Effects of low glycemic index/high-fat, high-calorie diet compared to the high-fat, high-calorie diet on glycemic control, lipid profile, and inflammatory markers in children and adolescence with cystic fibrosis: A study protocol for a randomized double-blind controlled clinical trial

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Abstract

Background: Inflammation plays an important role in pathogenesis of cystic fibrosis, and diet is considered as an effective factor in controlling inflammation. Low glycemic index diet seems to be effective for improving glycemic control and reducing the inflammation and lipid profile. Hence, this study was planned to compare the effects of a low glycemic index/high fat, high calorie diet with the high fat, high calorie diet on glycemic status, lipid profile, and inflammatory markers in patients with cystic fibrosis.

Methods: A total of 60 children and adolescence with cystic fibrosis will be randomized to receive a high fat, high calorie diet (n=30) or low glycemic index/high fat, high calorie diet (n=30) with similar calorie and macronutrients composition, for three months. Patients in high fat, high calorie diet arm will be able to use all sources of carbohydrates with different glycaemic indices; while those in another arm will be received their carbohydrate from low glycemic index sources. Before and after the intervention, serum levels of lipid profile (triglyceride, total cholesterol, HDL cholesterol, LDL cholesterol), insulin, glucose, HbA1c, the homeostasis model assessment-estimated insulin resistance index, and some inflammatory markers (IL-6, IL-10, IL-17A, and IFNγ) will be measured.

Discussion: To our knowledge, this study will be the first to examine the effects of a low glycemic index/high fat, high calorie diet compared to the high fat, high calorie diet in cystic fibrosis patients. The findings of this study might provide evidence to improve the nutritional status and immune system as well as to prevent the disease complications in these patients.

Background

Cystic fibrosis (CF) is an autosomal recessive inherited disorder which caused by a defect in the CF transmembrane conductance regulator (CFTR) on chromosome 7 [1]. This
autosomal genetic disease occurs in a raising trend worldwide and it is incidence is reported to be approximately 1 in 3000 births [2, 3]. Individuals with CF develop a wide spectrum of complications, of which, exocrine pancreatic dysfunction is one of the most common manifestations, found in 85% of patients [4]. This type of dysfunction leads to fat malabsorption and in turn results in fat-soluble vitamin inadequacy, malnutrition, and growth failure [5, 6]. CF-related diabetes (CFRD) is another hallmark of CF manifestations which directly linked to increased morbidity and mortality of CF [7]. This complication has been shown to be related to a decline in lung function [8] and also increased risk of bacterial chest infections [9]. Moreover, patients with both CF and CFRD have higher levels of inflammatory markers, and it has been shown that inflammatory biomarkers might be associated with complications of both CF and CFRD [10-13]. Hence, the strategies to reduce inflammatory markers are expected to be beneficial in prevention of CF-related complications [14].

Diet is considered as an effective factor in controlling inflammation. It has been shown that high glycemic index (GI) diet increase oxidative stress and inflammation [15-17], while low GI diet has been indicated potential for improving glycemic control and lipid profile in other forms of diabetes [18-21]. Also, previous studies have shown a beneficial effect of low GI diet on glycemic control and lipid profile in individuals with different health conditions [22, 23]. However, the evidence for the use of low GI diet in CFRD is scare. Given that hyperglycemia exacerbates the clinical situation in CF patients by induction of immune responses, it seems that low GI diet can be an effective factor in modulating the immune response in CF patients by controlling glucose homeostasis [14]. Moreover, given that 16% of CF children may suffer from diabetes and hyperlipidemia, administration of low GI diet might be benefit for these children [24].

As the primary goal of nutrition therapy in CF is to achieve optimal weight gain and
growth, a high fat and high calorie diet has been recommended for these patients [25]. Regarding a definite necessity for determination of the standard diet with the optimal effect on growth, survival rate, quality of life, and the reduction of complications in CF patients, we designed a study to compare the effects of low GI/high fat, high calorie diet with the high fat, high calorie diet on glycemic status, lipid profile, and inflammation markers in patients with CF. We hypothesized that low GI diet will improve the glycemic status and reduce the inflammation and lipid profile in children and adolescence with CF.

Methods/design

Study design

This is a randomized, double-blinded, parallel group, clinical trial. The study protocol was registered in the Iranian Registry of Clinical Trial (registration No, IRCT2017102325267N5).

