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## Physical Activity and Nonalcoholic Fatty Liver Disease: A Roundtable Statement from the American College of Sports Medicine

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## Abstract

Although physical activity (PA) is crucial in the prevention and clinical management of nonalcoholic fatty liver disease (NAFLD), most individuals with this chronic disease are inactive and do not achieve recommended amounts of PA. There is a robust and consistent body of evidence highlighting the benefit of participating in regular PA, including a reduction in liver fat and improvement in body composition, cardiorespiratory fitness, vascular biology and health-related quality of life. Importantly, the benefits of regular PA can be seen without clinically significant weight loss. At least 150 minutes of moderate or 75 minutes of vigorous intensity PA are recommended weekly for all patients with NAFLD, including those with compensated cirrhosis. If a formal exercise training program is prescribed, aerobic exercise with the addition of resistance training is preferred. In this roundtable document, the benefits of PA are discussed, along with recommendations for 1) PA assessment and screening; 2) how best to advise, counsel and prescribe regular PA and 3) when to refer to an exercise specialist.

## Keywords

EXERCISE; STEATOHEPATITIS; STEATOSIS; LIFESTYLE MODIFICATION; PREVENTION

## INTRODUCTION

Nonalcoholic fatty liver disease (NAFLD) has become a widespread pandemic driven by the increasing prevalence and incidence of obesity. Conservative estimates suggest 25–30% of the global population has NAFLD.(1, 2) In the US, almost 100 million adults have NAFLD, 25 million of whom have the progressive type called nonalcoholic steatohepatitis (NASH).(3, 4) In the absence of a regulatory-agency approved drug therapy or cure, disease progression to cirrhosis or primary liver cancer with hepatocellular carcinoma (HCC) is not uncommon and may necessitate life-saving liver transplantation. In fact, NASH-cirrhosis continues to grow as a leading reason for liver transplantation.(5, 6) Although regular

physical activity (PA) has many well-established benefits and may prevent or delay NAFLD and its complications, most individuals with NAFLD are inactive and fail to achieve recommended amounts of PA.(7–15) In this roundtable document, the benefits of PA are discussed, along with recommendations for 1) PA assessment and screening; 2) how best to advise, counsel and prescribe regular PA and 3) when to refer a patient with NAFLD to an exercise specialist.

### **Diagnosis, classification and etiology of NAFLD**

Currently, the American Association for the Study of Liver Diseases (AASLD) requires the following to diagnose NAFLD: 1) at least 5% hepatic steatosis in the absence of an alternative cause of steatosis (e.g., steatogenic medication use, starvation, monogenic disorders) and 2) the absence of significant alcohol consumption, defined as <20g/day for women and <30g/day for men.(16) NAFLD comprises two histological entities - nonalcoholic fatty liver (NAFL) and NASH. NASH, which can occur with or without liver fibrosis, is the progressive type of disease which may lead to cirrhosis or HCC. While NAFLD remains a complex disease process, it is generally agreed that a “multiple hit” pathogenesis, where environmental factors act on a genetically predisposed individual, is required to develop NAFLD and progress to NASH.(17) NAFLD occurs frequently in individuals with other metabolic conditions, such as dyslipidemia, metabolic syndrome, obesity, polycystic ovary syndrome and/or type 2 diabetes.

### **Treatment goals in NAFLD**

The primary goal of treatment in people with NAFLD/NASH is to prevent mortality and major adverse liver outcomes. It is generally accepted that reversing, arresting, or preventing liver fibrosis will lead to decreased risk for these clinical outcomes, and that a reduction in hepatic steatosis may be beneficial as well. Patients with NAFLD can decrease their liver fat by enacting a lifestyle intervention with Mediterranean-informed dietary change and increased PA. When coupled with clinically significant body weight loss of 7% or greater, histologic improvement with liver fibrosis reversal can be expected.(18) As lifestyle intervention is often viewed as a vehicle for clinically significant body weight loss, dietary change and PA remain crucial in the prevention and clinical management of NAFLD. Because most patients with NAFLD will not die from a major adverse liver outcome but rather from extrahepatic cancers or a cardiovascular disease (CVD) event,(19) therapeutic approaches that also decrease oncologic and CVD risk are ideal. Importantly, regular PA may also play a role in these extrahepatic treatment goals. Additionally, while there is no current regulatory agency approved drug therapy for NASH, novel treatments seem possible in the near future. Regardless of when a drug therapy is approved for NASH, it should augment lifestyle intervention rather than replace it.

## **METHODOLOGY**

To develop a consensus regarding PA and NAFLD, we carried out a Delphi study with a multidisciplinary international panel of 19 academic experts from four countries and 18 academic institutions.(20) The core group of five members developed the initial consensus statements following individual and group systematic reviews. An iterative process was then

undertaken with three rounds of review. The first round (R1) was survey-based, followed by an online consensus meeting [American College of Sports Medicine (ACSM) International Multidisciplinary Roundtable on PA and NAFLD](20) and discussion of each statement and the corresponding levels of evidence (Table 1) for the second round (R2) and a final survey round (R3) of the consensus statements and recommendation. We achieved response rates of 89% in the R1, 100% in R2 and 94% in R3. A super majority (75%) was used for final agreement and inclusion of each statement in this position statement. No consensus could be achieved about the relationship between PA and change in liver fibrosis. The resulting consensus statements and recommendations (Table 2) can serve as a strong guidance for routine clinical management in patients with NAFLD with regards to PA.

### **Role of Physical Activity in NAFLD Pathogenesis**

PA and NAFLD are intricately linked. There is a robust and consistent body of epidemiological evidence demonstrating regular PA reduces the risk of NAFLD, especially if guideline-based amounts of activity are achieved.(7–13, 21) Over time, consistently performing regular moderate-to-vigorous PA (MVPA) can lead to lower mortality overall and from CVD,(8) even if performed in short, continuous bouts of at least ten minutes in length.(21–23) Patients with NAFLD spend more time in sedentary behavior.(15, 24, 25) For those individuals who are more sedentary, the risk of liver fibrosis and disease progression to cirrhosis may be greater.(10) PA is also closely related to oncologic risk. Multiple systematic reviews and meta-analyses have shown at the population level both hepatic and extra-hepatic malignancy risk to be less in individuals who perform regular MVPA or leisure-time activity, including primary cancers which are more commonly found in patients with NAFLD, including as breast, colorectal, and pancreatic.(14, 26, 27)

While the mechanisms underpinning the complex interplay between PA and NAFLD pathogenesis are not yet fully elucidated, regular PA impacts multiple pathways simultaneously,(28–30) including uncoupling protein-1, peroxisome proliferator-activated receptor gamma, adipocytokines, branched chain amino acids, which feedback to reduce insulin resistance, and AMP-activated protein kinase (AMPK).(29, 31–42) Regular PA may also influence gene expression(43, 44) and reverse gut-liver-axis dysfunction.(28, 45, 46)

#### **Evidence statements:**

1. Regular physical activity prevents the development of NAFLD (LOE 2B, Individual Cohort)
2. Emerging evidence suggests individuals who are less physically active may be at increased risk for disease progression to NASH with liver fibrosis and/or cirrhosis (LOE 2B, Individual Cohort)
3. Individuals who are less physically active have greater risk of primary liver cancer (e.g., hepatocellular carcinoma) and other extrahepatic cancers commonly associated with NAFLD (e.g., breast, colorectal, esophageal, gastric, pancreatic, prostate and kidney) (LOE 2B, Individual Cohort)

