciences*, **2008**, 7, 34-37; DOI: 10.4314/njhbs.v7i2.11674.
- 73. Horsfall, A.U.; Olabiyi, O.A.; Aiyegbusi, A.; Noronha, C.C.; Okanlawon, A.O. *Morinda citrifolia* fruit juice augments insulin action in Sprague-Dawley rats with experimentally induced diabetes. *Nig. Q. J. Hosp. Med.* **2008**, 18, 162-165; DOI: 10.4314/nqjhm.v18i3.45020.
- 726 74. Su, C.X.; Jensen, C.J.; Zhou, B.N. Chapter 11, Morinda citrifolia (noni): its effect on insulin secretion by G 727 protein coupled receptor systems. In ACS Symposium Series, Vol. 987; Ho, C.T., Simon, J.E., Shahidi,
 728 F., Shou, Y., Eds.; American Chemical Society: Washington D.C., USA, 2008; pp. 172-172; DOI: 10.1021/bk 729 2008-0987.ch011.
- 730 75. Nerurkar, P.V.; Hwang, P.W.; Saksa, E. Anti-diabetic potential of noni: the yin and the yang. *Molecules* **2015**, 20, 17684-17719; DOI: 10.3390/molecules201017684.
- 732 76. Bartnicka, A.; Seidel, A.; John, M.; Basar-Maurer, S.; Westendorf, J.; Algenstaedt, P. Nonisaft senkt signifikant den HbA1c bei patienten mit typ-2-diabetes. *Diabetologie und Stoffwechsel* **2013**, 8, P121; DOI: 10.1055/s-0033-1341781.
- 735 77. Graz, B.; Kitalong, C.; Yano, V. Traditional local medicines in the republic of Palau and non-communicable diseases (NCD), signs of effectiveness. *J. Ethnopharm.*, **2015**, 23, 161-233; DOI: 10.1016/j.jep.2014.11.047.
- 737 78. Inada, A.C.; Figueiredo, P.S.; Santos-Eichler, R.A.D.; Freitas, K.C.; Hiane, P.A.; Castro, A.P; Guimarães, C.A. *Morinda citrifolia* Linn. (Noni) and its potential in obesity-related metabolic dysfunction. *Nutrients* 2017, 9, article ID 540; DOI: 10.3390/nu9060540.
- 740 79. Palu, A.K.; West, B.J.; Jensen, J. Noni-based nutritional supplementation and exercise interventions influence body composition. *N. Am. J. Med. Sci.* **2011**, 3, 552-556; DOI: 10.4297/najms.2011.3552.
- 80. Bogdanov, A.R.; Derbeneva, S.A.; Bogdanova, A.A.; Feofanova, T.B.; Panfilova, N.V.; Nesierova, V.E. Low calorie diet influence optimization on body composition at obese patients with secondary diastolic heart failure. *Voprosy Pitaniia* **2015**, 84, 37-46; PMID: 26852530.
- Langford, J.; Doughty, A.; Wang, M.; Clayton, L.; Babich, M. Effects of *Morinda citrifolia* on quality of life and auditory function in postmenopausal women. *J. Altern. Complement. Med.* 2004, 10, 737-739; DOI: 10.1089/acm.2004.10.737.
- Palu, A.K.; Santiago, R.A.; West, B.J.; Kaluhiokalani, N.; Jensen, J. Chapter 39: The effects of *Morinda citrifolia*L. noni on high blood pressure: a mechanistic investigation and case study. In *American Chemical Society*Symposia Series No. 993, Functional Food and Health. Shibamoto, T., Kanazawa, K., Shahidi, F., Ho, C.T., Eds.;
 American Chemical Society: Washington D.C., USA, 2008; pp. 446-453; DOI: 10.1021/bk-2008-0993.ch039.
- 752 83. Almeida-Souza, F.; de Souza Cda, S.; Taniwaki, N.N.; Silva, J.J.; de Oliveira, R.M.; Abreu-Silva, A.L.; 753 Calabrese Kda, S. *Morinda citrifolia* Linn. fruit (Noni) juice induces an increase in NO production and death of Leishmania amazonensis amastigotes in peritoneal macrophages from BALB/c. *Nitric Oxide* 2016, 58, 51-58; DOI: 10.1016/j.niox.2016.06.004.
- 756 84. Cabrera-Vique, C.; Navarro-Alarcón, M.; Rodríguez Martínez, C.; Fonollá-Joya, J. Hypotensive effect of an extract of bioactive compounds of olive leaves: preliminary clinical study. *Nutricion Hospitalaria* **2015**, 32, 242-249; DOI: 10.3305/nh.2015.32.1.8644.
