



Effect of a Probiotic Preparation (VSL#3) on Cardiovascular Risk Parameters in Critically-Ill Patients

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ABSTRACT

Introduction: Cardiovascular disease (CVD) counts for a major portion of morbidity and mortality globally mostly accompanied by lipid abnormalities. Being at increased risk of cardiac injury, critically ill patients suffer from various lipid disorders. Lipid homeostasis has been efficiently restored by the introduction of probiotics. The aim of this trial was to determine the effect of probiotics on inflammation, antioxidant capacity and lipid peroxidation in critically ill patients.

Methods: Forty patients admitted to the intensive care unit were randomized to receive placebo or probiotic for 7 days. Serum levels of triglyceride (TG), total cholesterol, HDL-C, LDL-C and high-sensitivity C-reactive protein (hs-CRP) were measured before initiation of the study and on the 7th day.

Results: There was a significant difference between two groups regarding the levels of TG, HDL and hs-CRP at the end of the study ($P=0.043$, <0.001 and 0.003 , respectively); however, there was not a significant difference in total cholesterol and LDL-C levels.

Conclusion: Administration of probiotics in critically ill patients reduced the levels of TG and hs-CRP and increased HDL-C levels. However, no significant change was detected in levels of total cholesterol or LDL-C.

Introduction

Cardiovascular disease (CVD) counts for a major portion of morbidity and mortality globally mostly accompanied by lipid abnormalities.¹ Abnormal levels of high density lipoprotein-cholesterol (HDL-C), low density lipoprotein-cholesterol (LDL-C) and triglyceride (TG) could accompany a higher prevalence of CVD.² Elevated LDL-C and reduced HDL-C levels are already-confirmed CVD risk factors with evidence suggesting that significant increase in HDL-C could be set as an important therapeutic goal.³ While critically ill patients frequently suffer from metabolic disturbances such as various lipid disorders, elevated triglyceride levels followed by increase in very low-density lipoprotein (VLDL), and low circulating HDL-C are their main characteristic in critically ill patients. LDL-C levels are also decreased.^{4,5} Commonly seen in critically ill patients, systemic inflammation and sepsis also accompany severe metabolic imbalances including decreases in HDL-C and LDL-C and high levels of TG.⁶⁻⁹ Association between systemic inflammation and

lipid metabolism has been clearly established in many studies.¹⁰ In sepsis, enhanced production of hepatic VLDL and/or inhibited peripheral and hepatic VLDL clearance lead to increase in plasma TG within VLDL. In contrast, sepsis decreases plasma cholesterol within LDL-C and mainly HDL-C.¹¹ Furthermore, lipid metabolism disorders are accompanied by worse prognosis in critically ill patients.¹² Indices of lipid metabolism have been found to be related to the severity of illness, the occurrence of sepsis and survival in critically ill patients.¹³ In addition, ICU patients are at increased risk of cardiac disorders due to the underlying presence of coronary circulation atherosclerosis and other non-cardiac factors including increased tissue oxygen demands, anemia, sepsis, mechanical ventilation, and hemodynamic instability. Furthermore, assessment of myocardial injury as an independent determinant of hospital mortality would make it possible to recognize ICU patients at augmented risk of death.¹⁴

The incidence of myocardial injury is usually defined by elevated levels of cardiac troponin I.15 Levels

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of high-sensitivity C-reactive protein (hs-CRP), a marker of systemic inflammation and a mediator of atherothrombotic disease, is in significant correlation with cardiovascular disease risk; inflammation is a major factor in atherothrombotic disease.¹⁶ Global risk assessment uses hs-CRP as an index in the primary prevention of cardiovascular disease.¹⁷

Probiotics, live microorganisms providing a health benefit on their host when used in adequate amounts¹⁸, are of beneficial effects in the prevention and treatment of different disease. Heterogeneous results have been given on the effects of probiotics consumption on the plasma lipid profile in different studies; yet, positive changes in lipid profile have been observed with the use of probiotics.^{19,20} In addition, probiotics not only are of useful effects on cellular immunity but also help to preserve the balance between pro- and anti-inflammatory cytokines.²¹ Despite the introduction of numerous pharmacologic lipid lowering therapies, there are known side effects.²² Hence, the use of probiotics considering their natural and safe properties and low ability of triggering adverse effects²³ seems logical for serum lipid improvement.

