Supplementation of Low Ratio N-6:N-3 Pufa Reduces Body Fatness In Young Obese Balinese Women: A Randomized Study with Optimized Energy Regulation

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Abstract
Background and Objectives: Obese Indonesians, including Balinese, are at higher risk of comorbidities like, CVD, diabetes, non-alcoholic fatty liver disease. Urban Balinese are changing their diet from traditional to fast foods and the like that are high in saturated fats. Nutritional modification, such as increasing n-3 PUFA content in the diet, may aid in managing body fat accumulation-related diseases. This study investigated the effects of supplementation of n-6:n-3 PUFA with ratio of 2:1 on body fat reduction in young obese Balinese women.

Methods and Study Design: Sixty-six young obese Balinese women, aged 18-25, were randomly assigned equally into Intervention and Control groups, supplemented with 2100 mg:1100 mg and 240 mg:100 mg of n-6:n-3 PUFAs, respectively. Data were collected at baseline, 6, and 12 weeks of intervention. BMI, waist circumference (WC), waist-to-height ratio (WHR), conicity index, triglyceride concentrations, and Lipid Accumulation Product (LAP) were measured. Participants were advised to maintain <1500-Kcal daily energy intake and participate in a guided low-impact aerobics once a week.

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Results: After 12-weeks supplementation, all body fat indices (BMI, WC, WHtR, conicity index, and LAP), decreased significantly whether unadjusted or adjusted by the reduced of energy intake in the Intervention group. Otherwise, in the Control group only some of WC indices (WC, WHtR, and conicity index) decreased statistically. BMI and WC of the Intervention group decreased significantly more than the Control group.

Conclusion: Twelve-week intervention with high-dose of low n-6:n-3 PUFAs ratio and restriction energy intake, reduced body fatness selectively in young obese Balinese women.

Introduction
The prevalence of obesity along with its complications, is increasing rapidly, particularly in developing countries such as Indonesia. Among obese people, adipocyte accumulation is considered a critical event because these fat cells function as active endocrine organs influencing many crucial functions of various systems of the body. As adipocytes enlarge, their metabolic roles change and the expression of proinflammatory cytokines increases. Therefore, obesity has been proposed as a chronic low-grade inflammatory disease. Severely obese people are at a high risk of obesity-related comorbidities, such as CVD, diabetes, and non-alcoholic fatty liver disease. In addition to BMI, other parameters are commonly used for obesity determination, such as lipid accumulation product (LAP)—an adiposity index for adults correlated with abdominal fat storage and adiposity dysfunction—is calculated using waist circumference (WC) and triglyceride (TG) level. Along with waist-to-height ratio (WHtR) and the conicity index (C-Index) (which is calculated using WC, body weight (BW), and height (BH)), LAP is strongly correlated with visceral fat accumulation. A combination of environmental, lifestyle, and genetic factors determines the individual risk for developing comorbidities and disease progression; in addition, nutrition, drug intake, metabolic disarrangement, toxin exposure, and infection can be modifying environmental factors. Therefore, careful nutritional modification may aid in managing the aforementioned conditions caused by overaccumulation of body fat, particularly visceral fat. Elizondo et al. reported that the content of n-3 PUFA and long chain (LC)-PUFA of the -erythrocyte phospholipids was 30% and 35% lower in obese people than in nonobese people, respectively. Furthermore, the content of n-3 PUFA [i.e., linolenic acid (ALA)] and LC-PUFA (i.e. EPA and DHA) was 56% and 59% lower in obese people than in nonobese people, respectively; in other words, the levels of n-6 PUFA [i.e., linoleic acid (LA)] and LC-PUFA (i.e. arachidonic acid) were relatively higher in obese people than in their nonobese counterparts. The n-6:n-3 LC-PUFA ratio was consequently 144% higher in obese people than in nonobese controls. Therefore, food intake with a high n-6:n-3 ratio may increase the risk of various metabolic and degenerative diseases, because n-6 PUFA is associated with proinflammatory markers. Modifying nutrition by increasing the dietary content of n-3 PUFA is an approach that may aid in reducing the risk of metabolic diseases; the dietary inclusion of canola oil (n-6:n-3 PUFA ratio, 2:1) can be suitable for this intervention. Because canola oil is high in n-3 PUFA (ALA) content, it can potentially modulate the inflammatory response and reduce the risk of obesity-associated metabolic syndrome.

