

RESEARCH

Open Access

Effects of Ramadan fasting on cardiovascular risk factors: a prospective observational study

Mohsen Nematy¹, Maryam Alinezhad-Namaghi², Masoud Mahdavi Rashed², Mostafa Mozhdehifard², Seyedeh Sania Sajjadi², Saeed Akhlaghi³, Maryam Sabery⁴, Seyed Amir R Mohajeri², Neda Shalaey², Mohsen Moohebaty⁵ and Abdolreza Norouzy^{2*}

Abstract

Background: Previous research has shown that Ramadan fasting has beneficial effects on cardiovascular risk factors, however there are controversies. In the present study, the effect of Ramadan fasting on cardiovascular risk factors has been investigated.

Method: This is a prospective observational study that was carried out in a group of patients with at least one cardiovascular risk factor (including history of documented previous history of either coronary artery disease (CAD), metabolic syndrome or cerebro-vascular disease in past 10 y). Eighty two volunteers including 38 male and 44 female, aged 29–70 y, mean 54.0 ± 10 y, with a previous history of either coronary artery disease, metabolic syndrome or cerebro-vascular disease were recruited. Subjects attended the metabolic unit after at least 10 h fasting, before and after Ramadan who were been fasting for at least 10 days. A fasting blood sample was obtained, blood pressure was measured and body mass index (BMI) was calculated. Lipids profile, fasting blood sugar (FBS) and insulin, homocysteine (hcy), high-sensitivity C-reactive protein (hs-CRP) and complete blood count (CBC) were analyzed on all blood samples.

Results: A significant improvement in 10 years coronary heart disease risk (based on Framingham risk score) was found (13.0 ± 8 before Ramadan and 10.8 ± 7 after Ramadan, $P < 0.001$, t test). There was a significant higher HDL-c, WBC, RBC and platelet count (PLT), and lower plasma cholesterol, triglycerides, LDL-c, VLDL-c, systolic blood pressure, body mass index and waist circumference after Ramadan ($P < 0.05$, t test). The changes in FBS, insulin, Homeostasis Model Assessment Insulin Resistance (HOMA-IR), hcy, hs-CRP and diastolic blood pressure before and after Ramadan were not significant ($P > 0.05$, t test).

Conclusions: This study shows a significant improvement in 10 years coronary heart disease risk score and other cardiovascular risk factors such as lipids profile, systolic blood pressure, weight, BMI and waist circumference in subjects with a previous history of cardiovascular disease.

Keywords: Ramadan, Fasting, hs-CRP, Homocysteine, FBS, Insulin resistance, Cardiovascular diseases, Risk factors

Introduction

Ramadan is a month for Muslims during which they abstain from eating, drinking and smoking from dawn to sunset.

During month of Ramadan there are changes in the quality of food and eating patterns. It might be due to the consumption of more carbohydrates and sweet foods, mainly in the form of two large meals at dawn

and sunset [1]. It has been established that a given nutrient ingested at an unusual time can induce different metabolic effects [2]; however, the physiological changes in Ramadan are not well known.

Cardiovascular diseases are the leading cause of death in the world [3]. Fasting during Ramadan is essentially a radical change in lifestyle for the period of one lunar month that may affect cardiovascular risk in patients with coronary artery disease (CAD) and cerebro-vascular disease (CVD) [4,5].

The most common risk factors associated with increased risk of atherosclerotic heart disease or stroke

* Correspondence: norouzya@mums.ac.ir

²Biochemistry and Nutrition Research Center and Department of Nutrition, Mashad, University of Medical Sciences, Mashad 91779-48564, Iran
Full list of author information is available at the end of the article

are abnormalities in plasma lipids and some coagulation and haemostatic factors, hypertension and smoking [6].

Lipids profile are affected by factors such as changes in the dietary habits, using different dietary fats, increased consumption of refined sugar, and reduced physical activity [7].