4. Physical activity offers promise to reverse NAFLD pathogenesis by impacting multiple mechanistic pathways simultaneously and possibly synergistically (LOE 2B Individual cohort studies including low-quality RCT)

### **Assessment and screening patients with NAFLD for physical activity**

Routine healthcare visits offer opportunities for all clinicians to screen patients with NAFLD for PA. While several validated tools exist in research and clinical practice,(47–49) the two question Physical Activity Vital Sign(50–52) is most easily performed and can readily be incorporated into electronic health records. At the population level, implementation of the Physical Activity Vital Sign has been reported to lead to significant loss of body weight in individuals with or at risk for metabolic disease.(53)

Because most individuals with NAFLD do not achieve recommended amounts of weekly PA, screening for barriers preventing routine PA can be helpful and may identify areas to intervene. This intervention should follow an individualized approach, including addressing common self-reported barriers such as cost, energy, fear of injury, motivation, time, skill or social support.(15, 54) While no one optimal tool has been identified, the screening, brief intervention, and referral to a treatment framework (SBIRT) may be considered(55) and used as a scaffolding to design a personalized exercise prescription. NAFLD-specific questionnaires do exist as well for the busy clinician, however, their efficacy remains unknown.(15)

#### **Evidence statements:**

1. All patients with NAFLD should be screened for physical activity during routine healthcare visits with primary care and specialist (e.g., hepatologist) healthcare providers (LOE 2B cohort/low-quality RCT)
2. Tools that provide standardized measures of physical activity (e.g., ACSM Physical Activity Vital Sign) should be administered regularly at routine healthcare visits (LOE 5 Expert opinion)
3. Barriers to physical activity should be assessed systematically. While no one optimal tool has been identified, the Screening, Brief Intervention and Referral to Treatment (SBIRT) framework or NAFLD-specific questionnaires may be considered (LOE 5 Expert opinion)

### **Advising and counseling patients with NAFLD about physical activity**

Because regular PA is important across all stages of NAFLD, all patients with NAFLD should be counseled about the benefits of PA. When approaching this conversation, it is important for the clinician to understand factors which determine whether a person engages in or avoids a certain behavior. Unfortunately, behavioral medicine is a field that is largely ignored across NAFLD research and recommendations in this area are adapted from non-NAFLD populations with metabolic disease. Nevertheless, the importance of multidisciplinary teams, use of patient-centered language and avoidance of stigmatizing words about body weight and body image have all been successful.(56, 57) Using these tools may help encourage NAFLD patients to choose help seeking behavior more often.

Motivational interviewing is perhaps the most widely studied behavioral change technique in patients with NAFLD. A feasible approach can empower patients with NAFLD to make health-related decisions(58) within the context of a short clinic visit and may also lead to a reduction in body weight through dietary change and increased PA.(59) Other techniques which have been studied in patients with NAFLD and may also lead to greater amounts of PA completion include cognitive behavioral therapy(60) and social-cognitive therapy.(61, 62)

Another tool available to the busy clinician comes from widely available print and web-based educational materials, which can be provided to reinforce the knowledge delivered as a part of an in-person conversation. When elected, these education materials, including that available from the ACSM Exercise is Medicine initiative, may lead to success in promoting health behavior change in patients with NAFLD,(63) and should be provided to all patients with NAFLD.

**Evidence statements:**

1. All patients with NAFLD, regardless of disease stage, should be counseled on the benefits of physical activity by their primary care and specialist (e.g., hepatologist) healthcare providers (LOE 5 Expert opinion)
2. Patient-centered language should be used at all counseling sessions. Stigmatizing language about body weight and body image should be avoided (LOE 5, Expert opinion)
3. Behavioral approaches such as motivational interviewing may be beneficial in helping patients with NAFLD become more physically active (LOE 5, Expert opinion)
4. Print or web-based educational materials specific to patients with NAFLD, including those from the ACSM Exercise is Medicine initiative, should be provided to all patients with NAFLD to reinforce in-person counseling by healthcare providers (LOE 5, Expert opinion)

**Benefits of physical activity in patients with NAFLD**

There are many well-established benefits of regular PA and, in particular, exercise training in patients with NAFLD and NASH, including loss of liver fat on imaging, improvement in NASH activity on histology, reversal of endothelial dysfunction, change in body composition, gain in physical fitness and improvement in health-related quality of life (HRQOL). Importantly, many of these benefits appear independent of clinically significant body weight loss.(45, 64) Whether weight loss is required for liver fibrosis improvement with exercise training remains unclear, however, in general it is accepted that at least 7% body weight loss is required to lead to improvement in liver fibrosis across all treatment interventions.(18, 65, 66)

The most widely studied and established benefit of exercise training in patients with NAFLD is improvement in liver fat measured by magnetic resonance imaging (MRI) scan.(35, 67–81) This benefit appears to be independent of significant body weight loss and is greatest for



aerobic exercise training. Other modalities of exercise which have also been shown to reduce liver fat include resistance training, high-intensity interval training (HIIT) and combined aerobic and resistance training. A recent systematic review and meta-analysis(82) suggests that exercise training achieves rates of clinically significant reduction in MRI-measured liver fat at the threshold which may be a surrogate for histologic improvement in both NASH activity and liver fibrosis at thresholds adapted from NASH drug trials.(83, 84) This is important because to date, there is inconsistent evidence that exercise training independently improves liver fibrosis when liver histology is directly examined, despite multiple studies showing histologic NASH activity can be improved, especially in the presence of significant body weight loss.(85–88)

CVD is a leading cause of death in patients with NAFLD.(19) Fortunately, exercise training has been shown to improve biomarkers of CVD, including reversal of endothelial dysfunction(89) and reduction in serum plasminogen activator inhibitor (PAI) 1 concentration.(45) Exercise training also improves body composition favorably, by reducing adipose tissue volume and possibly increasing lean body mass. Both visceral adipose tissue and subcutaneous adipose tissue have been shown to be reduced by aerobic exercise training.(32, 45, 90–94) The evidence supporting improvement in lean body mass is less consistent, where a statistically significant improvement in lean body mass has not been demonstrated, perhaps owing to short study durations.(45, 95–99)

Exercise training consistently improves cardiorespiratory fitness. There is a robust body of evidence demonstrating that peak oxygen uptake ( $VO_{2peak}$ ), which is universally poor in patients with NAFLD(24, 100) and related to liver fibrosis stage,(24, 101, 102) can be improved with regular exercise training.(68, 80, 95) This is consistent across multiple exercise types including both moderate-intensity aerobic exercise and HIIT protocols.(45, 68, 77, 80, 95) HRQOL, which is also poor in patients with NAFLD,(103, 104) may also be improved with aerobic exercise based on a recent single-center RCT.(45)

#### **Evidence statements:**

1. Established benefits of physical activity in patients with NAFLD include:
  - a. Loss of liver fat on imaging (LOE 1A Systematic Review with Homogeneity of RCTs)
  - b. Improvement in histologic NASH activity (LOE 2B, Low-quality RCT)
  - c. Reversal of endothelial dysfunction (LOE 2B, Low-quality RCT)
  - d. Change in body composition (loss of adipose tissue and gain of lean muscle mass) (LOE 2B, Low-quality RCT)
  - e. Gain in physical fitness (LOE 1A Systematic Review with Homogeneity of RCTs)
  - f. Improvement in health-related quality of life across multiple domains of health (LOE 2B, Low-quality RCT)

