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Differences between current clinical guidelines for screening, diagnosis and management of non-alcoholic fatty liver disease and real-world practice: a targeted literature review

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ABSTRACT

Introduction: Non-alcoholic fatty liver disease (NAFLD) is the most common liver disease and is associated with obesity and metabolic comorbidities. Liver steatosis can progress to non-alcoholic steatohepatitis (NASH) exhibiting a relevant risk of fibrosis and ultimately liver failure. To date, no approved treatment for NASH to reduce its clinical and humanistic burden has been developed.

Areas covered: We undertook a literature review to identify English language, national and international clinical guidelines for NAFLD regarding diagnosis, assessment and management, and determined their points of agreement and difference. Additionally, we investigated published literature relating to real-world management of NAFLD and NASH.

Expert opinion: National (China, England/Wales, Italy, the USA) and international society (Asia-Pacific, Europe, World Gastroenterology Organisation) guidelines were identified and analyzed. All guidelines addressed identifying and diagnosing subjects with likely NAFLD, as well as assessment and management of individuals with risk factors for advanced disease, including fibrosis. Real-world practice reveals widespread, suboptimal awareness and implementation of guidelines. In the absence of proven therapeutics, such gaps risk failure to recognize patients in need of specialist care and monitoring, highlighting the need for clear, easy-to-apply care pathways to aid in reducing the clinical and humanistic burden of NAFLD and NASH.

Keywords:

Non-alcoholic fatty liver disease; non-alcoholic steatohepatitis; cClinical guidelines; literature review; liver fibrosis; real-world practice

Article highlights

- Since the mid-2010s, national clinical guidance for the diagnosis, assessment and management of NAFLD has been published for China, England/Wales, Italy and the USA, as well as internationally from the Asia-Pacific, European and global perspectives.
- All guidelines we reviewed provided recommendations for targeted, risk-based screening of individuals deemed likely to have NAFLD. Tests to screen patients who may have NAFLD included blood-based markers, imaging and elastography techniques to exclude those at low risk of NASH and advanced fibrosis.
- To assess liver fibrosis, a key determining factor in risk of disease progression, FIB-4 and/or NFS scores were recommended in guidelines from China, Europe, Italy and the USA.
- All guidelines provided recommendations for NAFLD management through lifestyle changes, principally weight loss through restriction of calorie intake and increased regular exercise.
- To date, no drugs have received regulatory approval for the treatment of NAFLD or NASH. However, all guidelines included weak recommendations for pharmacological support where considered necessary, although restricted to patients diagnosed with NASH with, or at risk of, fibrosis or disease progression.
- Despite the widespread availability of clinical guidelines, awareness and understanding of NAFLD, its diagnosis and management remain poor outside of specialists in hepatology. A lack of guidance and education is leading to low rates of diagnosis and therefore missed cases of serious, potentially life-threatening NASH and missed opportunities for intervention.

- The innovation of clear algorithms for sequential screening and diagnosis of NAFLD could help improve rates of identification and referral of at-risk individuals and improve standards of care.

1. Introduction

Non-alcoholic fatty liver disease (NAFLD) is a major cause of chronic liver disease that is often underdiagnosed [1]. Although manageable and potentially reversible in its early stages, progressive liver steatosis can lead to non-alcoholic steatohepatitis (NASH) with advanced fibrosis (AF) and end-stage liver disease, which is rapidly becoming the leading indication for liver transplantation [1]. Importantly, patients with NASH also experience substantially impaired quality of life [2-4].

The incidence of NAFLD is increasing worldwide, with a prevalence of around 25%, and is associated with higher mortality rates than those in the general population [5-10]. Although adults are most commonly affected by NAFLD, with occurrence increasing with age, rates of childhood NAFLD are increasing, particularly in association with the occurrence of obesity [1,8,11,12].

Cardiometabolic comorbidities share epidemiological, pathophysiological and behavioural/lifestyle features with NAFLD, suggesting a close association with aspects of metabolic syndrome and insulin resistance [1,8,13,14,15, (Niederseer, 2021)]. Management of cardiometabolic conditions such as type 2 diabetes mellitus (T2DM) and metabolic syndrome using lifestyle modification and pharmacotherapy is a key component of NAFLD care [6,16, (Niederseer, 2021)].

Genetic background also plays a key role in determining the development and severity of NAFLD and NASH that may explain inter-individual variation in patterns of disease.

Genome-wide association and large candidate gene studies have identified the I148M variant in the *patatin-like phospholipase domain-containing protein 3 (PNPLA3)* gene as a common, strong genetic determinant of NAFLD, NASH and hepatocellular carcinoma (HCC).

Other single nucleotide polymorphisms (SNPs) that appear to play a moderate role in the NAFLD spectrum include *TM6SF2*, *MBOAT7* and *GCKR* (Eslam, 2020).

NAFLD is characterized by evidence of hepatic fat accumulation (steatosis), with or without chronic, mild inflammation, and where the role of other causative etiologies, such as viral hepatitis or excessive alcohol consumption, have been excluded [8]. Although steatosis itself is usually considered benign, evidence is accumulating that suggests it may contribute to NASH progression [1]. NASH is characterized by steatosis, ballooning degeneration and lobular inflammation, which can progress to fibrosis, which is found in over half of patient with NAFLD, predisposing some individuals to cirrhosis and end-stage liver disease as well as being a HCC risk factor) [1,8,9,17,18]. Although the incidence of NAFLD- and NASH-related HCC is low (0.44 and 5.29 per 1,000 person years, respectively), the high number of patients with NAFLD has led to HCC being the fifth most common cancer worldwide, as well as one of the two most frequent causes of cancer-related death (Loomba, 2020; Huang, 2021).

Compared to NAFLD without NASH, a diagnosis of NASH is associated with shorter survival times, more cardiovascular events, and greater liver cancer mortality [17]. As well as being a leading cause of liver cirrhosis, NASH is associated with a substantial humanistic and economic burden that increases with advancing fibrosis levels. For example, end-stage liver disease accounts for between 56% and 90% of economic and wellbeing costs, and most NASH-related costs are due to use of secondary healthcare and the need for extensive diagnostic testing [4,18,19].