Many studies reported a significant reduction in type 2 diabetic patients' weight during Ramadan [8,9], some other reported a non-significant reduction in weight of these patients [10]. Bouguerra et al. showed Ramadan fasting has a moderate effect on glycaemia and lipoprotein levels in type 2 diabetic patients when previous metabolic control was quite good; but fasting induced more deterioration when previous control was poor [9]. Although, another study in Iran showed a deterioration of glycaemic control after Ramadan fasting in Type 2 diabetic patients that was more evident in patients using oral hypoglycaemic medication than diet-controlled patients [11]. Shariatpanahi et al. showed that the combined change in the number and timing of meals and portioning of the entire intake into only two meals per day may increase insulin sensitivity in Ramadan [8]. Homocysteine (Hcy) is a well-known mixed amino acid, intermediary on the metabolic pathway between methionine and cysteine [12]. Mild to moderate elevation of plasma hcy concentration has been known as carrying an increased risk for the development of atherosclerotic vascular disease [13]. Hyperhomocysteinemia is thus an independent risk factor for atherosclerosis, and impaired arterial endothelial function is detectable in healthy adults with hyperhomocysteinaemia [13].

High-sensitivity C-reactive protein (hs-CRP), a plasma inflammation marker, has been known to play a role in the development of cardiovascular diseases and is considered as a biomarker that is useful in the prediction of early cardiovascular risk [14]. Several prospective epidemiological studies have demonstrated a consistent relationship between higher C-reactive protein (CRP) levels and increased risk of cardiovascular events, including myocardial infarction, stroke, and cardiovascular death [15,16]. CRP may reflect a greater burden of atherosclerosis or alternatively may identify a high-risk atherosclerosis phenotype with an active inflammation and atherosclerotic plaque that is vulnerable to rupture [17,18].

Fasting is not meant to create excessive hardship for individuals who are not able to do that, the Islamic practice exempts many people from fasting, including those who are ill and fasting is harmful for them [19]. One of the common questions for patients with a history of cardiovascular disease is whether it is safe for them to fast during Ramadan. Yet, studies on the effects of Ramadan fasting on blood lipids, blood pressure, anthropometric parameters and other cardiovascular risk factors are scarce, and have given inconclusive results [9,10,20].

In current research, the effect of Ramadan fasting on cardiovascular risk factors including biochemical indices, blood pressure and main anthropometric parameters has been investigated.

Subjects and methods

Subjects

This was a prospective observational study that was carried out in a group of patients with at least one cardiovascular risk factor (including history of documented previous history of either coronary artery disease (CAD), metabolic syndrome or cerebro-vascular disease in past 10 y). Subjects' risk factors were compared before and after Ramadan month to see the effects of fasting on it.

This study was conducted according to the guidelines laid down in the Declaration of Helsinki and all procedures involving human subjects were approved by the Research Ethics Committee of Mashad University of Medical Sciences (approval number 87224). Written informed consent was obtained from all subjects. This study was conducted in the month of Ramadan during September/October 2008 in city of Mashad, Iran.

Volunteers were recruited from outpatient cardiology clinics in Imam Reza and Ghaem teaching hospitals via poster advertisement or direct invitation. Inclusion criteria were male and female between 29 and 70 years old with a history of documented previous history of either coronary artery disease (CAD), metabolic syndrome or cerebro-vascular disease in past 10 y. 76 patients were taking medications including 52 patients on antihypertensive agents, 47 patients on statins, 26 patients on oral antidiabetic medications including metformin and sulphonylureas and 46 patients on antiplatelet aggregation agents. Six patients did not take any medications.

Diagnosis of cardiovascular disease was based of cardiologist judgment.

Exclusion criteria were pregnant and lactating women, subjects with the age of below 18 y old and those who fasted for less than 10 d. Subjects were nonsmoker and volunteers with any type of acute inflammation (hs-CRP >5) were excluded. One hundred and one subjects participated in the first part of the study; however, 82 subjects completed the study. Seventeen patients were excluded because of fasting for less than 10 d and 2 subjects withdrew their consent for personal reasons.

Past medical history, drug and smoking history were obtained prior the study.

Blood sampling and data gathering

Subjects attended the metabolic unit after 10–12 h fasting, in two stages; between 7 d before and 2 first days of starting Ramadan and from the 27th of Ramadan till 6 d after end of this month. An 8 mL fasting blood sample was collected from the median cubital vein by using

vacuum sampling method. The blood samples were analyzed in an University affiliated laboratory.

Systolic and diastolic blood pressures were measured in duplicate by sphygmomanometers (Omron M3, M4, Omron Corporation, the Netherlands). If difference between two measurements was more than 5%, a third reading was performed and average of two near readings was considered as mean blood pressure. Food frequency questionnaire (FFQ) was completed in both stages before and after Ramadan at the same Ramadan in the matched BMI from the control group.