Material and Methods
Research Design, Participants, and Intervention
A double-blind, randomised-controlled trial was conducted during May–September 2013. A total of 80 young women (aged 18–25 years) who felt obese and showed their interests to participate in the study were screened. Out of these 80 women, 74 women met the following entry criteria: BMI $\geq$ 25kg/m², unmarried (single) status, ready as a subject and comply fully with the study protocol and procedure, and living in Denpasar for several years and at least next one year.
Women who were on body weight control program using pharmacologic or herbal regimen, on routine routine using supplementation of antioxidants, vitamins, or minerals, on treatment of chronic diseases using steroid or non steroid anti inflammation, and alcoholic (daily consumption of more than 20 g alcohol) were excluded from the study.

These 74 women were equally allocated to either the Intervention or Control group (n= 37 in each group). Informed consent was obtained before commencing the study, and the participants' anonymity was maintained throughout. The study protocol was approved by the Research Ethics Committee of Udayana University/Sanglah Hospital (Ethical Clearance No. 787/UN.14.2/Litbang /2012, 17 September 2012). The study was registered as Clinical trial ID ACTRN12615000757516.

The Intervention group received daily supplementation of 30 mL of emulsion containing either 10 g of canola oil (700 mg SFA, 6100 mg MUFA, 2100 mg LA, and 1100 mg ALA, and the Control group received daily supplementation of 2 g palm oil (66 mg SFA, 1000 mg MUFA, 240 mg LA, and 100 mg ALA; control group) for 12 weeks (June–September 2013).

Participants in Intervention and Control groups were also recommended to restrict their daily energy intake to less than 1500 Kcal/day. To maintain the participants’ compliance, we conducted weekly meetings every Sunday, where they performed 1-h exercise with an aerobic instructor weekly meetings with clinical investigators were done every
Sunday, wherein subjects performed 1-h exercise with an aerobic instructor. During this meeting, the participants were provided 1 week’s supply (250 mL) of emulsion. Any complaints or adverse effects potentially caused by the supplementation was also monitored and reported. To ensure that the emulsion was consumed as instructed, the emulsion remaining in a used bottle container was measured weekly. The remaining emulsion in a used bottle container was measured weekly in order to ensure that the emulsion was consumed as instructed. Eight participants (10.8%) dropped out during the course of the study (4 from each group), leaving a total of 66 participants (33 in each group) who completed the study and included in the final analyses. Summary of recruitment and participation during the trial is summarised in Figure 1.

Data collection
Assessment of Food Intake
Dietary intake was assessed twice at the baseline and the endline, by two trained dietitians, using Food Frequency Questionnaire (FFQ) and Semi-Quantitative FFQ (SQ-FFQ) methods. Food patterns were calculated from FFQ as number of foods frequency consumed per day. Conversion of food to nutrient intakes per day from SQ-FFQ data was done using Nutrisurvey software developed by Erhardt in 2007 and distributed by SEAMEO RECFON, Universitas Indonesia resulting in the quantity of macronutrients (energy, protein, fat, carbohydrates, fiber, and cholesterol).

Anthropometric and Biochemical Assessments
Anthropometric and biochemical variables were assessed three times: on the first day of week-1 (baseline), on the last day of week-6 (midline), and on the last day of week-12 (endline).

All anthropometric variables were measured twice, and for the final analyses mean values were used. Body weight (BW) was assessed using a digital scale (Omron HBF-362 model, Kyoto, Japan), with a precision of 0.1 kg. Body height (BH) was assessed using a stature meter (General Care No 26SM), with a precision of 0.1 cm. Waist circumference (WC) was measured using a flexible non-elastic tape, exactly at the middle level of the abdomen, with a precision of 0.1 cm.