Screening and stratification of severity in large numbers of individuals are key components of efforts to combat disease progression. However, the complexity of NAFLD diagnosis and management is such that primary and specialist healthcare professionals play important roles, and multidisciplinary care is essential. Several international and national guidelines for diagnosing and managing NAFLD have been developed or updated in recent years [20-27]. Previously, inconsistencies between individual published guidelines for the diagnosis and

treatment of NAFLD have been reported [28,29], as well as a paucity of country-specific guidance and supporting documentation to encourage guideline implementation [30]. This may go some way towards accounting for poor rates of NAFLD diagnosis and lack of screening for disease severity [5].

1.1. Objective of this review

The objective of this literature search and review was to determine areas of consistency in NAFLD clinical practice guidance and identify points of disagreement. A secondary objective was to investigate real-world practice patterns in NAFLD management based on published data.

2. Methods

The current review follows on from an earlier literature search and review [29] which explored publications between 2005 and 2019. An update was undertaken employing the same methodology and included publications up to October 2020.

In the original literature search, the PubMed, Embase, and DynaMed publication databases were screened for potential publications. For the update described here, PubMed, MEDLINE, Embase, and Cochrane databases were screened. Targeted, iterative literature searches were conducted to identify relevant English-language publications related to key themes in the objectives and the predefined research questions described below. Search terms included at least one of the following: “non-alcoholic fatty liver disease”, “non-alcoholic steatohepatitis,” “fatty liver,” “liver fibrosis,” and “cirrhosis.” Additional search terms were included to identify guidelines, consensus statements, and relevant research relating to all stages of the identification, evaluation and management of NAFLD. Search results included a range of publication types: research papers, systematic literature reviews, narrative reviews, qualitative/quantitative research, clinical guidelines or consensus statements.

The final publications for inclusion were reviewed and summarized in the context of predefined research questions. Study findings are described qualitatively; quantitative data synthesis was not feasible owing to the high levels of heterogeneity across the study/publication types, patient characteristics and data presented.

To aid evaluation and synthesis of the narrative from this literature review, the authors analyzed the results in the context of the following topics relating to real-world practice in NAFLD.

- *Disease awareness among patients diagnosed with NAFLD.*
- *Discrepancies between routine diagnosis and screening practice and the implementation of recommendations in NAFLD guidelines.*
- *Lifestyle modification and pharmacotherapeutic management of NAFLD in practice.*

Initially identified articles of relevance were supplemented through hand search, including review of 'cited in' articles in PubMed and with additional publications known to the authors or cited in the papers considered.

3. Results

3.1. Guidelines included in this review

Seven sets of national or international society guidelines or consensus statements were identified. Three of these (European Association for the Study of the Liver [EASL] [20], National Institute for Health and Care Excellence [NICE] [25] and Asia-Pacific Working Party [22,23]) were based on systematic literature reviews. Literature review and/or consensus among experts supported recommendations in other guidelines (Table 1) [21,24,26,27]. The American Association for the Study of Liver Diseases (AASLD) guideline was an update to an earlier document published in 2012 [31]. The earliest published guidelines were from Italy (2014) [26], with the rest published in the years 2016 to 2018 [20-25,27].

Only three of the guidelines use an evidence-based approach (the Grading of Recommendations Assessment, Development and Evaluation [GRADE] system) to develop their recommendations: EASL [20], Asia-Pacific Working Party [22,23] and the Chinese National Consensus Workshop [27] (Table 1). The AASLD [31] differentiates its guidance developed from expert consensus from evidence-based recommendations. Quality assessment used and considered ranking of credibility.

3.2. Comparative overview of guidelines and consensus statements for NAFLD

3.2.1. Summary and overview

Seven guidelines and consensus statements published in English were identified for inclusion (Table 1).

All included guidelines provided recommendations for screening of individuals deemed likely to have NAFLD (Table 2), their diagnosis and where there is a risk of advanced disease, an assessment of severity. In patients with NAFLD with or at risk of severe and/or progressive NASH, recommendations for follow-up and monitoring are included (Tables 1 and 3). They also provided recommendations for management through lifestyle changes and where deemed necessary, off-label prescription of pharmacological agents and use of dietary supplements and plant-derived preparations that may protect the liver against damage (hepatoprotectors), as well as surgical interventions (Table 4).

Screening patients with NAFLD risk factors in primary care is essential to identify those in need of further evaluation in specialist hepatology services for risk of advanced disease [30,32-35]. However, there was a general lack of detailed guidance on referral to hepatologists and implementation of multidisciplinary models of care in the guidelines. In particular, screening, diagnostic and follow-up pathways and recommendations for primary care management and referral pathways were not clearly stated.

3.2.2. Risk groups

All seven guidelines define at-risk populations in which the exclusion of other causes, such as viral hepatitis or excessive alcohol consumption has to be considered (Table 1). All guidelines except those from NICE in England and Wales [25] included a lower limit for hepatic fat accumulation of 5%.

Although all guidelines acknowledged multiple risk factors for NAFLD, particularly T2DM, metabolic syndrome or obesity, there was considerable variation in the number and types of other risk factors between sets of guidelines, most of which appeared in three guidelines or fewer (Table 2). All except the Asia-Pacific guidelines, which refer to overnutrition as 'invariable' in this context [23], included T2DM and metabolic syndrome as risk factors (Table 2).

All except guidelines issued by NICE in England and Wales did not include the presence of obesity as a risk factor (Table 2); However, NICE does identify this condition as a target for intervention to reduce overweight via its specific obesity recommendations (CG189) [25,36]. In the context of bodyweight, it should be noted that managing 'lean NAFLD' (i.e., without obesity/elevated body mass index [BMI <30 kg/m² [20,27]]) is highlighted as an issue in some populations, in particular Asian subjects and those with metabolic disturbances [20,23,25,27].

Insulin resistance, which is considered to have a central role in the development of NAFLD, was only mentioned in guidelines issued by the World Gastroenterology Organisation (WGO) [26], EASL [20] and the Asia-Pacific region [23]. Other metabolic risk factors mentioned included hypertension [26], dyslipidemia [21,26], weight gain [23,24] and sleep apnea [21,27] (Table 2).

Although the role of genetics, particularly the SNPs *PNPLA3*, *TM6SF2* and *MBOAT7*, in NAFLD, NASH and HCC risk are recognized (Carlsson, 2020), current guidelines do not recommend routine screening for these mutations (Table 1). Although genetic targets for therapeutic interventions offer the possibility of precision medicine, this aspect of NAFLD medicine must be considered in its infancy currently (Carlsson, 2020).