Anthropometric parameters such as height, body weight and waist circumference were measured in both phases. Body mass index (BMI) was calculated as body weight (kg) divided by squared height in meters (m²). Height and weight were measured, using standardized procedures. Height was measured using a portable stadiometer (OTM, Tehran, Iran). Height measurement was taken to the nearest 0.1 cm, without shoes, with the subject stretching to the maximum height and the head positioned in the Frankfort plane. Weight was measured using Rassa weight scale (Rassa, Tehran, Iran) to the nearest 100 g with subjects removing shoes and in light clothing. Waist circumference was measured to the nearest 0.1 cm at the mid-point between the lower rib and the upper margin of the iliac crest [21] in a horizontal plane by using a no stretching tape with an insertion buckle at one end.

Assays

Lipids profile, fasting blood sugar (FBS), fasting insulin, hs-CRP, hcy and complete blood count (CBC) were analyzed using standard methods. Homocysteine measurement was being performed on blood plasma and the other variables on blood serum.

Triglyceride (TG) and cholesterol were measured by GPO-PAP and CHOD-PAP methods respectively (Parsazmun kit, Karaj, Iran, for both). The direct method was used for determining the HDL and LDL by Human kits (Human Gesellschaft für Biochemica und Diagnostica GmbH, Max-Planck-Ring 21 – D-65205 Wiesbaden – Germany) and fasting blood glucose was measured by glucose-oxidize, GOD-PAP method [22] (Parsazmun kit, Karaj, Iran). Total hcy measurement was performed with immunoturbidometry, based on assays to co-substrate conversion product [23,24] (Demeditec Diagnostic GmbH, Lise-Meitner-strabe 2, D-24145 Kiel Germany). Insulin was measured, using radioimmunoassay methods (ELISA) with IRMA kit (Immunotecha.s. Prague, Czech Republic). Hs-CRP concentration was measured by immunoturbidometry method by auto analyzer [25] (Prestige 24i, Tokyo Boeki, Japan). Insulin resistance was estimated from the homeostasis model assessment of insulin resistance (HOMA-IR) originally described by

Matthews et al. [26] as calculated from the following equation:

$$\begin{aligned} &[\text{Fasting insulin concentration}(\mu\text{U/mL}) \\ &\times \text{fasting glucose concentration}(\text{mmol/L})]/22.5 \end{aligned}$$

[22] Cell blood counts were measured by KX-21 N Sysmex autoanalyser system (Sysmex Shanghai Ltd, Lujiazui, China).

Framingham coronary heart disease risk score was used to calculate the risk of CAD [27].

Data analysis

The Kolmogorov-Smirnov test was performed to assess the normal distribution. Differences between results were analyzed, using paired samples *t* test for normally distributed parameters and Wilcoxon Signed Ranks test for not normally distributed parameters (LDL, TG, FBS, HOMA-IR and hs-CRP). Relationship between dichotomous and quantitative variables was assessed by independent sample *t* test and correlation between quantitative and quantitative variables was evaluated by Pearson's correlation. Two-way repeated measures ANOVA were used to assess interactions among variables seem to be effective on parameter differences ($P < 0.15$ in Pearson's correlation or independent sample *t* test) on variables before and after Ramadan. Quantitative data were expressed as the mean \pm SD for all parameters. Statistical significance was considered at $P < 0.05$ for all tests. Relationship among cholesterol serum level and existing CAD, using statin family medication and hypertension (based on history) were evaluated by using two-way repeated measures ANOVA. These statistical analyses were conducted by using SPSS statistical software (version 11.5, SPSS Inc. Chicago, IL).

Results

Duration of daily fasting during Ramadan 2008 was between 14 h and 42 min in first day and 13 h and 35 min in last day in Mashad, Iran. The mean days subjects fasted in this study was 26.3 ± 5 (range 10–35 d).

Eighty two volunteers including 38 male and 44 female, age 29–70 y, mean 54.0 ± 10 y, with a previous history of either coronary artery disease (CAD) (60.9%), metabolic syndrome (67%) or cerebro-vascular disease (2.4%) in past 10 y completed the study. Twenty eight percent of them (23 volunteers) were suffering from both CAD and metabolic syndrome.

10 years coronary heart disease risk

This study shows a significant improvement in 10 years coronary heart disease risk, based on Framingham risk score (13.0 ± 8 before Ramadan and 10.8 ± 7 after Ramadan, $P < 0.001$, *t* test).

