January 2013

Are we ready for a new epidemic of under recognized liver disease in South Asia especially in Pakistan? Non alcoholic fatty liver disease

Om Parkash  
Aga Khan University, om.parkash@aku.edu

Saeed Hamid  
Aga Khan University, saeed.hamid@aku.edu

Follow this and additional works at: https://ecommons.aku.edu/pakistan_fhs_mc_med_gastroenterol

Part of the Gastroenterology Commons

Recommended Citation
Available at: https://ecommons.aku.edu/pakistan_fhs_mc_med_gastroenterol/164
Abstract
Nonalcoholic fatty liver disease (NAFLD) is increasingly recognized as an important public health problem nowadays. NAFLD encompass a variety of liver pathologies including simple steatosis, NASH, fibrosis, cirrhosis and finally cancer. It is associated with obesity, metabolic syndrome, dyslipidaemia, Insulin resistance (IR) and type 2 diabetes.

It is the most common chronic liver disease in USA and considered to be increasing in Asia Pacific region including South Asia however there is no community based study from Pakistan. Customarily NAFLD had been regarded as a benign disease; however clinical as well as epidemiological studies had contradicted this belief because approximately 20% of the patients with NAFLD had NASH which has propensity to develop cirrhosis and ultimately to HCC.

The diagnosis of NAFLD is made most of the times incidentally on abdominal imaging which is done for other purposes.

Despite its prevalence, treatment options are very limited. However modification of risk factors such as dyslipidemia, diabetes control and weight reduction does help in NAFLD. Fatty liver results due to lack of physical activity; hence foremost step to manage such patients would be to develop the healthy life style. We need population based studies in our country so that we can protect our population from a new epidemic.

Keywords: NAFLD, NASH, Fatty liver, Physical activity, Hepatocellular carcinoma.

Introduction
Nonalcoholic fatty liver disease (NAFLD) is recognized as an important public health problem nowadays. NAFLD encompass a variety of liver pathologies including simple steatosis, Nonalcoholicsteatohepatitis (NASH), fibrosis, cirrhosis and finally cancer.

Approximately three decades ago Ludwig et al described liver lesions similar to alcohol associated lesions within liver in patients who did not take alcohol and called it NAFLD. NAFLD is defined as fatty liver (Liver fat > 5-10% of liver weight) which is not due to excess alcohol or other cause of steatosis. NAFLD is associated with obesity, metabolic syndrome, dyslipidemia, insulin resistance (IR) and type 2 diabetes. However, exact pathogenesis of NAFLD is incompletely understood and factors that determine the severity are still to be clearly understood.

NAFLD is the most common chronic liver disease in USA and considered to be increasing in Asia pacific region including South Asia. NAFLD affects approximately 15-40% of general population and its prevalence is increasing worldwide. The community prevalence of NAFLD in South Asia and South East Asia ranges from 5-30%. Recently a hospital based study in Pakistan had shown a frequency of approximately 14% however, there is no community based study from Pakistan to the best of our knowledge. In India it varies from 5-28% in general population especially those who are undergoing healthy checkups. Indians have increased propensity for visceral fat accumulation which may present from birth. In China, prevalence ranges from 5-24% higher in urban areas, compared to rural areas. Fatty liver on ultrasound was detected in 18% of white collar people in Beijing and 11% in labourers and similar figures were reported from other areas of Shanghai. Other countries like Japan, Hong Kong, Taiwan, Indonesia have a prevalence of 9-14%,16%,11-41% and 30% respectively.

Customarily NAFLD had been regarded as a benign disease; however clinical as well as epidemiological studies have contradicted this belief and have shown that approximately 20% of the patients with NAFLD had Nonalcoholic steatohepatitis (NASH). This NASH group constituted approximately 80% of cryptogenic cirrhosis and progressed to advanced fibrosis in 32-37%. Certain risk factors are associated with NAFLD like obesity, type II diabetes mellitus (Type 2 DM) and insulin resistance. These risk factors increase the risk of fibrosis progression. Hence, about 5-20% of patients with Non cirrhotic NASH would develop cirrhosis in 10 years and about one in...
every 200 NASH subject would develop hepatocellular carcinoma (HCC) over a 7 years period. Second most common cause of death in patients with NASH is liver disease (cirrhosis related complications) causing 12% deaths in 10 years of follow up. Hence, it is clear that NASH cannot be considered as benign disease anymore.