3.2.3. Modalities used for screening for NAFLD and assessing severity

There was agreement across all guidelines that universal screening of the general population is not recommended. Six of the seven guidelines, but not those issued in the USA [21], included recommendations for screening of high-risk populations defined by the presence of the risk factors, including obesity with or without T2DM [20,23,25,27], metabolic syndrome and abnormal liver enzyme profiles [20], NASH cirrhosis [24], and insulin resistance [26] (Table 1). Subsequently, EASL has issued updated its specific clinical practice guidelines on the use of non-invasive tests (NITs) for assessing severity of (Berzigotti, 2021).

Although evaluation of liver biopsy is the reference standard method for the grading and staging of NAFLD it is not practical or affordable for large-scale, routine use [25,32,34,37]. Alternative, non-invasive and cost-effective tests that are easy to implement are therefore necessary to sequentially screen and assess the large number of patients who may have NAFLD, exclude those with low risk of advanced disease from further assessment, and identify those at risk of severe disease [32,34].

For screening of patients who may have NAFLD, all seven of the guidelines we analyzed highlighted use of NITs, including blood-based, imaging and elastography techniques to exclude those at low risk of AF (who can be managed using diet and lifestyle modification in the first instance) and help identify those in need of further assessment for liver fibrosis. There is ongoing evaluation of NIT selection in routine clinical practice and choice of methodology may be influenced by a range of factors, including proven diagnostic ability, operator competence, ease of access, clinical setting (primary versus specialist care), patient preference and cost [32–37]. A detailed evaluation of these factors is beyond the scope of this review, and the recent review by Campos-Murguía *et al* [37] is recommended for test-specific technical details, together with the latest practice guidance from EASL for the use of NITs in liver disease assessment (Berzigotti, 2020).

Beyond individual measures of liver status/performance and patient characteristics that are known risk factors (Table 2), composite scores based on risk factors assessment (T2DM, obesity etc.) and serological measurements are recommended. In particular the Enhanced Liver Fibrosis (ELF) test, the Fatty Liver Index (27), Fibrosis-4 (FIB-4) Index, Hepatic Steatosis Index, and NAFLD Fibrosis Score (NFS) can be used for assessment of disease severity (including fibrosis) and risk of progression [32,34,37].

Guidelines from NICE in England and Wales [25], the Asia-Pacific region [23], China [27], Italy [24] and the WGO [26] recommend ultrasound, supported by measurement of liver enzyme levels in the Chinese National Consensus Workshop and WGO guidelines [26,27] and transient elastography (TE) according to those for the Asia-Pacific region [23] (Table 1).

For patients with suspected NAFLD requiring an assessment of severity, including the presence and extent of fibrosis (see below), the WGO recommends measuring blood markers, anthropometric features (weight, BMI, waist circumference, height), blood pressure and imaging, including ultrasound and abdominal computed tomography scan where required [26]. TE using FibroScan is recommended in the EASL and Italian guidelines as confirmation where other tests are inclusive [20,24]. TE is also recommended for assessment of NAFLD in the guidelines from China along with NFS [27], and may also be used in accordance with AASLD recommendations [21].

Liver fibrosis plays a major role in the progression to severe liver disease and associated mortality [32,37], so its assessment is vital in patients with NAFLD deemed to be at risk of, or suspected to have, fibrosis. Liver fibrosis assessment is recommended using FIB-4 and/or NFS in guidelines from AASLD [21], China [27], EASL [20], and Italy [24]. However, the NFS may not be suitable for use in assessing patients with NAFLD and T2DM [32]. In England and Wales, ELF is recommended test for AF [25].

Liver biopsy is the acknowledged reference standard, but all guidelines agree that it is not suitable for routine screening and assessment. Where specified, liver biopsy is to be

reserved for cases of diagnostic uncertainty after NITs [26] and in cases where patients are considered at risk of having or likely to have NASH or AF [20,21,26,27].

3.2.4. Follow-up recommendations

Guidance on follow-up to monitor risk and progression of disease varies between guidelines, with no recommendations included in those from the Asia-Pacific region [22], the AASLD [21], and the WGO [26]. EASL recommends monitoring in specialist care, including NITs and repeat liver biopsy after at least 5 years in patients at high risk of liver disease progression [20]. Repeat assessment of disease severity every 3 years are recommended by NICE [25] and every 2 years in Italy [24], rising to every 6 months when cirrhosis is present. In China, regular follow-up of 'lean' patients with NAFLD is advised [27]. No generally accepted model of care has yet been published, although NICE in England/Wales has issued a pathway (available at <http://pathways.nice.org.uk/pathways/non-alcoholic-fatty-liver-disease>; see discussion).

Recommendations for HCC screening are not universal. Guidelines from Asia-Pacific [23], China [27] and the USA [21] recommend screening when cirrhosis is present, whereas those from Italy recommend screening specifically in proven cases of NAFLD with AF [24] (Table 3). European guidelines acknowledge the risk of HCC in NAFLD, they do not make any specific recommendation about timing of surveillance [20]. Although outside of the scope of this literature search, it is worth noting that some national guidelines include pre-cirrhotic NASH as a condition for HCC screening, and the American Gastroenterology Association has recently published an evidence-based review and best practice advice for HCC screening in people with NAFLD (Loomba, 2020).

3.2.5. Algorithms in guidelines

Algorithms for diagnosis and/or follow-up were absent in four of the seven guidelines (Table 3) [21,23,24,27]. The guidelines did not include information to support decisions on patient referral from primary care to specialist hepatology care, although the EASL guidelines did

include an algorithm for fibrosis assessment [20]. It is also noteworthy that differentiation and integration of care across primary and secondary care, relating to screening and referral, as well as long-term monitoring, is currently lacking in the guidelines reviewed here. This is reflected in clinical practice, where care pathways are often lacking or, where they do exist, are not standardized based on best-practice principles (Lazarus, 2021a). The paucity of easy-to-follow care pathways developed from evidence-based best practice therefore remains a major hurdle to translating the knowledge base for NAFLD and NASH into improved outcomes for patients.

3.2.6. *Treatment using lifestyle modification*

All guidelines recommend the use of lifestyle modification to manage NAFLD. Interventions feature structured programs incorporating diet and regular physical activity, often with personalization and expert nutritional support, to facilitate long-term adherence [20-22,25-27]. However, no clear and consistent guidance was provided regarding the care pathway through which lifestyle modification should be delivered.

