

Understanding the socioeconomic and genetic risk factors for MASH



Metabolic dysfunction-associated steatohepatitis (MASH), a severe form of metabolic dysfunction-associated steatotic liver disease (MASLD) is a chronic, metabolic liver disease that is associated with certain cardiometabolic risk factors, including¹:



Obesity



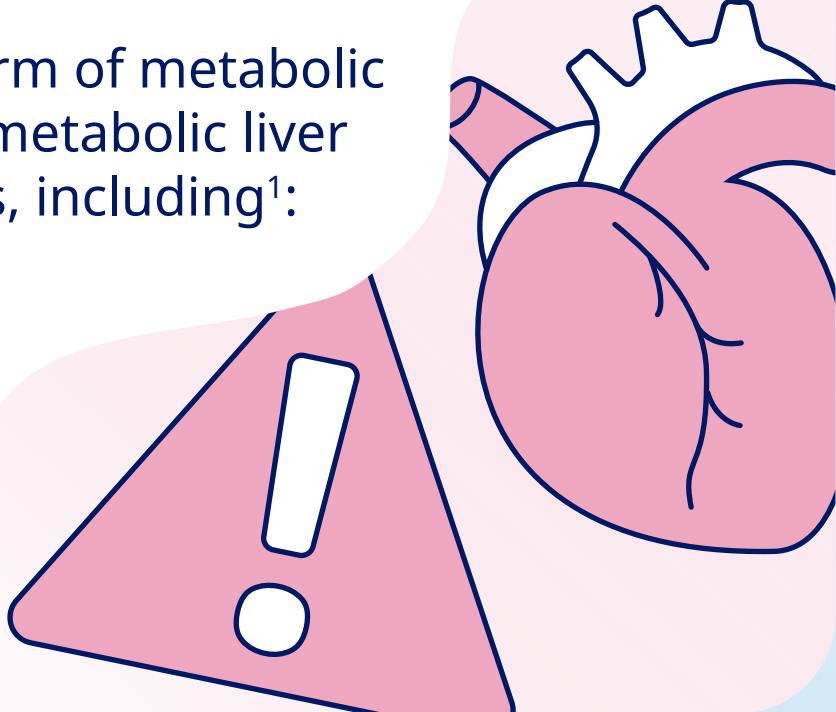
Type 2 diabetes



Hypertension



Dyslipidemia
Hypertriglyceridemia and/or low HDL

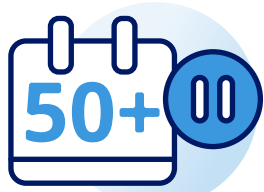


MASLD and MASH disproportionately impacts populations based on sex and socioeconomic group

Sex differences in MASLD and MASH prevalence:

Men are **19%** more likely to have MASLD than women²

Both sexes have similar risk of MASH, the progressive form of MASLD, however women have a **37%** greater risk of advanced fibrosis²



Women aged >50 are at an increased risk of MASH and advanced fibrosis²

- Lower levels of estrogen in post-menopausal women may contribute to the development of advanced fibrosis²

Socioeconomic disparities contribute to MASLD disease burden:



Poverty



Poor access to **nutritionally adequate food**

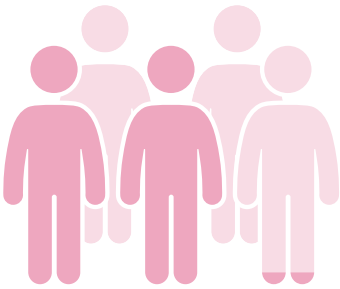
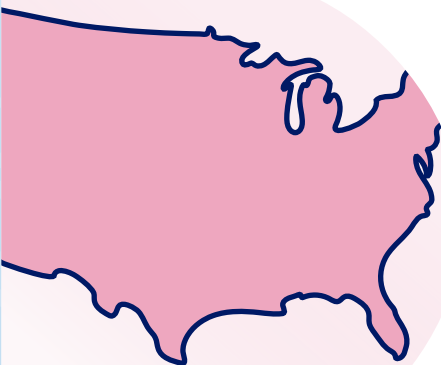


Food insecurity

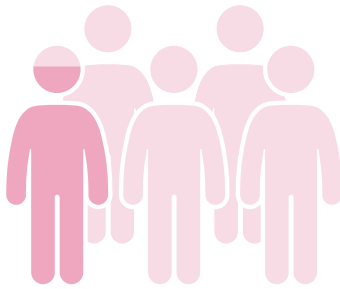


MASLD and MASH burden is higher in certain racial and ethnic populations

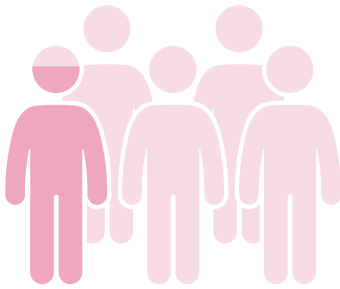
Prevalence of MASLD among different racial and ethnic groups in the United States³⁻⁵



