

Article



Correlation of Adiponectin and Leptin with Anthropometrics and Behavioral and Physical Performance in Overweight and Obese Chinese College Students

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Simple Summary: We studied a population of overweight and obese Chinese university students. Obesity, which was previously limited to the Western world, has emerged in Asia and China in recent years. Owing to the massive population of the Asian region, even effects on a small percentage of this population correspond to substantial problems. University students are an at-risk group for obesity due to the sedentary lifestyle imposed by their heavy study schedule. We measured the physical capacities, as well as several biochemical and behavioral variables, of Chinese students. We measured the blood levels of two multifunctional hormones, namely leptin (LEP) and adiponectin (ADPN), and levels of physical activity. We found that in obese students, physical capacities are linked with the hormone ADPN, which has been hypothesized to be linked with muscle development. We also found that sleep was associated with the blood levels of LEP and ADPN. To the best of our knowledge, our study is the first work to provide data on the association of LEP and ADPN with behavioral, physical, and biochemical variability in overweight and obese Asian populations. Such an association has potential for use as a comparison tool.

Abstract: The aim of this study is to assess the relationship of leptin (LEP) and adiponectin (ADPN) with other circulating fat markers, physical capacity, behaviors, and anthropometric indices in a population of overweight and obese Chinese university students. LEP and ADPN levels, as well as behavioral, anthropometric, biochemical, and performance characteristics, were measured. Method: A total of 17 anthropometric parameters, 8 questionnaires (investigating quality of life, sleep, eating, perceived functioning, stress, and depression), 9 biochemical parameters, and 12 functional parameters were investigated. Results: In contrast to previous studies, our work found an unusually strong relationship between LEP and ADPN (r = 0.961, p = 0.000) that can be related to ethnicity. We also found that LEP and ADPN were associated with stress and bodily pain. A total of 12 anthropometric measures were also associated with LEP/ADNP levels. Moreover, LEP and ADPN were found to be related to lower limb, hand, and abdominal strength; blood pressure; and basic metabolism. However, we did not find associations with sleep; eating habits; or cardiovascular fitness, which was measured in the form of resting heart rate and VO_{2max}. Conclusion: This study reveals new relationships of LEP and ADPN with selected anthropometric and behavioral parameters in obese Chinese college students.

Keywords: leptin; adiponectin; obese Chinese college students

1. Introduction

Leptin [1,2] (LEP) and adiponectin [3] (ADPN) are two adipocytokines that have been widely studied in connection with obesity [4]. LEP increases and ADPN decreases with increasing body fat. The ADPN:LEP ratio is a prognostic factor for the development of



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Copyright: © 2024 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). obesity [5], cardiovascular disease [6], insulin resistance [7], and diabetes [8], as well as several other conditions, due to the ubiquity of LEP and ADPN in the human body [1,2]. Given that they are produced by adipose tissues, plasma LEP and ADPN have been connected with different anthropometric measures in the Caucasian population, particularly with waist and hip circumference, waist:hip ratio, BF%, and body fat mass [9]. By contrast, in the Asian population, plasma LEP and ADPN are connected with abdominal fat [10–14]. The relationship of LEP and ADPN with anthropometric measures is of clinical interest because it may aid in the diagnosis of hyperleptinemia, which has been widely explored [15,16]. Considering the proximity of LEP receptors to neuropeptide Y receptors in the hypothalamus [17], LEP has been studied in connection with anxiety, stress [18–20], depression [21–23], sleep [24], and eating behaviors [23,25]. LEP and ADPN levels have also been associated with behavioral factors, such as physical activity levels [25,26]. Although LEP and ADPN, as ergogenic molecules, have been suggested as markers of training status [27] in athletes, only a few studies on the possible link of LEP with the capacity for physical performance exist. In children, LEP is inversely correlated with the amount of physical activity [28] and positively correlated with VO_{2max} [29]. However, the relationships of LEP and ADPN with physical performance in obese adults have been poorly investigated [30]. To the best of our knowledge, no studies exist on the relationship of LEP and ADPN with basic physical performance (lower limb, hand, and abdominal muscle strength; balance; and flexibility) in overweight/obese Asian college students. Furthermore, studies investigating behavioral, physical, biochemical, and anthropometric indices in the same cohort of overweight and obese subjects do not exist.

