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# Association between coffee consumption and functional disability in older US adults

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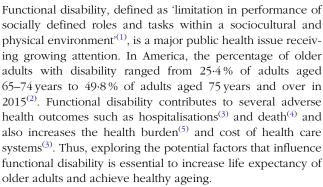
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#### **Abstract**

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The effect of coffee consumption on functional disability has been scarcely investigated. Thus, this study aimed to examine the association between coffee consumption and functional disability in older American adults. Participants ( $\geq$ 60 years old, n 7704) were from the National Health and Nutrition Examination Survey 2007–2016. Coffee consumption was assessed through two 24-h dietary recall interviews. Five domains of functional disability including lower extremity mobility (LEM), general physical activity (GPA), leisure and social activities (LSA), activities of daily living (ADL) and instrumental activities of daily living (IADL) were self-reported. Age- and multivariate-adjusted logistic regression models and restricted cubic spline analyses were used. Total coffee consumption was inversely associated with LEM, GPA, LSA and IADL disability. Compared with non-drinkers of total coffee, those who consumed  $\geq$ 2 cups/d reported lower odds of LEM (OR 0·67, 95 % CI 0·50, 0·91), GPA (OR 0·65, 95 % CI 0·47, 0·88), LSA (OR 0·61, 95 % CI 0·45, 0·83) and IADL (OR 0·59, 95 % CI 0·44, 0·78) disability. The dose—response analyses confirmed these relationships. Intake of  $\geq$ 2 cups/d caffeinated coffee was also inversely linked to GPA (OR 0·67, 95 % CI 0·48, 0·92), LSA (OR 0·66, 95 % CI 0·46, 0·93) and IADL (OR 0·57, 95 % CI 0·43, 0·75) disability, whereas the inverse association of 2+ cups/d decaffeinated coffee was only on LEM (OR 0·43, 95 % CI 0·23, 0·81) and LSA (OR 0·39, 95 % CI 0·16, 0·94) disability. The present study suggested that coffee consumption was inversely associated with functional disability in older American adults. Those associations of diverse coffee types differed across domains of functional disability.

Key words: Coffee consumption: Functional disability: National Health and Nutrition Examination Survey: Cross-sectional studies

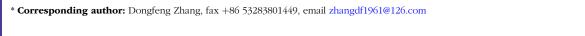


Previous epidemiological literature has examined the effects of dietary factors on functional disability, such as dairy consumption<sup>(6,7)</sup>, fruit and vegetables<sup>(8)</sup>. Healthy dietary patterns could contribute to prevention or delay of functional disability<sup>(9)</sup>.

Coffee, as one of the most popular consumed beverages worldwide, is considered as part of a healthy diet<sup>(10)</sup> due to its numerous underlying benefits to reduce the risk of type 2 diabetes<sup>(11)</sup>, CVD<sup>(12)</sup>, sarcopenia<sup>(13)</sup>, falls<sup>(14)</sup> and mortality<sup>(15)</sup>, which predispose individuals to functional disability. However, few studies have investigated the association between coffee consumption and functional disability. To our knowledge, only a cohort study has investigated the association in older Spanish adults and found the protective association between coffee consumption and functional impairment just among women, hypertensive, diabetic and obese patients<sup>(16)</sup>.

Therefore, we conducted a cross-sectional study using a large population-based sample derived from the National Health and Nutrition Examination Survey (NHANES) (2007–2016) to estimate the association between coffee consumption and functional disability in older American adults.

**Abbreviations:** ADL, activities of daily living; GPA, general physical activity; IADL, instrumental activities of daily living; LEM, lower extremity mobility; LSA, leisure and social activities; NHANES, National Health and Nutrition Examination Survey.





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#### Methods

## Study sample

NHANES is an ongoing biennial cross-sectional survey of the non-institutionalised, civilian US population. Based on the complex stratified multistage probability sampling design, NHANES collected demographic, socio-economic, health and nutritional data on approximately 5000 participants every 2 years. Participants completed health interviews at home and then performed the health examination at mobile examination centres that included medical, dental and physiological measurements. Details about survey design and data collection procedures were available elsewhere<sup>(17)</sup>. The Centers for Disease Control and Prevention National Center for Health Statistics institutional review board approved the NHANES, and all participants provided written informed consent.

Data from NHANES 2007–2016 were combined for the present study, including 50 588 participants. The study sample was restricted to adults aged  $\geq 60$  years (n 9760). About 2053 participants who had ineligible dietary recall status including 'not reliable or not met the minimum criteria' and 'not done' were excluded. After excluding the three participants with incomplete functional disability data, 7704 individuals were included in this study (Fig. 1).

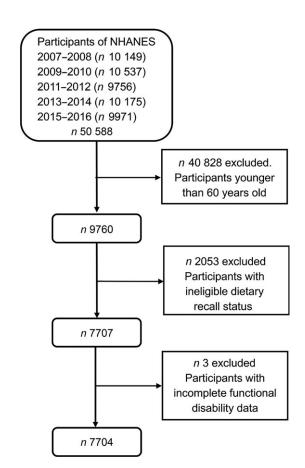


Fig. 1. Study participant flow. NHANES, National Health and Nutrition Examination Survey.