The purpose of this randomized clinical trial was to determine the effect of probiotic containing lactobacillus, bifidobacterium and streptococcus thermophilus on lipid profile and hs-CRP levels in critically ill patients in the intensive care unit.

Methods

After approval of ethics committee of Tabriz University of Medical Sciences and obtaining written informed consent from the patients or their legal guardians, 40 patients who were admitted between December of 2011 and October of 2012 to the ICU of the Shohada Hospital (Tabriz, Iran) were enrolled in this trial. Inclusion criteria were critically ill patients who were expected to stay in ICU for at least 7 days aged 18 to 40 years old receiving enteral nutrition. Exclusion criteria were patients who could not tolerate enteral nutrition, those with unstable hemodynamic, immune disorders, history of diabetes mellitus and hyperlipidemia, intestinal obstruction or ischemia, cancer and patients who were expected to expire in the next 24 hours. Patients were randomized to two 20-person groups; the first group received standard treatment plus placebo and the second group received standard treatment plus VSL#3, 2 sachets daily for 7 days. Each sachet of probiotics (VSL#3; VSL Pharmaceuticals, Ft Lauderdale, FL) contained 450 billion viable lyophilized bacteria consisting of 4 strains of *Lactobacillus* (*L. casei*, *L. plantarum*, *L. acidophilus*, and *L. delbrueckii* subsp. *Bulgaricus*), 3 strains of *Bifidobacterium* (*B. longum*, *B. breve*, and *B. infantis*) and *Streptococcus salivarius* subsp. *Thermophilus*.

All patients received enteral nutrition with Fresubin original fibre (*Fresenius Kabi, Homburg, Germany*) at the first 24-hour of admission via nasogastric tube.

Energy requirements were calculated as 25–30 kcal/kg. Fasting blood samples were obtained from each patient to evaluate lipid profile and hs-CRP on days 1 and 7. Serum triglyceride, total cholesterol, and HDL-cholesterol concentrations were determined by routine clinical assays using commercial kits on an automated analyzer. Serum LDL-cholesterol concentrations were calculated based on the Friedewald calculation ($LDL = TC - HDL - TG/5$). Data were analyzed by SPSS 16.

P values < 0.05 were considered significant for all statistical tests.

Results

There were 20 patients in each group. No significant differences in patients' demographic data between two groups were observed (Table 1). Levels of TG, total cholesterol, LDL-C, HDL-C and hs-CRP at the baseline and at the end of the study are shown in Table 2. There was a significant difference between two groups in levels of TG, HDL and hs-CRP at the end of the study ($P = 0.043$, <0.001 and 0.003, respectively); however, there was no significant difference in total cholesterol and LDL-C levels. In addition, levels of TG significantly decreased while levels of HDL-C significantly increased after the treatment period in probiotic group ($P = 0.001$ and 0.041, respectively).

Levels of hs-CRP in both groups are shown in Figure 1.

Table 1. Demographic data of patients

	Control group	Probiotic group	P value
Age (year)	35.60±5.03*	33.60±5.50	0.238
Male/Female	14/6	13/7	0.500
BMI	24.70±3	24.30±2.92	0.677

*mean±SD, BMI: Body Mass Index

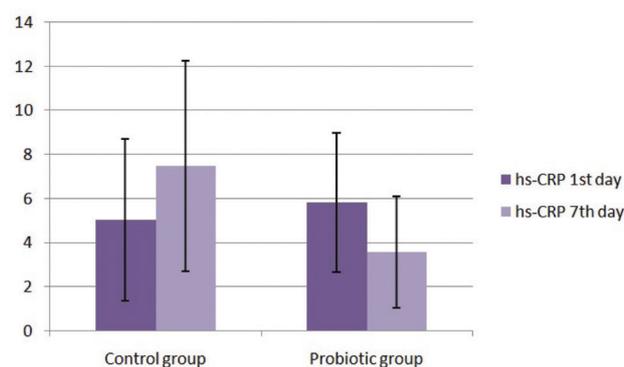


Figure 1. Mean (±SD) hs-CRP levels in two groups of patients.