As suggested by the literature [10–14], the associations of LEP and ADPN with anthropometrics in Asians are different from those in Caucasians. Comparing different populations within the same BMI range revealed that ADPN concentrations were lower in the Indonesian population than in the Dutch population [31]. In obese Caucasians, LEP and ADPN showed no or low correlations with each other [32]. LEP has been hypothesized to be connected with different behaviors [33]. It regulates food intake, reward, and motivation. Reduced LEP impairs cognition [33].

To the best of our knowledge, only a few studies have linked LEP and ADPN with behavioral characteristics in Chinese populations. Behavior and LEP levels have been studied in Chinese schoolchildren. LEP was found to be associated with low math scores and high verbal scores. Low ADPN was found to be correlated with family-related behavioral problems [34]. LEP has been proposed as a biomarker of sleep disorders given that it acts as a mediating factor between obesity and poor sleep [35]. High ADPN and low LEP levels were associated with anxiety [36] and stress [37]. Therefore, our aim is to assess the relationship of LEP and ADPN with other circulating fat markers, physical capacity, behaviors, and anthropometric indices in a population of overweight and obese Chinese university students.

2. Materials and Methods

2.1. Participants

This study was a randomized, controlled trial. Obese and overweight adults were recruited from Tongji University through workshops, leaflets, social media, official web pages, and paper bulletins disseminated on campus.

The ethics committee of Tongji University approved all interventions (tjdxsr046). A total of 40 respondents were invited to visit the research department. Prior to the collection of eligibility data, researchers carefully explained the purpose and protocol of the study in detail. Prior to registration, participants signed a written informed consent letter. All participants completed an initial screening by filling out questionnaires on background information (including brief sedentary questions, medical history, current health status, a food intake questionnaire [FFQ25 [38]], the International Physical Activity Questionnaire [IPAQ [39]], and a physical examination). A total of 8 participants did not meet the inclusion criterion of sedentary obesity, and out of the 32 participants who were invited to enroll in

this study, 6 refused. This study included 26 subjects (3 females) from different colleges of Tongji University in Shanghai (age 19.23 ± 1.39 years) (Table 1). Obesity was defined in accordance with Chinese reference values [40] as BMI > 28.

Table 1. Anthropometrics, biochemical, cardiorespiratory indices, and functional measures (mean and standard deviation).

	Mean	SD	Range
Anthropometrics			
Height (cm)	176.92	± 6.62	167.00-180.7
Weight (kg)	86.75	± 20.46	65.60-142.10
BMI	29.28	± 5.48	24.100-40.40
Fat-free mass (kg)	61.43	± 13.60	41.30-87.90
Fat mass (kg)	26.75	± 11.81	12.10-61.10
Muscle mass (kg)	58.4	± 425	54.80-53.10
Body fat %	24.2	± 6.53	17.30-30.30
Waist:hip ratio	0.94	± 0.04	0.91-0.98
Chest circum (cm)	95.143	± 19.73	101-123.00
Waistline (cm)	93	± 13.12	78.50-129.00
Minimum waist circum (cm)	88.98	± 10.16	73.20-116.00
Hipline (cm)	99.98	± 20.87	91.00-106.00
Thigh circum (cm)	62.9	± 7.16	53.00-78.00
Calf circum (cm)	41.75	± 4.14	36.00-52.00
Biceps contract circum (cm)	32.07	± 4.32	24.4-44.00
Biceps relaxed circum (cm)	30.91	± 3.42	24.80-37.00
Abdominal skinfold (mm)	48.90	± 19.64	19.50-75.00
Shoulder crease skinfold (mm)	33.89	± 10.97	13.00-53.00
Biceps skinfold (mm)	39.51	± 13.75	12.00-73.00
Thigh crease skinfold (mm)	50.14	± 19.26	8.00-75.00
Iliac skinfold (mm)	24.53	± 12.52	5.00-51.00
Calf skinfold (mm)	34.093	± 13.81	12.00-56.00
Biochemical parameters			
TC (mM/L)	4.03	± 0.92	1.87-5.67
TG (mM/mL)	1.14	± 0.57	0.27-2.84
HDL (mM/L)	1.35	± 0.25	0.79-1.85
LDL (mM/L)	2.59	± 0.78	0.70-4.28
Glucose (mM/L)	4.76	± 0.34	3.88-5.59
LEP (ng/mL)	3.83	± 0.82	2.50 - 5.42
ADPN (mg/L)	8.69	± 1.82	5.58-12.31
ADPN/LEP ratio	2.30	± 0.11	2.07 - 2.48
HGB (mM/L)	82.66	± 8.44	66.55-100.38
HbA1c (mM/L)	32.28	± 16.14 19.72–98.49	
Cardiorespiratory indices			
Basal VO ₂ (L/min)	1822	± 157.04	1688–1995
$VO_2 \max (L/min)$	3374.27	± 1016.87	1222-5889
$VO_2 \max (mL/Kg/min)$	40.46	± 5.88	30.90-53.40
METS	11.71	± 2.47	4.40-17.50
Physical performance parameters			
Systolic pressure (mmHg)	128.96	± 22.35	95.00-157.00
Diastolic pressure (mmHg)	76.38	± 10.01	63.00–97.00
Resting heart rate (beats/min)	80.44	± 13.73	54.00-101.00
Hand grip strength (right hand) (kg)	37.64	±9.23	19.30-61.90
Vital capacity (mL)	3878.69	± 913.54	2674-7140
Sit-and-reach (cm)	+14.05	± 8.58	-2.90-(+)29.00
I min sit-ups (number)	30.81	±6.22	18.00-45.00
Standing long jump (cm)	195.71	2 ± 5.71	144.00-243.00
1 min squats (number)	48.38	9 ± 0.71	31.00-65.00
Standing on one leg with eyes closed (s)	33.95	± 32.37	1.20–131.53