## Study variables

Functional disability. In NHANES, five domains of functional disability<sup>(18)</sup>, including lower extremity mobility (LEM), general physical activity (GPA), leisure and social activities (LSA), activities of daily living (ADL) and instrumental activities of daily living (IADL), were evaluated by asking participants about their ability to perform nineteen routine tasks in the Physical Functioning Questionnaire without special equipment (online Supplementary Table S1). Participants were provided five options: 'no difficulty', 'some difficulty', 'much difficulty', 'unable to do' or 'don't do this activity'. For each domain, disability status was defined as having any difficulty in performing one or more activities within a given domain<sup>(19,20)</sup>.

Coffee consumption. The dietary interview component of NHANES was conducted as a partnership between the US Department of Agriculture and the US Department of Health and Human Services. All participants were eligible for two 24-h dietary recalls administered by trained interviewers. The first recall was collected in person through mobile examination centres, and the second one was collected via telephone 3–10 d later. The food intake data, which were collected and processed by the automated methods described elsewhere<sup>(21)</sup>, were used to measure the type and quantity of consumed food and beverages as well as the daily total consumption of nutrients, energy and other components from those food and beverages.

Caffeinated and decaffeinated coffee consumption was assessed according to the food code, and total coffee consumption was the sum of those two. Caffeine consumption was estimated from the intake of all caffeine-containing foods and beverages. As the high agreement of two dietary recalls (interclass correlation coefficient 0.71 for total coffee, 0.71 for caffeinated coffee, 0.65 for decaffeinated coffee and 0.67 for caffeine, P < 0.001), we chose the average coffee intake of two dietary recalls for our main analysis. In order to visualise the coffee consumption, total, caffeinated and decaffeinated coffee consumption (in g) was divided by one cup size (10 oz or 283.5 ml)<sup>(22)</sup> and classified into four groups: none, <1 cup/d, 1 to <2 cup/d and  $\ge 2$  cup/d. Caffeine intake was divided into quartile.

Other variables. Covariates recorded in this study included age, sex (male and female), race and ethnicity, marital status ('single' and 'married'), educational level, annual household income, BMI, physical activity, cigarette smoking, alcohol drinking, energy intake, tea intake, soft drink intake, depression, hypertension, diabetes, arthritis, gout, stroke and cancer. Race and ethnicity were self-reported and allocated into five categories: Mexican American, other Hispanic, non-Hispanic White, non-Hispanic Black and another race. Education was grouped as 'below high school', 'high school' and 'above high school'. Annual household income was categorised as '<\$20 000', '\$20 000 to <\$45 000', '\$45 000 to <\$75 000' and '≥\$75 000'. BMI was grouped into four categories: underweight (<18·5 kg/m²), normal (18·5–24·9 kg/m²), overweight (25–29·9 kg/m²) and obesity (≥30 kg/m²). Energy



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intake, tea intake and soft drink intake were calculated based on the two 24-h dietary recalls. Based on the Global Physical Activity Questionnaire (23), physical activity was calculated and classified as 'none', 'not meeting WHO recommendations' and 'meet WHO recommendations'. Cigarette smoking was categorised as 'never', 'former' and 'current'. Alcohol drinking was defined as 'none', 'light', 'moderate' and 'heavy' (24). Depressive symptom was assessed by using the Patient Health Questionnaire<sup>(25)</sup>. Hypertension, diabetes, arthritis, gout, stroke and cancer were dichotomised as 'yes' or 'no' based on answers of the question 'Has a doctor ever diagnosed you with "this problem"?"

## Statistical analysis

Participants' characteristics were presented as numbers and percentages for categorical variables and mean values and standard deviations for continuous variables according to total coffee consumption groups. The  $\chi^2$  test or ANOVA was used to determine descriptive statistics differences across groups. Age-adjusted and multivariate-adjusted binary logistic regression models were used to explore the associations of total, caffeinated, decaffeinated coffee and caffeine consumption with each functional disability's domain. Linear trend across categories of coffee consumption was evaluated by Stata's 'contrast' command assigning orthogonal polynomial coefficients (26,27). Moreover, the multivariable-adjusted models were repeated with strata of sex. After 1% abnormal values before and after were rejected, non-drinkers occupied 28.0%; thus, we used restricted cubic spline analyses with three knots located at the 30th, 50th and 70th percentiles of the exposure distribution to assess doseresponse relationships between total coffee consumption and functional disability. The multivariate-adjusted and restricted cubic spline models were adjusted for these potential confounders: age, sex, race and ethnicity, marital status, educational level, annual household income, BMI, physical activity, cigarette smoking, alcohol drinking, energy intake, tea intake, soft drink intake, depression, hypertension, diabetes, arthritis, gout, stroke and cancer. When caffeinated coffee was tested, decaffeinated coffee was adjusted as covariates and vice versa.

Considering that diseases might influence the association between coffee consumption and functional disability and adjusting them as covariates might not completely remove residual confounding, we conducted the sensitivity analyses after excluding participants with CVD and cancer.

All analyses were accounted for the complex survey design of NHANES conducted on Stata version 15 (StataCorp LP). NHANES 10-year sampling weights, primary sampling unit and strata information were used to obtain unbiased, nationally representative estimation. Statistical significance was defined as twosided P values  $\leq 0.05$ .