- 759 85. Khayyal, M.T.; el-Ghazaly, M.A.; Abdallah, D.M.; Nassar, N.N.; Okpanyi, S.N.; Kreuter, M.H. Blood pressure lowering effect of an olive leaf extract (*Olea europaea*) in L-NAME induced hypertension in rats. *Arzneimittelforschung* 2002, 52, 797-802; DOI; 10.1055/s-0031-1299970.
- 762 86. Glang, J.; Falk, W.; Westendorf, J. Effect of *Morinda citrifolia* L. fruit juice on gingivitis/periodontitis. *Modern* 763 *Research in Inflammation* **2013**, 2, 21-27; DOI: 10.4236/mri.2013.22003.
- 764 87. Issell, B.F.; Gotay, C.C.; Pagano I.; Franke, A.A. Using quality of life measures in a phase I clinical trial of noni in patients with advanced cancer to select a phase II dose. *J. Diet. Suppl.* **2009**, 6, 347-359; DOI: 10.3109/19390210903280272.

- 767 88. Prapaitrakool, S.; Itharat, A. *Morinda citrifolia* Linn. for prevention of postoperative nausea and vomiting. 768 J. *Med. Assoc. Thai.* **2010**, 93, S204-S209; PMID: 21294416.
- 769 89. Nima, S.; Kasiwong, S.; Ridtitid, W.; Thaenmanee, N.; Mahattanadul, S. Gastrokinetic activity of *Morinda citrifolia* aqueous fruit extract and its possible mechanism of action in human and rat models. *J. Ethnopharm.* 771 2012, 142, 354-361; DOI: 10.1016/j.jep.2012.04.044.
- 772 90. Cornejo, F.M.P.; Asmat, A.A.S.; Ruiz, R.S.G. Post exodontia analgesic effect of *Morinda citrifolia* (noni): a randomized trial in parallel groups. *Int. J. Odontostomat.* **2014**, 8, 433-438; DOI: 10.4067/S0718-774 381X2014000300018.
- 91. Seely, D; Singh, R. Adaptogenic potential of a polyherbal natural health product: report on a longitudinal clinical trial. *Evid. Based Complement. Alternat. Med.* **2007**, 4, 375-380; DOI: 10.1093/ecam/nel101.
- 777 92. Lee, J.N.; Kim, S.W.; Yoo, Y.K.; Lee, G.T.; Lee, K.K. Anti-wrinkle effect of *Morinda citrifolia* (noni) extracts. *J. Soc. Cosmet. Sci. Korea*, **2006**, 32, 227-231.
- 779 93. West, B.J.; Deng, S.; Palu, A.K.; Jensen, C.J. *Morinda citrifolia* Linn. (Rubiaceae) leaf extracts mitigate UVB induced erythema. *J. Nat. Med.* **2009**, 63, 351-354; DOI: 10.1007/s11418-009-0327-7.
- Palu, A.K.; West, B.J.; Jensen, C.J. Noni seed oil topical safety, efficacy, and potential mechanisms of action.
 J. Cosmetics Dermatol. Sci. Appl. 2012, 2, 74-78; DOI: 10.4236/jcdsa.2012.22017.
- 783 95. West B.J.; Sabin, R. Efficacy of a *Morinda citrifolia* based skin care regimen. *Curr. Res. J. Biol. Sci.* **2012**, 4, 310-314.
- 96. Uwaya, A.; Isami, F.; Abe, Y.; Jensen, C.J.; Deng, S.; West, B.J. Topical antiaging activity of a combination of *Morinda citrifolia* (noni) seed concentrate, fruit juice, leaf juice, and seed oil. *Pharma Science Monitor* **2017**, 8, 8-24.
- 788 97. Wang M.Y.; Su, C. Cancer preventive effect of *Morinda citrifolia* (Noni). *Ann. N. Y. Acad. Sci.* **2001**, 952, 161-168; DOI: 110.1111/j.1749-6632.2001.tb02737.x.
- Wang, M.Y.; G.; Nowicki, D.; Jensen, J. Hepatic protection by noni fruit juice against CCl(4)-induced chronic liver damage in female SD rats. *Plant Foods Hum. Nutr.* **2008**, 63, 141-145; DOI; 0.1007/s11130-008-0083-y.
