Lipid profile in patients with alcohol dependence syndrome

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BACKGROUND: Alcohol dependence syndrome (ADS) has become a global public health challenge because of its high prevalence and the concomitant increase in risk of liver disease, cardiovascular disease and premature death. Influence of alcohol use on lipid metabolism is well recognized. Investigations had been carried out in the earlier period on abnormal lipid profile as a risk factor for Coronary Heart disease (CHD). Patients of alcohol dependence usually have a consumption pattern of more heavy use. Therefore it is useful to study the lipid profile in patients of alcohol dependence, to understand the effects of increasing levels of consumption.

METHODS: This cross-sectional study was conducted in TU Teaching Hospital. ADS patients were screened by the consultant psychiatrist using the Alcohol Use Disorder Identification Test (AUDIT) questionnaire. A total of 89 patients scored positive on the AUDIT as having alcohol-related problems and were included in the study. 89 ADS patients and 89 healthy controls both male and female were enrolled as participants. Blood Pressure and other anthropometric parameters were measured while fasting blood samples were analyzed for serum lipid profile. SPSS program was used to analyze data, t-test & Spearman's correlation coefficient was used to find correlation.

RESULTS: Among the ADS cases 95% were current smokers. Mean age of cases and controls was $35.42\pm5.6 \& 34.53\pm3.5$ years respectively. The mean total cholesterol levels were found to be higher in cases (5.41 ± 0.70) than controls (3.79 ± 0.74) with a strong statistical significance (p<0.001). Also, Mean triglyceride (TG) levels (2.09 ± 0.72), along with the mean HDL-cholesterol (1.66 ± 0.40) and LDLcholesterol levels (2.79 ± 0.81) were also elevated in cases when compared to the control samples (p<0.001).

CONCLUSION: This study has demonstrated definitive lipid profile changes in patients of alcohol dependence, with some correlation to the liver dysfunction. Alcohol causes alteration in various parameters of lipid metabolism including those which predispose to CHD. Low to moderate alcohol use over prolonged periods has been linked to have protective influence for development of coronary heart disease (CHD), through increase in high density lipoprotein cholesterol (HDL-C) levels. Key words: ADS (Alcohol dependence syndrome), CHD (Coronary heart disease), HDL-C (high density lipoprotein cholesterol), TG (Triglyceride), LDL-C (Low density lipoprotein cholesterol)

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Introduction

Alcohol consumption might be the cause of several diseases, and it is, well known, the high burden of its consumption over mortality around the world [1]. Alcohol is the only psychoactive drug that provides energy (7.1 kcal/g). However, its calories are considered "empty," because alcohol ingestion does not provide vitamins and minerals [2] and its use may cause alterations to the nutritional state [3]. Alcohol dependence "A syndrome is defined as cluster of physiological, behavioral, and cognitive phenomena in which the use of a substance or a class of substances takes on a much higher priority for a given individual than other behaviors that once had greater value [4]. Hazardous alcohol intake and related disorders are a major health issue. A World Health Organization (WHO) project on psychological problems in general practice has shown that alcohol dependence or harmful alcohol use is present in about 6% of patients attending primary care, ranking third in frequency after major depression and generalized anxiety [5]. Harmful or heavy alcohol drinking makes a substantial contribution to the burden of disease and premature mortality [6]. Among persons admitted to general hospitals, 20 to 40 percent have alcohol-related problems, and among the elderly, alcohol-related hospitalizations are as numerous as those due to myocardial infarction [7]. The liver is the organ most severely affected by alcoholism. In some urban areas, cirrhosis (usually a complication of alcoholism) is the fourth most frequent cause of death among people 25 to 64 years of age [8]. Different studies have been done in alcoholic liver diseases but only few studies have been done in alcohol dependence syndrome. Alcohol abuse is a major

¹Department of Biochemistry, Maharajgunj Medical campus, IOM, TU, Nepal. ²Nepal Medical College, Jorpati, Kathmandu, Nepal. ³Department of Psychiatry and Mental Health, TU teaching hospital, Nepal Correspondence to: Mithileshwer Raut, Department of Biochemistry, Maharajgunj Medical Campus, IOM, TU Email: clinbio.mraut@gmail.com health problem as well as social problem in the community. Patients of alcohol dependence usually have a consumption pattern of more heavy use. Therefore it is useful to study the lipid profile in patients of alcohol dependence, to understand the effects of increasing levels of consumption. Alcoholism may lead to the different health consequences, like alcoholic liver diseases, cardiovascular diseases, and kidney diseases. So, the early diagnosis of ADS can help the patient to prevent from these major health problems.

Methods

This cross-sectional study was conducted in TU Teaching Hospital. ADS patients were screened by the consultant psychiatrist using the Alcohol Use Disorder Identification Test (AUDIT) questionnaire. The aim of the study and the questions in the questionnaire form were fully explained to the patients. A total of 89 patients scored positive on the AUDIT as having alcoholrelated problems and were included in the study. 89 ADS patients and 89 healthy controls both male and female were enrolled as participants. Blood Pressure and other anthropometric parameters were measured while fasting blood samples were analyzed for serum lipid profile. Serum is used for analysis of lipid profile and traditional marker of alcoholism. Total cholesterol was estimated by enzymatic method as described by Allain et al [9]. Serum triglyceride was estimated by Fossati and Prencipe method [10] associated with Trinder reaction [11]. HDL-Cholesterol was estimated by precipitation method. in which chylomichron, LDLcholesterol and VLDL were precipited and the supernatant fluid containing HDL-cholesterol were estimated by cholesterol method. LDLcholesterol was calculated using the Friedewald formula [12]. Serum glutamate pyruvate (SGPT), Serum glutamate transferase oxaloacetate transferase (SGOT) and gamma glutamyl transferase (GGT) were estimated by enzymatic method. The test was performed by reagent manufactured by Human, Germany, in the fully automated chemistry analyzer, BT 3000, Italy.