#### Results

After filtering, 7704 participants aged 60 years and above were eligible for this study (Fig. 1). Total, caffeinated and decaffeinated coffee consumption weighted mean and standard deviations were 1.35 (sp 1.46) cups/d, 1.10 (sp 1.41) cups/d and 0.25 (sp 0.70) cups/d, respectively. Online Supplementary Table \$2 summarises the characteristics according to the categories of total coffee consumption.

Table 1 shows the weighted OR (95 % CI) of each functional disability's domain according to categories of total coffee consumption. In the age-adjusted models, higher total coffee consumption was associated with decreased odds of all domains of functional disability. After adjusting the multivariate confounders, the lower odds of LEM disability were observed among total coffee drinkers of 0 to <1 cups/d (OR 0.74, 95 % CI 0.57, 0.96) and ≥2 cups/d (OR 0.67, 95 % CI 0.50, 0.91) with respect to non-coffee drinkers ( $P_{\text{trend}} = 0.03$ ). Compared with nondrinkers of total coffee, those who consumed 2 cups or more per d had OR of 0.65 (95% CI 0.47, 0.88) and 0.61 (95% CI 0.45, 0.83) for disability of GPA ( $P_{\text{trend}} = 0.02$ ) and LSA  $(P_{\text{trend}} = 0.006)$ , respectively. And the intakes of 1 to <2cups and ≥2 cups of coffee/d were linked to reduced odds of IADL disability (OR<sub>1 to <2cups/d</sub> 0.72, 95 % CI 0.54, 0.95; OR<sub>>2 cups/d</sub>: 0.59, 95 % CI 0.44, 0.78,  $P_{\text{trend}} = 0.001$ ). Although total coffee drinkers ≥2 cups/d were also inversely associated with lower odds of reporting ADL disability (OR 0.70), the CI (0.50, 1.01) was wide as a result of the small number of participants.

The weighted associations between caffeinated coffee and functional disability are displayed in Table 2. In the age-adjusted models, participants with higher caffeinated coffee consumption tended to have lower odds of all domains of functional disability except LEM. In the fully adjusted models, compared with nondrinker of caffeinated coffee, drinkers ≥2 cups/d showed lower odds of reporting disability in GPA (OR 0.67, 95 % CI 0.48, 0.92,  $P_{\text{trend}} = 0.01$ ) and LSA (OR 0.66, 95 % CI 0.46, 0.93,  $P_{\text{trend}} = 0.02$ ). The odds of IADL disability was decreased across all other categories of caffeinated coffee consumption with respect to non-coffee drinkers (OR $_0$  to <1 cups/d 0.71, 95 % CI 0.55, 0.93;  $OR_{1 \text{ to } < 2 \text{ cups/d}} \text{ } 0.69, 95 \% \text{ CI } 0.52, 0.91; OR_{\geq 2 \text{ cups/d}} \text{ } 0.57, 95 \% \text{ CI }$  $0.43, 0.75, P_{\text{trend}} < 0.001$ ). A lower odds of ADL disability was also observed among caffeinated coffee drinkers ≥2 cups/d with respect to non-coffee drinkers, although the association was not statistically significant (OR 0.72, 95 % CI 0.52, 1.01). In Table 3, we report the weighted age- and multivariate-adjusted associations between categories of decaffeinated coffee intake and functional disability. The higher decaffeinated coffee intake was only related to disability in LEM and LSA after adjusting all confounders. Comparing drinkers of 0 to <1 and  $\ge 2 \text{ cups/d}$ to non-drinkers of decaffeinated coffee consumption, the corresponding OR was 0.65 (95 % CI 0.47, 0.89) and 0.43 (95 % CI 0.23, 0.81) for LEM disability ( $P_{\text{trend}} = 0.02$ ) and 0.64 (95% CI 0.50, 0.82) and 0.39 (95% CI 0.16, 0.94) for disability in LSA  $(P_{\text{trend}} = 0.08).$ 

The weighted associations between caffeine intake (mg/d) and functional disability are presented in online Supplementary Table S3. Participants in the highest quartile of caffeine intake had lower odds of reporting disability in GPA (OR 0.76, 95% CI 0.59, 0.99,  $P_{\text{trend}} = 0.04$ ) than those in the lowest quartile. Also, the odds of reporting disability of ADL was reduced among participants in the third quartile of intake



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Table 1. Age- and multivariate\*-adjusted associations between categories of total coffee intake and functional disability, National Health and Nutrition Examination Survey (NHANES) (2007–2016)

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(Numbers and percentages; weighted odds ratios and 95 % confidence intervals)