Discussion

The present study used a double-blind, placebo-controlled, randomized design to determine the effects of probiotics on lipid profile and inflammation in critically ill, enterally

Table 2. Levels of lipid profile and hs-CRP at baseline and at the end of the study

		Control group	Probiotic group	P value
TG(mg/dl)	1 st day	146.85±68.38 [*]	178.70±57.98	0.120 [†]
	7 th day	171.70±89.97	126.70±34.07	0.043
Cholesterol(mg/dl)	1 st day	129.85±30.00	150.65±47.88	0.108
	7 th day	152.55±32.46	137.45±31.32	0.143
LDL(mg/dl)	1 st day	61.48±26.47	76.79±44.10	0.191
	7 th day	86.96±31.90	68.61±31.38	0.075
HDL(mg/dl)	1 st day	39.00±14.29	38.00±11.63	0.810
	7 th day	31.25±8.40	43.50±10.01	<0.001
hs-CRP(μg/ml)	1 st day	5.02±3.69	5.82±3.16	0.471
	7 th day	7.48±4.79	3.56±2.52	0.003

^{*}mean±SD
[†]Independent-samples t-test

fed patients. A significant effect on the levels of TG and HDL-C was observed following adding probiotics to the enteral feeding of the patients; however, no significant effect was attained for total cholesterol and LDL-C in our study. In a study performed on 32 healthy subjects, it was reported that administration of probiotics was associated with a significant reduction in serum total cholesterol, LDL-C and TG and increase in HDL-C.²⁴ In another study on type 2 diabetic patients, no noteworthy changes were reported in TG and HDL-C, while probiotic group had significantly decreased total cholesterol levels: HDL-C ratio and LDL-C: HDL-C ratio.²⁵ In a meta-analysis, it was indicated that a probiotics rich diet decreases total cholesterol and LDL-C in participants with high, borderline high and normal cholesterol levels.¹⁹ The varied effects of probiotics on lipid profile observed in the studies might be as a result of the type of the patient population and also the used probiotic. In our study, probiotics could not have applied their LDL-C lowering effects, as we assessed their effects in critically ill population who generally present Low LDL-C levels.⁴

The patients who received probiotic showed a significant reduction in hs-CRP levels. Hs-CRP as a marker of systemic inflammation is a strong independent predictor of risk of future myocardial infarction, stroke, peripheral arterial disease, and vascular death.²⁶⁻³⁶ In the present study, a significant reduction in hs-CRP concentrations following the treatment course with probiotics was observed. Similar findings were reported by Asemi et al. who studied the effect of daily consumption of probiotic yoghurt vs. conventional yogurt on inflammatory factors in pregnant women. Their results showed that the probiotic yogurt decreased serum hs-CRP level, whereas there was no significant change in serum hs-CRP level in the conventional yogurt group.³⁷ In another study which compared a multifunctional (active) diet (AD) including a probiotic strain (*Lactobacillus plantarum*) with a control diet (CD) in healthy adults, CD did not alter the measured metabolic variables; however, AD changed hs-CRP levels significantly.³⁸ Yet, the results of this study cannot be credited to the sole probiotic; as a mixed multifunctional diet was used and the obtained

results might have been due to the use of antioxidant-rich foods, oily fish or wholegrain products.

In conclusion, the results of this randomized trial suggest that administration of probiotics in critically ill patients compared to placebo-treated patients significantly reduces the levels of TG and hs-CRP and increases HDL-C levels. However, it does not significantly affect the levels of total cholesterol and LDL-C. Hence, further studies with larger sample sizes are required to clarify their usefulness in this group of patients.

Ethical issues: The study was approved by the institutional review board, and written informed consent was obtained from all participants.

Competing interests: The authors had no competing interests to declare in relation to this article.