Triglyceride (TG) concentrations were measured from blood serum using a colorimetric method (Cobas 6000; Roche Diagnostics, Mannheim, Germany), with a precision of 1 mg/dL.

Calculation of body fatness
BMI was calculated as BW (kg)/(BH (m))^2. WHtR was calculated as WC (cm)/BH (cm), and C-Index was calculated as WC (cm)/(0.109 × √[BW (kg)/BH (cm)]). Finally, LAP for women was calculated as [WC (cm) − 58] × TG (mmol).

Statistical Analyses
Statistical analyses were performed using Stata 12.1 (Stata Corp, College Station, TX, USA). The normal distribution of continuous data is presented as mean ± standard error of the mean (SE). Comparison within groups were tested using general linear model, and changes between groups were compared using an independent t test. Statistical significance was set at p<0.05.

Results
All participants, in the intervention and the control-groups were comparable in baseline characteristics, namely; age, occupations, ethnicities, selected nutrients intake (except fiber), anthropometric and body fat index values. However, there were extreme values for certain nutrients intake, such as for cholesterol and fiber. Cholesterol intake of the both group were relatively high (more than 250 mg), especially in the Intervention group (319 mg), although it was not different between group. On the contrary, both groups had a very low fiber consumption when compared with Indonesian RDA (2014) for that of age and gender group (32 g). The Intervention group (8.1 g) had a particularly lower fiber intake than those the Control group (11.9 g) (p=0.024) (Table 1). Most of the energy intake came from traditional Balinese food such as, rice, chicken, tempeh, and tofu, with additional energy from street and fast food (Table 2).
### Table 1: Baseline characteristics of participants

| Parameters                        | Intervention | Control   | p     |
|-----------------------------------|--------------|-----------|-------|
| **Age (years) (mean±SE)**         | 20.9±1.8     | 20.6±1.5  | 0.413 |
| **Occupation (f(%))**             |              |           |       |
| Student                           | 30(90.9)     | 30 (90.9) | 1.000 |
| Officer staff                     | 2(6.1)       | 2 (6.1)   |       |
| Unemployment                      | 1(3.0)       | 1 (3.0)   |       |
| **Ethnicity (f(%))**              |              |           |       |
| Balinese                          | 30 (90.9)    | 30 (90.9) | 0.613 |
| Javanese                          | 2 (6.0)      | -         |       |
| Chinese                           | -            | 1 (3.0)   |       |
| Indian                            | 1 (3.0)      | 2 (6.0)   |       |
| **Selective Nutrients Intake (mean±SE)** |          |           |       |
| Energy (Kcal)                     | 1803±107     | 1812±118  | 0.956 |
| Fat (g)                           | 78.6±6.67    | 66.0±7.08 | 0.679 |
| PUFA (g)                          | 18.9±2.50    | 16.1±2.50 | 0.832 |
| Cholesterol (mg)                  | 319±25.9     | 265±25.3  | 0.109 |
| Fiber (g)                         | 8.1±0.68     | 11.9±1.60 | 0.024 |
| **Anthropometric (mean±SE)**      |              |           |       |
| Body weight (kg)                  | 76.5±2.71    | 76.3±2.14 | 0.929 |
| BMI (kg/m²)                       | 30.3±0.97    | 30.5±0.75 | 0.766 |
| Waist circumference (cm)          | 92.3±2.15    | 92.7±2.15 | 0.899 |

Presented in mean±SE, analysed using independent student t-test.
Presented in f (%), tested using chi-square. PUFA= poly unsaturated fatty acid, BMI= body mass index. All characteristics variables between both were comparable, except it was very low intake of fiber in both group, especially in intervention group.

