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#### RESEARCH LETTER

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#### Abstract

**Background:** Following the adoption of new nomenclature for steatotic liver disease, we aimed to build consensus on the use of International Classification of Diseases codes and recommendations for future research and advocacy.

**Methods:** Through a two-stage Delphi process, a core group (n = 20) reviewed draft statements and recommendations (n = 6), indicating levels of agreement. Following revisions, this process was repeated with a large expert panel (n = 243) from 73 countries.

**Results:** Consensus ranged from 88.8% to 96.9% (mean = 92.3%).

**Conclusions:** This global consensus statement provides guidance on harmonizing the International Classification of Diseases coding for steatotic liver disease and future directions to advance the field.

### INTRODUCTION

Nomenclature changes in the field of steatotic liver disease (SLD) were recently proposed and are currently being adopted by a wide range of stakeholders.<sup>[1]</sup> Among the suggested modifications, the change from NAFLD to metabolic dysfunction–associated steatotic liver disease (MASLD) reflects a drop of the "nonalcoholic" label, enabling the inclusion of positive diagnostic criteria while removing potentially stigmatizing classification. The intake of alcohol as a disease contributor is also acknowledged in the new nomenclature, with the introduction of the term "MASLD and alcohol-associated liver disease (ALD)", abbreviated as MASLD and ALD (MetALD).<sup>[1]</sup> Moreover, the nomenclature process introduced new defining criteria for MASLD and MetALD. Whereas studies have demonstrated almost complete overlap between populations defined by NAFLD and MASLD criteria, the cutoffs for alcohol use in the MetALD definition remain to be clarified by future studies.<sup>[2–4]</sup>

As a consequence of these nomenclature changes and to aid in their implementation, administrative coding will need to be adjusted. Globally, the International Classification of Diseases (ICD) coding system is the

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Abbreviations: ALD, alcohol-associated liver disease; ICD, International Classification of Diseases; MASH, metabolic dysfunction–associated steatohepatitis; MASLD, metabolic dysfunction–associated steatotic liver disease; MetALD, MASLD and ALD; SLD, steatotic liver disease.

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I ABLE 1	LE 1 Final consensus statements and recommendations								
State	Statements	Grade	A (%)	SA (%)	(%) A+SA	SD (%)	D (%)	NQ (%)	z
-	MASLD is currently best coded using the ICD-10 code for NAFLD (K76.0).	в	67.4	21.5	88.8	5.6	5.6	4.1	233
2	MASH is currently best coded using the ICD-10 coding for NASH (K75.8 or K75.81, depending on the setting).	۷	70.8	21.0	91.8	4.3	3.9	4.1	233
ო	ALD is best coded using the "alcohol-associated liver disease" spectrum of ICD-10 codes (K70).	A	84.6	12.3	96.9	2.2	0.9	6.6	227
4	As no appropriate MetALD ICD-10 code exists, clinical research and health care professionals should use ICD coding for the more relevant part of MASLD/ALD on an individual basis while awaiting ICD-10/11 definition changes by the World Health Organization.	ш	62.3	26.8	89.2	6.9	3.9	4.9	231
Reco	Recommendations								
2ı	Research should focus on identifying how to best distinguish between MASLD, MetALD, and ALD when using historical data sources (eg, register-based data).	۲	68.5	22.7	91.2	5.9	2.9	2.1	238
9	International societies should advocate for a global update of ICD terminology by the World Health Organization to better reflect the nomenclature change, including separate diagnostic codes for MASLD, MASH, MetALD, ALD, and cryptogenic steatotic liver disease.	۷	86.4	9.5	95.9	2.9	1.2	0.4	242
Mea	Mean % agreement	73.3	19.0	92.3	I	I	I	I	Ι
Notes: agreen	Notes: Percentages may add up to more than 100 due to rounding. Grades are based on the percentage of combined agreement (agree+somewhat agree): A, 90%–99% combined agreement; B, 78%–89% combined agreement. Responses to each statement and recommendation are presented as percentages of the total responses.	eement (agre	e+somew	hat agree): A	۰, 90%–99% col	mbined agre	ement; B, 7	.8%—89% co	mbined

Abbreviations: A, agree; ALD, alcohol-associated liver disease; D, disagree; ICD, International Classification of Diseases; MASH, metabolic dysfunction-associated steatohepatitis; MASLD, metabolic dysfunction-associated steatotic liver disease; MetALD, MASLD and ALD; N, total number of responses; NQ, percentage of participants that indicated that they were not qualified to respond; SA, somewhat agree; SD, somewhat disagree vears.

# most used. We thus aimed to build consensus on the appropriateness of using current ICD NAFLD and NASH codes to code MASLD and metabolic dysfunctionassociated steatohepatitis (MASH), respectively. We also sought to develop recommendations to guide research and advocacy on amending future ICD codes for SLD. While ICD systems vary at the local level (eg, which version is in use), ICD-10 is currently the dominant system. Nonetheless, following its release in 2022, ICD-11 use will be gradually introduced over the coming

# METHODS

We performed a two-stage Delphi process whereby, first, a core group of people (n = 20) indicated their agreement or disagreement with statements and recommendations (n = 6) (Supplemental Table 1, http://links.lww.com/HC9/A809) using "yes" to agree and "no" to disagree, through Microsoft Forms, from July 23 to August 6, 2023. Respondents were also invited to provide qualitative feedback on each item and overall, which was considered during item revisions. This group included individuals who had previously contributed to a consensus statement on the use of NAFLD ICD codes in research<sup>[5]</sup> and key opinion leaders involved in the nomenclature change.