BMI: body mass index; TC: total cholesterol; TG: triglyceride; HDL: high-density lipoprotein; LDL: low-density lipoprotein cholesterol; HGB: hemoglobin; LEP: leptin; ADPN: adiponectin. In the sit-and-reach test, 0 is the feet line, + is beyond the feet line, - is before the feet line.

2.2. Inclusion and Exclusion Criteria

The inclusion criteria were (1) being apparently sedentary in accordance with living a sedentary lifestyle (spending more than 8 h awake in a sitting position daily and continuously sitting still for over 90 min); (2) age between 18 and 25 years; (3) BMI \geq 25; (4) has not engaged in regular exercise for more than 30 min for 3–5 months; and (5) IPAQ score < 600 metabolic equivalents min/week. Out of 26 subjects, 15 were obese (BMI > 28).

The exclusion criteria included (1) severe cardiovascular or skeletal muscle disease; (2) diabetes complications (such as postexercise hypoglycemia or autonomic neuropathy or peripheral neuropathy); (3) diagnosed with mental illnesses; and (4) other sports/exercise contraindications (diagnosed with heart problems). All the students regularly consumed their meals at the campus canteen.

2.3. Measurements

2.3.1. Anthropometric Measures

A total of 16 anthropometric measures were taken in accordance with standard procedures [41]. In total, 11 circumferences (chest; minimum and maximal waist line; contracted and relaxed hip, thigh, calf, and biceps brachii), and 6 skinfolds (shoulder and tight crease, biceps, abdominal, iliac, and calf) were measured. Weight, fat, fat-free and muscle masses, and body fat% were measured with a X-SCAN PLUS (Selvas, Gyeong, Republic of Korea) bioelectric impedance scale. Height was measured by using a Hengkang Jiaye (Shenzhen, China) anthropometer. Physical tests, including 1 min sit-ups and squats, standing on one leg with eyes closed, and the sit-and-reach test, were performed in accordance with ACSM guidelines [42].

2.3.2. Biochemical Parameters

Biochemical parameters were measured in duplicate after overnight fasting for at least 10 h. The following parameters were measured with the fully automated biochemical analyzer Dimension EXL 200, (Siemens Healthineers, Munchen, Germany): total cholesterol, triglycerides, high-density lipoprotein cholesterol, low-density lipoprotein, glucose, hemoglobin, and HbA1c. ADPN and LEP were quantified by using an E-EL-H0004-96 T human ADP/Acrp30 (ADPN) ELISA kit (BLKAZ5BYNR, Elabscience, Wuhan, China) and E-EL-H0113-96 T human LEP ELISA kit (EKF8PFBUAG, Elabscience, Wuhan, China) in accordance with the manufacturer's instructions.