	Total coffee (cups/d)											
	0		0 to <1			1 to <2			2+			
	OR		OR		95 % CI	OR		95 % CI	OR		95 % CI	P <sub>trend</sub>
LEM ( <i>n</i> 7699)												
Cases/participants		932/2082		906/2065			808/2098			559/1454		
Weighted prevalence (%)		41.2		39.4			37⋅1			32.7		
Age adjusted	Ref.		0.83		0.67, 1.02	0.76		0.62, 0.94	0.71		0.57, 0.89	0.004
Multivariate adjusted	Ref.		0.74		0.57, 0.96	0.79		0.59, 1.05	0.67		0.50, 0.91	0.028
GPA (n 7699)		1050/0001		1000/0004			1000/0000			007/4455		
Cases/participants		1353/2081		1363/2064			1299/2099			897/1455		
Weighted prevalence (%)	Ref.	65⋅0	0.90	65⋅0	0.74, 1.10	0.80	61.9	0.64.0.00	0.70	55.8	0.55, 0.88	0.004
Age adjusted Multivariate adjusted	Ref.		0.86		0.74, 1.10	0.82		0.64, 0.99 0.62, 1.08	0.70		0.47, 0.88	0.004
LSA ( <i>n</i> 7704)	i ici.		0.00		0.07, 1.10	0.02		0.02, 1.00	0.03		0.47, 0.00	0.013
Cases/participants		573/2083		593/2066			488/2100			307/1455		
Weighted prevalence (%)		22.7		24.4			21.1			16.3		
Age adjusted	Ref.		1.03		0.83. 1.28	0.86		0.67. 1.11	0.68		0.53, 0.86	0.001
Multivariate adjusted	Ref.		0.93		0.68, 1.26	0.96		0.69, 1.34	0.61		0.45, 0.83	0.006
ADL ( <i>n</i> 7704)					,			,			-,	
Cases/participants		553/2083		574/2066			458/2100			296/1455		
Weighted prevalence (%)		22.3		22.9			20.5			16.3		
Age adjusted	Ref.		0.97		0.77, 1.21	0.85		0.67, 1.08	0.69		0.52, 0.91	0.008
Multivariate adjusted	Ref.		0.88		0.65, 1.19	0.95		0.69, 1.30	0.70		0.50, 1.01	0.088
IADL ( <i>n</i> 7704)												
Cases/participants		749/2083		757/2066			638/2100			416/1455		
Weighted prevalence (%)		32.6		32.0			28.0			23.4		
Age adjusted	Ref.		0.88		0.70, 1.11	0.74		0.59, 0.93	0.65		0.51, 0.82	<0.001
Multivariate adjusted	Ref.		0.77		0.57, 1.03	0.72		0.54, 0.95	0.59		0.44, 0.78	0.001

LEM, lower extremity mobility; Ref., reference; GPA, general physical activity; LSA, leisure and social activities; ADL, activities of daily living; IADL, instrumental activities of daily living.

(OR 0·71, 95 % CI 0·53, 0·97,  $P_{\rm trend}$  = 0·08). Compared with the participants with lowest quartile of caffeine intake, those with the second and third quartiles had OR of 0·65 (95 % CI 0·48, 0·87) and 0·63 (95 % CI 0·47, 0·84) of IADL disability ( $P_{\rm trend}$  < 0·001).

The associations between total, caffeinated, decaffeinated coffee and functional disability stratified by sex are displayed in online Supplementary Tables S4 and S5. For males, total, caffeinated coffee and caffeine intakes were inversely linked to four domains of functional disability (adjusted OR ranged from 0.48 to 0.67) except LSA disability, and the intake of decaffeinated coffee was negatively associated with four domains of functional disability (adjusted OR ranged from 0.35 to 0.64) except IADL disability. While for females, there was inverse association of total coffee intake with LSA and IADL disability (adjusted OR ranged from 0.51 to 0.53), the inverse association was observed between caffeinated coffee consumption and LSA, ADL and IADL disability (adjusted OR ranged from 0.47 to 0.75), decaffeinated coffee consumption was negatively linked to LEM, GPA and LSA disability (adjusted OR ranged from 0.58 to 0.67) and the caffeine intake was inversely associated with IADL disability (OR 0.56, 95 % CI 0.40, 0.79). In the sensitivity analysis, the results were almost unchanged when we excluded participants with CVD and cancer (online Supplementary Tables **S6** and **S7**).

Fig. 2 depicts dose–response relationship between total coffee consumption and each domain of functional disability except ADL. We found that with increasing intakes of total coffee consumption, the OR of the disability of LEM ( $P_{\rm for\ non-linearity}=0.06$ ), GPA ( $P_{\rm for\ non-linearity}=0.36$ ), LSA ( $P_{\rm for\ non-linearity}=0.44$ ) and IADL ( $P_{\rm for\ non-linearity}=0.11$ ) decreased in dose relationship manner. The does–response relationship of total coffee consumption was not carried out because no statistically significant association was observed between total coffee consumption and ADL in the multivariate logistic regression model.

## Discussion

In this large-scale cross-sectional study, after adjusting confounders, older adults with high total coffee consumption had lower odds of reporting disability in LEM, GPA, LSA, as well as IADL, and the dose–response relationship was detected. For different types of coffee, higher intake of caffeinated coffee was associated with reduced prevalence of GPA, LSA and IADL disability. And the decaffeinated coffee showed meaningful inverse association only on the disability of LEM and LSA. The association between different types of coffee and five domains of functional disability was discrepant in males and females.

Limited studies have evaluated the association between coffee consumption and functional disability. Machado-Fragua



<sup>\*</sup> Adjusted for age, sex, race and ethnicity, marital status, educational level, annual household income, BMI, physical activity, cigarette smoking, alcohol drinking, energy intake, tea intake, soft drink intake, depression, hypertension, diabetes, arthritis, gout, stroke and cancer.