All seven guidelines reviewed included recommendations for weight loss, with only those from NICE not quantifying therapeutic weight loss [25]. Five of the seven propose a standard target weight loss 5%–10% [20,23,24,26,27], whereas the AASLD recommends weight loss stratified by NAFLD severity: 3%–5% for steatosis and 7%–10% for NASH [21]. To achieve recommended weight loss, guidelines from Asia-Pacific, Europe, China and the USA recommend a target daily energy *intake deficit* of between 500 and 1000 kcal [20-22,27], whereas in Italy a target daily *intake limit* of 1200 to 1600 kcal is stated [24]. WGO guidelines indicate reducing by 25% the normally recommended daily calorie intake to achieve weight loss [26].

3.2.7. *Treatment using pharmacological agents and supplements*

Despite the universally acknowledged absence of an approved pharmacological therapy in the explored countries, all of the guidelines identified in this review included consideration of

the use of pharmacological agents and nutritional supplements. However, their use was restricted to patients with NASH [20-22,24,27], and/or fibrosis [20,22,25,27] or at high risk of disease progression [20,24] (Table 4). No guidelines made strong, evidence-based recommendations for the use of any pharmacological agent, with three stating that specific agents ‘may be used’ [21,22,27]. EASL states that ‘no firm recommendation’ can be made [20], and the WGO states that any use is to be ‘considered experimental’ [26] (Table 4).

Several of the recommended agents described in Table 4 target, and are appropriately indicated for, NAFLD risk factors and comorbidities. Specifically, in the context of NAFLD and NASH, they have been evaluated for their effects on blood markers and histology (reviewed in the development of the European guidelines) [20]. However, other than in India no drugs have yet been approved specifically for the treatment of NAFLD or NASH. A systematic review undertaken in the development of the EASL guidelines of randomized controlled trials in NAFLD that included histological outcomes showed poor, inconsistent efficacy across drug types in small or medium-sized trials [20]. It is also noteworthy that most of the trials of antidiabetic drugs that are considered potentially beneficial in managing NAFLD, in particular glucagon-like peptide I receptor antagonists (GLP-1 RAs) and sodium-glucose cotransporter 2 (SGLT-2) inhibitors, were ongoing and unpublished when the guidelines under consideration were being developed (Dougherty, 2021). Although not suitably indicated, GLP-1 RAs are favorably viewed for evaluation as pharmacotherapy in NAFLD, due to their combined effects on markers of steatosis, weight and cardiometabolic parameters (Dougherty, 2021; Niederseer, 2021; Wilding, 2021). SGLT-2 inhibitors also show promise, although the risk of urinary tract infections may limit their use (Niederseer, 2021). Both classes of antidiabetic appear to have better benefit–risk profiles than pioglitazone, which features in NAFLD guidelines from EASL [20], NICE [25] and the AASLD [21] (Niederseer, 2021). Other, novel agents with mechanisms of action that target metabolic, inflammatory or fibrotic processes have also been studied, some in late-phase trials, but development has stalled due to benefit–risk profiles perceived as inadequately

favorable [38] (Xanthakos, 2021). For patients with multiple cardiovascular risk factors, statins have been viewed favorably due to their proven safety profile and their strong efficacy for the reduction of the cardiovascular risk, but not for benefits in NAFLD *per se* [28].

Many studies of dietary supplements (e.g., vitamin E and omega-3 fatty acids) and plant-derived hepatoprotectors (e.g., silymarin, phosphatidylcholine and ursodeoxycholic acid [UDCA]) have been performed, with summaries of evidence and outcomes provided in guidelines from EASL [20], the USA [21] and the Asia-Pacific region [22]. In the Chinese guidelines, silymarin, bicyclol, polyene and phosphatidylcholine, along with vitamin E, are noted as having broad utility as add-on treatments for liver injury in patients with chronic liver diseases [27]. However, alack of evidence, the overall low quality of source data where assessments have been made and the need for data from clinical trials were noted.

3.2.8. *Bariatric interventions*

Bariatric surgery is recommended considered in the absence of cirrhosis to reduce obesity (for example, where dietary interventions, lifestyle modifications and/or supplements are unsuccessful) [20-22,26,27] or as an adjunct to liver transplant [20]. No BMI cut-offs for bariatric interventions were specified in guidelines from EASL [20], AASLD [21] or WGO [26]. Those for the Asia-Pacific region [22] define a qualifying BMI greater than 30 kg/m² as eligible for bariatric surgery, whereas guidelines for China [27] differentiate obesity by BMI level (severe obesity: at least 40 kg/m²; moderate obesity: 35 to 39.9 kg/m², stipulating a 2.5 kg/m² reduction in threshold for Asian populations). Patients with NAFLD having a BMI between 30 and 34.0 kg/m² (adjusted accordingly for Asian populations) might also be considered for bariatric surgery when cardiometabolic risk factors are present [27]. Evidence is accumulating in support of bariatric surgery for patients with NASH in need of rapid weight loss, leading to reduction in hepatic steatosis and even resolution of NAFLD in some cases [39-41]. There were no recommendations for endoscopic bariatric devices such as gastric balloons.

3.2.9. Liver transplantation

Liver transplantation is a recognized option for patients with end-stage liver disease due to decompensated cirrhosis or the onset of HCC (Table 4) according to regional standards. However, thorough assessment of patient status, in particular age and cardiovascular and renal disease risk factors, is necessary [20-22,27,42]. High BMI ($>40 \text{ kg/m}^2$) may also present challenges in performing liver transplant [21,22,26]. Risks associated with post-transplant infections should also be borne in mind [20,22,42].

3.3. Clinical practice, awareness and application of NAFLD guidelines in the real world

3.3.1. Disease awareness among patients diagnosed with NAFLD

Patients with significant risk factors for NAFLD repeatedly report low levels of awareness of potential liver disease [43-47]. A recent study of awareness around NAFLD among 30 patients with T2DM revealed that just half were familiar with the concept of ‘fatty liver’ [46]. Although disease awareness among patients with NAFLD was found to be low, it has increased slightly over the past two decades [45]. It is worth noting that just 16 of 667 (2.4%) with imaging-confirmed NAFLD had been made aware of their diagnosis through physician and/or nurse communication [44]. Regarding high-risk comorbidities, NAFLD awareness rates of between 19% and 38% have been reported among those with risk factors such as obesity or T2DM [43]. There was also a poor level of understanding of the therapeutic nature of lifestyle modification and the interplay between NAFLD and T2DM (including insulin resistance), as well as the possible consequences of NAFLD, including risk of progression. Experience of post-diagnosis follow-up examinations, including monitoring of disease progression, was reported by just half of the subjects [46].

Further details on disease awareness among patients and practitioners are provided in the Appendix.