Study population

This study will be conducted on children and adolescents with CF who are referred to the Cystic Fibrosis Clinic of the Children’s Medical Center, affiliated to Tehran University of Medical Sciences, Tehran, Iran. All children and adolescents, aged 6–18 years old, who are diagnosed with CF by a sweat test or genetic test by a clinical specialist, will be included. Also, we will check oral glucose tolerance test (OGTT) levels from participants’ medical records and those with normal OGTT levels and with no CFRD will be included in the study. The participants will be excluded if: 1) have other chronic diseases such as diabetes and thyroid disease, liver diseases (i.e., hepatitis and cirrhosis), 2) were hospitalized at the beginning of the study or one month earlier, 3) had a history of surgery one month prior to the intervention or were supposed to undergo surgery (except for dental surgery) in the
next three months, and 4) take steroids. Moreover, the participants will be excluded if they will not willing to follow the prescribed diets, unable to adhere to the diets, hospitalize or require surgery during the study.

Sample size

Sample size was determined by formula suggested for randomized clinical trials, with type I error of 5%, type II error of 20%, study power of 80%, and serum interleukin-17A (IL-17A) level as a key variable [26]. The number of needed samples was calculated as 21 participants. To get a more confident result with a 30% dropout rate, we will consider 30 participants in each group.

Randomization

All eligible participants will be enrolled by convenience sampling method and will be randomly (1:1 ratio) allocated by an assistant to high fat, high-calorie diet (HFHC; n = 30) or low GI/high fat, high calorie diet (LG; n = 30), using permuted block randomization method, stratified by age and sex.

Blinding

Given the nature of the intervention (diet), investigators will not be blinded to the intervention. However, all outcomes will be measured by an independent assessor who will be blind to the group allocation. Also, patients and data analysts will be blinded to the group allocation.

Procedure and intervention

In both group, the intervention will be done during three months and both groups will be visited at baseline and after three months of study. All patients will be received a leaflet, presenting the number of portions per meal, food exchange list, food recommendations, and allowed and forbidden foods (for LG diet only).
To design a diet for CF children and adolescents, we first calculate the required energy intake of participants according to the Paediatrics Clinics formula [27]. Then, we added further 300 to calculated energy. The patients in the HFHC diet arm will be received a high-fat, high-calorie diet containing 40% fat, 20% protein, and 40% carbohydrate. This group will be able to use all sources of carbohydrates with different GIs. The patients in the LG diet arm will be received a high-fat, high-calorie diet with macronutrients percentage similar to that in HFHC group; however, the difference will be that the carbohydrate sources were supplied from low GI sources. LG and high glycemic index (HGI) were defined as GI < 50 and GI ≥ 50, respectively. The low GI diet group will be instructed to select carbohydrate containing foods from a list of low GI grains, fruits, vegetables, and dairy, which was provided by the study researchers. Moreover, a list of prohibited HGI foods will be given to members of this group.

The criterion for compliance in the low GI group will be GI<50, calculated from the food diaries. To extract GI values, we will refer to the Iranian GI table [28], and the international table of GI [29] will be used for GI values not reported on the Iranian-specific table. The GI for foods not included in the Iranian or the international table will estimate using the GI for the most similar food. To calculate the mean GI for diet, we will use the reported formula [30].

**Measurement**

**General information**

Information regarding socio-demographic background, current use of medications and supplements, other medical interventions related to the treatment, and lack of allergy or intolerance to certain foods will be recorded by participants’ medical record and also interviews at baseline.
Outcomes

The primary outcomes are changes in the following parameters at the end of the study in comparison with the baseline values: Hemoglobin A1c (HbA1c); fasting blood glucose (FBS); insulin; homeostasis model assessment-estimated insulin resistance (HOMA-IR) index; lipid profile such as triglyceride (TG), total cholesterol (TC), high-density lipoprotein (HDL) cholesterol, low-density lipoprotein (LDL) cholesterol; and inflammatory markers such as interleukin 6 (IL-6), interleukin 10 (IL-10), IL-17A, and interferon-gamma (IFNγ).

The primary variables will be measure at the beginning and the end of the study. For this purpose, venous blood sample (10 mL) will be drawn after 12-h overnight fasting by trained nurses in seated position to measure biochemical markers. Then, blood samples will be centrifuged at 3000 rpm for 10 min at 4 °C to obtain serum which will be stored at −80 °C until biochemical analyses. Lipid profiles and serum levels of glucose will be measured by the enzymatic colorimetric method using standard kits. HbA1c concentrations will be measured by using an auto-analyzer (Selectra E, Vitalab, Holliston, the Netherlands). Serum insulin will be measured using enzyme-linked immunosorbent assay (ELISA) kit. Insulin resistance will be determined using the HOMA-IR equation. Serum levels of IL-6, IL-10, IL-17A, and IFNγ will be measured using ELISA kits.