### Table 2: Food consumption frequency pattern (time/day) of total participants (n=66)

| Food Sources          | f(%)  | Mean±SE   |
|-----------------------|-------|-----------|
| **Carbohydrates**     |       |           |
| - Rice                | 66(100)| 2.23±0.12 |
| - Noodle              | 64(97.0)| 0.19±0.03 |
| - Bread               | 63(95.5)| 0.47±0.13 |
| - Table sugar         | 45(68.2)| 1.39±0.36 |
| - Snack (biscuit, commercial snack) | 40(60.6)| 0.81±0.14 |
| - Soft drink          | 43(65.2)| 0.19±0.06 |
| **Proteins**          |       |           |
| - Beef                | 35(53.0)| 0.20±0.05 |
| - Pork                | 45(68.2)| 0.17±0.04 |
| - Lamb                | 20(30.3)| 0.26±0.09 |
| - Chicken             | 65(98.5)| 1.13±0.17 |

Presented in f (%), tested using chi-square.
Overall, the energy intake of both the intervention and control groups significantly decreased from the baseline measurements \((p<0.001\) and \(p=0.001\), respectively; Table 3). The decrease energy intake was most probably due to the reduction of rice consumption.

Among the unadjusted or adjusted by the reduced energy intake, after intervention compared with the baseline. Specifically, BMI, WC, WHtR and C-Index significantly decreased at the midline \((\text{unadjusted}: p=0.007, p<0.001, p<0.001, \text{and} p=0.001; \text{adjusted}: p=0.08, p<0.001, p<0.001 \text{and} p=0.001,\) respectively) and end line \((\text{same} p\text{ value among unadjusted and adjusted}: p=0.005, p<0.0001, p<0.001, \text{and} p<0.001,\) respectively). Notably, in the control group, only the parameters calculated considering WC decreased at the end line, compared with that at the baseline weather that with unadjusted or adjusted by reduced energy intake \((p=0.007 \text{ or} p=0.06; \text{and} 0.006 \text{ or} 0.005 \text{ for WC and WHtR, respectively}); \text{however, the C-Index decreased at midline as well as end line} (p=0.019 \text{ and} 0.001,\) respectively), and the reduction in BMI was non-significant \((p=0.961)\). We next observed that TG concentration in the intervention-group participants significantly decreased at the end line, compared with that at the baseline with unadjusted and adjusted by reduced energy intake \((p=0.021,\) and \(p=0.022); \text{by contrast, TG concentration demonstrated no significant changes in the control group} \((p=0.033)\) and end line \((p<0.001),\) compared with that at the baseline; by contrast, the control group demonstrated no significant change in LAP \((p=0.001)\). Furthermore, LAP also decreased significantly in the intervention group at the midline \((p=0.033)\) and end line \((p<0.001),\) compared with that at the baseline; by contrast, the control group demonstrated no significant change in LAP \((p=0.001)\).

Between group analyses, the decline of energy intake after 12 weeks supplementation did not differ among two groups \((p=0.961)\). BMI, WC and WHtR declined statistically different from the baseline to the endline in the Intervention group compared to the Control group \((p=0.038, p=0.047,\) and \(p=0.052,\) respectively). No differences in changes of C-index, TG concentrations, and LAP were observed between Intervention and Control groups \((p=0.019)\).
Table 3: Changes of research variables values were adjusted by energy intake based on groups of intervention and time of assessments at baseline, midline, and endline of the participants