2.3.3. Cardiorespiratory Indices

Basal VO₂, VO_{2max}, and basal metabolism (METS) were measured on the basis of the mean of the FatMax test results. FatMax is the optimal heart rate for fat utilization [43]. Subjects were asked to refrain from vigorous exercise for 24 h and fast for 10 h before the test. All FatMax tests were conducted in the morning between 8 a.m. and 12 a.m. to avoid circadian variance. A graded treadmill exercise test was used to perform the FatMax test. In brief, a walking warm-up was performed at 3.0 km/h with an incline of 1% for 3 min. Subsequently, the first stage of exercise was performed at a speed of 6.0 km/h for 2 min. Each increase in speed had a duration of 2 min. The protocol was modified to adapt to the physical capacities of the sample from a previous study employing FatMax on obese adults [44]. The speeds of the following steps were 7.0, 8.0, 9.0, and 10 km/h until the respiratory exchange ratio of 1.15 was reached or the subject became exhausted. VO₂ and VCO₂ were measured by using a portable metabolic system (K5, COSMED, Rome, Italy) and Polar H10 (Polar Electro, Tampere, Finland) heartrate band. The test was stopped immediately if the subject showed symptoms of dizziness, nausea, or dyspnea during the test. Vital capacity was measured with an electronic activity tester (Hengkang Jiaye, Shenzhen, China).

2.3.4. Physical Performance Parameters

Diastolic and systolic blood pressure levels were measured with a standard sphygmomanometer, and resting heart rate was measured with a Polar H10 HR monitor. The grip strength of both hands was measured with a Jamar dynamometer. Progressive maximal contractions were performed for both hands, and the best value of 3 attempts was retained. All subjects were right-handed (Table 1).

2.3.5. Scale Evaluation

SF36 [45] quality of life; the IPAQ scale [39], which was also validated for blood pressure [46]; HAMA for anxiety and depression [47,48] (mental and body anxiety and total scores); PSS for perceived stress levels [49]; SDS for anxiety [50]; TFEQR21 for eating behaviors [51,52]; FS14 [53] (physical and mental fatigue and total scores); and PSQI [54] for the assessment of sleep were used. The SF-36 measures 8 scales: physical functioning (PF), physical role, bodily pain (BP), general health, vitality, social functioning (SF), emotional role, and mental health. Questionnaires were submitted in the form of the Chinese version validated in previous studies [45–54]. All measurements were performed in the morning, and the testing order was randomized between subjects for each test by using a random number sorter.

2.3.6. Statistical Analysis

Descriptive statistic and correlation analyses (Spearman) were performed by using SPSS IBM, SPSS Statistics 25. Levine and Shapiro–Wilk tests were used to check the homogeneity of variance and normality of data, respectively. All data were normally distributed. The extracted data included the mean and SD of participant characteristics, cardiorespiratory indices, and questionnaire scores.

3. Results

3.1. Sample Descriptive Statistics

Table 1 reports the means and standard deviations of the observed variables.

3.2. Questionnaires Scores

The questionnaire results of overweight and obese Chinese college students are reported in Table 2.

	Mean	SD	Range	
SF-36				
PF	66.12	± 39.85	70-100	
RP	77.93	± 17.61	25-100	
BP	52.38	± 13.54	41-90	
GH	65.2	± 21.19	30-92	
VT	82.98	± 21.45	15-90	
SF	41.51	± 32.35	44.44-100.00	
RE	67.28	± 21.08	33-100	
MH	56.01	± 30.58	32–96	
HT	54.46	± 25.51	25-100	
IPAQ				
JOB	1057	± 1024.55	330-3816	
Transport	1083.16	± 1029.71	198–3312	
House work	139.82	± 231.91	60-1080	
Recovery	714.17	± 1026.03	23-3876	
MET	2642.8	± 1864.69	442.2-8544.0	
Walk	1402.63	± 1447.35	82.5-5841.0	
Medium intensity	991.96	± 760.62	60-2940	
High intensity	779.43	± 653.57	120-2520	

Table 2. Questionnaires scores (mean and standard deviation).