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Table 2. Age- and multivariate\*-adjusted associations between categories of caffeinated coffee intake and functional disability, National Health and Nutrition Examination Survey (NHANES) (2007-2016)

(Numbers and percentages; weighted odds ratios and 95 % confidence intervals)

	Caffeinated coffee (cups/d)											
	0		0 to <1			1 to <2			2+			
	OR		OR		95 % CI	OR		95 % CI	OR		95 % CI	$P_{trend}$
LEM (n 7699)												
Cases/participants		1327/3002		794/1840			621/1688			463/1169		
Weighted prevalence (%)		40.2		38.4			36-2			33.6		
Age adjusted	Ref.		0.87		0.71, 1.06	0.82		0.66, 1.01	0.82		0.65, 1.03	0.084
Multivariate adjusted	Ref.		0.79		0.61, 1.01	0.77		0.59, 1.02	0.75		0.55, 1.02	0.084
GPA ( <i>n</i> 7699)												
Cases/participants		1977/3000		1198/1840			1018/1689			719/1170		
Weighted prevalence (%)		64.8		65⋅1			60.0			55⋅3		
Age adjusted	Ref.		0.95		0.78, 1.17	0.79		0.65, 0.95	0.72		0.56, 0.92	0.005
Multivariate adjusted	Ref.		0.92		0.72, 1.18	0.79		0.61, 1.01	0.67		0.48, 0.92	0.014
LSA (n 7704)							000/1000					
Cases/participants		813/3003		502/1842			392/1689			254/1170		
Weighted prevalence (%)		21.9		24.2			20.9			16.4		
Age adjusted	Ref.		1.09		0.87, 1.37	0.93		0.71, 1.20	0.74		0.56, 0.98	0.024
Multivariate adjusted	Ref.		1.07		0.79, 1.43	0.96		0.71, 1.32	0.66		0.46, 0.93	0.016
ADL (n 7704)		705/0000		400/4040			0.40/4.000			045/4470		
Cases/participants		795/3003		498/1842			343/1689			245/1170		
Weighted prevalence (%)	Ref.	22.3	0.00	21.7	074 115	0.05	19-9	0.05 1.10	0.71	15.7	0.55.000	0.010
Age adjusted	Ref.		0.93 0.88		0.74, 1.15	0.85 0.88		0.65, 1.10	0.71 0.72		0·55, 0·93 0·52, 1·01	0.013 0.066
Multivariate adjusted IADL (n 7704)	nei.		0.00		0.66, 1.18	0.00		0.65, 1.19	0.72		0.52, 1.01	0.000
Cases/participants		1087/3003		644/1842			496/1689			333/1170		
Weighted prevalence (%)		32.9		29.6			27.5			22.5		
Age adjusted	Ref.	32.3	0.80	23.0	0.65, 0.99	0.75	21.5	0.59. 0.94	0.63	22.0	0.49, 0.81	<0.001
Multivariate adjusted	Ref.		0.71		0.55, 0.93	0.73		0.59, 0.94	0.57		0.43, 0.75	<0.001
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LEM, lower extremity mobility; Ref., reference; GPA, general physical activity; LSA, leisure and social activities; ADL, activities of daily living; IADL, instrumental activities of daily living.

et al. (16) conducted a cohort study with 3289 Spanish adults aged ≥60 years and found that although without significant interaction of coffee intakes and subgroups, coffee consumption was associated with lower risk of functional disability in women and adults with hypertension, obesity or diabetes. But the protective effect of coffee was not significant in all adults. Whereas our study suggested that older American adults who consumed higher total, caffeinated and decaffeinated coffee were associated with reporting lower odds of different domains of functional disability after adjusting the risk factors like hypertension, diabetes and so on. The inconsistent conclusions could possibly due to different study designs, studied samples, types of coffee consumed and measurements of functional disability.

Our findings supported that coffee consumption was associated with functional disability. What remains unknown is what components of coffee confer such benefits. We examined the relationship of caffeinated, decaffeinated coffee and caffeine intake on the functional disability and found that although caffeinated coffee and decaffeinated coffee were both inversely related to functional disability, those relationships differ across domains of functional disability. Caffeinated coffee and caffeine ntake both have beneficial associations on GPA and IADL disability, which means the caffeine might be the beneficial component of coffee. In consistent with the present results, previous studies have demonstrated that the protective association between caffeinated coffee and falls might be due to the caffeine<sup>(14)</sup>. A randomised controlled trial assessed the effect of caffeine ingestion on functional performance and found that acute caffeine ingestion positively enhanced functional performance in apparently healthy older adults<sup>(28)</sup>. While we also found that higher decaffeinated coffee was related to the lower odds of LEM and LSA disability. Decaffeinated coffee contains many non-caffeine compounds, such as chlorogenic acid and diterpenoids. Chlorogenic acid, as the most common compounds, improves many aspects of cognition such as sustained attention, reaction time and alertness<sup>(29)</sup>; those ability was essential for older adults to maintain their physical function<sup>(30)</sup>. Therefore, the beneficial effects of decaffeinated coffee might be partially due to chlorogenic acid which enhanced sustained attention and reaction time.