The second stage involved inviting a panel of individuals with SLD experience to indicate their level of agreement ("agree," "somewhat agree," "somewhat disagree," or "disagree") with the modified items (n=6)(Table 1), using the described methodology,<sup>[6]</sup> through Qualtrics XM, from October 6-23, 2023. Respondents were also invited to provide qualitative feedback on each item and overall, which was considered during manuscript writing. Invitees who were not familiar with ICD codes and their use could opt out. Respondents who did not feel qualified to indicate their level of agreement with a survey item could choose the option "not qualified to respond." For the purposes of this study, we defined reaching consensus as having > 80% agreement on each item, with overall agreement being the sum of the "agree" and "somewhat agree" categories in stage 2.

# **Ethical considerations**

This study received ethical review exemption from the Hospital Clínic of Barcelona, Spain, ethics committee on October 4, 2023. All research was conducted in accordance with the Declarations of Helsinki and Istanbul. Respondents consented to participating, and data were anonymized for all analyses.

A total of 479 individuals were invited to participate in stage 2, of whom 269 (56.2%) responded. Of these, 26 (9.7%) opted out as they were not familiar with ICD codes and their use. The 243 respondents (90.3%) who completed the survey worked in 73 countries and had a mean age of 53.9 (SD: 9.4). Most respondents were male (65.4%), worked in high-income countries (66.3%) and in the Europe and Central Asia World Bank region (41.2%), and primarily worked in academia (67.9%) and as clinicians/medical doctors (72.8%) (Supplemental Table 2, http://links.lww.com/HC9/A809, contains further panelist details).

In stage 2, consensus ranged from 88.8% to 96.9% (mean = 92.3%). Four items had < 80% "agree" responses and relied more heavily on the "somewhat agree" category to reach a consensus. A total of 351 qualitative comments were provided across items. There was  $\geq$  88.8% consensus that MASLD, MASH, and ALD are currently best coded with K76.0, K75.8, or K75.81, and the K70 spectrum of ICD-10 codes, respectively. As for MetALD, which has no ICD code as it was newly introduced, 89.2% agreed that using ICD coding for the perceived dominant disease driver (MASLD or ALD) on an individual basis was preferable while awaiting updates to the ICD system. In terms of recommendations, 91.2% of participants agreed that research should prioritize how best to distinguish between MASLD, MetALD, and ALD when using historical data. Furthermore, the consensus that international societies should advocate for a global update of ICD terminology to better reflect the SLD nomenclature changes was 86.4%.

# DISCUSSION

This study found that, among a large panel of experts working across 73 counties, there was a high degree of consensus that NAFLD and NASH ICD codes can be updated to reflect the new MASLD and MASH names and definitions, respectively, without the need for new codes. Renaming the administrative terms across various systems and countries to reflect the nomenclature change should be a priority. This is important, as introducing coding changes may lead to considerable difficulties in comparing study results and interpreting disease epidemiology patterns across settings and over time. It should be noted that definition and ICD code modifications will not mitigate the challenge of correctly calculating the amount of alcohol consumed by patients, but we hope that the recommendation of focusing research on identifying how to best distinguish between MASLD, MetALD, and ALD will promote investigations around this topic. Further work to introduce novel ICD codes to specifically define HEPATOLOGY COMMUNICATIONS

MetALD is needed, which may be achieved through discussions with national and regional norm-setting bodies and the World Health Organization, which maintains and updates the ICD system.

# CONCLUSIONS

This global expert consensus statement recommends that the currently available ICD codes for NAFLD and NASH can be used to define MASLD and MASH, respectively, although advocacy is needed to update ICD terminology to better reflect the nomenclature change and introduce new codes for MetALD specifically.

## AUTHOR CONTRIBUTIONS

Hannes Hagström and Jeffrey V. Lazarus: Study conception and design; MVR: Statistical analysis; Hannes Hagström, Marcela Villota-Rivas, HEM, and Jeffrey V. Lazarus: Drafting of the manuscript; All: Acquisition of data; analysis and interpretation of data; and critical revision

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#### REFERENCES

- Rinella ME, Lazarus JV, Ratziu V, Francque SM, Sanyal AJ, Kanwal F, et al. A multi-society Delphi consensus statement on new fatty liver disease nomenclature. J Hepatol. 2023;79: 1542–56.
- Ratziu V, Boursier J. Confirmatory biomarker diagnostic studies are not needed when transitioning from NAFLD to MASLD. J Hepatol. 2023:S0168-8278(23)04996-6. doi:10.1016/j.jhep.2023. 07.017.
- Song SJ, Che-To Lai J, Lai-Hung Wong G, Wai-Sun Wong V, Cheuk-Fung Yip T. Can we use old NAFLD data under the new MASLD definition? J Hepatol. 2023:S0168-8278(23)05000-6. doi:10.1016/j. jhep.2023.07.021.
- Hagström H, Vessby J, Ekstedt M, Shang Y. 99% of patients with NAFLD meet MASLD criteria and natural history is therefore identical. J Hepatol. 2023:S0168-8278(23)05080-8. doi:10.1016/ j.jhep.2023.08.026.
- Hagström H, Adams LA, Allen AM, Byrne CD, Chang Y, Grønbæk H, et al. Administrative coding in electronic health care record-based research of NAFLD: An expert panel consensus statement. Hepatology. 2021;74:474–82.
- Lazarus JV, Mark HE, Allen AM, Arab JP, Carrieri P, Noureddin M, et al. A global research priority agenda to advance public health responses to fatty liver disease. J Hepatol. 2023;79: 618–34.