Secondary outcomes

The secondary outcomes are changes in weight and body mass index (BMI) which will be measure at the beginning and the end of the study. Height will be measured using the SECA stadiometer to the nearest 0.1 cm and weight will be measured using a SECA electronic scale to the nearest 0.1 kg. BMI will be calculated as weight (kg) divided by height squared (m^2).
Adherence

All patients will be trained to follow the instructions of the diets. At the beginning and the end of the study, dietary information of all participants will be collected using a 24-hour dietary recall in three non-consecutive days, and will be analysed by Nutritionist IV software (First Databank, San Bruno, CA). To assess participants adherence during study period, a dietitian will be assigned to check them weekly using telephone. Patients will be monitored weekly during the study period and any occurrence of adverse events will be recorded.

Statistical analysis

Continues and categorical variables will be reported as mean ± standard deviation and number and percentages, respectively. Kolmogorov-Smirnov test will be applied to explore the normality distribution of variables. Paired sample t-test will be performed for within-group changes comparisons (baseline vs. post-intervention values). The analysis of covariance (ANCOVA), will be adjusted for baseline values and will be used for evaluation of between-group differences. All outcome variables will be assessed based on intention-to-treat and per protocol analysis. Missing data will be entered using “regression imputation” method. The level of significance will set at a probability of <0.05 for all analyses. All data analyses will be conducted using the statistical package for social sciences, version 23.0 (SPSS, Chicago, IL, U.S.A).

Discussion

CF is an autosomal recessive inherited disorder which affects many organs, including the lungs, the digestive tract, and various epithelial tissues [31]. Generally, CF is an inflammatory disorder and since inflammation is associated with an increased risk of mortality, the strategies to reduce inflammatory markers is expected to be beneficial in
prevention of this chronic disease [14].

Diet is an effective factor in controlling inflammation in many diseases. Many interventional and observational studies have emphasized the effects of different types of carbohydrate on acute and chronic inflammation [15, 17]. It has been shown that high GI foods increase oxidative stress and inflammation [15–17], while low GI diets improve glycemic control and lipid profile [18–21]. However, the evidence for the use of low GI diet in CF is scare. According to our knowledge, there is no previous study that compared the effects of low GI/high fat, high calorie diet compared to the high fat, high calorie diet in patients with CF. The findings of this study might provide evidence to improve the nutritional status and immune system as well as to prevent the disease complications in these patients.

Trail status

Authors confirmed that the recruitment has been begun in 15th Dec, 2018 and expected that recruitment will be completed in 20th Dec, 2019. The flow chart of the study protocol and the time schedule of enrolment, interventions, assessments, and visits for participants are indicated in Figures 1 and 2.

Abbreviations

ANCOVA: Analysis of covariance; BMI: Body mass index; CF: Cystic fibrosis; CFRD: Cystic fibrosis-related diabetes mellitus; CFTR: CF transmembrane conductance regulator; ELISA: Enzyme-linked immunosorbent assay; GI: Glycemic index; FBG: Fasting blood glucose; HbA1c: Hemoglobin A1c; HDL: High-density lipoprotein; HFHC: High-fat, high-calorie diet; HGI: High glycemic index; HOMA-IR: Homeostasis model assessment-estimated insulin resistance index; IFNγ: interferon-gamma; IL: Interleukin; LDL: Low-density lipoprotein; LG: Low-glycemic index/high-fat, high-calorie diet; OGTT: Oral glucose tolerance test; TG:
Triglyceride; TC: Total cholesterol.

Declarations

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Authors’ contributions

ZG designed the initial idea of this work, which was further developed by MM and MM. SY advised on statistical analysis. ZG organized participant management and data collection. ZG and MM drafted the manuscript. The manuscript has been read and approved by all authors.

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Availability of data and materials

Not applicable.

Ethics approval and consent to participate

The local Ethics Committee of Tehran University of Medical Sciences has approved study protocol (IR.TUMS.VCR.REC.1396.3171). A written informed consent form will be obtained before participating in the study by investigator. Whenever a person is unable to continue supplementation, he/she will be excluded from the study.

Consent for publication

Not applicable.
Competing interests

The authors declare that they have no competing interests.

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Figures
Figure 1. Study’s flow diagram
| TIMEPOINT**     | Enrolment | Allocation | Post-allocation | Close-out |
|----------------|-----------|------------|----------------|-----------|
| **ENROLMENT:** |           |            |                |           |
| Eligibility screen | X         |            |                |           |
| Informed consent | X         |            |                |           |
| Demographic measurements | X    |            |                |           |
| Anthropometric measurements | X    |            |                |           |
| Randomization |            |            | X              |           |
| Allocation     |            |            | X              |           |
| **INTERVENTIONS:** |         |            |                |           |
| [Intervention A] |           |            |                |           |
| [Intervention B] |           |            |                |           |
| **ASSESSMENTS:** |         |            |                |           |
| [List baseline variables] | X   | X          |                |           |
| [List outcome variables] |       |            | X              | X         |
| [List other data variables] |       |            | X              | X         |

**Figure 2.** Time schedule of enrolment, interventions, assessments, and visits for participants.

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