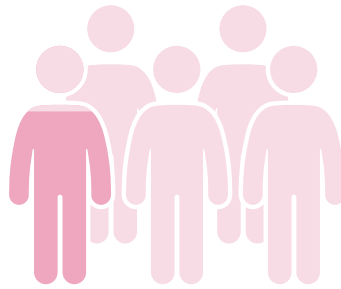
Hispanic Americans:
41.0%*



Asian Americans:
18.1%†

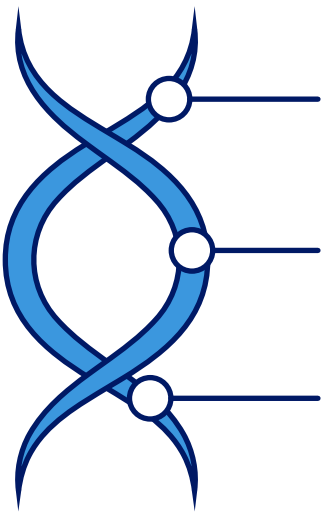


Black Americans:
18.0%†



White Americans:
14.4%‡

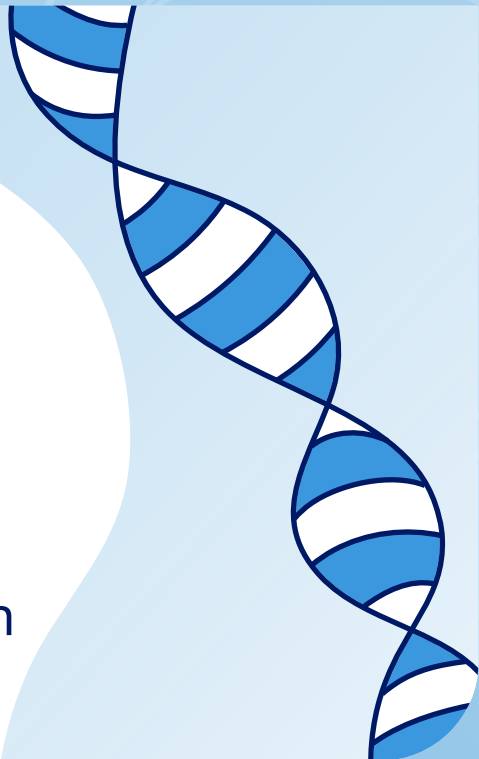
Certain genetic determinants can increase risk of MASH, regardless of cardiometabolic health



PNPLA3 variant is strongly linked to an increased risk of MASH associated with more advanced liver disease in Hispanics and lean MASLD in Asian populations^{6,7}

TM6SF2 increases liver triglycerides and ALT levels contributing to MASLD risk⁸

HSD17B13 loss of function variant may be associated with protection against MASH development⁷



There are many factors that can increase risk of MASLD and MASH, including sex, genetic factors, socioeconomic group and racial or ethnic subgroups, making it crucial to increase awareness about populations at risk for MASH



* Data for Hispanic American population is pooled prevalence data from a systematic review and meta-analysis quantifying MASLD disease burden to characterize health disparities experienced by adult Hispanic individuals in the United States. † Data for Asian and Black Americans from the National Health and Nutrition Examination Survey 2011-2016 (n=4538). ‡‡ Data for White American population is pooled prevalence data from a systematic review and meta-analysis of patients with MASLD, diagnosed using biochemical, radiologic, or histologic criteria per AASLD guidelines.⁴
ALT, alanine aminotransferase; HDL, high-density lipoprotein; HSD17B13, hydroxysteroid 17-beta dehydrogenase 13; MASH, metabolic dysfunction-associated steatohepatitis; MASLD, metabolic dysfunction-associated steatotic liver disease; PNPLA3, patatin-like phospholipase domain-containing protein 3; TM6SF2, transmembrane 6 superfamily member 2.
1. Younossi ZM et al. Hepatology. 2023;77:1335-1347; 2. Kardashian A et al. Hepatology. 2023;77:1382-1403; 3. Tesfai K et al. Clin Gastroenterol Hepatol. 2025;23:236-249; 4. Le MH et al. J Int Med. 2020;287:711-22; 5. Rich NE et al. Clin Gastroenterol Hepatol. 2018;16:198-210; 6.Yip TCF et al. J Hepatol 2022;76(3):726-734; 2024;119:1089-1101; 7. Riazi K et al. Nutrients. 2022;14:4556; 8. Luo F et al. Hepatol Commun. 2022;6:448-460.