Lipids profile

The values of total cholesterol, triglycerides, VLDL-c, LDL-c, cholesterol/HDL and LDL/HDL ratio were significantly decreased ($P = 0.02$ for cholesterol and $P < 0.001$ for rest, Wilcoxon test for triglycerides and LDL, and paired t test for rest) and HDL-c increased significantly ($P < 0.001$, paired t test) after Ramadan fasting (Table 1).

It appears that there was no significant effects of CAD, hypertension and taking statin drugs on cholesterol mean level variations before and after Ramadan ($P = 0.12$, $P = 0.09$ and $P = 0.71$ respectively). Effect of Ramadan fasting on cholesterol was independent of existing CAD, hypertension and taking statins ($P = 0.04$, ANOVA).

Blood pressure

There was a significant decrease in systolic blood pressure (132.9 ± 16 mmHg, 129.9 ± 17 mmHg, $P = 0.03$, paired t test). Although, no significant change in diastolic blood pressure was found (80.2 ± 9 , 78.6 ± 11 mmHg, $P = 0.14$, paired t test).

FBS, insulin, HOMA-IR, homocysteine, hs-CRP

As it is shown in Table 2, FBS, insulin, HOMA-IR, hcy and hs-CRP concentrations showed no significant difference after Ramadan fasting ($P = 0.33$, $P = 0.58$, $P = 0.76$, $P = 0.06$ and $P = 0.07$ respectively, paired t test for insulin and hcy and Wilcoxon test for rest).

Anthropometric parameters

Body weight, BMI and waist circumference reduced significantly after Ramadan fasting (73.5 ± 13 , 71.7 ± 13 kg, 28.4 ± 4 , 27.7 ± 4 kg/m² and 98.3 ± 9 , 96.7 ± 9 cm respectively and $P < 0.001$, paired t test).

Hematological parameters

Table 3 illustrates the values regarding hematological parameters before and after Ramadan. The mean concentrations of all these parameters (except hemoglobin and Hematocrit) showed significant changes when compared before and after Ramadan ($P = 0.25$ and $P = 0.32$

Table 1 Effect of Ramadan fasting on lipids profile before and after Ramadan (n = 82)

Biochemical variables	Before Ramadan	After Ramadan	p-values
	$\bar{x} \pm SD^3$	$\bar{x} \pm SD^3$	
Cholesterol(mg/dL)	193.4 ± 51	184.3 ± 42	0.023
Triglycerides (mg/dL)	224.8 ± 129	182.9 ± 112	0.000
HDL-c (mg/dL)	43.0 ± 9	47.9 ± 8	0.000
Cholesterol/HDL(ratio)	4.5 ± 1	3.9 ± 1	0.000
VLDL-c (mg/dL)	38.1 ± 14	33.8 ± 17	0.000
LDL-c (mg/dL)	109.96 ± 46	96.83 ± 35	0.000
LDL/HDL (ratio)	2.5 ± 0.8	2.0 ± 0.6	0.000

³SD = standard deviation.

Table 2 Effect of Ramadan fasting on some clinical and biological parameters before and after Ramadan (n = 65 for hs-CRP and 82 for rest)

Biochemical variables	Before Ramadan	After Ramadan	p-values
	$\bar{x} \pm SD^3$	$\bar{x} \pm SD^3$	
FBS ¹ (mmol/l)	124.7 ± 48	128.1 ± 54	0.327
insulin (μU/mL)	12.4 ± 9.5	11.8 ± 7.6	0.583
HOMA-IR (μUmol/l ²)	3.8 ± 3	3.6 ± 3	0.093
Hcy ² (μmol/l)	10.5 ± 3.8	11.5 ± 3.9	0.065
hs-CRP (mg/L)	1.0 ± 1	0.9 ± 1	0.160

¹FBS, fasting blood sugar.

²Hcy, homocysteine.

³SD = standard deviation.

respectively, paired t test). There was a significant higher WBC, RBC, platelet count (PLT) ($P = 0.01$, $P < 0.001$, $P < 0.001$ respectively, paired t test), and significant lower MCV and MCH ($P < 0.001$, paired t test) after Ramadan.

There was no significant correlation among number of fasting days and studied parameters in this study.

Food records

FFQ results are shown in Table 4. The data show a statistically non-significant reduction in energy and other macronutrients intake during fasting in month of Ramadan.