| Parameter          | Intervention (n=33) | Control (n=33) |
|--------------------|---------------------|----------------|
|                    | Baseline (mean±SE)  | Midline (mean±SE) | Endline (mean±SE) | Baseline (mean±SE)  | Midline (mean±SE) | Endline (mean±SE) | p  |
| Energy Intake (Kcal) | 1803±107            | NA             | 1351±110          | <0.001             | 1812±118           | NA             | 1368±97           | 0.001      |
| BMI (kg/m²)        | 30.3±0.97†‡          | 29.9±0.94†     | 29.8±0.97‡        | 0.007†             | 0.005‡             | 0.005‡         | 0.005‡       | NS          |
| WC (cm)            | 92.3±2.15†‡          | 89.1±2.08†§    | 86.9±2.08 ‡§      | <0.001†            | <0.001‡            | <0.001‡        | <0.001‡      | NS          |
| WHtR               | 0.58±0.01 †‡         | 0.56±0.01 †§   | 0.55±0.01 †       | <0.001†            | <0.001†            | <0.001†        | <0.001†      | NS          |
| C Index            | 1.22±0.02†‡          | 1.19±0.01†§    | 1.16±0.01†§       | 0.001†             | 0.001†             | 0.001†         | 0.001†       | NS          |
| TG (mg/dL)         | 115±9.99†            | 109±9.18       | 103±7.05‡         | 0.021‡             | 0.022‡             | 0.021‡         | 0.021‡       | NS          |
| LAP                | 47.9±7.01†‡          | 41.6±6.02†§    | 36.3±4.97 ‡§      | 0.033‡             | 0.036‡             | 0.033‡         | 0.036‡       | NS          |

p within-group, comparison of baseline, midline and endline mean tested by general linear model, adjusted by the reduced energy intake from baseline to end-line.
Baseline week-0, midline week-6, endline week-12.
†; baseline vs midline;§;midline vs endline;‡; baseline vs endline
BMI; body mass index, WC;waist circumference, WHtR; waist to height ratio, CIndex; conicity index, TG; triglyceride, LAP; lipid accumulation product.
Table 4: Comparison of changes of research variables values between intervention vs control groups during the intervention

| Parameters | Time Interval (Week) | Groups – value changes |  |
|------------|----------------------|------------------------|--|
|            |                      | Intervention (mean±SE) | Control (mean±SE) | p       |
| Energy intake | 0-12 | -451±115 | -443±116 | 0.961 |
| BMI         | 0-6 | -0.40±0.14 | 0.21±0.29 | 0.100 |
|             | 6-12 | -0.11±0.09 | 0.03±0.07 | 0.275 |
|             | 0-12 | -0.50±0.17 | 0.24±0.29 | 0.038* |
| WC          | 0-6 | -3.21±0.78 | -1.58±0.81 | 0.153 |
|             | 6-12 | -2.20±0.67 | -1.16±0.70 | 0.283 |
|             | 0-12 | -5.41±0.92 | -2.74±0.95 | 0.047* |
| WHtR        | 0-6 | -0.02±0.00 | -0.01±0.00 | 0.161 |
|             | 6-12 | -0.01±0.00 | -0.01±0.00 | 0.295 |
|             | 0-12 | -0.03±0.01 | -0.02±0.01 | 0.052* |
| C Index     | 0-6 | -0.04±0.01 | -0.03±0.01 | 0.639 |
|             | 6-12 | -0.03±0.01 | -0.02±0.01 | 0.329 |
|             | 0-12 | -0.06±0.01 | -0.04±0.01 | 0.329 |
| TG          | 0-6 | -6.27±7.50 | -8.24±6.01 | 0.838 |
|             | 6-12 | -6.06±6.45 | 2.82±5.11 | 0.285 |
|             | 0-12 | -12.33±5.09 | -5.42±8.11 | 0.473 |
| LAP         | 0-6 | -6.32±2.84 | -5.66±2.29 | 0.871 |
|             | 6-12 | -5.28±2.36 | -0.99±2.40 | 0.209 |
|             | 0-12 | -11.6±2.70 | -6.65±4.52 | 0.351 |

BMI: body mass index; WC: waist circumference; WHtR: waist to height ratio; C Index: conicity index; TG: triglyceride; LAP: lipid accumulation product.

Discussion
In present study, we try to identify and elaborate the ways to control body fatness and prevent its comorbidities, through optimized the role of nutrition intervention, such as energy restriction and supplementation high dose of low n-6:n-3 PUFA ratio, especially in young obese women.