References

1. WHO. The Global Burden of Disease: 2004 Update. Genève, Switzerland: World Health Organization; 2008.
2. Lozano JV, Pallarés V, Cea-Calvo L, Listerri JL, Fernández-Pérez C, Martí-Canales JC, et al. Serum lipid profiles and their relationship to cardiovascular disease in the elderly: the PREV-ICTUS study. *Curr Med Res Opin* 2008; 24: 659-70.
3. Evans M, Roberts A, Davies S, Rees A. Medical lipid-regulating therapy: current evidence, ongoing trials and future developments. *Drugs* 2004; 64:1181-96.
4. Gordon BR, Parker TS, Levine DM, Saal SD, Wang JC, Sloan BJ, et al. Low lipid concentrations in critical illness: implications for preventing and treating endotoxemia. *Crit Care Med* 1996; 24: 584-9.
5. Barlage S, Gnewuch C, Liebisch G, Wolf Z, Audebert FX, Glück T, et al. Changes in HDL-associated apolipoproteins relate to mortality in human sepsis and correlate to monocyte and platelet activation. *Intensive Care Med* 2009; 35: 1877-85.
6. Mahmoodpoor A, Eslami K, Mojtahedzadeh M, Najafi A, Ahmadi A, Dehnadi-Moghadam A, et al. Examination of Setarud(IMOD) in the management of patients with severe sepsis. *DARU* 2010; 18: 23- 8.
7. Hamishehkar H, Beigmohammadi MT, Abdollahi M, Ahmadi A, Mahmoodpour A, Mirjalili MR, et al. Identification of enhanced cytokine generation following sepsis. Dream of Magic bullet for mortality prediction and therapeutic evaluation. *DARU* 2010; 18: 155- 62.
8. Eslami K, Mahmoodpoor A, Ahmadi A, Abdollahi M, Kamali K, Mousavi S, et al. Positive effect of septimebTM on mortality rate in severe sepsis: a novel non antibiotic strategy. *DARU J Pharm*

Sci 2012; 20: 40.

9. Tabeeefar H, Beigmohammadi MT, Javadi MR, Abdollahi M, Mahmoodpoor A, Ahmadi A, et al. Effects of pantoprazol on systemic and gastric pro- and anti-inflammatory cytokines in critically ill patients. **Iranian Journal of Pharmaceutical Research** 2012, 11: 1051-58.

10. Wendel M, Paul R, Heller AR. Lipoproteins in inflammation and sepsis. II. Clinical aspects. **Intensive Care Med** 2007; 33:25-35.

11. Berbée JF, Havekes LM, Rensen PC. Apolipoproteins modulate the inflammatory response to lipopolysaccharide. **J Endotoxin Res** 2005; 11: 97-103.

12. Cappi SB, Noritomi DT, Velasco IT, Curi R, Loureiro TC, Soriano FG. Dyslipidemia: a prospective controlled randomized trial of intensive glycemic control in sepsis. **Intensive Care Med** 2012; 38: 634-41.

13. Lind L, Lithell H. Impaired glucose and lipid metabolism seen in intensive care patients is related to severity of illness and survival. **Clin Intensive Care** 1994; 5:100-105.

14. Quenot JP, Le Teuff G, Quantin C, Doise JM, Abrahamowicz M, Masson D, et al. Myocardial injury in critically ill patients: relation to increased cardiac troponin I and hospital mortality. **Chest** 2005; 128: 2758-64.

15. Guest TM, Ramanathan AV, Tuteur PG, Schechtman KB, Ladenson JH, Jaffe AS. Myocardial injury in critically ill patients. A frequently unrecognized complication. **JAMA** 1995; 273:1945-9.

16. Ridker PM. High-sensitivity C-reactive protein and cardiovascular risk: rationale for screening and primary prevention. **Am J Cardiol** 2003; 92:17K-22K.

17. Ridker PM. High-Sensitivity C-Reactive Protein: Potential Adjunct for Global Risk Assessment in the Primary Prevention of Cardiovascular Disease Paul M. Ridker, MD, MPH) **Circulation**. 2001; 103:1813-8.

18. Food and Agriculture Organization (FAO). Health and nutritional properties of probiotics in food including powder milk with live lactic acid bacteria: report of a joint FAO/WHO expert consultation on evaluation of health and nutritional properties of probiotics in food including powder milk with live lactic acid bacteria. Geneva: Food and Agriculture Organization; 2001.

19. Guo Z, Liu XM, Zhang QX, Shen Z, Tian FW, Zhang H, et al. Influence of consumption of probiotics on the plasma lipid profile: a meta-analysis of randomised controlled trials. **Nutr Metab Cardiovasc Dis** 2011; 21: 844-50.

20. Sadrzadeh-Yeganeh H, Elmadfa I, Djazayeri A, Jalali M, Heshmat R, Chamary M. The effects of probiotic and conventional yoghurt on lipid profile in women. **Br J Nutr** 2010; 103: 1778-83.