2. Most benefits of regular physical activity in patients with NAFLD are independent of clinically significant weight loss (LOE 1B, individual RCT with narrow confidence intervals)
3. Emerging evidence suggests that sustained physical activity may improve liver histology (NASH activity and liver fibrosis) independent of significant weight loss, however, at this time it remains accepted that weight loss may lead to the greatest impact on liver histology (LOE 1A Systematic Review with Homogeneity of RCTs).
4. In individuals with NAFLD who are overweight or obese, 7–10% weight loss may improve liver histology (LOE 1A Systematic Review with Homogeneity of RCTs).
5. Modest weight loss between 3–5% may lead to the same benefits in lean individuals as seen for patients who are overweight or obese. (LOE 1A Systematic Review with Homogeneity of RCTs)

### **Referring a patient with NAFLD to an exercise specialist**

Because physicians are not routinely trained in exercise prescription during medical education and may not feel comfortable delivering this information,(105) the exercise specialist remains a crucial member of the multidisciplinary NAFLD treatment team and may be found across various exercise society databases. Exercise specialists are encouraged to assess a patient's understanding of NAFLD and how lifestyle impacts this common disease and also to establish goals for treatment. Exercise specialists have an additional role in assessing: 1) current and historical PA level; 2) physical fitness in the context of cardiometabolic risk factors and other pertinent medical comorbidities and; 3) barriers and facilitators to PA completion.

Published ACSM guidelines exist to help the exercise specialist determine when medical referral should be sought prior to initiating an exercise training program.(106) Because many patients with NAFLD are inactive and have a history of either CVD or metabolic disease, medical guidance from a treating physician may be necessary prior to beginning an exercise program, especially if a moderate-intensity program is to be prescribed. For individuals who are actively engaged in a moderate-intensity training program already, medical clearance is recommended prior to progressing to a vigorous intensity program.(106) Progressive exercise, starting with low-intensity activities, should also be considered in individuals with NAFLD who have signs or symptoms suggestive of CVD, metabolic and/or renal disease. No physician clearance is required for low intensity activity, which is performed at <50% VO<sub>2</sub>peak (e.g., slowly walking).

#### **Evidence statements:**

1. Referral to an exercise specialist should be considered for all patients with NAFLD to support the clinician in designing and implementing an exercise training program to allow patients with NAFLD to become more physically active (LOE 5, Expert opinion)



2. For patients not meeting recommended activity amounts or intensities, referral to an exercise specialist is strongly recommended for individual exercise prescription feedback and development (LOE 5, Expert opinion)
3. Routine use of exercise specialist databases, including those maintained by the ACSM, may assist the clinician in finding the most appropriate exercise professional for each individual patient (LOE 5, Expert opinion)
4. Prior to starting a moderate to vigorous exercise training program, patients with NAFLD who are inactive, are encouraged to seek medical clearance. For those who are actively engaged in a moderate exercise training program, medical clearance is recommended if seeking to begin a vigorous exercise training program (LOE 1A, Systematic Review with Homogeneity of RCTs).
5. In patients with signs or symptoms suggestive of cardiovascular, metabolic and/or renal disease, starting with low-intensity exercise and progressing in intensity as physiologic acclimation occurs may be considered (LOE 5, Expert opinion).
6. An exercise specialist with additional training or expertise in metabolic disease and obesity is preferred and may offer the ability to expertly address physical conditions unique to patients with NAFLD that may prevent certain types of activity (e.g., osteoarthritis, sarcopenia, frailty and deconditioning) (LOE 5, Expert opinion)

### **Physical activity recommendations in patients with NAFLD**

Exercise training remains a key component in the clinical management of patients with NAFLD. To date, there is a clear and consistent body of evidence supporting this statement despite the heterogeneity across the individual clinical trials which vary widely in each of the ACSM FITT principles of exercise prescription. Because exercise frequency (3–7 days/wk.), intensity (low, moderate, moderate-vigorous and vigorous), time (20–60 minutes) and type (aerobic, resistance, aerobic and resistance combined, HIIT, Pilates, yoga) are variable in the published literature,(35, 67, 77, 107–110) no single optimal exercise prescription has been defined for patients with NAFLD. However, it has been suggested that the combination of aerobic and resistance training is likely more effective than either independently,(107) although this remains controversial and in need of further validation.

What appears to be more important than each individual FITT component, is the total dose of exercise delivered. Exercise dose is routinely reported in metabolic equivalent of task (MET) minutes and is calculated by intensity (METs) by time and aggregated each week. An exercise dose of 750 MET-min/wk., which is the equivalent of 150 min/wk. of moderate-intensity activity or 75 min/wk. of vigorous intensity activity, appears sufficient to improve clinical outcomes in patients with NASH.(82) Additionally, while the majority of the published literature uses direct, in-person supervision of exercise, emerging evidence suggests that remote monitoring with telehealth or unsupervised mobile health delivered lifestyle intervention programs may be equally feasible, safe, acceptable and possibly as efficacious as traditional, supervised in-person exercise training.(111–115)

Patients with advanced liver disease, including those with cirrhosis, represent a unique patient population that has traditionally been viewed independently of patients with earlier stage disease. Several exercise intervention trials challenge this historical perspective and show that exercise training is feasible, safe and highly-efficacious in patients with cirrhosis, including those whose primary etiology of liver disease is NASH.(116–119) Collectively, these studies have shown that regular exercise training decreases portal hypertension,(116) and may improve physical performance, frailty and HRQOL.(117, 118) It is important to recognize that these studies did not include patients with decompensated cirrhosis and therefore no recommendation can be made in this patient population.(20, 120)

**Evidence statements:**

1. At least 150 minutes per week of moderate intensity physical activity or 75 minutes per week of vigorous physical activity are recommended each week for all adult patients with NAFLD (LOE 1B, Individual RCT with small CI)
2. A graded, “staircase” approach is encouraged for patients with NAFLD to progress and achieve moderate-vigorous intensity activity. (LOE 1B- Individual RCT with small CI).
3. If a formal exercise training program is to be prescribed, aerobic exercise with the addition of resistance training is the preferred type of exercise (LOE 5, Expert opinion).
4. In select patients, supervised high-intensity interval training (HIIT) can be considered as an alternative to traditional aerobic exercise and resistance training. Inclusion of low-impact exercises may lessen the risk of injury with HIIT (LOE 2B, Low-quality clinical trial).
5. Supervised and unsupervised physical activity appear to be equally safe and feasible, although efficacy data directly comparing the two are lacking (LOE 5, Expert opinion)
6. An individualized approach where components of ACSM’s FITT exercise principles (e.g., frequency, intensity, time and type) are varied in order to achieve recommended amounts of weekly physical activity may help improve adherence (LOE 5, Expert opinion)
7. Emerging evidence suggests that telehealth, including mobile health applications, may be considered as a safe, feasible and effective way to increase physical activity in patients with NAFLD (LOE 2B, Low quality trial)
8. Patients with NAFLD and advanced liver disease, including those with compensated cirrhosis, may safely perform regular physical activity at amounts similar to what is recommended for all patients with NAFLD (LOE1B, Individual RCT with small CI).