The mechanism about the inverse association of coffee on functional disability is unclear. Previous findings, along with those from our study, may suggest that the inverse association of coffee consumption against functional disability might be due to the beneficial effects of coffee on muscle strength (31) cognition<sup>(32,33)</sup> and anti-inflammatory effect<sup>(34)</sup>. Guo et al.<sup>(31)</sup> examined the effect of coffee on the skeletal muscles used aged mice in vivo and detected that total coffee treatment was benefit to the prevention of age-related sarcopenia in mice. These



Adjusted for age, sex, race and ethnicity, marital status, educational level, annual household income, BMI, physical activity, cigarette smoking, alcohol drinking, energy intake, tea intake, soft drink intake, depression, hypertension, diabetes, arthritis, gout, stroke, cancer and decaffeinated coffee consumption.

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**Table 3.** Age- and multivariate\*-adjusted associations between categories of decaffeinated coffee intake and functional disability, National Health and Nutrition Examination Survey (NHANES) (2007–2016) (Numbers and percentages; weighted odds ratios and 95 % confidence intervals)

	Decaffeinated coffee (cups/d)											
	0		0 to <1			1 to <2			2+			
	OR		OR		95 % CI	OR		95 % CI	OR		95 % CI	$P_{trend}$
LEM (n 7699)												
Cases/participants		2582/6195		386/892			168/416			69/196		
Weighted prevalence (%)		38.4		36.1			34.3			30.9		
Age adjusted	Ref.		0.80		0.62, 1.02	0.75		0.52, 1.06	0.74		0.45, 1.19	0.237
Multivariate adjusted	Ref.		0.65		0.47, 0.89	0.73		0.50, 1.06	0.43		0.23, 0.81	0.022
GPA (n 7699)												
Cases/participants		3924/6195		594/893			263/415			131/196		
Weighted prevalence (%)		61.8		62.6			61.6			61.9		
Age adjusted	Ref.		0.92		0.67, 1.26	0.89		0.60, 1.32	1.04		0.68, 1.59	0.911
Multivariate adjusted	Ref.		0.75		0.52, 1.07	0.80		0.50, 1.27	0.67		0.40, 1.14	0.180
LSA ( <i>n</i> 7704)												
Cases/participants		1590/6199		243/893			92/416			36/196		
Weighted prevalence (%)		21.5		20.1			19.8			15.2		
Age adjusted	Ref.		0.84		0.66, 1.07	0.84		0.60, 1.17	0.67		0.34, 1.29	0.244
Multivariate adjusted	Ref.		0.64		0.50, 0.82	1.03		0.69, 1.54	0.39		0.16, 0.94	0.082
ADL ( <i>n</i> 7704)												
Cases/participants		1511/6199		232/893			97/416			41/196		
Weighted prevalence (%)		20.7		20.8			18-2			18-1		
Age adjusted	Ref.		0.93		0.72, 1.20	0.80		0.56, 1.13	0.87		0.52, 1.45	0.492
Multivariate adjusted	Ref.		0.73		0.53, 1.00	0.92		0.60, 1.39	0.57		0.30, 1.09	0.157
IADL (n 7704)												
Cases/participants		2036/6199		331/893			138/416			55/196		
Weighted prevalence (%)		28.8		31.7			29.1			24.8		
Age adjusted	Ref.		1.03		0.80, 1.33	0.93		0.66, 1.30	0.84		0.48, 1.46	0.484
Multivariate adjusted	Ref.		0.82		0.61, 1.10	0.86		0.60, 1.23	0.55		0.28, 1.06	0.107

LEM, lower extremity mobility; Ref., reference; GPA, general physical activity; LSA, leisure and social activities; ADL, activities of daily living; IADL, instrumental activities of daily living.

results provided the evidence to support the protective effect of coffee on functional disability that might be caused by the loss of skeletal muscles. Moreover, cognition is essential for planning and monitoring performance. Lower levels of cognition have been reported to be related to self-reported disability<sup>(33)</sup>. Meanwhile, coffee drinkers at midlife had better cognition later in life with respect to those drinking no or only little coffee<sup>(32)</sup>. Thus, the inverse associations of coffee in our study might be related to better cognition. Additionally, antiinflammatory effects of coffee might attenuate many healthrelated outcomes such as disability. Many previous studies have reported that coffee consumption was associated with lower levels of inflammatory markers such as IL-8<sup>(35)</sup>, fetuin-A<sup>(36)</sup> and 8-isoprostane levels<sup>(35)</sup>, which were risk factors of incident mobility limitation<sup>(37)</sup> and ADL disability<sup>(38)</sup>. The mechanism of coffee on functional disability needs to be explored by more research considering the potential hypothesis of muscle strength, diabetes and anti-inflammatory effects.

A major strength of this study is the large number of participants from the NHANES, which could generate nationally representative estimates of older American adults. Furthermore, we used five domains to measure the functional disability, including LSA which has not been investigated in the previous studies. Moreover, the correlation coefficients of two 24-h dietary recalls are 0.71, which proved the high reliability of

habitual coffee consumption in this study. In addition, we detected the dose-response relationships between total coffee consumption and functional disability. Finally, our study investigated the association of diverse type of coffee on each functional disability domain. However, there were also some limitations in our study. First, owing to the cross-sectional design of this study, the causal inferences of coffee consumption and functional disability cannot be inferred, the possibility of reverse causality also cannot be excluded. Participants may drink less coffee due to the reduced mobility (unable to prepare or buy coffee) and reduced social activities (less reason to drink coffee). Therefore, longitudinal cohort study is necessary to investigate the prospective association between coffee consumption and functional disability in older American adults in the future. Second, the coffee consumption and functional disability were self-reported, which might introduce recall bias. Finally, although numerous potential confounders have been adjusted, some residual confounding might still persist. Yet, we hope these results may stimulate interest in achieving a better understanding of the associations between coffee consumption and functional disability.