3.3.2. Discrepancies between routine diagnosis and screening practice and the implementation of recommendations in NAFLD guidelines

National healthcare policy and support is essential to facilitate necessary changes in clinical practice to implement clinical guidelines as they become available. Lazarus et al identified very low levels of health policy support and awareness programs for the implementation of NAFLD guidelines across Europe [30]. Surveying experts across Europe and reviewing official documents relating to policies, clinical guidelines, awareness and monitoring showed that necessary guidance and information is missing on a country level. In the 29 countries studied, there were no written strategies or action plans for NAFLD despite over 40% having policies on obesity, cardiovascular disease, T2DM and/or healthy living and nutrition. Two countries incorporated NAFLD/NASH into obesity and alcohol strategies. Subsequent analysis of these data and evaluation of a 'European Preparedness Index' for meeting the NAFLD challenge shows that none of the countries involved in the study have a high level of preparedness (Lazarus, 2021b). Only the UK could be considered to have above a low level of preparedness in this scenario.

A third of countries had issued national guidelines, all of which recommended screening of risk factors and liver cirrhosis. Specifically, in the UK, only one-fifth of 84 specialist gastroenterology/hepatology centers reported having access to local NAFLD guidelines [48]. Specialist care (gastroenterology and hepatology) was responsible for NAFLD management in over 80% of countries, and around half involved primary care [30]. However, in keeping with the observations on the published guidelines, just five countries (17%) had developed algorithms for primary follow-up and referral to specialist care [30].

Given the underdiagnosis of NAFLD and NASH, implementing guidelines should help to identify cases in need of monitoring and/or referral. However, there is concern that attempts to optimize diagnosis could result in a rapid increase in the number of patients being referred to specialized care [49-51]. Over-referral from primary to specialist care has been identified as a potential problem, particularly among patients with T2DM [50]. On the other hand,

referral rates may often be higher among specialists than physicians in primary care [52-55]. Moreover, if appropriately applied, adhering to current guidelines could avoid unnecessary rates of invasive liver biopsy [56]. For example, patients in specialist care for HIV mono-infection suspected of having NAFLD are not subjected to unnecessarily high rates of referral to hepatologists when EASL guidelines are applied [57], further supporting the application of current clinical guidance. Developments in nurse-led screening could further help redirect patients from specialist back to primary care, thereby avoiding or limiting the impact and increased workload due to higher numbers of referrals [58].

Development of clear algorithms and training for primary care physicians would also facilitate selection of patient types for screening, including a recognition of fibrosis and its implications for advanced disease [35,58,59]. For example, applying local processes such as the UK's Camden and Islington NAFLD Pathway can reduce unnecessary referrals to specialist care by 80%, offering both clinical and economic efficiencies [60,61]. However, there are currently widespread deficiencies in the number and standardization of care pathways for NAFLD (Lazarus, 2021a).

Specifically, regarding fibrosis evaluation, the available data indicated that assessment of fibrosis using non-invasive techniques is not consistently conducted according to guideline-defined recommendations, and many physicians are unsure of the assessments and interpretations required or do not apply them rigorously [52,54,62]. Limited use of non-invasive scoring systems and infrequent referral of high-risk patients to specialists as described above likely result in underdiagnosis and missed opportunities to identify patients at risk of progression to severe, advanced NASH [55,63].

3.3.3. Lifestyle modification and pharmacotherapeutic management of NAFLD in practice

Adherence to lifestyle modifications has repeatedly been observed to be suboptimal [64,65]. In a study of patients with NAFLD in Russia, 86% of those assigned to strict diets were not adherent to strict dietary recommendations on more than a few occasions per month [64].

Among patients with T2DM, levels of disease awareness and understanding of the therapeutic importance of lifestyle modification were found to be low [46]. Moreover, a qualitative study revealed low levels of patient education, support and follow-up related to their diagnosis and implementation of lifestyle modification [47]. Poor understanding and inadequate support have been associated with low adherence to lifestyle modification programs among patients with NAFLD across Europe [65-67].

Improved patient education about NAFLD and the therapeutic role of lifestyle modification to reduce future risk of clinically burdensome disease could help to bridge this gap in awareness [66,68]. However, an evaluation of online material developed for patients with liver cirrhosis highlighted that the available information from health platforms and specialist hepatology centers in the USA is often overly lengthy and complex, hindering broad understanding [69].

Per guidelines, analysis of real-world practice reveals that most physicians recognize and adhere to recommendations for lifestyle modification through physical activity and diet to manage NAFLD. These studies also show that pharmacotherapy is commonly used despite the lack of approved drugs and minimal, generally low-quality, evidence base for agents cautiously mentioned in guidelines. The number of individual drugs and rates of use vary across studies, which may be partly driven by local practices. Overall, use of drugs otherwise indicated for the management of risk factors and that may improve liver pathology (e.g., metformin, glitazones and statins), or other agent, including vitamin E, silymarin, phosphatidylcholine and UDCA, was reported repeatedly: Asia [70], France [71], Germany [67], Poland [54], Romania [72], Russia [64] and the USA [73].

Current guidelines emphasize that supportive evidence for some treatments is lacking or inadequate (e.g., metformin and UDCA). However, usage of products such as vitamin E, UDCA, phosphatidylcholine and silymarin) and drugs classes such as antidiabetics (glitazones, GLP-1 antagonists) and statins is observed in real-life practice. Clinical evidence from multiple sources reviewed in guidelines from EASL [20], the Asia-Pacific region [22],

and the USA [21] does not indicate a clear benefit in terms of reducing fibrosis, but does suggest that tolerability is generally acceptable. On the other hand, a number of small but promising clinical trials, the results of which were not available when the guidelines were being developed, suggest that GLP-1 RAs and SGLT-2 inhibitors offer promise in the management of NAFLD across NASH and cardiometabolic endpoints (Niederseer, 2021; Dougherty, 2021). Further investigations in clinical trials are needed to better understand the therapeutic effects of these treatments on steatohepatitis and fibrosis in NAFLD [27].

4. Discussion and conclusion

This literature review identified the availability of current clinical guidelines that offer recommendations to for the identification, evaluation and management of the growing number of patients with NAFLD or at risk of NASH in the absence of approved pharmacological interventions. However, despite broad consistencies in their approach, inconsistencies across guidelines and gaps in recommendations that support their application risk suboptimal clinical practice, including the identification of those individuals most at risk of severe disease who require referral to specialist care. This is supported by observational data from routine clinical practice.