The available literature shows that clinical and social burden of NAFLD continues to increase.

Pathogenesis:
The pathogenesis of NAFLD is still incompletely understood and factors which determine disease are still not very clear nevertheless, overweight/obesity and insulin resistance are considered as risk factors. Typically, patients at the time of diagnosis are of middle age. The prevalence of other diseases being risk factors for NAFLD, are also increasing. For example majority of patients (>80%) are overweight with approximately 30% being obese, 30-70% are hypertensive and 20% have Type 2 DM. Therefore NAFLD has been considered as part of metabolic syndrome.

NAFLD may present with a variety of signs and symptoms including insulin resistance (IR), peripheral lipolysis, increased hepatic uptake of fatty acids, hormonal abnormalities (elevated leptin), release of pro-inflammatory cytokines and elevated insulin level induced mitochondrial function abnormality. All these abnormalities start the cascade of NAFLD from simple fatty liver to NASH and result in decompensated cirrhosis or HCC.

Diagnosis:
The diagnosis of NAFLD is made on abdominal imaging for evaluation of elevated liver enzymes or for some other indications. Most of the patients are asymptomatic. If symptoms develop, these are nonspecific such as fatigue or upper abdomen pain. Majority patients seek medical consultation because of deranged liver functions seen as elevated liver enzymes. The population based data such as 3rd National Health and Nutrition Examination Survey (NHADES III) from North America has shown prevalence of deranged liver enzymes in 8% and in 2/3 cases centre could not be determined. Most of these unexplained abnormal liver functions are strongly associated with metabolic syndrome and hence represent possible NAFLD. However, transaminase levels are typically normal or elevated by <5 times of upper limit of normal. In contrast to those with alcoholic hepatitis, most patients with NAFLD had ratio of aspartate aminotransferase (AST) to alanine aminotransferase (ALT) of less than 1. As the disease progresses, AST level increases more than ALT level, and if the ratio is greater than 1, more advanced liver disease may be suspected.

Recent studies suggest that upper limit of normal for serum ALT (generally between 40 and 55 IU/L) should be lowered to ≤30 IU/L for men and ≤19 IU/L for women.

The assessment of patients with NAFLD includes confirmation of diagnosis, disease severity assessment to differentiate between NAFLD/NASH and to determine the concomitant metabolic abnormalities including insulin resistance and cardiovascular abnormalities.

The current guidelines recommend hepatic ultrasound (US) as first step of diagnostic evaluation of NAFLD. The ultrasound features of NAFLD include increased hepatic echogenicity, vascular blurring and deep attenuation of US signals. These three US features have good accuracy in detecting fatty liver and had good correlation with visceral obesity and metabolic syndrome (MetS). NAFLD diagnosis also requires exclusion of other liver conditions such as viral hepatitis (HBV, HCV) and alcoholic liver disease. Liver histology remains the gold standard for assessing disease severity in NAFLD. Being invasive, biopsy is unsuitable for community studies and particularly for studying hepatic fibrosis progression, and also histological assessment of NAFLD is associated with sampling error and can lead to underestimation of the fibrosis score especially when specimen is small.

It is now however, clear that NAFLD and insulin resistance (IR) have a ‘chicken and egg’ type relationship where each supports the other. Insulin resistance has been directly linked to development of diabetes mellitus, hypertension, and atherosclerosis and all these lead to significant morbidity and mortality.