	Mean	SD	Range
PSS	43.85	± 4.24	34–56
SDS	29.46	± 6.87	20-47
TFEQR-21			
Uncontrolled eating	18.46	± 4.90	11–31
Restrain	15.07	± 4.89	8–26
Emotional eating	12.30	± 5.88	4-25
FS-14			
Physical fatigue	3.73	± 2.06	2-8
Mental fatigue	2.57	± 1.83	1–6
Total score	7.30	± 3.26	1–12
PSQI	5.5	± 2.70	1–13
HAMA			
Mental anxiety	5.84	± 4.64	0–16
Body anxiety	4.69	± 2.71	0-12
Total score	10.53	± 6.15	2–2618

Table 2. Cont.

SF-36: The 36-Item Short Form Health Survey questionnaire; PF: physical function; RP: physical role; BP: bodily pain; GH: general health; VT: vitality; SF: social functioning; RE: emotional role; MH: mental health; HT health changes in the past year; PSS: perceived stress scale; SDS: stress depression scale; IPAQ: international physical activity scale; TFEQR-21: Three Factor Eating Questionnaire-R21; FS-14: Fatigue Scale-14; PSQI: Pittsburgh Sleep Quality Index; HAMA: Hamilton Anxiety Scale.

The mean PSQI score of our sample was within the range of normal sleep (0–5 points). However, we observed 10 subjects with a score between 6 and 10 (mild insomnia) and 1 subject with moderate insomnia (>10). The HAMA total cut-off point for anxiety disorders was 14. The mean of our subjects was below the anxiety threshold (6 subjects were beyond the threshold). The maximum FS14 fatigue score was 14. Our group average showed that our subjects, with the exception of 8 subjects who scored >10, did not perceive considerable fatigue. Our sample had a score of <50 on the SDS scale (no depression symptoms).

3.3. Correlations among Measured Variables

Table 3 reports the results of the correlation analysis. In contrast to several other studies [32,55–57], our work found that LEP and ADPN were strongly correlated with each other (r = 0.961, p = 0.000). As expected, LEP and ADPN were correlated with all other biochemical indices of fat metabolism that we measured (Table 3), with LEP showing an inverse correlation and ADPN a strong positive correlation with total cholesterol. LEP is correlated with LDL and HbA1c, and LEP and ADPN were correlated with HGB. In agreement with another study on obesity that found a CC of 0.670 (p = 0.0001) [58], this work discovered that ADPN was correlated with HDL (Table 3). ADPN showed a correlation with BMI; height; hip, bicep (contracted and relaxed), waist, and calf circumferences; shoulder crease skinfolds; and waist:hip ratio and muscle mass. LEP presented correlations with the previous variables, except for muscle mass, fat-free mass, waist circumferences, and shoulder crease skinfold. However, in contrast to ADPN, LEP showed a correlation with thigh circumference (Table 2). ADPN was correlated with muscle and fat-free masses, 1 min squat (reps), long jump, 1 min sit-up (reps), and hand grip but not with sit-and-reach test results and body balance. LEP exhibited the same correlations with physical performance (Table 2).

Although previous studies also observed a correlation between LEP and blood pressure in obese individuals with high blood pressure [59], our sample did not appear to have hypertension. METS was also correlated with LEP and ADPN. LEP was correlated with the total score on the PSQI questionnaire and with the PF and BP subscales of SF36. LEP and ADPN were correlated with the depression and stress scales, as well as with the BP and physical function scales (Table 3).