Although extensive clinical guidelines on the screening, diagnosis and management of NAFLD have been published, the real-world evidence that we have discussed reveals substantial shortfalls in disease awareness and management, patient education and adherence to therapeutic programs, as well as physician practice, referral and the off-label use of pharmacological agents. Rates of awareness were lowest among primary care physician and non-gastroenterology/hepatology specialists [43,52,62]. The risk of under-recognition due to low awareness of patients with or at risk of NASH/cirrhosis is compounded by low rates of referral from primary care and 'other' specialists to gastroenterologists or hepatologists [52-55]. This is a particular clinical concern in cases

where patients with normal liver enzymes who are, nevertheless, at high risk of NAFLD are overlooked in primary care [52,71]. Consequently, rates of NAFLD and the associated clinical, healthcare system and economic burden continue to increase.

Education about NAFLD and relevant guidelines would help primary care physicians overcome the reported lack of comfort dealing with liver diseases [48,63,74,75]. When asked, general physicians expressed requirements for improved levels of awareness, knowledge and confidence to allow improved identification and management of liver disease [74,75]. As patients may be referred back to primary care for lifestyle modification therapy in some systems [48], there remains an unmet need for development of skills and the use of multidisciplinary care to optimize early management of NAFLD in community and primary health services, and streamline specialist hepatology care accordingly [33,60,61,63].

Although the majority of clinicians recognize and adhere to recommendations for lifestyle modification through physical activity and diet, as we describe here, patients often appear not to understand the therapeutic intent of such interventions and have poor adherence to them. Consequently, the potential benefits of dietary control and physical activity may be limited.

Dietary supplements and hepatoprotective agents are commonly used, despite variable evaluations in clinical guidelines. Guidelines allow for some off-label use of drugs to aid management of NAFLD, specifically reducing impact of risk factors and, based on limited evidence, improving disease pathology. However, pharmacotherapy usage is commonly reported in observational studies. Despite substantial investment in clinical development, pharmacological treatment has yet been approved for advanced disease, despite the existence of evidence that interventions can reduce steatosis and dysmetabolism. The absence of clear, evidence-based recommendations for pharmacotherapy of NAFLD is partly due to the practical challenges of designing and implementing trials of new drugs in NAFLD impeding progress [38] and partly due to the timing of data availability versus guideline development (Dougherty, 2021).

In advanced disease, including end-stage liver disease, surgical interventions involving highly specialized, targeted care can offer major clinical benefits. We did not examine data relating to bariatric surgery in the publications we reviewed. Likewise, there were no evaluations of the outcomes of liver transplantation, although there is growing evidence that it can be beneficial in suitably selected patients [39-42]. However, the cost-effectiveness and long-term effectiveness of bariatric surgery for weight loss and liver transplantation to correct liver failure require more evidence in the context of the increasing burden of NAFLD and its consequences.

In conclusion, despite the availability of clinical recommendations and guidelines for the diagnosis and management of NAFLD, real-world practice reveals substantial disparities in levels of application and outcomes. The evidence indicates a need for education, training and development of algorithms to close the gap between current and optimal screening, diagnosis and management of NAFLD. This needs to be supplemented by healthcare policy and associated structural developments to turn recommendations into actionable patient care.

5. Expert opinion

Since the mid-2010s, multiple national clinical guidelines for the diagnosis, assessment and management of NAFLD and NASH has been published (in English) . The need for such clinical guidance reflects the growing prevalence of NAFLD and its increasing health, humanistic and economic burden, particularly among those who have or are at risk of developing NASH and/or fibrosis. However, challenges persist in the optimal identification of these patients and their direction into appropriate specialist care must be prioritized to reduce the impact of NAFLD on individual and public health.

Application of guideline recommendations for promotion of lifestyle changes is often suboptimal. To date, no drugs have received regulatory approval for the treatment of NAFLD or NASH. All guidelines included weak recommendations for pharmacological support,

restricted to patients diagnosed with NASH with or at risk of fibrosis or disease progression. Investigational drugs have reached late phases of development, but none to date have demonstrated benefit–risk profiles likely to be acceptable to health regulators. Overall, NAFLD clinicals are therefore limited in terms of recommendations for management of progressive, advanced disease.

In practice, awareness of NAFLD, its diagnosis and management remain poor outside of specialist hepatology. We suggest that poor guidance and education may result in suboptimal rates of diagnosis, and potentially leading to cases of serious, potentially life-threatening NASH being overlooked. There is a clear need for education and for integration across healthcare disciplines to improve identification of at-risk individuals and to optimize referral pathways. Development and adoption of algorithms for sequential screening and diagnosis of NAFLD and risk assessment based on these guidelines could help address this need.

Although early intervention may help to minimize the health impact of NAFLD, patients frequently appear not to understand the therapeutic intent of such interventions and rates of adherence to such recommendations are low . In this context, the recent recommendations from the international Liver Forum's Standard of Care Working Group proposing standardized core features of lifestyle modification assessment and implementation in the management of NAFLD, principally to refine measurement of outcomes in clinical trials, are important to highlight [76].

We believe that harmonizing clinical guidelines and developing clear, easily-applied screening and assessment algorithms will help establish optimum pathways of care for patients with NAFLD. In the first instance, this will help to streamline referral and monitoring of individuals according to disease status and severity as well as risk of progression, making the most of current options for clinical care. It would also aid efficient allocation of health resources and budgets. By optimizing current resources, the true extent of unmet needs in NAFLD and NASH management can be fully understood, which will help the development of

new pharmacological interventions on a background of best practice in both primary care and specialist hepatology services. Identification of clinical trial populations, definitions of care in the control arms of randomized clinical trials, and selection of endpoints and appropriate size effects for assessment of efficacy will be central to improving drug development and assessment of agents approved for other, physiologically relevant conditions. Additionally, the development and routine implementation of precision medicine based on genetic screening will help to direct efficient and optimal management using available resources and agents as they evolve.

There is encouraging evidence that this optimization of care and identification of trial-ready patient populations is beginning to be realized. Published models of care, although limited in number, demonstrating impacts on outcomes and cost efficiencies are becoming available. Likewise, testable, algorithms for sequential assessment of patients at risk of advanced NASH fibrosis are in the literature. To fully assess standards of care, there is a need for large, long-term, international observational studies to fully understand the state of current practice such as the European NAFLD Registry [77] and the TARGET-NASH study [78]. The results of these investigations can help identify and support the development of new paradigms to inform updated, refined guidelines in the light of current challenges. Improved standards of care will help to define patient cohorts and clinical outcomes for the investigation and development of new interventions aimed at alleviating the growing burden of NAFLD and NASH.