In conclusion, our results show that coffee consumption is inversely associated with the lower odds of functional disability in older American adults. The inverse associations of diverse coffee type differ across the different domains of functional



<sup>\*</sup> Adjusted for age, sex, race and ethnicity, marital status, educational level, annual household income, BMI, physical activity, cigarette smoking, alcohol drinking, energy intake, tea intake, soft drink intake, depression, hypertension, diabetes, arthritis, gout, stroke, cancer and caffeinated coffee consumption.

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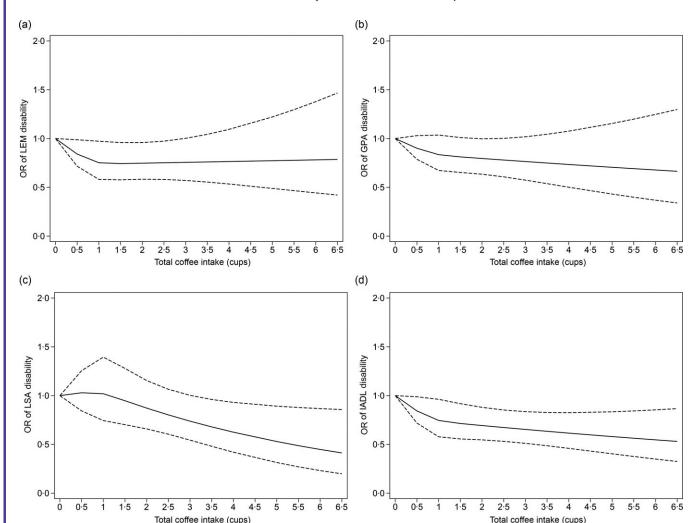


Fig. 2. (a) Examination of the dose-response relationship between total coffee consumption and lower extremity mobility (LEM) disability (Pfor non-linearity = 0.06); (b) examination of the dose-response relationship between total coffee consumption and general physical activity (GPA) disability (Pfor non-linearity = 0.36); (c) examination of the dose–response relationship between total coffee consumption and leisure and social activities (LSA) disability (P<sub>for non-linearity</sub> = 0.44) and (d) examination of the dose-response relationship between total coffee consumption and instrumental activities of daily living (IADL) disability (Pfor non-linearity = 0-11). No total coffee consumption was used as the reference group. The solid line and dashed line represent the estimated odds ratio and its 95 % confidence interval.

disability. Further research is essential to confirm these results in prospective design in older American adults and explore the mechanism of the inverse associations.

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## Supplementary material

For supplementary material referred to in this article, please visit https://doi.org/10.1017/S0007114520003153

## References

- 1. Guralnik JM & Ferrucci L (2003) Assessing the building blocks of function: utilizing measures of functional limitation. Am J Prev Med 25, 112-121.
- Erickson W, Lee C & von Schrader S (2016) 2015 Disability Status Report: United States. Ithaca, NY: Cornell University Yang Tan Institute on Employment and Disability (YTI).



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- Fried TR, Bradley EH, Williams CS, et al. (2001) Functional disability and health care expenditures for older persons. Arch Intern Med 161, 2602–2607.
- Lyyra TM, Leskinen E & Heikkinen E (2005) A cohort study found good respiratory, sensory and motor functions decreased mortality risk in older people. J Clin Epidemiol 58, 509–516.
- McGrath R, Al Snih S, Markides K, et al. (2019) The burden of functional disabilities for middle-aged and older adults in the United States. J Nutr Health Aging 23, 172–174.
- Kim J & Lee Y (2011) Frequency of dairy consumption and functional disability in older persons. J Nutr Health Aging 15, 795–800
- Yoshida D, Ohara T, Hata J, et al. (2019) Dairy consumption and risk of functional disability in an elderly Japanese population: the Hisayama Study. Am J Clin Nutr 109, 1664–1671.
- Ribeiro SM, Morley JE, Malmstrom TK, et al. (2016) Fruit and vegetable intake and physical activity as predictors of disability risk factors in African-American middle-aged individuals. J Nutr Health Aging 20, 891–896.
- Parsons TJ, Papachristou E, Atkins JL, et al. (2019) Healthier diet quality and dietary patterns are associated with lower risk of mobility limitation in older men. Eur J Nutr 58, 2335–2343.
- Grosso G, Godos J, Galvano F, et al. (2017) Coffee, caffeine, and health outcomes: an umbrella review. Annu Rev Nutr 37, 131–156.
- 11. Mirmiran P, Carlstrom M, Bahadoran Z, et al. (2018) Long-term effects of coffee and caffeine intake on the risk of pre-diabetes and type 2 diabetes: findings from a population with low coffee consumption. Nutr Metab Cardiovasc Dis 28, 1261–1266.
- 12. Rodriguez-Artalejo F & Lopez-Garcia E (2018) Coffee consumption and cardiovascular disease: a condensed review of epidemiological evidence and mechanisms. *J Agric Food Chem* **66**, 5257–5263.
- Dirks-Naylor AJ (2015) The benefits of coffee on skeletal muscle. Life Sci 143, 182–186.
- Machado-Fragua MD, Struijk EA, Ballesteros JM, et al. (2019)
   Habitual coffee consumption and risk of falls in 2 European cohorts of older adults. Am J Clin Nutr 109, 1431–1438.
- Ding M, Satija A, Bhupathiraju SN, et al. (2015) Association of coffee consumption with total and cause-specific mortality in 3 large prospective cohorts. Circulation 132, 2305–2315.
- Machado-Fragua MD, Struijk EA, Graciani A, et al. (2019) Coffee consumption and risk of physical function impairment, frailty and disability in older adults. Eur J Nutr 58, 1415–1427.
- National Center for Health Statistics & Centers for Disease Control and Prevention (2017) About the National Health and Nutrition Examination Survey. https://www.cdc.gov/ nchs/nhanes/about\_nhanes.htm (accessed October 2019).
- 18. Chen H & Guo X (2008) Obesity and functional disability in elderly Americans. *J Am Geriatr Soc* **56**, 689–694.
- Kalyani RR, Saudek CD, Brancati FL, et al. (2010) Association of diabetes, comorbidities, and A1C with functional disability in older adults: results from the National Health and Nutrition Examination Survey (NHANES), 1999–2006. Diabetes Care 33, 1055–1060.
- Egede LE (2004) Diabetes, major depression, and functional disability among U.S. adults. *Diabetes Care* 27, 421–428.