		LEP		ADPN	
		r	р	r	р
Biochemical parameters	Glucose	0.607	0.001	0.626	0.000
	TC	-0.501	0.000	0.812	0.000
	TG	0.377	0.006	0.547	0.003
	HDL	n.s.	n.s.	0.498	0.007
	LDL	0.602	0.000	n.s	n.s.
1	HGB	0.709	0.000	0.545	0.003
	HbA1c	0.283	0.042	n.s.	n.s.
	BMI	0.419	0.026	0.438	0.02
	Hip circum	0.487	0.009	0.471	0.11
	Height	0.531	0.004	0.549	0.002
	Muscle mass	n.s.	n.s.	0.388	0.041
	Fat-free mass	n.s.	n.s.	0.39	0.04
	Biceps control	0.414	0.016	0.474	0.011
Anthropometry	Biceps relaxed	0.491	0.008	0.502	0.006
	Calf circum	0.5	0.007	0.504	0.006
	Shoulder crease skin	n.s.	n.s.	0.384	0.044
	Waist:hip	0.585	0.001	0.608	0.001
	Waist circum	n.s.	n.s.	0.52	0.005
	Thigh circum	0.533	0.003	n.s.	n.s.
Functional indices	1 min-squat (reps)	0.375	0.049	0.416	0.028
	Long jump	0.45	0.016	0.49	0.008
	Hand grip	0.347	0.071	0.398	0.036
	1 min sit-ups	0.362	0.058	0.389	0.041
	Systolic pressure	0.581	0.002	0.599	0.001
	Diastolic pressure	0.597	0.001	0.575	0.002
	METS	0.416	0.028	0.428	0.023
	SDS total	0.47	0.012	0.486	0.009
Questionnaires	PSS total	n.s.	n.s.	0.538	0.003
	SF36/bp	0.546	0.003	0.546	0.003
	SF36/pf	0.379	0.047	0.416	0.028
	PSOI	0.288	0.039	0.538	0.003

Table 3. Correlations of LEP and ADPN with biochemical parameters, anthropometry, functional indices, and questionnaires.

TC: total cholesterol; TG: triglyceride; HDL: high-density lipoprotein; LDL: low-density lipoprotein cholesterol; HGB: hemoglobin; SDS: stress depression scale; PSS: perceived stress scale; SF36/bp: bodily pain; SF36/pf: physical functioning; PSQI: Pittsburgh Sleep Quality Index. n.s. = nonsignificant.

4. Discussion

Low (or decreased) levels of LEP and high (or increased) levels of ADPN are beneficial in terms of reducing weight and inflammation and desirable in overweight and obese individuals [3]. The present study aims to explore the relationships between LEP and ADPN in obese Chinese university students. Considering the absence of comprehensive studies exploring the relationships of LEP and ADPN with the anthropometric indices and behavioral and physical performances of Chinese college students, our present study is the first work to provide such information. LEP and ADPN are inversely correlated in obesity. The major finding of our study is a quasiperfect (r = 0.961, p = 0.000) correlation between LEP and ADPN. This result is surprising because LEP and ADPN have an inverse relationship in humans [32,55–57,60]. Given that we did not find any study on a similar cohort (age and ethnicity), we can hypothesize that this result can be attributed to diet. In fact, all of our subjects attend the same university and may share common dietary factors that influence ADPN levels. Although our sample size is relatively small, the subjects are highly homogeneous, all being college students of the same age and gender (with the exception of three females). They also consume their meals at the same place and sleep in dormitories under similar conditions. We can hypothesize that some factors can influence the strong correlation that we found between LEP and ADPN. For example, the consumption of green tea is common among Chinese students and could reduce LEP and increase ADPN considerably [61]. Moreover, the students live in Shanghai, wherein the local diet is rich in fish omega-3, which can also reduce LEP [62]. Another possible explanation for the strong correlation between LEP and ADPN may be the distribution of body fat in the Asian biotype [4]. Such a distribution is associated with the differing secretion of LEP and ADPN in the blood stream. Asians accumulate more visceral fat tissue and less subcutaneous fat than Caucasians. Although the association of LEP and ADPN with muscle mass (fat-free mass) and lower limb strength was previously observed in young tennis players [63] and Chinese high school students [64], it remains controversial. A previous work found that LEP was positively associated with muscle density (it would decrease skeletal muscle lipid content and promote lipid oxidation) [65], whereas another study discovered that LEP was negatively correlated with muscle density and relative hand grip strength [66]. In contrast to a study on young Caucasians [66] and Afro-Americans [67], our work found that hand strength was positively correlated with LEP. Our study also discovered a relationship between LEP/ADPN and abdominal strength. The correlation with muscle mass and thus with strength can be explained by the involvement of ADPN in the phosphorylation of adenosine monophosphate, which acts as a signaling factor in skeletal muscles [68].

