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LETTERS TO THE EDITOR

# Alcohol consumption and fatty liver disease

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# **Abstract**

Hamaguchi *et al* recently reported some interesting observations on alcohol consumption and risk of fatty liver disease from a large population. However, we feel that it might be necessary to discuss some concerns in this study. As the alcohol consumption categorization was defined by the same criteria in both men and women, which might affect their results. As another factor is soft drinks consumption. It has been proved that soft drinks, especially fructose, contributes to the development of obesity, diabetes, metabolic syndrome, and nonalcoholic fatty liver disease. However, this confounding factor was not adjusted or discussed in this article. The third is the genetic background, for some genetic factors are related with the development of fatty liver disease, which was also not considered yet.

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**Key words:** Alcohol; Fatty liver disease; Obesity; Diabetes; Metabolic syndrome

Core tip: Modest alcohol consumption was significantly inversely associated with fatty liver disease in recent studies. However, some studies did not consider some important potential confounding factors when they conclude their findings. Herein, we raised and discussed these important factors in this letter.

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## TO THE EDITOR

We read with great interest the article by Hamaguchi *et al*<sup>11</sup> published in January 2012 at *World J Gastroenterol*. This cross-sectional study reported some interesting observations on alcohol consumption and risk of fatty liver disease from a large population. However, we feel that it might be necessary to discuss some concerns in this study.

The authors clearly indicated that alcohol consumption was significantly inversely associated with fatty liver disease, especially in men. However, they did not find this association in their previous cohort study. The reason for this contradiction might be that some important confounding factors were not considered. As the alcohol consumption categorization was defined by the same criteria in both men and women, only 84 female subjects (1.1%) were defined as excess alcohol consumers and 207 (2.7%) were defined as moderate alcohol consumers, the numbers being much lower than those in men (13.5% and 14.7%, respectively). Although the authors analyzed the data in both men and women, the initial categorization was not separated, which might affect their results.



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Another factor is soft drinks consumption. It has been proved that soft drinks, especially fructose, contributes to the development of obesity, diabetes, metabolic syndrome, and nonalcoholic fatty liver disease<sup>[3]</sup>. However, this confounding factor was not adjusted or discussed in this article. The last point is about the genetic background. Although the mechanisms of this inverse association between alcohol consumption and fatty liver disease are still unclear, some genetic factors are related with the development of fatty liver disease<sup>[4-6]</sup>, such as peroxisome proliferator-activated receptor gamma and hemochromatosis gene polymorphisms<sup>[7,8]</sup>. Therefore, as some genetic factors might interact with alcohol consumption in fatty liver disease, it could be an interesting topic for further investigations.

#### REFERENCES

- Hamaguchi M, Kojima T, Ohbora A, Takeda N, Fukui M, Kato T. Protective effect of alcohol consumption for fatty liver but not metabolic syndrome. World J Gastroenterol 2012; 18: 156-167 [PMID: 22253522 DOI: 10.3748/wjg.v18.i2.156]
- 2 Hamaguchi M, Kojima T, Ohbora A, Takeda N, Fukui M, Kato T. Aging is a risk factor of nonalcoholic fatty liver disease in premenopausal women. World J Gastroenterol 2012;

- 18: 237-243 [PMID: 22294826 DOI: 22294826]
- Nseir W, Nassar F, Assy N. Soft drinks consumption and nonalcoholic fatty liver disease. World J Gastroenterol 2010; 16: 2579-2588 [PMID: 20518077 DOI: 10.3748/wjg.v16.i21.2579]
- 4 Li YY. Genetic and epigenetic variants influencing the development of nonalcoholic fatty liver disease. World J Gastroenterol 2012; 18: 6546-6551 [PMID: 23236228 DOI: 10.3748/wjg.v18.i45.6546]
- 5 Sookoian S, Pirola CJ. PNPLA3, the triacylglycerol synthesis/hydrolysis/storage dilemma, and nonalcoholic fatty liver disease. World J Gastroenterol 2012; 18: 6018-6026 [PMID: 23155331 DOI: 10.3748/wjg.v18.i42.6018]
- 6 Nagarajan P, Mahesh Kumar MJ, Venkatesan R, Majundar SS, Juyal RC. Genetically modified mouse models for the study of nonalcoholic fatty liver disease. World J Gastroenterol 2012; 18: 1141-1153 [PMID: 22468076 DOI: 10.3748/wjg. v18.i11.1141]
- 7 Raszeja-Wyszomirska J, Kurzawski G, Lawniczak M, Miezynska-Kurtycz J, Lubinski J. Nonalcoholic fatty liver disease and HFE gene mutations: a Polish study. World J Gastroenterol 2010; 16: 2531-2536 [PMID: 20503453 DOI: 10.3748/wjg.v16. i20.2531]
- 8 Rey JW, Noetel A, Hardt A, Canbay A, Alakus H, Zur Hausen A, Dienes HP, Drebber U, Odenthal M. Pro12Ala polymorphism of the peroxisome proliferator-activated receptor γ2 in patients with fatty liver diseases. World J Gastroenterol 2010; 16: 5830-5837 [PMID: 21155004 DOI: 10.3748/wjg.v16. i46.5830]

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