21. Isolauri E, Kirjavainen P, Salminen S. Probiotics: a role in the treatment of intestinal infection and inflammation? **Gut** 2002; 50: 54-9.

22. Nawarskas JJ. HMG-CoA reductase inhibitors and coenzyme Q10. **Cardiol Rev** 2005; 13: 76-9.

23. Roberfroid M. Prebiotics: The Concept Revisited. **J Nutr** 2007; 137: 830-7.

24. Xiao JZ, Kondo S, Takahashi N, Miyaji K, Oshida K, Hiramatsu A, Iwatsuki K, et al. Effects of milk products fermented by *Bifidobacterium longum* on blood lipids in rats and healthy adult male volunteers. **J Dairy Sci** 2003; 86: 2452-61.

25. Ejtahed HS, Mohtadi-Nia J, Homayouni-Rad A, Niafar M, Asghari-Jafarabadi M, Mofid V, et al. Effect of probiotic yogurt containing *Lactobacillus acidophilus* and *Bifidobacterium lactis* on lipid profile in individuals with type 2 diabetes mellitus. **J Dairy Sci** 2011; 94: 3288-94.

26. Ridker PM, Hennekens CH, Buring JE, Rifai N. C reactive protein and other markers of inflammation in the prediction of cardiovascular disease in women. **N Engl J Med** 2000; 342: 836-43.

27. Kuller LH, Tracy RP, Shaten J, Meilahn EN. Relationship of C-reactive protein and coronary heart disease in the MRFIT

nested case-control study. Multiple Risk Factor Intervention Trial. **Am J Epidemiol** 1996; 144: 537-47.

28. Ridker PM, Cushman M, Stampfer MJ, Tracy RP, Hennekens CH. Inflammation, aspirin, and the risk of cardiovascular disease in apparently healthy men. **N Engl J Med** 1997; 336: 973-79.

29. Tracy RP, Lemaitre RN, Psaty BM, Ives DG, Evans RW, Cushman M, et al. Relationship of C-reactive protein to risk of cardiovascular disease in the elderly: results from the Cardiovascular Health Study and the Rural Health Promotion Project. **Arterioscler Thromb Vasc Biol** 1997; 17: 1121-7.

30. Ridker PM, Buring JE, Shih J, Matias M, Hennekens CH. Prospective study of C-reactive protein and the risk of future cardiovascular events among apparently healthy women. **Circulation** 1998; 98: 731-3.

31. Ridker PM, Cushman M, Stampfer MJ, Tracy RP, Hennekens CH. Plasma concentration of C-reactive protein and risk of developing peripheral vascular disease. **Circulation** 1998; 97: 425-8.

32. Koenig W, Sund M, Fröhlich M, Fischer HG, Löwel H, Döring 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: 237-42.

33. Roivainen M, Viik-Kajander M, Palosuo T, Toivanen P, Leinonen M, Saikku P, et al. Infections, inflammation, and the risk of coronary heart disease. **Circulation** 2000; 101: 252-57.

34. Harris TB, Ferrucci L, Tracy RP, Corti MC, Wacholder S, Ettinger WH Jr, et al. Associations of elevated interleukin-6 and C-reactive protein levels with mortality in the elderly. **Am J Med** 1999; 106: 506-12.

35. Danesh J, Whincup P, Walker M, Lennon L, Thomson A, Appleby P, et al. Low grade inflammation and coronary heart disease: prospective study and updated meta-analyses. **BMJ** 2000; 321: 199-204.

36. Mendall MA, Strachan DP, Butland BK, Ballam L, Morris J, Sweetnam PM, et al. C-reactive protein: relation to total mortality, cardiovascular mortality and cardiovascular risk factors in men. **Eur Heart J** 2000; 21:1584-90.

37. Asemi Z, Jazayeri S, Najafi M, Samimi M, Mofid V, Shidfar F, et al. Effects of daily consumption of probiotic yoghurt on inflammatory factors in pregnant women: a randomized controlled trial. **Pak J Biol Sci** 2011; 14:476-82.

38. Tovar J, Nilsson A, Johansson M, Ekesbo R, Aberg AM, Johansson U, et al. A diet based on multiple functional concepts improves cardiometabolic risk parameters in healthy subjects. **Nutr Metab (Lond)** 2012; 9: 29.