- Raper N, Perloff B, Ingwersen L, et al. (2004) An overview of USDA's dietary intake data system. J Food Compos Anal 17, 545–555.
- Xiao Q, Sinha R, Graubard BI, et al. (2014) Inverse associations of total and decaffeinated coffee with liver enzyme levels in National Health and Nutrition Examination Survey 1999–2010. Hepatology 60, 2091–2098.
- Armstrong T & Bull F (2006) Development of the World Health Organization Global Physical Activity Questionnaire (GPAQ). *Journal of Public Health* 14, 66–70.
- Gay IC, Tran DT & Paquette DW (2018) Alcohol intake and periodontitis in adults aged >/=30 years: NHANES 2009–2012. J Periodontol 89, 625–634.
- Zhang Z, Jackson S, Merritt R, et al. (2019) Association between cardiovascular health metrics and depression among U.S. adults: National Health and Nutrition Examination Survey, 2007–2014. Ann Epidemiol 31, 49–56.e42.
- Brighthaupt SC, Schneider KE, Johnson JK, et al. (2019) Trends in adolescent heroin and injection drug use in nine urban centers in the U.S., 1999–2017. J Adolesc Health 65, 210–215.
- Frenk SM, Kit BK, Lukacs SL, et al. (2016) Trends in the use of prescription antibiotics: NHANES 1999–2012. J Antimicrob Chemother 71, 251–256.
- Duncan MJ, Clarke ND, Tallis J, et al. (2014) The effect of caffeine ingestion on functional performance in older adults. J Nutr Health Aging 18, 883–887.
- Camfield DA, Silber BY, Scholey AB, et al. (2013) A randomised placebo-controlled trial to differentiate the acute cognitive and mood effects of chlorogenic acid from decaffeinated coffee. PLOS ONE 8, e82897.
- 30. Wang T, Wu Y, Li W, *et al.* (2019) Weak grip strength and cognition predict functional limitation in older Europeans. *J Am Geriatr Soc* **67**, 93–99.
- 31. Guo Y, Niu K, Okazaki T, *et al.* (2014) Coffee treatment prevents the progression of sarcopenia in aged mice *in vivo* and *in vitro*. *Exp Gerontol* **50**, 1–8.
- Eskelinen MH, Ngandu T, Tuomilehto J, et al. (2009) Midlife coffee and tea drinking and the risk of late-life dementia: a population-based CAIDE study. J Alzheimers Dis 16, 85–91.
- Buchman AS, Boyle PA, Leurgans SE, et al. (2011) Cognitive function is associated with the development of mobility impairments in community-dwelling elders. Am J Geriatr Psychiatry 19, 571–580.
- Loftfield E, Shiels MS, Graubard BI, et al. (2015) Associations of coffee drinking with systemic immune and inflammatory markers. Cancer Epidemiol Biomarkers Prev 24, 1052–1060.
- Kempf K, Herder C, Erlund I, et al. (2010) Effects of coffee consumption on subclinical inflammation and other risk factors for type 2 diabetes: a clinical trial. Am J Clin Nutr 91, 950–957.
- Wedick NM, Brennan AM, Sun Q, et al. (2011) Effects of caffeinated and decaffeinated coffee on biological risk factors for type 2 diabetes: a randomized controlled trial. Nutr. J 10, 93.
- 37. Penninx BW, Kritchevsky SB, Newman AB, et al. (2004) Inflammatory markers and incident mobility limitation in the elderly. *J Am Geriatr Soc* **52**, 1105–1113.
- 38. Sujarwoto S & Tampubolon G (2015) Inflammatory markers and physical performance in middle-aged and older people in Indonesia. *Age Ageing* **44**, 610–615.