Discussion

This study shows a significant improvement in 10 years coronary heart disease risk, based on Framingham risk score. To the best of our knowledge, there was no similar study among cardiovascular patients to assess the effect of Ramadan fasting on cardiovascular risks. Most of the studies evaluated changes of some cardiovascular risk factors among metabolic syndrome, diabetic, hyperlipidemic patients and or healthy adults.

There was a significant higher HDL-c and lower cholesterol, triglycerides, LDL-c, VLDL-c levels, cholesterol/

Table 3 Effect of Ramadan fasting on hematological factors before and after Ramadan (Mean values with standard deviation) (n = 82)

Hematological factors	Before Ramadan	After Ramadan	p-values
	$\bar{x} \pm SD^3$	$\bar{x} \pm SD^3$	
WBC (1000/μl)	6.6 ± 1.5	6.9 ± 1.5	0.011
RBC (mil/μl)	4.7 ± 0.3	4.9 ± 0.4	0.000
HGB ¹ (g/dL)	13.6 ± 1.2	13.7 ± 1.0	0.254
MCV (fl)	87.6 ± 5.0	83.7 ± 4.2	0.000
MCH (pg)	28.8 ± 2.1	27.9 ± 1.8	0.000
PLT ² (1000/μl)	219.2 ± 54.1	242.7 ± 56.2	0.000

¹HGB, Hemoglobin.

²PLT, Platelet count.

³SD = standard deviation.

Table 4 Effect of Ramadan fasting on energy and nutrient intake of control group before and after Ramadan (Mean values with standard deviation) (n =82)

Consumption	Before Ramadan	After Ramadan	p-values
	$\bar{x} \pm SD^1$	$\bar{x} \pm SD^1$	
Energy (kcal)	2588.6 ± 1485	2449.3 ± 1179	0.337
Carbohydrates(g/day)	327.8 ± 179	321.0 ± 171	0.746
Protein (g/day)	96.7 ± 59	85.4 ± 45	0.071
Fat (g/day)	100.0 ± 64	92.6 ± 42	0.263
Fiber(g/day)	22.3 ± 14	20.2 ± 11	0.180

¹SD = standard deviation.

HDL and LDL/HDL ratios after Ramadan in the present study. In Helsinki Heart Study a 10% reduction in TC, LDL-c (11%), 35% decrease in triglycerides and 11% increase in HDL-c after the intervention with pharmacological agent was associated with a 34% reduction in cardiovascular event [28]. A 2 unit increment of TC/HDL adds an excess of 68% to CHD risk [27,29]. Our study demonstrates significant improvement (-0.6) in TC/HDL and other lipids profile after Ramadan fasting which might be associated with a nearly 20% reduction in the risk of CHD.

A reduction in cholesterol [30,31], and triglycerides levels [30-32], has been reported in previous studies; however, The result of some other reports showed no significant changes in cholesterol [1,12,32-34] and triglycerides [1,2,33,34]. In a study in Kuwait on sixteen sedentary healthy adult males, there was no significant change in TG level after Ramadan [33]. This discrepancy was possibly due to small sample size or different Ramadan diet pattern in Kuwait study which includes more sweetened foods. Also, there are some reports about increased cholesterol [10,35] and triglycerides [10] after Ramadan fasting. There is a tendency for higher simple carbohydrate consumption during Ramadan [36].

Improvement in HDL-c has been shown in some studies after fasting in Ramadan [2,30,31,34], while some other studies reported no significant change [1,32] or decrease [9,10] in HDL-c after Ramadan.

Significant decline in LDL-c has been reported in some studies after Ramadan fasting[2,30,31]. Also, no significant change [1,12,32,34] or significant increase [9,10,35] of LDL-c has been reported in other investigations.

It seems that the effect of Ramadan fasting on serum lipid levels may be closely related to the dietary habits and other life style changes [2,7].

There were significant decrease in systolic blood pressure and no significant change in diastolic blood pressures after Ramadan fasting in the present study. However, there are reports about significant decrease [1], no significant change [9,20,30,35] and significant increase [33] in systolic blood pressure.

There was no significant change in FBS, fasting insulin and insulin-resistance (HOMA-IR) after Ramadan fasting in the present study. Some studies showed no significant changes in the serum level of glucose [1,33,35]. While, the others reported higher [9] or lower [2,10,32] FBS after Ramadan fasting. These controversies may be explained by different food habits, amount of calorie intake [30], the number of fasting days, period of daily fasting, time of sampling, genetic tendency and daily activity in different reports. There is a report about significant increase in insulin sensitivity (1/HOMA IR) in subjects with the metabolic syndrome [8] and significant decrease in insulin and insulin resistance among men patients with type II diabetes after Ramadan fasting [35]. The lack of improvement in HOMA-IR or FBS despite reduction in body weight may be due to decrease in total body water content instead of body fat content after Ramadan fasting.