Recently Asia pacific society of liver diseases has incorporated the anthropometric measurements as regional criteria based on multiple Asian publications. Majority of patients with NAFLD are overweight or obese, therefore small increases in weight such as 2-3kg in body weight may lead to NAFLD, but more importantly this may occur within the normal ranges of weight. The World Health Organization (WHO) has redefined the criteria of obesity for Asians when body mass index (BMI) is >25 Kg/m² and overweight when BMI is between 23-25Kg/m².

Unlike hospital based studies, population based studies have defined NAFLD by biochemical criteria (increased serum transaminases or gamma glutamyltranspeptidase) or by hepatic imaging (liver US, Computerized tomography i.e. CT, magnetic resonance imaging) or both. Therefore, an operational definition of NAFLD based on liver imaging, supported by appropriate exclusion, appears sustainable and can overcome some of the problems associated with liver biopsy. Hence, deciding for operational definition of
NAFLD in South Asia, cost and availability should be considered before defining for clinical as well as research purpose. Literature shows that CT may not carry more weightage than US in terms of diagnosis and CT may only be useful in identifying focal fatty change in liver. Magnetic resonance imaging (MRI) can be robust for NAFLD because it can quantify the triglycerides stores in liver that can be useful in assessing the efficacy of therapeutic intervention, but this modality is very expensive and not available everywhere in this part of world (South Asia). In Japan US liver has been used for NAFLD as a standard of care. In one of the comparative study from Cleveland clinic, which used liver histology (at least 30% fat) as the gold standard, sensitivity and specificity of liver US for NAFLD was good (89% and 93% respectively), but somewhat less for liver fibrosis (77% and 89% respectively).

**Treatment:**

Despite its prevalence, treatment options are very limited. However, modification of risk factors such as dyslipidaemia, control of diabetes and weight reduction helps in control of NAFLD. Fatty liver occurs due to lack of physical activity, excess calorie intake in comparison to normal person without fatty liver, hence, the foremost step to advise such patients to develop a healthy life style. When such intervention proves to be ineffective then one should go for drugs which would be the second line strategy. Often, it is difficult to encourage people to modify their unhealthy behaviour to adopt a healthier life style because life style modifications depend largely on personal beliefs and values. Therefore psychological/behavior strategies are essential to produce a change in behaviour.

**Lifestyle modification:**

**Behavioral therapy:**

Behavioral therapy is an area of research widely applied to the treatment of obesity and associated metabolic diseases. This type of therapy is a global therapeutic approach which delivers patients the practical approach to achieve their eating and exercise goals. Subjects are administered with a set of principles and techniques for modifying diet and exercise such as keeping records of their daily food intake along with physical activity, counting food calories and the time spent in exercise, being more active in their daily life, avoiding situations which could lead to incidental eating and correcting unrealistic goals about weight loss and body image. This type of therapy is usually offered in groups of 10-15 subjects or occasionally individually but preferable in groups; in weekly sessions of 60-90 minutes for a period of 3-12 months. Patients learn to calculate food calories, manage their nutritional changes and are encouraged to do physical exercise for 30-40 min daily for at least 5 days a week. This type of intensive programme often leads to a mean weight loss of 10% of initial weight in 6 months in subjects who complete the treatment. Unfortunately some patients regain part of their lost weight within one year but majority maintain healthy weight loss of ≤ 5% bodyweight by following treatment and weight maintenance strategies. Recently a meta-analysis also confirmed that lifestyle intervention is at least as effective as pharmacotherapy. This was also seen in a study done in India which revealed that moderate physical activity and diet was effective in preventing diabetes in Indians despite having low BMI. Behavioral therapy requires a well-defined programme, must be supported by a manual for both operators and patients, and must be carried out by a multidisciplinary team, including physicians, dieticians, psychologists, trained nurses and physical exercise experts. There are several techniques that aim at enhancing patient compliance in lifestyle changes, for example formulating realistic goals with patients, empathetic communication by physicians, being sensitive to the general stigma of obesity, planning individualized weight loss and physical activity programs, encouraging self-efficacy, explaining treatment and its benefits, motivating patients to self-monitor diet and physical activity and arranging regular follow-up clinic. This strategy also includes reinforcement methods and relapse prevention techniques to ensure long term sustainability of change.

