



#### PREMISE

- To promote optimal health and well-being, adults aged 18-60 years are recommended to sleep at least 7 hours each night.
- Sleep disruption has been associated with metabolic diseases, such as obesity and diabetes in murine models.
- Sleep disruption can interrupt the circadian rhythm and thus the body metabolism.
- It can be speculated that sleep-related problems may trigger several pathophysiologic processes associated with nonalcoholic fatty liver disease (NAFLD).

**Aim:** We hypothesized an association between sleep duration, steatosis, and advanced fibrosis.

### METHODS

- Using the NHANES database, we identified all patients aged 18 and older from 2017 to March 2020 prepandemic surveys.
- The presence of fatty liver was determined using vibration - controlled transient elastography (VCTE).
- Sleep duration was grouped by short sleep duration  $(\leq 7 \text{ hours})$  and normal sleep duration ( >7 hours).
- Linear regression models were used to examine the relationship between continuous CAP score and sleep duration, after adjusting for potential confounders.

# Association between sleep duration and hepatic steatosis in the USA

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Table 1. Cohort characteristics			
		Sleep hours	
	Overall	≤7 hours	>7 hours
Age, (mean ±)	47 ± 16	46.6 ± 14	47 ± 16
Sex, female (%)	51	46	54
Race/ethnicity, (%)			
White	62	59	64
Black	11	13	9
Hispanic	16	17	15
Asian	5	5	6
Other	4	4	3
BMI, (mean ±)	29.6 ± 6.4	30 ± 7	29 ± 6
Diabetes, (%)	10	11	10
FibroScan®			
CAP, (mean ±)	264 ± 56	269 ± 56	260 ± 56
kPa, (mean)	5.8 ± 4	5.9 ± 4	5.7 ± 4

**BMI**, body mass index; **CAP**, controlled attenuation parameter; **kPa**, kilopascales

#### CONCLUSIONS

Overall, FibroScan<sup>®</sup> data in NHANES 2017-2020 support a positive association between short sleep duration and hepatic steatosis independently of metabolic factors such as BMI.

### RESULTS







### Model 1:

Adjusted for age and body mass index

#### Model 2:

Adjusted for age, body mass index and sex