In the present data, we provide the details of the cross-sectional study, from the Washington Heights-Inwood Community Aging Project (WHICAP) that examined the association between Apolipoprotein E (APOE-ε4) and snoring/sleep apnea in non-demented older adults. A total of 1944 non-demented older adults constituted our sample. Sleep dysfunction was measured using sleep categories derived from the Medical Outcomes Study Sleep Scale. Stratified analyses were conducted in order to examine the association between APOE-ε4 and sleep variables by ethnic group. For further analyses and enhanced discussion, see “Examining the association between Apolipoprotein E (APOE) and self-reported sleep disturbances in non-demented older adults” by Tsapanou et al. (2015) [1].

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### Specifications Table

| Subject area          | Neuropsychology, Neurology, Genetics     |
|-----------------------|-----------------------------------------|
| More specific subject area | Neurogenetics                           |
| Type of data          | Questionnaire, Tables                   |
| How data was acquired | Quantitative data of 1944 non-demented elderly, using a self-reported sleep questionnaire and their \( APOE-\varepsilon 4 \) status |
| Data format           | Raw data, Analyzed                      |
| Experimental factors  | \( APOE-\varepsilon 4 \) and sleep questions were the main variables used. Age, sex, ethnicity, years of education, and body mass index were used as covariates. |
| Experimental features | Characterization of the \( APOE-\varepsilon 4 \) status |
| Data source location  | New York, New York, USA                 |
| Data accessibility    | Data is in this article                 |

### Value of the data

- Identifying the sleep measures associated with \( APOE-\varepsilon 4 \).
- Providing novel information about the differences among different ethnic groups.
- Providing the frequencies of the \( APOE \) alleles in a large sample of older adults.

1. **Data, experimental design, materials and methods**

   In the present data, we present the details of the cross-sectional study that examined the association between Apolipoprotein E \( (APOE-\varepsilon 4) \) and snoring/sleep apnea in a large group of older adults. We provide the demographic characteristics of our sample, the frequencies of the \( APOE \) alleles in our population, the sleep scale we used, and the results of the association analyses between \( APOE-\varepsilon 4 \) and sleep variables stratified by ethnic group [1].

1.1. **Sample collection**

   Participants were drawn from the Washington Heights–Inwood Community Aging Project (WHICAP) at Columbia University Medical Center [2,3]. WHICAP is a community based research study designed to identify risk factors and biomarkers for aging and Alzheimer's disease in a multi-ethnic cohort that includes Whites, African-Americans, Caribbean-Hispanics, and Other [4]. All participants were over 65 years old, and non-demented at the time of the evaluation. For details, see supplementary Table 1.

1.2. **Sleep measures**

   Sleep quality was assessed using the Sleep Scale from the RAND Medical Outcomes Study. This scale is a self-report 12-item questionnaire [5,6]. Each of the questions has a possible rating of 0-6, based on the frequency of the sleep problem. Using the sleep questionnaire manual [6], we used the five clustered sleep categories to define our analyses phenotype: 1. Sleep disturbance, 2. Snoring, 3. Sleep short of breath/awaking with a headache, 4. Sleep adequacy, and 5. Daytime somnolence. Additionally, categories 2 and 3 were combined into a single variable ‘sleep apnea’ (see Supplementary Fig. 1).
1.3. APOE genotyping

WHICAP participants were APOE genotyped as previously described [7]. APOE genotypes were transformed into a dichotomous trait based on the number of APOE-ε4 alleles: 0 if the individual does not carry any copy of the ε4 allele (non-ε4 carriers) or 1 if the individual carries 1 or 2 copies of the ε4 allele (ε4 carriers). Carriers of ε2ε4 alleles were not included in the initial sample due to the opposite effect of these two alleles [8–10]. More details about the APOE genotyping can be found in previous studies [7]. For the frequencies of the APOE alleles, see supplementary Table 2.

1.4. Analysis

All statistical analyses were performed using SPSS 22 (SPSS, Chicago, Illinois). Nominally significant p values were defined as p < 0.05. Unadjusted Linear Regression Analyses: We used linear regression models with APOE-ε4 at baseline evaluation as the predictor and the previously described sleep score as the independent variable. Adjusted Linear Regression Analyses: Secondary analyses were performed adjusting for: age, sex, ethnicity, education, and body mass index (BMI). Ethnicity was ascertained based on self-report using the format of the 1990 census [11]. Participants were then assigned to one of the three groups: White, African-American, and Caribbean-Hispanic. To further examine any possible differences among the ethnic groups, we stratified the sample and perform analyses within each ethnic group independently (Whites n = 431, African Americans n = 465, and Caribbean-Hispanics n = 1048) (see Supplementary Table 3).

Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at doi:http://dx.doi.org/10.1016/j.dib.2015.09.014.

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