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## REFERENCES

1. Allen AM, Therneau TM, Larson JJ, Coward A, Somers VK, Kamath PS. Nonalcoholic fatty liver disease incidence and impact on metabolic burden and death: a 20 year-community study. *Hepatology*. 2018;67(5):1726–36. [PubMed: 28941364]
2. Younossi ZM, Golabi P, Paik JM, Henry A, Van Dongen C, Henry L. The global epidemiology of nonalcoholic fatty liver disease (NAFLD) and nonalcoholic steatohepatitis (NASH): a systematic review. *Hepatology*. 2023;77(4):1335–47. [PubMed: 36626630]
3. Rinella ME. Nonalcoholic fatty liver disease: a systematic review. *JAMA*. 2015;313(22):2263–73. [PubMed: 26057287]
4. Stine JG, Rinella ME. Editorial: age and non-invasive markers of fibrosis in patients with nonalcoholic fatty liver disease: time to adjust the clock? *Am J Gastroenterol*. 2017;112(5):752–4. [PubMed: 28469217]
5. Nouredin M, Vipani A, Bresee C, et al. NASH leading cause of liver transplant in women: updated analysis of indications for liver transplant and ethnic and gender variances. *Am J Gastroenterol*. 2018;113(11):1649–59. [PubMed: 29880964]

6. Parikh ND, Marrero WJ, Wang J, et al. Projected increase in obesity and non-alcoholic-steatohepatitis-related liver transplantation waitlist additions in the United States. *Hepatology*. 2019;70(2):487–95. [PubMed: 28833326]
7. Kim D, Konyon P, Cholankeril G, Ahmed A. Physical activity is associated with nonalcoholic fatty liver disease and significant fibrosis measured by FibroScan. *Clin Gastroenterol Hepatol*. 2022;20(6):e1438–e55. [PubMed: 34214678]
8. Kim D, Murag S, Cholankeril G, et al. Physical activity, measured objectively, is associated with lower mortality in patients with nonalcoholic fatty liver disease. *Clin Gastroenterol Hepatol*. 2021;19(6):1240–7.e5. [PubMed: 32683103]
9. Zelber-Sagi S, Nitzan-Kaluski D, Goldsmith R, et al. Role of leisure-time physical activity in nonalcoholic fatty liver disease: a population-based study. *Hepatology*. 2008;48(6):1791–8. [PubMed: 18972405]
10. Schneider CV, Zandvakili I, Thaiss CA, Schneider KM. Physical activity is associated with reduced risk of liver disease in the prospective UK Biobank cohort. *JHEP Rep*. 2021;3(3):100263. [PubMed: 33898961]
11. Schnurr TM, Katz SF, Justesen JM, et al. Interactions of physical activity, muscular fitness, adiposity, and genetic risk for NAFLD. *Hepatology Commun*. 2022;6(7):1516–26. [PubMed: 35293152]
12. Kwak JH, Jun DW, Lee SM, et al. Lifestyle predictors of obese and non-obese patients with nonalcoholic fatty liver disease: a cross-sectional study. *Clin Nutr*. 2018;37(5):1550–7. [PubMed: 28918170]
13. Sung K-C, Lee M-Y, Lee J-Y, et al. Natural course of fatty liver in 36,195 South Korean adults. *Sci Rep*. 2019;9(1):9062. [PubMed: 31308382]
14. DiJoseph K, Thorp A, Harrington A, Schmitz KH, Chinchilli VM, Stine JG. Physical activity and risk of hepatocellular carcinoma: a systematic review and meta-analysis. *Dig Dis Sci*. 2023;68(3):1051–9. [PubMed: 35767138]
15. Stine JG, Soriano C, Schreiber I, et al. Breaking down barriers to physical activity in patients with nonalcoholic fatty liver disease. *Dig Dis Sci*. 2021;66(10):3604–11. [PubMed: 33098023]
16. Rinella ME, Neuschwander-Tetri BA, Siddiqui MS, et al. AASLD practice guidance on the clinical assessment and management of nonalcoholic fatty liver disease. *Hepatology*. 2023;77(5):1797–1835. [PubMed: 36727674]
17. Buzzetti E, Pinzani M, Tsochatzis EA. The multiple-hit pathogenesis of non-alcoholic fatty liver disease (NAFLD). *Metabolism*. 2016;65(8):1038–48. [PubMed: 26823198]
18. Vilar-Gomez E, Martinez-Perez Y, Calzadilla-Bertot L, et al. Weight loss through lifestyle modification significantly reduces features of nonalcoholic steatohepatitis. *Gastroenterology*. 2015;149(2):367–78.e5; quiz e14–5. [PubMed: 25865049]
19. Simon TG, Roelstraete B, Khalili H, Hagström H, Ludvigsson JF. Mortality in biopsy-confirmed nonalcoholic fatty liver disease: results from a nationwide cohort. *Gut*. 2021;70(7):1375–82. [PubMed: 33037056]
20. Stine JG LM, Corey KE, Sallis RE, et al. American College of Sports Medicine (ACSM) International Multidisciplinary Roundtable report on physical activity and nonalcoholic fatty liver disease. *Hepatology Commun*. 2023;7(4):e0108. [PubMed: 36995998]
21. Stine JG. Editorial: twenty minutes of moderate-to-vigorous physical activity a day keeps the NAFLD away. *Aliment Pharmacol Ther*. 2022;55(1):116–7. [PubMed: 34907567]
22. Tsunoda K, Kitano N, Kai Y, Jindo T, Uchida K, Arao T. Dose-response relationships of accelerometer-measured sedentary behaviour and physical activity with non-alcoholic fatty liver disease. *Aliment Pharmacol Ther*. 2021;54(10):1330–9. [PubMed: 34633105]
23. Long MT, Pedley A, Massaro JM, et al. Hepatic steatosis is associated with lower levels of physical activity measured via accelerometry. *Obesity (Silver Spring)*. 2015;23(6):1259–66. [PubMed: 25959049]
24. Krasnoff JB, Painter PL, Wallace JP, Bass NM, Merriman RB. Health-related fitness and physical activity in patients with nonalcoholic fatty liver disease. *Hepatology*. 2008;47(4):1158–66. [PubMed: 18266250]

