

Nonalcoholic Fatty Liver Disease in the U.S.: Clinical Characteristics of Patients Enrolled in TARGET-NASH

A. Sidney Barritt¹, Norman Gitlin², Samuel Klein³, Anna S. Lok⁴, Rohit Loomba⁵, Laura Malahias⁶, George E. DeMuth⁶, Julie M. Crawford⁶, K. Rajender Reddy⁷, Huy N. Trinh⁸, Miriam B. Vos⁹, Michael Weiss¹⁰, Kenneth Cusi¹¹, Brent A. Neuschwander-Tetri¹², Arun J. Sanyal¹³

¹. University of North Carolina, Chapel Hill, NC, United States ². Atlanta Gastroenterology Associates, Atlanta, GA, United States ³. Washington University School of Medicine, St. Louis, MO, United States ⁴. University of Michigan, Ann Arbor, MI, United States ⁵. University of California at San Diego, La Jolla, CA, United States ⁶. TARGET PharmaSolutions, Chapel Hill, NC, United States ⁷. University of Pennsylvania, Philadelphia, PA, United States ⁸. San Jose Gastroenterology, San Jose, CA, United States ⁹. Emory University Children's Healthcare of Atlanta, Atlanta, GA, United States ¹⁰. Gastro Florida, Clearwater, FL, United States ¹¹. University of Florida, Gainesville, FL, United States ¹². Saint Louis University, St. Louis, MO, United States ¹³. Virginia Commonwealth University, Richmond, VA, United States

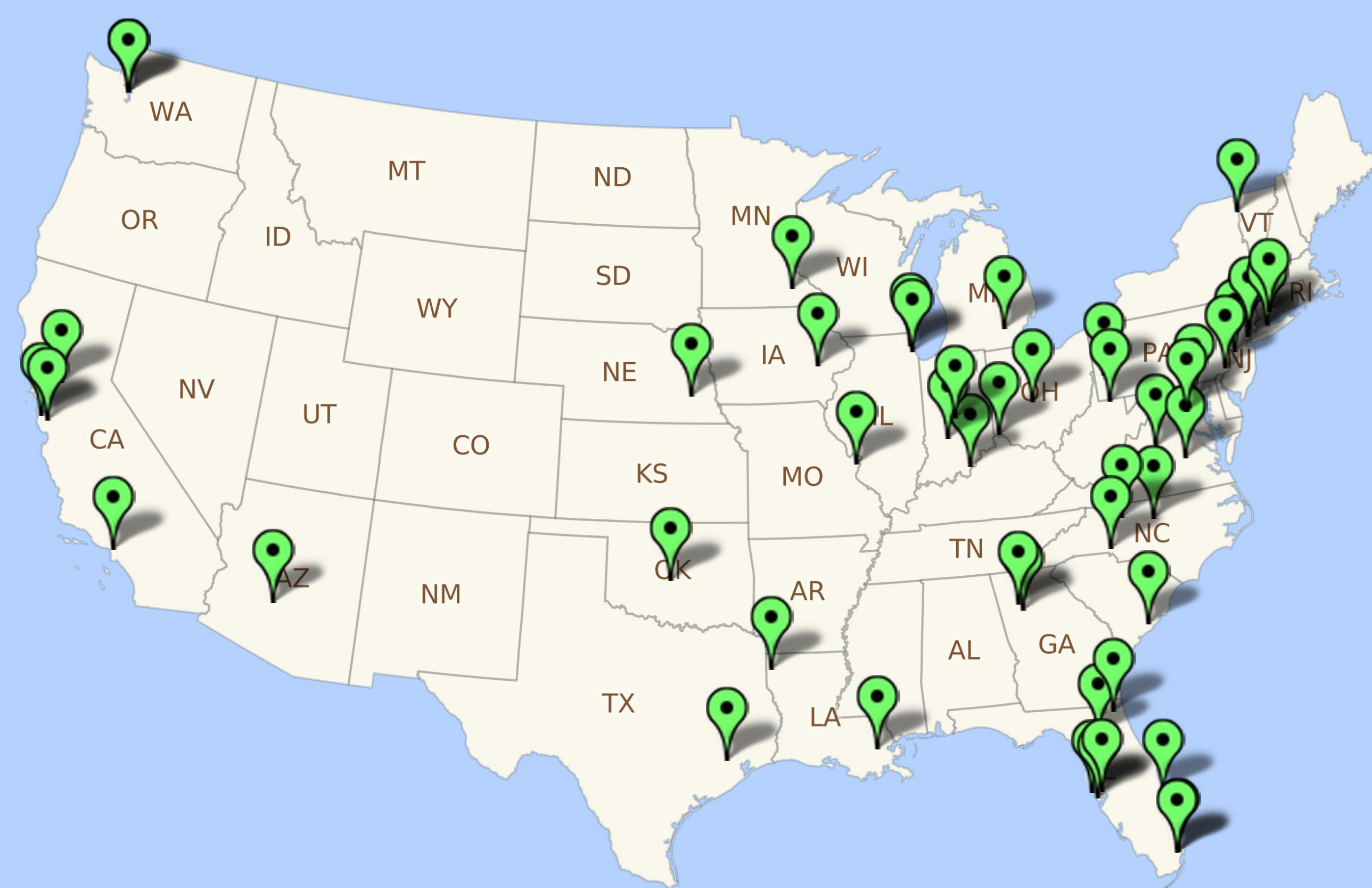
INTRODUCTION

- Nonalcoholic fatty liver disease (NAFLD) is highly prevalent in both children and adults.
- NAFLD can lead to cirrhosis, hepatocellular carcinoma and death from liver disease.
- NAFLD is also associated with increased risk of type II diabetes and cardiovascular events.
- Current treatment goals, limited to exercise and weight loss, are difficult for many patients to achieve and maintain. Thus, pharmacological therapies are greatly needed; many are in various stages of development.
- Large, observational cohorts are needed to better understand the spectrum of NAFLD by obtaining real-world data that avoids ascertainment bias from studies in tertiary care centers alone and allows for further validation of histology and non-invasive biomarkers.
- The overall aim of TARGET-NASH is to determine the natural history of NAFLD and to evaluate treatment regimens used in real world clinical practice.
- This current analysis describes the baseline characteristics of consecutive patients enrolled to date in TARGET-NASH.