We investigated the behavioral characteristics associated with LEP/ADNP. Although an association exists between feeding and LEP levels [69], we did not find such an association in our subjects. The relationship between high levels of LEP and sleep found in previous studies [70] could not be observed in our work. The LEP level in our sample was five times lower than that normally present in the obese Caucasian population [71]. Although a certain degree of this difference can be explained by the sensitivity of measurement methods, our Chinese samples had very low LEP levels. Our work found that LEP and ADPN were associated with depression and stress, confirming the findings in the Caucasian population [72]. Given that LEP and ADPN are also ergogenic [73], their association with the PF index measured by using the SF questionnaire is unsurprising. We hypothesize that the association with BP can be attributed to LEP-associated inflammation in obese individuals, as shown in a previous study [74]. However, the association of ADPN with pain remains under debate [75]. These factors could have contributed to the low levels of LEP and strong association between LEP and ADPN in our study. We discovered a correlation between ADPN and selected body dimensions (BMI; height; hip, biceps (contracted and relaxed), waist, and calf circumferences; shoulder crease skinfolds; waist:hip ratio and muscle mass). This result is in agreement with the hypertrophic role of ADPN [76]. In fact, ADPN was also correlated with muscle and fat-free masses, 1 min squat (reps), long jump, 1 min sit-up (reps), and hand grip but not with sit-and-reach test results and body balance. LEP presented correlations with previous variables, except for muscle mass, fat-free mass, waist circumferences, and shoulder crease skinfold. However, in contrast to ADPN, LEP showed a correlation with thigh circumference in agreement with the secretion of LEP by subcutaneous fat tissues [1]. A limitation of our study is that we did not measure visceral fat, which is highly correlated with LEP and inversely correlated with ADPN. In addition, diet was not assessed in detail, and we did not include a control group (which could have been helpful in strengthening the results). Our sample is also unbalanced in gender distribution. We are aware that our number of participants is limited and imbalanced. However, the incidence of obesity in China is considerably lower than in Western countries, and our sample included almost all of the obese students of the campus.

5. Conclusions

Our study found new relationships of LEP and ADPN with selected anthropometric and behavioral variables in obese Chinese college students. These correlations are different from those observed previously in the Caucasian population and deserve further investigation. We hypothesize that our results can be attributed to differences in diets. The relationships of LEP and ADPN with anthropometric and biochemical indices, performance, and mood indicate that LEP/ADPN has a relationship with selected anthropometric indices, mood states, and physical performance in overweight and obese Chinese college students.

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Institutional Review Board Statement: This study was conducted in accordance with the Declaration of Helsinki, and approved by the Institutional Ethics Committee) of Tongji University (protocol code 2021tjdx024 of 2 March 2021).

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: Due to privacy reasons, the datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

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Conflicts of Interest: The authors declare that they have no competing interests.