25. Hallsworth K, Thoma C, Moore S, et al. Non-alcoholic fatty liver disease is associated with higher levels of objectively measured sedentary behaviour and lower levels of physical activity than matched healthy controls. *Frontline Gastroenterol.* 2015;6(1):44–51. [PubMed: 25580206]
26. Moore SC, Lee IM, Weiderpass E, et al. Association of leisure-time physical activity with risk of 26 types of cancer in 1.44 million adults. *JAMA Intern Med.* 2016;176(6):816–25. [PubMed: 27183032]
27. Mantovani A, Petracca G, Beatrice G et al. Non-alcoholic fatty liver disease and increased risk of incident extrahepatic cancers: a meta-analysis of observational cohort studies. *Gut.* 2022;71(4):778–88. [PubMed: 33685968]
28. Hughes A, Dahmus J, Rivas G, et al. Exercise training reverses gut dysbiosis in patients with biopsy-proven nonalcoholic steatohepatitis: a proof of concept study. *Clin Gastroenterol Hepatol.* 2021;19(8):1723–5. [PubMed: 32882426]
29. Babu AF, Csader S, Männistö V, et al. Effects of exercise on NAFLD using non-targeted metabolomics in adipose tissue, plasma, urine, and stool. *Sci Rep.* 2022;12(1):6485. [PubMed: 35444259]
30. Thorp A, Stine JG. Exercise as Medicine: The Impact of Exercise Training on Nonalcoholic Fatty Liver Disease. *Curr Hepatol Rep.* 2020;19(4):402–11. [PubMed: 33767944]
31. Shi X, Yin H, Li J, et al. Circulating branch chain amino acids and improvement in liver fat content in response to exercise interventions in NAFLD. *Sci Rep.* 2021;11(1):13415. [PubMed: 34183740]
32. Zhang HJ, He J, Pan LL, et al. Effects of moderate and vigorous exercise on nonalcoholic fatty liver disease: a randomized clinical trial. *JAMA Intern Med.* 2016;176(8):1074–82. [PubMed: 27379904]
33. Cao H. Adipocytokines in obesity and metabolic disease. *J Endocrinol.* 2014;220(2):T47–59. [PubMed: 24403378]
34. Giby VG, Ajith TA. Role of adipokines and peroxisome proliferator-activated receptors in nonalcoholic fatty liver disease. *World J Hepatol.* 2014;6(8):570–9. [PubMed: 25232450]
35. Hashida R, Kawaguchi T, Bekki M, et al. Aerobic vs. resistance exercise in non-alcoholic fatty liver disease: a systematic review. *J Hepatol.* 2017;66(1):142–52. [PubMed: 27639843]
36. Yang X, Smith U. Adipose tissue distribution and risk of metabolic disease: does thiazolidinedione-induced adipose tissue redistribution provide a clue to the answer? *Diabetologia.* 2007;50(6):1127–39. [PubMed: 17393135]
37. Wajchenberg BL. Subcutaneous and visceral adipose tissue: their relation to the metabolic syndrome. *Endocr Rev.* 2000;21(6):697–738. [PubMed: 11133069]
38. Gehrke N, Biedenbach J, Huber Y, et al. Voluntary exercise in mice fed an obesogenic diet alters the hepatic immune phenotype and improves metabolic parameters – an animal model of life style intervention in NAFLD. *Sci Rep.* 2019;9(1):4007. [PubMed: 30850619]
39. Rector RS, Uptergrove GM, Morris EM, et al. Daily exercise vs. caloric restriction for prevention of nonalcoholic fatty liver disease in the OLETF rat model. *Am J Physiol Gastrointest Liver Physiol.* 2011;300(5):G874–83.
40. Cho J, Lee I, Kim D, et al. Effect of aerobic exercise training on non-alcoholic fatty liver disease induced by a high fat diet in C57BL/6 mice. *J Exerc Nutrition Biochem.* 2014;18(4):339–46.
41. Taniguchi H, Tanisawa K, Sun X, Kubo T, Higuchi M. Endurance exercise reduces hepatic fat content and serum fibroblast growth factor 21 levels in elderly men. *J Clin Endocrinol Metab.* 2016;101(1):191–8. [PubMed: 26562755]
42. Stine JG, Xu D, Schmitz K, Sciamanna C, Kimball SR. Exercise attenuates ribosomal protein six phosphorylation in fatty liver disease. *Dig Dis Sci.* 2020;65(11):3238–43. [PubMed: 32239376]
43. Piguat AC, Guarino M, Potaczek DP, Garn H, Dufour JF. Hepatic gene expression in mouse models of non-alcoholic fatty liver disease after acute exercise. *Hepatol Res.* 2019;49(6):637–52. [PubMed: 30811073]
44. Melo L, Bilici M, Hagar A, Klaunig JE. The effect of endurance training on non-alcoholic fatty liver disease in mice. *Physiol Rep.* 2021;9(15):e14926.
45. Stine JG, Schreiberman IR, Faust AJ, et al. NASHFit: A randomized controlled trial of an exercise training program to reduce clotting risk in patients with NASH. *Hepatology.* 2021.

46. Cheng R, Wang L, Le S, et al. A randomized controlled trial for response of microbiome network to exercise and diet intervention in patients with nonalcoholic fatty liver disease. *Nature Commun.* 2022;13(1):2555. [PubMed: 35538056]
47. Canadian Society for Exercise Physiology. Get Active Questionnaire. 2017. Available at [https://csep.ca/wp-content/uploads/2021/05/GETACTIVEQUESTIONNAIRE\\_ENG.pdf](https://csep.ca/wp-content/uploads/2021/05/GETACTIVEQUESTIONNAIRE_ENG.pdf).
48. Craig CL, Marshall AL, Sjöström M, et al. International physical activity questionnaire: 12-country reliability and validity. *Med Sci Sports Exerc.* 2003;35(8):1381–95. [PubMed: 12900694]
49. Warburton DE, Jamnik VK, Bredin SS, et al. Evidence-based risk assessment and recommendations for physical activity clearance: an introduction. *Appl Physiol Nutr Metab.* 2011;36 Suppl 1:S1–2. [PubMed: 21800937]
50. Bowen PG, Mankowski RT, Harper SA, Buford TW. Exercise is Medicine as a vital sign: challenges and opportunities. *Transl J Am Coll Sports Med.* 2019;4(1):1–7. [PubMed: 30828640]
51. <https://exerciseismedicine.org/wp-content/uploads/2021/04/EIM-Physical-Activity-Vital-Sign.pdf>.
52. Sallis R. Developing healthcare systems to support exercise: exercise as the fifth vital sign. *Br J Sports Med.* 2011;45(6):473–4. [PubMed: 21292925]
53. Grant RW, Schmittiel JA, Neugebauer RS, Uratsu CS, Sternfeld B. Exercise as a vital sign: a quasi-experimental analysis of a health system intervention to collect patient-reported exercise levels. *J Gen Intern Med.* 2014;29(2):341–8. [PubMed: 24309950]
54. Centers for Disease Control and Prevention. Overcoming Barriers to Physical Activity. Available at <https://www.cdc.gov/physicalactivity/basics/adding-pa/barriers.html>.
55. Office of National Drug Control Policy. Screening, brief intervention, and referral to treatment (SBIRT). [Washington, D.C.] : Office of National Drug Control Policy (ONDCP), Substance Abuse and Mental Health Services Administration (SAMHSA), [2012]; 2012.
56. Arora E, Babu AS, Vidhyasagar S, Maiya GA. Physical activity promotion program on physical activity and glycemic control in prediabetes. *Crit Rev Phys Rehabil Med.* 2018;30(2):181–5.
57. Lazarus JV, Kakalou C, Palayew A, et al. A Twitter discourse analysis of negative feelings and stigma related to NAFLD, NASH and obesity. *Liver Int.* 2021;41(10):2295–307. [PubMed: 34022107]
58. Hallsworth K, Adams LA. Lifestyle modification in NAFLD/NASH: facts and figures. *JHEP Rep.* 2019;1(6):468–79. [PubMed: 32039399]
59. Mazzotti A, Caletti MT, Brodosi L, et al. An internet-based approach for lifestyle changes in patients with NAFLD: two-year effects on weight loss and surrogate markers. *J Hepatol.* 2018;69(5):1155–63. [PubMed: 30290973]
60. Moscaticello S, Di Luzio R, Bugianesi E, et al. Cognitive-behavioral treatment of nonalcoholic Fatty liver disease: a propensity score-adjusted observational study. *Obesity (Silver Spring).* 2011;19(4):763–70. [PubMed: 20966900]
61. Montesi L, Caselli C, Centis E, et al. Physical activity support or weight loss counseling for nonalcoholic fatty liver disease? *World J Gastroenterol.* 2014;20(29):10128–36. [PubMed: 25110440]
62. Hallsworth K, McPherson S, Anstee QM, Flynn D, Haigh L, Avery L. Digital intervention with lifestyle coach support to target dietary and physical activity behaviors of adults with nonalcoholic fatty liver disease: systematic development process of VITALISE using intervention mapping. *J Med Internet Res.* 2021;23(1):e20491. [PubMed: 33448929]
63. de Sousa D, Fogel A, Azevedo J, Padrão P. The effectiveness of web-based interventions to promote health behaviour change in adolescents: a systematic review. *Nutrients.* 2022;14(6):1258. [PubMed: 35334915]
64. Stine JG DK, Pattison Z, Harrington A, Chinchilli VM, Schmitz KH, Loomba R. Exercise training is associated with treatment response in liver fat content by magnetic resonance imaging independent of clinically significant body weight loss in patients with nonalcoholic fatty liver disease: a systematic review and meta-analysis. *Am J Gastroenterol.* 2022;In press.
65. Wong VW, Chan RS, Wong GL, et al. Community-based lifestyle modification programme for non-alcoholic fatty liver disease: a randomized controlled trial. *J Hepatol.* 2013;59(3):536–42. [PubMed: 23623998]