METHODS

- TARGET-NASH is an observational study, initiated in 2016, of pediatric and adult patients with NAFLD managed at academic and community Hepatology, GI and Endocrinology practices.
- The medical record from consented patients including narratives, laboratory results, pathology, imaging data and patient-reported outcomes, is abstracted into a centralized data core.
- Detailed demographics, patient comorbidities, medications and disease progression are assessed as are adverse outcomes, including cardiovascular and neoplastic complications.
- Data from the first 1301 consecutively enrolled patients are presented. Descriptive statistics with ANOVA and Cochran-Armitage tests for trend are reported.

TARGET-NASH SITES



RESULTS

Characteristics of the adult cohort at enrollment

	All Subjects (N=1301)	NAFL (N=316)	NASH (N=504)	NAFLD Cirrhosis (N=481)	P-value ²
Median age (years) ¹	n=59	n=58	n=56	n=62	
18-39 years	10.2%	11.1%	16.1%	3.3%	<0.0001 ^a
40-64 years	55.6%	53.3%	58.7%	53.8%	
>=65 years	34.3%	35.6%	25.2%	42.9%	
Sex, female	57.8%	51.6%	60.9%	58.5%	0.0261 ^b
Median BMI (kg/m ²)	32.0	29.0	32.0	33.0	<0.0001 ^a
Diabetes	43.7%	21.5%	37.3%	64.9%	<0.0001 ^c
Cardiovascular disease	19.8%	14.2%	17.1%	26.4%	<0.0001 ^c
Hypertension	57.6%	49.7%	52.6%	68.0%	<0.0001 ^c
Anxiety or Depression	29.4%	19.0%	29.6%	36.0%	<0.0001 ^c
Osteoarthritis	20.9%	15.2%	17.5%	28.3%	<0.0001 ^c
Obstructive Sleep Apnea Syndrome	19.3%	6.7%	19.0%	27.7%	<0.0001 ^c
Medication: pioglitazone	3.5%	2.8%	3.4%	4.0%	0.4114 ^c
Medication: metformin	31.5%	17.8%	28.7%	42.8%	<0.0001 ^c
Supplement: vitamin E	14.1%	8.2%	15.6%	16.1%	0.0035 ^c

¹ Age calculated based on year of consent minus birth year² Tests of homogeneity or trend:^a ANOVA test^B Likelihood Ratio test^c Cochran-Armitage test

DISEASE CATEGORY DEFINITIONS

NAFL	Any participant not meeting criteria for clinical NASH or cirrhosis
NASH	Confirmed by biopsy: •Steatohepatitis by Brunt OR •NAS total score ≥ 4 Clinical diagnosis: •Adults: ALT > 19 U/L for female, > 30 U/L for men and; •Hepatic steatosis on biopsy or CT/US/MRI and; • ≥ 1 of the following: BMI ≥ 30 , type II diabetes, dyslipidemia
NAFLD	1) Liver biopsy with fibrosis stage = 4 OR
Cirrhosis	2) Liver biopsy with fibrosis stage = 3 and 1 \geq clinical signs of cirrhosis OR 3) 2 or more clinical signs of cirrhosis OR 4) FibroScan® stiffness result ≥ 11 kPa

RESULTS

Characteristics of the pediatric cohort at enrollment

- Number enrolled = 152
- Median age = 14
- Sex, female = 27.6%
- Median BMI = 31.5
- NAFL = 15.1%
- NASH = 80.2% (32% biopsy confirmed)
- NAFLD Cirrhosis = 4.6%

SUMMARY

- In the adult cohort, 76% of the first 1301 consecutive patients enrolled had NASH/NAFLD cirrhosis.
- Similarly, 85% of the first 152 pediatric patients enrolled had NASH/NAFLD cirrhosis.
- Metabolic syndrome risk factors like diabetes, hypertension and obesity are present in greater proportions in NASH and NAFLD cirrhosis.
- Comorbid conditions that make lifestyle intervention difficult (namely, anxiety / depression, osteoarthritis and cardiovascular disease) occur at significantly greater rates as disease severity increases.

CONCLUSIONS

- Participants enrolled in TARGET-NASH include populations, such as patients with cirrhosis and cardiovascular disease, not widely represented in clinical trials.
- Real-world clinical registries are important for obtaining unbiased natural history data and determining clinical effectiveness of new interventions.
- TARGET-NASH will be an important source of real world patient oriented outcome data.

STATEMENT AND DISCLOSURES

TARGET-NASH is a collaboration among academic & community investigators, the pharmaceutical industry, and NASH patient community advocates. TARGET-NASH is sponsored by TARGET PharmaSolutions, Inc. TARGET thanks the study staff, nurses, health care providers and patients at each study center for their contributions to this work.

Listings of Principal Investigators and Industry Partners are available upon request by emailing info@targetpharmasolutions.com.