- Wang, M.Y.; West, B.J.; Jensen, C.J.; Nowicki, D.; Su, C.; Palu, A.K.; Anderson, G. *Morinda citrifolia* (Noni):
 a literature review and recent advances in Noni research. *Acta Pharmacol. Sin.* 2002, 23, 1127-1141; PMID:
 12466051.
- 795 100. Comporti, M. Lipid peroxidation and cellular damage in toxic liver injury. *Lab. Invest.* **1985**, 53, 599-623; PMID: 3906270.
- 797 101. Recknagel, R.O.; Glende, E.A.; Dolak, J.A.; Waller, R.L. Mechanisms of carbon tetrachloride toxicity. *Pharmacol. Ther.* 1989, 43, 139-154; PMID: 2675128.
- 799 102. Zhang, J.J.; Wang, L.Y.; Ou, L.Y.; et al. Study on evaluation of antioxidant activity of Noni juice in vivo. *Science and Technology of Food Industry*, **2011**, 32, 392–393.
- 801 103. Butterfield, T.A.; T. M. Best, T.M.; Merrick, M.A. The dual roles of neutrophils and macrophages in inflammation: a critical balance between tissue damage and repair. *J. Train.* 2006, 41, 457-465; PMID: 17273473.
- 804 104. Dona, M.; Dell'Aica, I.; Calabrese, F.; Benelli, R.; Morini, M.; Albini, A.; Garbisa, S. Neutrophil restraint by green tea: inhibition of inflammation, associated angiogenesis, and pulmonary fibrosis. *J. Immunol.* 2003, 170, 4335-4341; DOI: 10.4049/jimmunol.170.8.4335.
- 807 105. Petrone, W.F.; English, D.K.; Wong, K.; McCord, J.M. Free radicals and inflammation: Superoxide-dependent activation of a neutrophil chemotactic factor in plasma. *Proc. Natl. Acad. Sci. U.S.A.* 1980, 77, 1159-1163; PMID: 6928666.
- 810 106. Feng, L.; Xia, Y.; Garcia, G.E.; Hwang, D.; Wilson, C.B. Involvement of reactive oxygen intermediates in cycloogygenase-2 expression induced by interleukin-1, tumor necrosis factor-α, and lipopolysaccharide. *J. Clin. Invest.* 1995, 95, 1669-1675; DOI: 10.1172/JCI117842.
- 813 107. O'Leary, K.A.; de Pascual-Teresa; S.; Needs, P.W.; Bao, Y.P.; O'Brien, N.M; Williamson, G. Effect of flavonoids and Vitamin E on cyclooxygenase-2 (COX-2) transcription. *Mutation Research*, **2004**, 551, 245-254; DOI: 10.1016/j.mrfmmm.2004.01.015.
- 816 108. de Pascual-Teresa, S.; Johnston, K.L.; DuPont, M.S.; O'Leary, K.A.; Needs, P.W.; Morgan, L.M.; Clifford, M.N.; Bao, Y.; Williamson, G. Quercetin metabolites downregulate cyclooxygenase-2 transcription in human lymphocytes ex vivo but not in vivo. *J. Nutr.* 2004; 134, 552-557; DOI: 10.1093/jn/134.3.552.
- 819 109. Hemler M.E.; Lands, W.E. Evidence for a peroxide-initiated free radical mechanism of prostaglandin biosynthesis. *J. Biol. Chem.* 1980, 255, 6253-6261; PMID: 6771268.

856

857

- 821 110. Wu, D. Hayek, M.G.; Meydani, S. Vitamin E and macrophage cyclooxygenase regulation in the aged. *J. Nutr.* **2001**, 131, 382S-388S; PMID: 11160566.
- 823 111. Werz, O.; Szellas, D.; Steinhilber, D. Reactive oxygen species released from granulocytes stimulate 5-lipoxygenase activity in a B-lymphocytic cell line. *Eur. J. Biochem.* **2000**, 267, 1263-1269; PMID: 10691962.
- 112. Yu, E.L.; Sivagnanam, M.; Ellis, L.; Huang, J.S. Acute hepatotoxicity after ingestion of *Morinda citrifolia*(Noni Berry) juice in a 14-year-old boy. *J. Pediatr. Gastroenterol. Nutr.* **2011**, 52, 222-2224, DOI: 10.1097/MPG.0b013e3181eb69f0.
- 828 113. West, B.J.; Deng, S. Ingredients other than noni may be culprits in case of acute hepatotoxicity in 14-year-old boy. *J. Pediatr. Gastroenterol. Nutr.* **2011**, 53, 469-470; DOI: 10.1097/MPG.0b013e3181eb69f0.