66. Harrison SA, Fecht W, Brunt EM, Neuschwander-Tetri BA. Orlistat for overweight subjects with nonalcoholic steatohepatitis: a randomized, prospective trial. *Hepatology*. 2009;49(1):80–6. [PubMed: 19053049]
67. Babu AF, Csader S, Lok J, et al. Positive effects of exercise intervention without weight loss and dietary changes in NAFLD-related clinical parameters: a systematic review and meta-analysis. *Nutrients*. 2021;13(9):3135. [PubMed: 34579012]
68. Baker CJ, Martinez-Huenschullan SF, D'Souza M, et al. Effect of exercise on hepatic steatosis: are benefits seen without dietary intervention? A systematic review and meta-analysis. *J Diabetes*. 2021;13(1):63–77. [PubMed: 32667128]
69. Battista F, Ermolao A, van Baak MA, et al. Effect of exercise on cardiometabolic health of adults with overweight or obesity: focus on blood pressure, insulin resistance, and intrahepatic fat - a systematic review and meta-analysis. *Obes Rev*. 2021;22 Suppl 4(Suppl 4):e13269.
70. Hens W, Taeyman J, Cornelis J, Gielen J, Van Gaal L, Vissers D. The effect of lifestyle interventions on excess ectopic fat deposition measured by noninvasive techniques in overweight and obese adults: a systematic review and meta-analysis. *J Phys Act Health*. 2016;13(6):671–94. [PubMed: 26694194]
71. Katsagoni CN, Georgoulis M, Papatheodoridis GV, Panagiotakos DB, Kontogianni MD. Effects of lifestyle interventions on clinical characteristics of patients with non-alcoholic fatty liver disease: a meta-analysis. *Metabolism*. 2017;68:119–32. [PubMed: 28183444]
72. Keating SE, Hackett DA, George J, Johnson NA. Exercise and non-alcoholic fatty liver disease: a systematic review and meta-analysis. *J Hepatol*. 2012;57(1):157–66. [PubMed: 22414768]
73. Khalafi M, Symonds ME. The impact of high intensity interval training on liver fat content in overweight or obese adults: A meta-analysis. *Physiol Behav*. 2021;236:113416.
74. Mohammad Rahimi GR, Attarzadeh Hosseini SR. Effect of aerobic exercise alone or in conjunction with diet on liver function, insulin resistance and lipids in non-alcoholic fatty liver disease. *Biol Res Nurs*. 2022;24(2):259–76. [PubMed: 35130757]
75. Musso G, Cassader M, Rosina F, Gambino R. Impact of current treatments on liver disease, glucose metabolism and cardiovascular risk in non-alcoholic fatty liver disease (NAFLD): a systematic review and meta-analysis of randomised trials. *Diabetologia*. 2012;55(4):885–904. [PubMed: 22278337]
76. Orci LA, Gariani K, Oldani G, Delaune V, Morel P, Toso C. Exercise-based interventions for nonalcoholic fatty liver disease: a meta-analysis and meta-regression. *Clin Gastroenterol Hepatol*. 2016;14(10):1398–411. [PubMed: 27155553]
77. Sabag A, Barr L, Armour M, et al. The effect of high-intensity interval training vs moderate-intensity continuous training on liver fat: a systematic review and meta-analysis. *J Clin Endocrinol Metab*. 2022;107(3):862–81. [PubMed: 34724062]
78. Sargeant JA, Gray LJ, Bodicoat DH, et al. The effect of exercise training on intrahepatic triglyceride and hepatic insulin sensitivity: a systematic review and meta-analysis. *Obes Rev*. 2018;19(10):1446–59. [PubMed: 30092609]
79. Smart NA, King N, McFarlane JR, Graham PL, Dieberg G. Effect of exercise training on liver function in adults who are overweight or exhibit fatty liver disease: a systematic review and meta-analysis. *Br J Sports Med*. 2018;52(13):834–43. [PubMed: 27317790]
80. Fernández T, Viñuela M, Vidal C, Barrera F. Lifestyle changes in patients with non-alcoholic fatty liver disease: a systematic review and meta-analysis. *PLoS One*. 2022;17(2):e0263931.
81. Golabi P, Locklear CT, Austin P, et al. Effectiveness of exercise in hepatic fat mobilization in non-alcoholic fatty liver disease: systematic review. *World J Gastroenterol*. 2016;22(27):6318–27. [PubMed: 27468220]
82. Stine JG, DiJoseph K, Pattison Z, et al. Exercise training is associated with treatment response in liver fat content by magnetic resonance imaging independent of clinically significant body weight loss in patients with nonalcoholic fatty liver disease: a systematic review and meta-analysis. *Am J Gastroenterol*. 2023. Online ahead of print.
83. Stine JG, Munaganuru N, Barnard A, et al. Change in MRI-PDFF and histologic response in patients with nonalcoholic steatohepatitis: a systematic review and meta-analysis. *Clin Gastroenterol Hepatol*. 2021;19(11):2274–83.e5. [PubMed: 32882428]