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New citations to add

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Tables

Table 1. Current NAFLD guidelines for the screening, diagnosis and management of NAFLD in adults.

Table 2. Comparison of NAFLD risk factors stated in the guidelines under consideration.

Table 3. Markers and measurements of liver fibrosis in current NAFLD guidelines.

Table 4. Drugs included in NAFLD guidelines for the management of NAFLD/NASH.

Table 1. Current NAFLD guidelines for the screening, diagnosis and management of NAFLD in adults.

	EASL [20]	NICE [25]	Asia-Pacific [22,23]	AISF [24]	AASLD [21]	CSH [27]	WGO [26]
<i>Year of publication</i>	2016	2016	2018	2017	2018	2019	2014
<i>Region/country</i>	Europe	England and Wales	Asia-Pacific	Italy	USA	China	Global
<i>Evidence base/origin</i>	Systematic literature review	Systematic literature review	Systematic literature review and author consensus	Literature review	Literature review and author consensus	Author consensus	Author consensus
<i>GRADE-based evidence assessment</i>	Yes	No	Yes	No	No	Yes	No
<i>Screening</i>							
• Systematic	No	No	No	No	No	No	No

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• High risk (subgroups)	Yes (Ob, met syn, abnormal liver enzymes)	Yes (Ob, T2DM)	Yes (Ob, T2DM)	Not stated	No	Yes (risk factors)	Yes (risk factors)
• Recommended methods	Liver enzymes	US	US, TE	-	-	US or liver enzymes	US, liver enzymes
• Genetic screening	Selected patients and in clinical trials (PNPLA3 I148M; TM6SF2 E167K)	No	No	No	No	No	No
<i>Diagnosis</i>							
• Criteria	Hepatic steatosis >5%*, insulin resistance, no other causes of steatosis	Excessive fat in liver, no other causes of steatosis	Hepatic steatosis*, no other causes of steatosis	Hepatic steatosis*, no other causes of steatosis	Hepatic steatosis*, no other causes of steatosis	Hepatic steatosis*, no other causes of steatosis	NASH: Hepatic steatosis >5%*, insulin resistance, no other causes

							of steatosis
• Alcohol limit (men)	30 g/d	30 g/d	140 g/wk (2 standard drink/d)	30 g/d	294 g/wk** (21 standard drinks/wk)	210 g/wk	30 g/d
• Alcohol limit (women)	20 g/d	20 g/d	70 g/wk (1 standard drink/d)	20 g/d	196 g/wk*** (14 standard drinks/wk)	140 g/wk	20 g/d

*Identified using either imaging or histology

**Equivalent to an average of 42 g/d

***Equivalent to an average of 28 g/d

AASLD: American Association for the Study of Liver Diseases; AISF: Italian Association for the study of the Liver; CSH: Chinese Society of Hepatology; d: day; EASL: European Association for the Study of the Liver; GRADE: Grading of Recommendations Assessment, Development and Evaluation; met syn: metabolic syndrome; NAFLD: non-alcoholic fatty liver disease; NASH: non-alcoholic steatohepatitis; NICE: National Institute for Health and Care Excellence; Ob: obesity; T2DM: type 2 diabetes mellitus; TE: transient elastography; wk: week; WGO: World Gastroenterology Organisation

Table 2. Comparison of NAFLD risk factors stated in the guidelines under consideration (green: included in at least four of the guidelines; grey: mentioned in three guidelines; red: included in one or two guidelines).

	EASL [20]	NICE [25]	Asia-Pacific [22,23]	AISF [24]	AASLD [21]	CSH [27]	WGO [26]
<i>Region/country</i>	Europe	England and Wales	Asia-Pacific	Italy	USA	China	Global
<i>T2DM</i>	✓	✓		✓	✓	✓	✓
<i>Metabolic syndrome</i>	✓	✓		✓	✓	✓	✓
<i>Obesity</i>	✓		✓	✓	✓	✓	✓
<i>Insulin resistance</i>	✓		✓				✓
<i>Overnutrition</i>			✓				
<i>Sleep Apnea</i>					✓	✓	
<i>Dyslipidemia</i>					✓		✓

<i>Polycystic Ovary Syndrome</i>					✓	✓	
<i>Hypopituitarism</i>					✓	✓	
<i>Hypertension</i>							✓
<i>Ethnicity</i>				✓			✓
<i>Age</i>	✓			✓			
<i>Weight gain</i>			✓	✓			

AASLD: American Association for the Study of Liver Diseases; AISF: Italian Association for the study of the Liver; CSH: Chinese Society of Hepatology;

EASL: European Association for the Study of the Liver; NAFLD: non-alcoholic fatty liver disease; NASH: non-alcoholic steatohepatitis; NICE: National Institute

for Health and Care Excellence; T2DM: type 2 diabetes mellitus; WGO: World Gastroenterology Organisation

Table 3. Screening guidance and use of markers and measurements of liver fibrosis in current NAFLD guidelines.

	EASL [20]	NICE [25]	Asia-Pacific [22,23]	AISF [24]	AASLD [21]	CSH [27]	WGO [26]
<i>Region/country</i>	Europe	England and Wales	Asia-Pacific	Italy	USA	China	Global
<i>Algorithm</i>	Yes*	Yes**	No	No	No	No	Yes
<i>Noninvasive tests</i>	NFS, FIB-4. TE where other tests are inconclusive	ELF for all patients with NAFLD,	Serum test and imaging (specific tests not specified)	NFS plus FIB-4. TE where other tests are inconclusive	NFS or FIB-4 to identify patients at high risk of AF. VCTE™ or MRE may also be used.	NFS, TE	Serum tests (specific tests not specified)
<i>Follow-up</i>	Progression followed in	No evidence of fibrosis repeat	Not stated	Negative test: repeat every 2 years;		Diagnosis of NASH: liver	Not stated

	specialist care	testing every 3 years Evidence of fibrosis: liver biopsy		fibrosis or abnormal liver enzymes: erases every year; cirrhosis: reassess every 6 months		biopsy	
HCC screening	No recommendation made per timing	Not stated	NASH with cirrhosis	NAFLD with AF and/or cirrhosis	NAFLD with cirrhosis	NASH with cirrhosis	Not stated

