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## Letters to the editor

**Correspondence: “A multisociety Delphi consensus statement on new fatty liver disease nomenclature”**

1 Dear Editor,

2 Steatotic liver disease (SLD) has been proposed by Rinella *et al.* as

3 the new overarching term to encompass various aetiologies of steato-

4 sis, including metabolic dysfunction-associated steatotic liver disease

5 (MASLD), MASLD and increased alcohol intake (MetALD), alcohol-

6 associated liver disease (ALD), specific aetiology SLD, and cryptogenic

7 SLD [1]. However, unlike metabolic dysfunction-associated fatty liver

8 disease (MAFLD) suggested by Eslam *et al.* in 2020, blood biomarkers

9 and scores were not explicitly outlined as one of the methods for

10 identifying hepatic steatosis [2]. These biomarkers or scores, such as

11 the fatty liver index, are deemed appropriate for extensive epidemio-

12 logical studies to detect hepatic steatosis in adults [2]. Indeed, the

13 European clinical practice guidelines stated that validated biomarkers

14 and scores are acceptable substitutes to diagnose fatty liver when

15 imaging methods are unavailable or impractical, such as in big epide-

16 miological surveys [3].

17 With increasing clinical and public health burdens from SLD, a

18 population-based study is currently being conducted to determine

19 its prevalence in Malaysia [4]. Biomarkers and scores are very

20 useful here because imaging is neither financially nor logistically

21 feasible in the nationwide survey [4]. Thus, biomarkers and scores

22 can be important for low- and middle-income countries like

23 Malaysia, where the availability and cost of imaging affect feasi-

24 bility [3]. Besides that, their use in primary care can increase

25 awareness and early diagnosis of SLD, which is essential for sec-

26 ondary disease prevention. Furthermore, the fibrosis-4 (FIB-4)

27 score can be used to screen for more severe SLD.[5] Those with

28 less severe SLD can be managed in the primary care setting. In

29 contrast, referrals for further evaluations are necessary for indi-

30 viduals with high or intermediate scores [5].

31 With a better understanding of the pathophysiology and epi-

32 demiology of SLD, it is clear that multi-disciplinary collaborative

33 efforts are vital to managing the complex disease [1,2,5]. Public

34 health-wise, determining the national prevalence of SLD is essen-

35 tial to gauge the issue's magnitude [4] accurately. The informa-

36 tion can raise SLD priority in health agenda setting at the

37 national level, catalyze evidence-based policymaking to prevent

38 and manage SLD, and incorporate SLD into the broader noncom-

39 municable disease prevention and control initiatives due to

40 shared common risk factors [4,5]. Public health practitioners play

41 a crucial role in raising awareness and knowledge of various

42 stakeholders on SLD, with healthcare professionals in primary

43 care and the general population being the key target groups [5].

44 In this sense, accepting and including readily available bio-

45 markers and scores for identifying hepatic steatosis is essential

and should be considered in future updates of the nomenclature 46  
or guidelines on the disease. 47

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**Ethical statement** 51

The Medical Research and Ethics Committee of the Ministry of 52  
Health Malaysia has granted ethical approval to conduct the study 53  
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**Declaration of competing interest** 55

Wah Kheong Chan has served as a consultant or advisory board 56  
member for Roche, Abbvie, Boehringer Ingelheim, and Novo Nordisk 57  
and as a speaker for Echosens, Viatrix, and Hisky Medical. Other 58  
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