84. Tamaki N, Munaganuru N, Jung J, et al. Clinical utility of 30% relative decline in MRI-PDFF in predicting fibrosis regression in non-alcoholic fatty liver disease. *Gut*. 2022;71(5):983–90. [PubMed: 33883248]
85. O’Gorman P, Naimimohasses S, Monaghan A et al. Improvement in histological endpoints of MAFLD following a 12-week aerobic exercise intervention. *Aliment Pharmacol Ther*. 2020;52(8):1387–98. [PubMed: 32717123]
86. Hickman IJ, Jonsson JR, Prins JB, et al. Modest weight loss and physical activity in overweight patients with chronic liver disease results in sustained improvements in alanine aminotransferase, fasting insulin, and quality of life. *Gut*. 2004;53(3):413–9. [PubMed: 14960526]
87. Eckard C, Cole R, Lockwood J, et al. Prospective histopathologic evaluation of lifestyle modification in nonalcoholic fatty liver disease: a randomized trial. *Therap Adv Gastroenterol*. 2013;6(4):249–59.
88. Promrat K, Kleiner DE, Niemeier HM, et al. Randomized controlled trial testing the effects of weight loss on nonalcoholic steatohepatitis. *Hepatology*. 2010;51(1):121–9. [PubMed: 19827166]
89. Green DJ, Maiorana A, O’Driscoll G, Taylor R. Effect of exercise training on endothelium-derived nitric oxide function in humans. *J Physiol*. 2004;561(Pt 1):1–25. [PubMed: 15375191]
90. Keating SE, Hackett DA, Parker HM, et al. Effect of aerobic exercise training dose on liver fat and visceral adiposity. *J Hepatol*. 2015;63(1):174–82. [PubMed: 25863524]
91. Bacchi E, Negri C, Targher G, et al. Both resistance training and aerobic training reduce hepatic fat content in type 2 diabetic subjects with nonalcoholic fatty liver disease (the RAED2 Randomized Trial). *Hepatology*. 2013;58(4):1287–95. [PubMed: 23504926]
92. Abdelbasset WK, Tantawy SA, Kamel DM, et al. Effects of high-intensity interval and moderate-intensity continuous aerobic exercise on diabetic obese patients with nonalcoholic fatty liver disease: a comparative randomized controlled trial. *Medicine (Baltimore)*. 2020;99(10):e19471.
93. Pugh CJ, Spring VS, Kemp GJ, et al. Exercise training reverses endothelial dysfunction in nonalcoholic fatty liver disease. *Am J Physiol Heart Circ Physiol*. 2014;307(9):H1298–306. [PubMed: 25193471]
94. Cuthbertson DJ, Shojaee-Moradie F, Sprung VS, et al. Dissociation between exercise-induced reduction in liver fat and changes in hepatic and peripheral glucose homeostasis in obese patients with non-alcoholic fatty liver disease. *Clin Sci (Lond)*. 2016;130(2):93–104. [PubMed: 26424731]
95. Gonzalez A, Valero-Breton M, Huerta-Salgado C, Achiardi O, Simon F, Cabello-Verrugio C. Impact of exercise training on the sarcopenia criteria in non-alcoholic fatty liver disease: a systematic review and meta-analysis. *Eur J Transl Myol*. 2021;31(1):9630. [PubMed: 33709647]
96. Hallsworth K, Fattakhova G, Hollingsworth KG, et al. Resistance exercise reduces liver fat and its mediators in non-alcoholic fatty liver disease independent of weight loss. *Gut*. 2011;60(9):1278–83. [PubMed: 21708823]
97. Houghton D, Thoma C, Hallsworth K, et al. Exercise reduces liver lipids and visceral adiposity in patients with nonalcoholic steatohepatitis in a randomized controlled trial. *Clin Gastroenterol Hepatol*. 2017;15(1):96–102.e3. [PubMed: 27521509]
98. Sullivan S, Kirk EP, Mittendorfer B, Patterson BW, Klein S. Randomized trial of exercise effect on intrahepatic triglyceride content and lipid kinetics in nonalcoholic fatty liver disease. *Hepatology*. 2012;55(6):1738–45. [PubMed: 22213436]
99. Cheng S, Ge J, Zhao C, et al. Effect of aerobic exercise and diet on liver fat in pre-diabetic patients with non-alcoholic-fatty-liver-disease: a randomized controlled trial. *Sci Rep*. 2017;7(1):15952. [PubMed: 29162875]
100. Sayiner M, Stepanova M, Pham H, Noor B, Walters M, Younossi ZM. Assessment of health utilities and quality of life in patients with non-alcoholic fatty liver disease. *BMJ Open Gastroenterol*. 2016;3(1):e000106.
101. Canada JM, Abbate A, Collen R, et al. Relation of hepatic fibrosis in nonalcoholic fatty liver disease to left ventricular diastolic function and exercise tolerance. *Am J Cardiol*. 2019;123(3):466–73. [PubMed: 30502049]
102. Dahmus J, Hummer B, Rivas G, et al. Patients with nonalcoholic steatohepatitis and advanced liver disease have the lowest cardiorespiratory fitness. *Dig Dis Sci*. 2023. Online ahead of print.

103. Golabi P, Otgonsuren M, Cable R, et al. Non-alcoholic fatty liver disease (NAFLD) is associated with impairment of Health Related Quality of Life (HRQOL). *Health Qual Life Outcomes*. 2016;14:18. [PubMed: 26860700]
104. Dan AA, Kallman JB, Wheeler A, et al. Health-related quality of life in patients with non-alcoholic fatty liver disease. *Aliment Pharmacol Ther*. 2007;26(6):815–20. [PubMed: 17767465]
105. Avery L, Exley C, McPherson S, Trenell MI, Anstee QM, Hallsworth K. Lifestyle behavior change in patients with nonalcoholic fatty liver disease: a qualitative study of clinical practice. *Clin Gastroenterol Hepatol*. 2017;15(12):1968–71. [PubMed: 28624648]
106. Thompson PD, Arena R, Riebe D, Pescatello LS. ACSM’s new preparticipation health screening recommendations from ACSM’s guidelines for exercise testing and prescription, ninth edition. *Curr Sports Med Rep*. 2013;12(4):215–7. [PubMed: 23851406]
107. Zhou BJ, Huang G, Wang W, et al. Intervention effects of four exercise modalities on nonalcoholic fatty liver disease: a systematic review and Bayesian network meta-analysis. *Eur Rev Med Pharmacol Sci*. 2021;25(24):7687–97. [PubMed: 34982430]
108. Buzzetti E, Linden A, Best LM, et al. Lifestyle modifications for nonalcohol-related fatty liver disease: a network meta-analysis. *Cochrane Database Syst Rev*. 2021;6(6):CD013156.
109. Singh AK, Kaur N, Kaushal S, et al. Partitioning of radiological, stress and biochemical changes in pre-diabetic women subjected to Diabetic Yoga Protocol. *Diabetes Metab Syndr*. 2019;13(4):2705–13. [PubMed: 31405697]
110. Keymasi Z, Sadeghi A, Pourrazi H. Effect of pilates training on hepatic fat content and liver enzymes in men with nonalcoholic fatty liver disease. *J Appl Health Studies Sport Physiol*. 2017;4(2):49–56.
111. Huber Y, Pfirrmann D, Gebhardt I, et al. Improvement of non-invasive markers of NAFLD from an individualised, web-based exercise program. *Aliment Pharmacol Ther*. 2019;50(8):930–9. [PubMed: 31342533]
112. Motz V, Faust A, Dahmus J, Stern B, Soriano C, Stine JG. Utilization of a directly supervised telehealth-based exercise training program in patients with nonalcoholic steatohepatitis: feasibility study. *JMIR Form Res*. 2021;5(8):e30239.
113. Faust A, Stine JG. Time to step it up: mobile health intervention for lifestyle modification in patients with nonalcoholic fatty liver disease. *Dig Dis Sci*. 2022;67(2):403–5. [PubMed: 33939148]
114. Tincopa M, Lyden, A, Wong, J, et al. . Impact of a pilot structured mobile technology based lifestyle intervention for patients with non-alcoholic fatty liver disease. *Dig Dis Sci*. 2022;67(2):4812–91.
115. Stine JG RG, Hummer BS, Duarte-Rojo A, et al. Mobile health lifestyle intervention program leads to clinically significant loss of body weight in patients with NASH. *Hepatol Commun*. 2023;7(4):e0052. [PubMed: 36930864]
116. Berzigotti A, Albillos A, Villanueva C, et al. Effects of an intensive lifestyle intervention program on portal hypertension in patients with cirrhosis and obesity: The SportDiet study. *Hepatology*. 2017;65(4):1293–305. [PubMed: 27997989]
117. Lai JC, Dodge JL, Kappus MR, et al. A multicenter pilot randomized clinical trial of a home-based exercise program for patients with cirrhosis: the Strength Training Intervention (STRIVE). *Am J Gastroenterol*. 2021;116(4):717–22. [PubMed: 33982941]
118. Duarte-Rojo A, Bloomer PM, Rogers RJ, et al. Introducing EL-FIT (Exercise and Liver FITness): a smartphone app to prehabilitate and monitor liver transplant candidates. *Liver Transpl*. 2021;27(4):502–12. [PubMed: 33232547]
119. Faust A, Stine JG. Leveraging the Coronavirus disease 2019 pandemic: is it time to consider incorporating mobile applications into standard clinical management of the liver transplantation patient? *Liver Transpl*. 2021;27(4):479–81. [PubMed: 33484229]
120. Johnston HE, Takefala TG, Kelly JT, et al. The effect of diet and exercise interventions on body composition in liver cirrhosis: a systematic review. *Nutrients*. 2022;14(16):3365. [PubMed: 36014871]