- 830 114. West, B.J.; Su, C.X.; Jensen, C.J. Prenatal toxicity test of *Morinda citrifolia* (noni) fruit. *J. Toxicol. Sci.* **2008**, 33, 647-649; DOI: 10.2131/jts.33.647.
- 832 115. Wang, M.Y.; Hurn, J.; Peng, L.; Nowicki, D.; Anderson, G. A multigeneration reproductive and developmental safety evaluation of authentic *Morinda citrifolia* (noni) juice. *J. Toxicol. Sci.* **2011**, 36, 81-85; DOI: 10.2131/jts.36.81.
- 835 116. West B.J.; Berrio, L.F. Tahitian Noni juice is not hepatotoxic. *Rev. Esp. Enferm. Dig.* **2007**, 99, 737-738; DOI: 10.4321/S1130-01082007001200017.
- 117. European Commission Scientific Committee on Food. Opinion of the Scientific Committee on Food on Tahitian
 Noni® juice expressed on 4 December 2002. Available online:
 https://ec.europa.eu/food/sites/food/files/safety/docs/sci-com_scf_out151_en.pdf (Accessed 18 January 2018).
- 841 118. European Food Safety Authority. Opinion on a request from the Commission related to the safety of noni 842 juice (juice of the fruits of *Morinda citrifolia*). *EFSA Journal*, **2006**, 4, article ID 376; DOI: 843 10.2903/j.efsa.2006.376.
- 844 119. European Food Safety Authority. Opinion on the safety of Tahitian Noni® 'Morinda citrifolia (noni) fruit 845 puree and concentrate' as a novel food ingredient. *EFSA Journal* **2009**, 7, article ID 998; DOI: 846 10.2903/j.efsa.2009.998.
- 847 120. Bussmann, R.W.; Hennig, L/; Giannis, A.; Ortwein, J.; Kutchan, T.M.; Feng, X. Anthraquinone Content in Noni (*Morinda citrifolia* L.). *Evid. Based Complement. Alternat. Med.* 2013, 2013, article ID 208378; DOI: 10.1155/2013/208378.
- 850 121. Y. Chan-Blanco, F. Vaillant, A. M. Perez, M. P. Belleville, C. Zuniga, and P. Brat, P. The ripening and aging of noni fruits (*Morinda citrifolia* L.): microbiological flora and antioxidant compounds. *J. Sci. Food Agric.* 2007, 87, 1710-1716; DOI: 10.1002/jsfa.2894.
- Yang, J.; Gadi, R.; Thomson, T. Antioxidant capacity, total phenols, and ascorbic acid content of noni (*Morinda citrifolia*) fruits and leaves at various stages of maturity. *Micronesica*, **2011**, 41, 167-176.
 - 123. Yang, J.; Paulino, R.; Janke-Stedronsky, S.; Abawi, F. Free-radical-scavenging activity and total phenols of noni (*Morinda citrifolia* L.) juice and powder in processing and storage. *Food Chem.* **2007**, 102, 302-308; DOI: 10.1016/j.foodchem.2006.05.020.
- 858 124. West, B.J.; Deng, S.; Jensen, C.J. Nutrient and phytochemical analyses of processed noni puree. *Food Res.* 859 *Int.* **2011**, 44, 2295-2301; DOI: 10.1016/j.foodres.2010.09.038.
- B60 125. Deng, S.; West, B.J.; Jensen, C.J. A quantitative comparison of phytochemical components in global noni fruits and their commercial products. *Food Chem.* **2010**, 122, 267-270; DOI: 10.1016/j.foodchem.2010.01.031.
- 126. Lachenmeier, K.; Mußhoff, F.; Madea, B.; Reusch, H.; Lachenmeier, D.W. Authentication of noni (*Morinda citrifolia*) juice. *Deutsche Lebensmittelrundschau* 2006, 102, 58-61.
- Potterat, O.; Von Felten, R.; Dalsgaard, P.W.; Hamburger, M. Identification of TLC Markers and
 Quantification by HPLC-MS of Various Constituents in Noni Fruit Powder and Commercial Noni-Derived
 Products. J. Agric. Food Chem. 2007, 55, 7489-7494; DOI: 10.1021/jf071359a.
- 867 128. Ghasemzadeh, A.; Jaafar, H.Z.; Rahmat, A. Phytochemical constituents and biological activities of different extracts of *Strobilanthes crispus* (L.) Bremek leaves grown in different locations of Malaysia. BMC Complement. Altern. Med. **2015**, 15, article ID 422; DOI: 10.1186/s12906-015-0873-3.