The value of hcy level did not change significantly after Ramadan in the present study. Although, Aksungar's study has been reported significant decrease in hcy after Ramadan fasting in 40 healthy volunteers [12].

Body weight and BMI decreased significantly in the present study (-1.4 kg and -0.7 kg/m² respectively, *P* <0.001). Most reports are similar to these results [31,32]. Although, there are some reports about non-significant changes in body weight [37] and BMI [37,38] after Ramadan fasting.

In another study, no change was seen concerning anthropometric parameters [7].

Limitations

There was no control group in this study. Pre Ramadan sampling in Ramadan was performed in the morning after an overnight fasting, while post Ramadan blood sampling was performed in the afternoon, after at least 10 h fasting during day time. This difference in the time of blood sampling was inevitable to have at least 10 h fasting. As the circadian rhythms of nutrition-related biological variables shows some degree of changes during Ramadan [39], there might be concerns about amount of daily activities or circadian rhythm of hormones like cortisol which might affect FBS levels. In the present study, we had only two measurements during the study, possibly evaluation in the middle of the month and after a time after Ramadan would reveal more details and shows us the trend of changes. Level of physical activity of the study population was not measured in this study. FFQ was completed for a control group not our subjects as this study was part of the bigger study that food data were completed as a separate study for a 500 volunteers to assess food pattern changes in Ramadan.

A multi-center study with larger sample size and other habitual changes controlled with matched control group is warranted.

Conclusion

This study shows a significant improvement in 10 years coronary heart disease risk score and other cardiovascular risk factors such as weight, BMI and waist circumference in subjects with a previous history of cardiovascular disease after an average of 26 d fasting during month of Ramadan. Result from this study suggested that Ramadan fasting may be useful to improve CVD risk factors.

Competing interests

None of the authors had a personal or financial conflict of interest.

Authors' contributions

MN, AN, MAN: were responsible for study concept and design, MAN, MMR, MMF, SA, MS, SAM, MM SSS and NS: contributed for the design and conduct of the study, acquired the data, analyzed and interpreted data, performed statistical analysis and contributed to the writing manuscript. All authors read and approved the submitted manuscript.

Source of support

This work was supported by a grant from Mashad University of Medical Sciences.

Acknowledgments

This study was supported by a research grant from Mashad University of Medical Sciences. We are most grateful to all subjects for their participation.

Author details

¹Department of Nutrition, Mashad, Biochemistry and Nutrition, Endoscopic & Minimally Invasive Surgery, and Cancer Research Centers, University of Medical Sciences, Mashad 91779-48564, Iran. ²Biochemistry and Nutrition Research Center and Department of Nutrition, Mashad, University of Medical Sciences, Mashad 91779-48564, Iran. ³Department of Community Medicine, Mashad, University of Medical Sciences, Mashad 91779-48564, Iran. ⁴Iranian Applied Research Center for Health and Sustainable Development (IRCPHD), North Khorasan University of Medical Sciences, North Khorasan, Iran. ⁵Department of Cardiology, Ghaem Teaching Hospital, Mashad, University of Medical Sciences, Mashad 91766-99199, Iran.