**Table 1.**

## Levels of Evidence

Level	Type of Evidence
1A	Systematic review (with homogeneity) of RCTs
1B	Individual RCT (with narrow CIs)
1C	All or none study
2A	Systematic review (with homogeneity) of cohort studies
2B	Individual cohort study (including low quality RCT)
2C	“Outcomes” research Ecological studies
3A	Systematic review (with homogeneity) of case-control studies
2B	Individual case-control study
4	Case series Poor quality cohort or case-control study
5	Expert opinion without explicit critical appraisal or based on physiology bench research or “first principles”

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**Table 2.**

## Consensus evidence statements and recommendations

**Role of physical activity in NAFLD pathogenesis:**

1. Regular physical activity prevents the development of NAFLD (LOE 2B, Individual Cohort)
2. Emerging evidence suggests individuals who are less physically active may be at increased risk for disease progression to NASH with liver fibrosis and/or cirrhosis (LOE 2B, Individual Cohort)
3. Individuals who are less physically active have greater risk of primary liver cancer (e.g., hepatocellular carcinoma) and other extrahepatic cancers commonly associated with NAFLD (e.g., breast, colorectal, esophageal, gastric, pancreatic, prostate and kidney) (LOE 2B, Individual Cohort)
4. Physical activity offers promise to reverse NAFLD pathogenesis by impacting multiple mechanistic pathways simultaneously and possibly synergistically (LOE 2B Individual cohort studies including low-quality RCT)

**Assessing and screening patients with NAFLD for physical activity:**

5. All patients with NAFLD should be screened for physical activity during routine healthcare visits with primary care and specialist (e.g., hepatologist) healthcare providers (LOE 2B cohort/low-quality RCT)
6. Tools that provide standardized measures of physical activity (e.g., ACSM Physical Activity Vital Sign) should be administered regularly at routine healthcare visits (LOE 5 Expert opinion)
7. Barriers to physical activity should be assessed systematically. While no one optimal tool has been identified, the Screening, Brief Intervention and Referral to Treatment (SBIRT) framework or NAFLD-specific questionnaires may be considered (LOE 5 Expert opinion)

**Advising and counseling patients with NAFLD about the benefits of physical activity:**

8. All patients with NAFLD, regardless of disease stage, should be counseled on the benefits of physical activity by their primary care and specialist (e.g., hepatologist) healthcare providers (LOE 5 Expert opinion)
9. Patient-centered language should be used at all counseling sessions. Stigmatizing language about body weight and body image should be avoided (LOE 5, Expert opinion)
10. Behavioral approaches such as motivational interviewing may be beneficial in helping patients with NAFLD become more physically active (LOE 5, Expert opinion)
11. Print or web-based educational materials specific to patients with NAFLD, including those from the ACSM Exercise is Medicine initiative, should be provided to all patients with NAFLD to reinforce in-person counseling by healthcare providers (LOE 5, Expert opinion)
12. Established benefits of physical activity in patients with NAFLD include:
  - a. Loss of liver fat on imaging (LOE 1A Systematic Review with Homogeneity of RCTs)
  - b. Improvement in histologic NASH activity (LOE 2B, Low-quality RCT)
  - c. Reversal of endothelial dysfunction (LOE- 2B, Low-quality RCT)
  - d. Change in body composition (loss of adipose tissue and gain of lean muscle mass) (LOE 2B, Low-quality RCT)
  - e. Gain in physical fitness (LOE 1A Systematic Review with Homogeneity of RCTs)
  - f. Improvement in health-related quality of life across multiple domains of health (LOE 2B, Low-quality RCT)
13. Most benefits of regular physical activity in patients with NAFLD are independent of clinically significant weight loss (LOE 1B, individual RCT with narrow confidence intervals)
14. Emerging evidence suggests that sustained physical activity may improve liver histology (NASH activity and liver fibrosis) independent of significant weight loss, however, at this time it remains accepted that weight loss may lead to the greatest impact on liver histology (LOE 1A Systematic Review with Homogeneity of RCTs).
15. In individuals with NAFLD who are overweight or obese, 7–10% weight loss may improve liver histology (LOE 1A Systematic Review with Homogeneity of RCTs).
16. Modest weight loss between 3–5% may lead to the same benefits in lean individuals as seen for patients who are overweight or obese. (LOE1A Systematic Review with Homogeneity of RCTs)

**Referring a patient with NAFLD to an exercise specialist:**

17. Referral to an exercise specialist should be considered for all patients with NAFLD to support the clinician in designing and implementing an exercise training program to allow patients with NAFLD to become more physically active (LOE 5, Expert opinion)
18. For patients not meeting recommended activity amounts or intensities, referral to an exercise specialist is strongly recommended for individual exercise prescription feedback and development (LOE 5, Expert opinion)
19. Routine use of exercise specialist databases, including those maintained by the ACSM, may assist the clinician in finding the most appropriate exercise professional for each individual patient (LOE 5, Expert opinion)

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**Role of physical activity in NAFLD pathogenesis:**

20. Prior to starting a moderate-to-vigorous exercise training program, patients with NAFLD who are inactive are encouraged to seek medical clearance. For those who are actively engaged in a moderate exercise training program, medical clearance is recommended if seeking to begin a vigorous exercise training program (LOE 1A, Systematic Review with Homogeneity of RCTs).

21. In patients with signs or symptoms suggestive of cardiovascular, metabolic and/or renal disease, starting with low-intensity exercise and progressing in intensity as physiologic acclimation occurs may be considered (LOE 5, Expert opinion).

22. An exercise specialist with additional training or expertise in metabolic disease and obesity is preferred and may offer the ability to expertly address physical conditions unique to patients with NAFLD that may prevent certain types of activity (e.g., osteoarthritis, sarcopenia, frailty and deconditioning) (LOE 5, Expert opinion)

**Physical activity recommendations in patients with NAFLD:**

23. At least 150 minutes per week of moderate intensity physical activity or 75 minutes per week of vigorous physical activity are recommended each week for all adult patients with NAFLD (LOE1B, Individual RCT with small CI)

24. A graded, “staircase” approach is encouraged for patients with NAFLD to progress and achieve moderate-vigorous intensity activity. (LOE 1B- Individual RCT with small CI).

25. If a formal exercise training program is to be prescribed, aerobic exercise with the addition of resistance training is the preferred type of exercise (LOE 5, Expert opinion).

26. In select patients, high-intensity interval training (HIIT) can be considered as an alternative to traditional aerobic exercise and resistance training. Inclusion of low-impact exercises may lessen the risk of injury with HIIT (LOE 2B, Low-quality clinical trial).

27. Supervised and unsupervised physical activity appear to be equally safe and feasible, although efficacy data directly comparing the two are lacking (LOE 5, Expert opinion)

28. An individualized approach where components of ACSM’s FITT exercise principles (e.g., frequency, intensity, time and type) are varied in order to achieve recommended amounts of weekly physical activity may help improve adherence (LOE 5, Expert opinion)

29. Emerging evidence suggests that telehealth, including mobile health applications, may be considered as a safe, feasible and effective way to increase physical activity in patients with NAFLD (LOE 2B, Low quality trial)

30. Patients with NAFLD and advanced liver disease, including those with cirrhosis, may safely perform regular physical activity at amounts similar to what is recommended for all patients with NAFLD (LOE1B, Individual RCT with small CI)

\* no consensus was achieved about physical activity and improvement in liver fibrosis stage