Majority of patients with NAFLD take more calories, do less exercise/physical activity which makes them prone to insulin resistance and overweight or obesity as compared to subjects who don't have NAFLD. The American gastroenterology Association recommends a weight loss of 10% of baseline weight as an initial goal if the patient is overweight. Palmer et al showed that weight reduction is associated with normalization of transaminases as well as liver enlargement. A Japanese study proved that moderate weight loss (approximately 6%) is associated with improvement in insulin resistance and fatty liver. An Israeli study on 48 subjects who underwent dietary restriction leading to moderate weight loss, showed improvement in liver enzymes in 96% of the subjects.

Variety of dietary combinations i.e either low carbohydrate or low fat are equally effective in weight reduction, and there is no long term difference in weight loss or its maintenance after treatment. Moreover, no specific recommendation favours Atkins, Ornish or South Beach diets. Therefore, diet should be modified on individual preferences. The current American
Physical Activity:
Physical activity (PA) plays a protective role in NAFLD. PA includes structured exercise, involves aerobic activities of moderate to vigorous intensity (e.g., jogging, brisk walking, bicycling, swimming, skiing, and ball games) and resistance training which comply with current exercise recommendations as well as other leisure time tasks performed at low intensity below current recommendations for improving cardio respiratory fitness (for example casual walking, bicycling, dancing and non-structural lifestyle activities such as gardening, housework, hobbies and yoga). It has been seen that weight reduction associated with dietary restriction are not long lasting and often the weight returns to its baseline once food restriction is stopped. Weight loss and improvement in physical activity shows histological improvement too. Diabetes prevention program research group suggest that exercise adherence appears to be more sustainable than weight loss over time therefore every effort should be made to encourage physical activity (exercise). Studies have also shown low prevalence of metabolic syndrome in subjects who adhere to increased physical activity and have higher muscle strength and higher cardio respiratory fitness. Studies have shown that physical activity leads to a reduction of risk of type 2 DM, IR, HTN and dyslipidaemia. Majority of these clinical effects might be mediated by changes in the release of adipocytokines (leptin, adiponectin and resistin), which are implicated in the development of hepatic steatosis, inflammation and fibrosis. A study from North America has shown that moderate intensity aerobic exercise has a beneficial effect on insulin resistance and alters substrate use in skeletal muscle. Another study revealed that physical activity enhances insulin sensitivity and glucose homeostasis through insulin sensitization along with facilitating glucose uptake, in the absence of weight loss. Physical activity (exercise) also decreases liver triglycerides within the development of hepatic steatosis, inflammation and fibrosis. A study from Australia on 141 patients with NAFLD has shown a significant improvement in liver enzymes in patients who adhered to PA of 150min/week or more and those who increased their fitness. This effect was independent of any weight change. Another study on obese patients showed moderate improvement in hepatic fat without losing weight, when aerobic exercise was done for 4 weeks only.

Pharmacological treatment:
At present there is no registered drug treatment for NAFLD. Earlier studies suggest that insulin sensitizers and antioxidants may confer some benefit whereas ursodeoxycholic acid (UDCA) and pentoxiphyline have not proved to be of any benefit in NAFLD in clinical trials.

Surgical treatment:
Bariatric surgery is only helpful in patients who are morbidly obese and had shown benefit in NAFLD with NASH.

Conclusion
NAFLD is an emerging problem and can cause significant morbidity and mortality if individuals don’t adopt healthy life style such as regular exercise and a weight maintenance diet. Since there is no approved treatment for NAFLD we need to educate Pakistani population about healthy life style. Also population based studies in our country are needed so that we can protect our population from a new epidemic.

References
7. Amarapurkar DN, Hashimoto E, Lesmana LA, Sollano JD, Chen PJ, Goh KL. How common is non-alcoholic fatty liver disease in the Asia-Pacific region and are there local differences? J Gastroenterol Hepatol 2007; 22: 788-93.