Received: 17 September 2011 Accepted: 30 July 2012

Published: 10 September 2012

References

- Sarraaf-Zadegan N, Atashi M, Naderi GA, Baghai AM, Asgary S, Fatehifar MR, et al: **The effect of fasting in Ramadan on the values and interrelations between biochemical, coagulation and hematological factors.** *Ann Saudi Med* 2000, **20**(5-6):377-381.
- Ziaee V, Razaee M, Ahmadinejad Z, Shaikh H, Yousefi R, Yarmohammadi L, et al: **The changes of metabolic profile and weight during Ramadan fasting.** *Singapore Med J* 2006, **47**(5):409-414.
- WHO: **The global burden of disease: 2004 update. THE REPORT IN SECTIONS. Part 2: Causes of death;** 2009. Internet: http://www.who.int/healthinfo/global_burden_disease/2004_report_update/en/index.html. (accessed 11 November 2009).
- Al SJ, Bener A, Hajar HA, Numan MT: **Does hospitalization for congestive heart failure occur more frequently in Ramadan: a population-based study (1991-2001).** *Int J Cardiol* 2004, **96**(2):217-221.
- Akhan G, Kutluhan S, Koyuncuoglu HR: **Is there any change of stroke incidence during Ramadan?** *Acta NeurolScand* 2000, **101**(4):259-261.
- Sarraaf-Zadegan N, Boshnam M, Rafiei M: **Risk factors for coronary artery disease in Isfahan, Iran.** *Eur J Public Health* 1999, **9**(1):20-26.
- Afrasiabi A, Hassanzadeh S, Sattarivand R, Mahboob S: **Effects of Ramadan fasting on serum lipid profiles on 2 hyperlipidemic groups with or without diet pattern.** *Saudi Med J* 2003, **24**(1):23-26.
- Shariatpanahi ZV, Shariatpanahi MV, Shahbazi S, Hossaini A, Abadi A: **Effect of Ramadan fasting on some indices of insulin resistance and components of the metabolic syndrome in healthy male adults.** *Br J Nutr* 2008, **100**(1):147-151.
- Bouguerra R, Jabrane J, Maatki C, Ben SL, Hamzaoui J, El KA, et al: **Ramadan fasting in type 2 diabetes mellitus.** *Ann Endocrinol (Paris)* 2006, **67**(1):54-59.
- Khaled BM, Bendahmane M, Belbraouet S: **Ramadan fasting induces modifications of certain serum components in obese women with type 2 diabetes.** *Saudi Med J* 2006, **27**(1):23-26.
- Norouzy A, Mohajeri SM, Shakeri S, Yari F, Sabery M, Philippou E, et al: **Effect of Ramadan fasting on glycaemic control in patients with Type 2 diabetes.** *J Endocrinol Invest* 2011, Oct 6.[Epub ahead of print].
- Aksungar FB, Topkaya AE, Akyildiz M: **Interleukin-6, C-reactive protein and biochemical parameters during prolonged intermittent fasting.** *Ann NutrMetab* 2007, **51**(1):88-95.
- Graham IM, Daly LE, Refsum HM, Robinson K, Brattstrom LE, Ueland PM, et al: **Plasma homocysteine as a risk factor for vascular disease. The European Concerted Action Project.** *JAMA* 1997, **277**(22):1775-1781.
- Shroff GR, Cen YY, Duprez DA, Bart BA: **Relationship between carotid artery stiffness index, BNP and high-sensitivity CRP.** *J Hum Hypertens* 2009, **23**(12):783-787.
- Rost NS, Wolf PA, Kase CS, Kelly-Hayes M, Silbershatz H, Massaro JM, et al: **Plasma concentration of C-reactive protein and risk of ischemic stroke and transient ischemic attack: the Framingham study.** *Stroke* 2001, **32**(11):2575-2579.
- Koenig W, Sund M, Frohlich M, Fischer HG, Lowel H, Doring A, et al: **C-Reactive protein, a sensitive marker of inflammation, predicts future risk of coronary heart disease in initially healthy middle-aged men: results from the MONICA (Monitoring Trends and Determinants in Cardiovascular Disease) Augsburg Cohort Study, 1984 to 1992.** *Circulation* 1999, **99**(2):237-242.
- Verma S, Wang CH, Li SH, Dumont AS, Fedak PW, Badiwala MV, et al: **A self-fulfilling prophecy: C-reactive protein attenuates nitric oxide production and inhibits angiogenesis.** *Circulation* 2002, **106**(8):913-919.
- Falk E, Shah PK, Fuster V: **Coronary plaque disruption.** *Circulation* 1995, **92**(3):657-671.
- The office of the supreme leader, Sayyid Ali Khamenei. **Practical lows of Islam.** Internet: 2009. <http://www.leader.ir/langs/en/> (accessed 10 November 2009).
- Perk G, Ghanem J, Amar S, Ben-Ishay D, Bursztyn M: **The effect of the fast of Ramadan on ambulatory blood pressure in treated hypertensives.** *J Hum Hypertens* 2001, **15**(10):723-725.
- Lohman TG, Roche AF, Martorell R: **Anthropometric Standardization Reference Manual.** IL: Kinetics Books Champaign; 1988.
- Burtis CA, Ashwood ER, Bruns DE: **Tietz Textbook of Clinical Chemistry AND Molecular Diagnostics.** Fourthth edition. Philadelphia: Elsevier Saunders; 2001.
- Frantzen F, Faaren AL, Alfheim I, Nordhei AK: **Enzyme conversion immunoassay for determining total homocysteine in plasma or serum.** *ClinChem* 1998, **44**(2):311-316.
- Tan Y, Tang L, Sun X, Zhang N, Han Q, Xu M, et al: **Total-homocysteine enzymatic assay.** *ClinChem* 2000, **46**(10):1686-1688.
- Azarpazhooh MR, Mobarra N, Parizadeh SM, Tavallaie S, Bagheri M, Rahsepar AA, et al: **Serum high-sensitivity C-reactive protein and heat shock protein 27 antibody titers in patients with stroke and 6-month prognosis.** *Angiology* 2010, **61**(6):607-612.
- Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF, Turner RC: **Homeostasis model assessment: insulin resistance and beta-cell function from fasting plasma glucose and insulin concentrations in man.** *Diabetologia* 1985, **28**(7):412-419.
- Wilson PW, D'Agostino RB, Levy D, Belanger AM, Silbershatz H, Kannel WB: **Prediction of coronary heart disease using risk factor categories.** *Circulation* 1998, **97**(18):1837-1847.
- Manninen V, Huttunen JK, Heinonen OP, Tenkanen L, Frick MH: **Relation between baseline lipid and lipoprotein values and the incidence of coronary heart disease in the Helsinki Heart Study.** *Am J Cardiol* 1989, **63**(16):42H-47H.
- Kinosian B, Glick H, Garland G: **Cholesterol and coronary heart disease: predicting risks by levels and ratios.** *Ann Intern Med* 1994, **121**(9):641-647.
- Fakhrzadeh H, Larjani B, Sanjari M, Baradar-Jalili R, Amini MR: **Effect of Ramadan fasting on clinical and biochemical parameters in healthy adults.** *Ann Saudi Med* 2003, **23**(3-4):223-226.

