

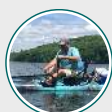
PREVALENCE OF LIVER DISEASE STATE FACT SHEET

American Liver Foundation (ALF) Liver Life Advocates—Raising Awareness, Advancing Change™, represents more than 100 million Americans affected by liver disease.

TENNESSEE



Tennessee Residents Affected by Liver Disease



Brad D.

Brad advocates for increased federal funding for rare liver disease research. Brad learned he had Gilbert's syndrome, a genetic disorder where the liver does not properly process bilirubin. Doctors monitored him and he had no symptoms, until January 2022, at age 50. Brad now has liver cirrhosis and is in need of a lifesaving liver transplant.



Patrick M.

Patrick advocates for increased liver health education and better access to specialized care. After months of trying to navigate the healthcare system, Patrick's wife was diagnosed with alcohol-associated cirrhosis and is now in need of a lifesaving liver transplant. Patrick believes her outcome may have been better had they seen a liver specialist, or received clear medical direction months earlier.



Carley V.

Carley, Miss Tennessee 2024, advocates to raise awareness about liver disease and help dispel negative stigma in the U.S. Diagnosed with autoimmune hepatitis at age 12, Carley has turned her diagnosis into a platform for good; recently placing third in the 2025 Miss America Pageant, Carley stands as a beacon of hope for the 100 million Americans affected by liver disease.

State Statistics

1,398	Tennessee residents died of liver disease/cirrhosis in 2022
264	liver transplants were performed in Tennessee
162	people are on the liver transplant waitlist
12	Tennesseans have died while on the liver transplant waitlist
13.1%	of adults in Tennessee have diabetes
39.1	of adults in Tennessee are overweight or obese
26.5%	of adults have no leisurely physical activity

ALF 2025 Legislative Priorities

- Increase funding for MASLD research and education
- Expand liver disease surveillance and prevention efforts at the CDC
- Finalize coverage for anti-obesity medication under Medicare Part D
- Support laws like the [Living Donor Protection Act](#), [Safe Step Act](#), [HOLD Act](#) and [HELP Copays Act](#)
- Improve access to liver disease treatments and reduce out-of-pocket costs

ALF is a national community of patients, caregivers and medical professionals dedicated to helping people improve their liver health. Providing guidance and life-saving resources, we are a beacon for the 100 million Americans affected by liver disease. We advocate for patients and families, fund medical research and educate the public about liver wellness and disease prevention. We bring people together through our educational programs and events and create a network of support that lasts a lifetime. ALF is the largest organization focused on all liver diseases and the trusted voice for patients and families living with liver disease. For more information visit liverfoundation.org or call 1-800-GO-LIVER (800-465-4837).