The energy restriction seemly has the main role to control body fatness. In both, the Intervention and the Control group were significantly decreased energy intake, in the endline compared to the baseline. The reduced energy intake of the Control group, was also followed by decreased of the sensitive body fatness, such as WC, WHtR and C-Index. The reduction of body fat could be optimized by supplementation high dose of low n-6:n-3 PUFA ratio. We observed the significant reduction of all body fat indexes, namely BMI, WC, WHtR, C-Index, TG concentration, and LAP within the Intervention group. Furthermore, between group analyses, some body fatness such as BMI, WC, and WHtR have been reducing more in the Intervention than the Control group.
The most prominent improvement was noted in BMI and WC after 12 weeks of the intervention. During the 12 weeks of intervention, LAP decreased as well. We propose that the supplementation for longer than 12 weeks will have more substantial positive effects. However, because we lost 10.8% of the participants to follow-up in this short-term study, we suggest that participants’ compliance should be ensured for studies exploring the effects of long-term intervention.

Most of studies, both in animals and human reported the positive effects of n-3 PUFA supplementation in weight management. A systematic review\textsuperscript{15} reported, four of five studies in adults, have concluded the changes in body weight by supplementation with n-3 PUFA. The evidence corroborates the role of n-3 PUFA, in controlling adiposity in obese people. This n-3 PUFA can control pathways involved in lipid metabolism by regulating biomolecular the gene transcription factors paroxysmal proliferator activated receptor (PPAR\textsubscript{α}), PPAR\textsubscript{γ}, sterol regulatory element binding protein-1 (SREBP-1), and carbohydrate-responsive element-binding protein (ChREBP).\textsuperscript{16,17} Two mechanisms involving n-3 PUFAs, particularly EPA and DHA, can control adiposity. The first involves endogenous lipid synthesis inhibition and oxidation stimulation; both EPA and DHA reduce endogenous lipid production by inhibiting the expression and processing of SREBP-1 involved in the stimulation of lipogenic gene transcription.\textsuperscript{16-19} In the second mechanism, n-3 PUFAs also function as potent PPAR\textsubscript{α} and PPAR\textsubscript{γ} activators and inhibitors; both proteins upregulate the expression of several genes involved in stimulation of fatty acid oxidation.\textsuperscript{16,17,20,21} Roriz \textit{et al.},\textsuperscript{9} reported that WHtR, C-Index, LAP, and visceral adiposity index (VAI; calculated using WC, BMI, and TG and HDL-cholesterol levels) were correlated with visceral adipose tissue area and assessed through computerised tomography; this is the gold standard for assessing men and women (aged 20–59 years), particularly their WHtRs and C-Index scores. The researchers concluded that for predicting visceral fat accumulation in adults and elderly people of both genders, the simple anthropometric measurements of WHtR and C-Index are more accurate regarding sensitivity and specificity than are those of LAP and VAI, both of which are anthropometric and biochemical measurements. Compared with those of the anthropometric parameters, the LAP and VAI are potentially attributable to the weaker correlations of biochemical parameters used for calculating them (e.g., TG and HDL-cholesterol levels) with visceral fat.\textsuperscript{22} Other studies have reported that high WHtR and C-Index values are associated with an increased risk of CHD, diabetes, hypertension, and dyslipidemia.\textsuperscript{22,23} A systematic review reported that in 91% of women, WHtR was more suitable for predicting CVDs than was WC and BMI.\textsuperscript{24} In addition, visceral fat, commonly responsible for abdominal obesity, has been reported to be the origin of metabolic and degenerative diseases. Thus, the simple parameters WHtR and C-Index are good predictors of abdominal obesity along with visceral fat accumulation.

In present study, supplementation this oil was strongly correlated with reduction in lipid accumulation, indicated by decreased BMI, WC, WHtR, and TG levels. Because adipose cells are considered endocrine organs, which commonly release proinflammatory mediators, this intervention potentially reduces inflammatory and oxidative stress. By limiting both the hypertrophy and hyperplasia of adipocytes, marine n-3 PUFA has an antidiapogenic effect during obesity development.\textsuperscript{25} Another study\textsuperscript{26} reported that decreasing the n-6:n-3 PUFA intake ratio to 3:1 reduces plasma TG levels in older people (aged 45–70 years).