Laboratory standard operation procedures were maintained for all laboratory analysis. Internal quality control sera, both normal and pathological, were also run for each lot of the test, for the validation of the results. SPSS program was used to analyze data, t-test &Spearman's correlation coefficient was used to find correlation.

Results

Mean age of the patients and control subjects was 35.42 ± 5.6 years and 34.53 ± 3.5 years respectively. Range of age was 25-47 years for both groups. The majority of subjects (78%) had begun voluntarily, while 22% claimed to have done so due to peer pressure. The average daily intake of alcohol as stated by the patients was 71.36 gm. The mean duration of drinking was 12.0 years (Range 5-28 years). The frequency of consumption was: daily in 48% patients, 3-5 times a week in 41 % patients. The majority of alcohol dependent patients (72%) consumed alcohol alone while 28% claimed to drink only in company.

Table 1. Comparison of mean age of ADS cases and healthy control

Subject class	Mean±SD	p-value
Cases	35.42 ± 5.6	0.201
Controls	34.53 ± 3.5	0.201

Applied one way ANOVA test, statistically significant at pvalue <0.05

Table 2.	Comparison	of mean	of Lipid	profile
between	cases and con	trols		

	Control Mean ± SD	Case Mean ± SD	p- value
Total Cholesterol	3.79±0.74	5.41±0.70	0.001
Triglyceride	1.23±0.60	2.09 ± 0.72	0.001
HDL-C	1.08 ± 0.24	1.66 ± 0.40	0.001
LDL-C	2.15 ± 0.90	2.79±0.81	0.001

Applied one way anova test, statistically significant at p-value < 0.05

Table 3. Comparison of mean of different traditional marker and liver enzymes between ADS patients and normal healthy control

·	Control Mean±SD	Case Mean±SD	p- value
Gamma-GT	41.80±10.56	181.02±78.16	0.001
MCV	89.77±2.18	97.22±4.6	0.001
SGOT	35.26±14.27	114.35±46.22	0.001
SGPT	26.30±10.6	60.28±13.12	0.001

Applied one way anova test, statistically significant at p-value < 0.05

Table 4. Spearman	's correlation	coefficient of
GGT with Lipid p	rofile between	1 (2365

	Spearman's rho	p-Values	
Total Cholesterol	0.081	0.449	
Triglyceride	-0.005	0.964	
HDL-C	-0.067	0.528	
LDL-C	0.105	0.323	

The table 4 shows the Spearman's correlation of GGT with lipid profile between the cases and controls. GGT was not significantly correlated with lipid profile.

Table 5. Spearman's correlation coefficient of SGOT with Lipid profile between cases

	Spearman's	p-Values
	rho	
Total Cholesterol	0.037	0.730
Triglyceride	-0.128	0.228
HDL-	0.128	0.231
Cholesterol		
LDL-cholesterol	0.020	0.849

The table 5 shows the Spearman's correlation of SGOT with lipid profile between the cases and controls. SGOT was not significantly correlated with lipid profile.

Table 6. Spearman's correlation coefficient of MCV with Lipid profile between cases

	Spearman's	p-Values
	rno	
Total Cholesterol	0.065	0.542
Triglyceride	0.004	0.969
HDL-	0.089	0.407
Cholesterol		
LDL-cholesterol	0.027	0.804

The table 6 shows the Spearman's correlation of MCV with lipid profile between the cases and controls. MCV was not significantly correlated with lipid profile.

Discussion

The mean total cholesterol levels were found to be higher in cases than controls with a strong statistical significance. Mean TG levels, along with the mean HDL-cholesterol and LDLcholesterol levels were also elevated in cases when compared to the control samples. Within the group analysis of cases, it revealed borderline high total cholesterol in about half of the cases and very high level of total cholesterol was seen in one-tenth of the patients.

Also, among the cases, majority had elevated triglycerides level. LDL cholesterol was normal in most of the patients. A elevated HDL cholesterol was noted in more than half of the cases. These features are in the line of the notion of cardio protective effect of alcohol consumption by maintaining the level of LDL and HDL with an expense of slightly raised triglyceride level. The present study needs to be seen in the light of earlier studies which appear to confirm the linear relationship between increasing amounts of alcohol use and lipid profile changes known to have protective role for CHD [13]. Alcohol consumption has been found to be associated with increased serum levels of Tg and high density lipoproteins (HDL) [14,15]. The increase in HDL cholesterol has been estimated to account for half of the beneficial effects of alcohol consumption on cardiovascular events.16 Alcohol has narrow therapeutic range and only the moderate drinking has beneficial effects on cardiovascular health [17]. Prolonged excessive drinking causes various structural and functional abnormalities of heart.

Conclusion

In conclusion, this study has demonstrated definitive lipid profile changes in patients of alcohol dependence, with some correlation to the liver dysfunction. Alcohol causes alteration in various parameters of lipid metabolism including those which predispose to CHD. Low to moderate alcohol use over prolonged periods has been linked to have protective influence for development of coronary heart disease (CHD), through increase in high density lipoprotein cholesterol (HDL-C) levels. Acessed May 2012.

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