*For assessing severity and risk of fibrosis

**Available at <http://pathways.nice.org.uk/pathways/non-alcoholic-fatty-liver-disease> (last accessed 16 July 2021)

AASLD: American Association for the Study of Liver Diseases; AF: advanced fibrosis; AISF: Italian Association for the study of the Liver; CSH: Chinese Society of Hepatology; EASL: European Association for the Study of the Liver; ELF: Enhanced Liver Fibrosis; FIB-4: Fibrosis-4 Index; HCC: hepatocellular carcinoma; MRE: magnetic resonance elastography; NAFLD: non-alcoholic fatty liver disease; NASH: non-alcoholic steatohepatitis; NFS: NAFLD Fibrosis

Score; NICE: National Institute for Health and Care Excellence; TE: transient elastography; VCTE: Vibration Controlled Transient Elastography; WGO: World Gastroenterology Organisation

Table 4. Patient groups and strength of recommendations for pharmacotherapy, and surgery recommendations included in clinical guidelines for the management of NAFLD/NASH (recommended drugs are not approved by regulatory bodies specifically for the treatment of NAFLD as of December 2020; shading: green, recommended in guidelines (off-label use form prescription-only medicines]; grey, may be considered; red, not recommended; white, not stated).

	EASL [20]	NICE [25]	Asia-Pacific [22,23]	AISF [24]	AASLD [21]	CSH [27]	WGO [26]
Patient groups considered suitable for pharmacotherapy	NASH, particularly with significant fibrosis (\geq F2); high risk of progression	AF	NASH and/or fibrosis	NASH at maximal risk of progression	Biopsy-proven NASH	Suspected of NASH; AF in the absence of biopsy	Not stated
Strength of recommendation for pharmacotherapy	'No firm recommendation can be made'	Not stated	Specific agents 'may be used'	Not stated	Specific agents 'may be considered'	'May be considered'	Any use 'should be considered experimental'
<i>Drugs approved for risk factors</i>							
Metformin	x	x	x	-	x	✓*	x

PPAR-gamma agonists (glitazones)	✓	✓	✗	✗*	✓	✓*	✗
Statins	✓*	✓*	✓*	✓*	✓*	✓*	✓
GLP-1 analogues	✗*	✗	✗	✗*	✗		
<i>Supplements and hepatoprotective agents</i>							
Vitamin E	✗	✓	✗	✗	✓	✗*	✗*
PUFA	✗	✗	✗	-	✗	-	-
Pentoxifylline	✗	-	✗	-	-	✗*	✗*
UDCA	✗	✗	-	-	✗	-	-
Phosphatidylcholine	-	-	-	-	-	✗*	
Obeticholic acid	✗	-	✗**	✗**	✗	-	-
Silymarin	-	-	✗*	-	-	✗*	-
Omega-3 fatty acid supplements	-	✗	✗	-	✗	✓*	✗

Bariatric surgery	Acceptable for improving obesity and T2DM No BMI limit stated	Not mentioned	Acceptable in patients with obesity BMI >30 kg/m ²	Not mentioned	As adjunct to liver transplant No BMI limit stated	To improve liver histology in obesity, metabolic syndrome and T2DM Stratified: ≥40 / ≥35 and ≤39.9 / ≤34.9 kg/m ² Adjustment for Asian patients: reduce BMI score by 2.5 kg/m ²	To reduce obesity No BMI limit stated
Liver transplant	Acceptable in NASH with end-stage liver disease	Not mentioned	Acceptable in NASH with end-stage liver disease	Not mentioned	Acceptable in NASH with end-stage liver disease	NASH-related decompensated cirrhosis, HCC	In patients meeting criteria for liver transplantation

Drugs/classes considered for use indicated by ✓. Where use is restricted to specific groups, these are stated in parentheses. Those where evidence is considered insufficient, or where the agent is not beneficial are indicated by ✕. ✓*: acceptable safety but not considered beneficial; ✕*: insufficient evidence, potentially useful; ✕**: no trial results

AASLD: American Association for the Study of Liver Diseases; AF: advanced fibrosis; AISF: Italian Association for the study of the Liver; BMI: body mass index; CSH: Chinese Society of Hepatology; EASL: European Association for the Study of the Liver; GLP-1: glucagon-like peptide 1; HCC: hepatocellular carcinoma; NAFLD: non-alcoholic fatty liver disease; NASH: non-alcoholic steatohepatitis; NICE: National Institute for Health and Care Excellence; PPAR: peroxisome proliferator-activated receptor; PUFA: polyunsaturated fatty acids; T2DM: type 2 diabetes mellitus; UDCA: ursodeoxycholic acid; WGO: World Gastroenterology Organisation

Appendix

Non-alcoholic fatty liver disease (NAFLD) awareness and implementation of clinical guidelines among primary care and specialist healthcare providers

Multiple publications of studies of disease awareness among healthcare professionals and patients and the implementation of clinical guidelines for NAFLD were noted in the literature search results. Although not a major focus of this review, their findings were considered instructive and key results are summarized here (**Table A1**).

Table A1. Awareness of NAFLD among patients and healthcare providers, and implementation of clinical guidelines by primary care physicians and specialists.

	Disease awareness/familiarity	Underestimate of prevalence	Awareness of guidelines	Diagnosis of NAFLD against guidelines	Diagnosis of NASH/fibrosis	Screening and diagnostic tools	Referral to specialists
Patients	19%/38% (obesity/T2MD) [53] 31% [43]						
Clinicians (various)		70% to 84% [43]		>80% based on liver enzymes of US [54]	50% MRI [54]		>75% [54]
Primary care/general physicians	18% [62]	51% [52]		76% [52]	>63% unsure about fibrosis	56% [43] 56%/53%/44% (NAFLD/NASH/T2DM) [62]	28% [53] 56% overall/ 71% with normal liver

					scores [52]		function tests [52]
Specialists: Gastroenterologists/ hepatologists	88% [62]				13% (lab) and 9% (US) for fibrosis [62]	71%/71%/68% (NAFLD/NASH/T2DM) [62]	70%/22% (increasing AT/normal AT) [71]
Specialists: other	6% (diabetologists) [55]	37% (obesity care practitioners) [79]	59% (obesity care practitioners) [79]	90% used US (obesity care practitioners) [79]			67% (internal medicine, family medicine, endocrinology, cardiology, obstetrics and gynecology) [53]

MRI: magnetic resonance imaging; NAFLD: non-alcoholic fatty liver disease; NASH: non-alcoholic steatohepatitis