Obesity and lipid accumulation are notable risk factors for metabolic and degenerative diseases; therefore, weight loss and body fat reduction (particularly visceral fat) evidently decrease the risk of these diseases. Because most diseases complicated by obesity are actually preventable, developing a simple and reliable strategy for early detection in participants with risks is essential. We suggest performing simple and applicable screenings using C-Index, WHtR, or LAP algorithms for early detection of increasing body fat composition and distribution. However, any confirmed participants at risk should promptly undergo follow-up and treatment with appropriate management, through modified nutrient intake and exercise. Appropriate detection and prevention of a complicated disease at earlier stages is more effective and efficient than is its treatment at advanced stages.
most metabolic and degenerative diseases lack definitive pharmacologic regimens, and those currently available in the market are expensive. Hence, developing a simple and applicable approach comprising restricted energy intake through a specific diet high in n-3 PUFA and increased physical activity, as the main appropriate prevention and curative strategy, particularly in obese participants, is warranted.

Adipose tissue not only roles as the energy deposit, but also an endocrine organs that release pro-inflammatory mediators, that increase oxidative stress of the body. This study may reduce inflammatory and oxidative stress by limiting both hypertrophy and hyperplasia of adipocytes. Ruzckova et al., reported, that marine n-3 PUFA has an anti-adipogenic effect during obesity development. Another study by Sander et al., reported that decreasing the n-6:n-3 PUFA intake ratio to 3:1 reduces plasma TG levels in older people (aged 45–70 years).

Obesity and lipid accumulation are notable risk factors for metabolic and degenerative diseases; therefore, weight loss and body fat reduction (particularly visceral fat) may decrease the risk of these diseases. Since the majority of obesity-related diseases are preventable, developing a simple and reliable strategy for early risk detection is essential. We suggest the adoption of these simple and applicable screenings by using BMI, WC, C-Index, WHtR, or LAP algorithms. These parameters can detect the increasing of body fat composition and distribution. Furthermore, any confirmed participants at risk should promptly undergo follow-up and treatment with appropriate management, through modified nutrient intake and exercise. Appropriate detection and prevention of a complicated disease at earlier stages is more effective and efficient than its treatment at advanced stages. Moreover, most metabolic and degenerative diseases are still lacking on definitive pharmacologic regimens, and those currently available in the market are expensive. Hence, developing a simple and applicable approach comprising of restricted energy intake, high n-3 PUFA diet consumption and increased physical activity, as the main appropriate prevention and curative strategy, particularly in obese participants, is warranted.

Diet recommendation should consider not only from the health point of view, but it should also take into account the sustainability of the environment. Plant based food that is produced by converting the forest into plantation, such as palm oil, is less sustainable than the naturally growth plant like coconut oil. Currently, most countries recommended plant rather than animal based food, due to it is not only healthier but also has lower environmental impact. Moreover, using plant (canola) oil supplementation in our study offers two advantages. Firstly, it contains high n-3 PUFA (ALA). This will be desaturated and elongated in the human body to form the LC-PUFA (EPA, DPA, and DHA). Secondly, it also contains n-6 PUFA (LA) that will be desaturated and elongated to form dihomo gamma linolenic acid (DGLA). EPA and DGLA will release prostaglandin 3 series and prostaglandin 1 series as anti-inflammatory eucosanoids. These eucosanoids are essential in reducing level of inflammation.

The reduction in lipid parameters after the supplementation indicated the ability of n-3 PUFA to decrease the health risk of fat accumulation, such as insulin resistance, dyslipidaemia, cardiovascular, fatty liver, and several degenerative diseases.

Conclusion Optimized energy restriction and daily supplementation of 3000mg PUFAs, n-6:n-3 ratio 2:1 (2100 mg LA and 1100 mg ALA) for 12 weeks decreases body fat accumulation in young obese women in Bali.

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Conflict of Interest The authors declare no conflict of interest.
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