31. Adlouni A, Ghalim N, Benslimane A, Lecerf JM, Saile R: **Fasting during Ramadan induces a marked increase in high-density lipoprotein cholesterol and decrease in low-density lipoprotein cholesterol.** *Ann NutrMetab* 1997, **41**(4):242–249.
32. Khatib FA, Shafagaj YA: **Metabolic alterations as a result of Ramadan fasting in non-insulin-dependent diabetes mellitus patients in relation to food intake.** *Saudi Med J* 2001, **25**(12):1858–1863.
33. Ramadan J: **Does fasting during Ramadan alter body composition, blood constituents and physical performance?** *Med PrincPract* 2002, **11**(Suppl 2):41–46.
34. Aksungar FB, Eren A, Ure S, Teskin O, Ates G: **Effects of intermittent fasting on serum lipid levels, coagulation status and plasma homocysteine levels.** *Ann NutrMetab* 2005, **49**(2):77–82.
35. Yarahmadi S, Larijani B, Bastanhagh MH, Pajouhi M, Baradar JR, Zahedi F, *et al*: **Metabolic and clinical effects of Ramadan fasting in patients with type II diabetes.** *J Coll Physicians Surg Pak* 2003, **13**(6):329–332.
36. Frost G, Pirani S: **Meal frequency and nutritional intake during Ramadan: a pilot study.** *Hum Nutr Appl Nutr* 1987, **41**(1):47–50.
37. Yucel A, Degirmenci B, Acar M, Albayrak R, Haktanir A: **The effect of fasting month of Ramadan on the abdominal fat distribution: assessment by computed tomography.** *Tohoku J Exp Med* 2004, **204**(3):179–187.
38. Aybak M, Turkoglu A, Sermet A, Denli O: **Effect of Ramadan fasting on platelet aggregation in healthy male subjects.** *Eur J Appl Physiol Occup Physiol* 1996, **73**(6):552–556.
39. Iraki L, Bogdan A, Hakkou F, Amrani N, Abkari A, Touitou Y: **Ramadan diet restrictions modify the circadian time structure in humans. A study on plasma gastrin, insulin, glucose, and calcium and on gastric pH.** *J Clin Endocrinol Metab* 1997, **82**(4):1261–1273.

doi:10.1186/1475-2891-11-69

Cite this article as: Nematy *et al.*: Effects of Ramadan fasting on cardiovascular risk factors: a prospective observational study. *Nutrition Journal* 2012 **11**:69.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

 **BioMed** Central