References

- 1. Friedman, J.M.; Halaas, J.L. Leptin and the regulation of body weight in mammals. Nature 1998, 395, 763–770. [CrossRef]
- 2. Zhang, Y.; Chua, S., Jr. Leptin function and regulation. *Compr. Physiol.* 2011, *8*, 351–369.
- 3. Fang, H.; Judd, R.L. Adiponectin regulation and function. *Compr. Physiol.* 2011, *8*, 1031–1063.
- 4. Aisike, G.; Kuerbanjiang, M.; Muheyati, D.; Zaibibuli, K.; Lv, M.X.; Han, J. Correlation analysis of obesity phenotypes with leptin and adiponectin. *Sci. Rep.* **2023**, *13*, 17718. [CrossRef] [PubMed]
- Zurita-Cruz, J.N.; Villasís-Keever, M.A.; Manuel-Apolinar, L.; Damasio-Santana, L.; Garrido-Magaña, E.; Rivera-Hernández, A.D.J. Leptin/adiponectin ratio as a prognostic factor for increased weight gain in girls with central precocious puberty. *Front. Endocrinol.* 2023, 14, 1101399. [CrossRef]
- 6. Hou, N.; Luo, J.D. Leptin and cardiovascular diseases. Clin. Exp. Pharmacol. Physiol. 2011, 38, 905–913. [CrossRef] [PubMed]
- Vatier, C.; Barbara, A.P.; Fellahi, S.; Vigouroux, C.; Capeau, J.; Bastard, J.P. Le rapport adiponectine sur leptine, un biomarqueur d'insulino-résistance et de risque cardiométabolique encore méconnu. *Ann. Biol. Clin.* 2020, 78, 265–268.
- Ye, Y.; Wu, P.; Wang, Y.; Yang, X.; Ye, Y.; Yuan, J.; Pan, A. Adiponectin, leptin, and leptin/adiponectin ratio with risk of gestational diabetes mellitus: A prospective nested case-control study among Chinese women. *Diabetes Res. Clin. Pract.* 2022, 191, 110039. [CrossRef]
- 9. Marques-Vidal, P.; Bochud, M.; Paccaud, F.; Mooser, V.; Waeber, G.; Vollenweider, P. Distribution of plasma levels of adiponectin and leptin in an adult Caucasian population. *Clin. Endocrinol.* **2010**, *72*, 38–46. [CrossRef]
- 10. Conroy, S.M.; Chai, W.; Lim, U.; Franke, A.A.; Cooney, R.V.; Maskarinec, G. Leptin, adiponectin, and obesity among Caucasian and Asian women. *Mediat. Inflamm.* 2011, 253580. [CrossRef]
- Guan, Y.; Zuo, F.; Zhao, J.; Nian, X.; Shi, L.; Xu, Y.; Wu, B. Relationships of adiponectin to regional adiposity, insulin sensitivity, serum lipids, and inflammatory markers in sedentary and endurance-trained Japanese young women. *Front. Endocrinol.* 2023, 14, 1097034. [CrossRef] [PubMed]
- 12. Sun, C.; Kovacs, P.; Guiu-Jurado, E. Genetics of body fat distribution: Comparative analyses in populations with European, Asian and African ancestries. *Genes* **2021**, *12*, 841. [CrossRef] [PubMed]
- Lear, S.A.; Humphries, K.H.; Kohli, S.; Chockalingam, A.; Frohlich, J.J.; Birmingham, C.L. Visceral adipose tissue accumulation differs according to ethnic background: Results of the Multicultural Community Health Assessment Trial (M-CHAT). *Am. J. Clin. Nutr.* 2007, *86*, 353–359. [CrossRef] [PubMed]
- 14. Tanaka, S.; Horimai, C.; Katsukawa, F. Ethnic differences in abdominal visceral fat accumulation between Japanese, African-Americans, and Caucasians: A meta-analysis. *Acta Diabetol.* **2003**, *40*, s302–s304. [CrossRef] [PubMed]
- Cicchella, A.; Jurimae, T.; Stefanelli, C.; Purge, P.; Saar, M. Correlations of skinfold thicknesses and circumferences at exactly defined body sites with leptin in 10–12-year-old boys with different BMIs. *Coll. Antropol.* 2014, 38, 459–465. [PubMed]
- Wada, N.; Hirako, S.; Takenoya, F.; Kageyama, H.; Okabe, M.; Shioda, S. Leptin and its receptors. J. Chem. Neuroanat. 2014, 61, 191–199. [CrossRef] [PubMed]
- 17. Roubos, E.W.; Dahmen, M.; Kozicz, T.; Xu, L. Leptin and the hypothalamo-pituitary–adrenal stress axis. *Gen. Comp. Endocrinol.* **2012**, 177, 28–36. [CrossRef] [PubMed]
- 18. Jin, H.; Meyer, J.M.; Mudaliar, S.; Jeste, D.V. Impact of atypical antipsychotic therapy on leptin, ghrelin, and adiponectin. *Schizophr. Res.* **2008**, *100*, 70–85. [CrossRef]

- Otsuka, R.; Yatsuya, H.; Tamakoshi, K.; Matsushita, K.; Wada, K.; Toyoshima, H. Perceived Psychological Stress and Serum Leptin Concentrations in Japanese Men. *Obesity* 2006, 14, 1832–1838. [CrossRef]
- 20. Tomiyama, A.J. Stress and obesity. Annu. Rev. Psychol. 2019, 70, 703–718. [CrossRef]
- 21. Carvalho, A.F.; Rocha, D.Q.; McIntyre, R.S.; Mesquita, L.M.; Koehler, C.A.; Hyphantis, T.N.; Berk, M. Adipokines as emerging depression biomarkers: A systematic review and meta-analysis. *J. Psychiatr. Res.* **2014**, *59*, 28–37. [CrossRef] [PubMed]
- Scott, S.R.; Millwood, S.N.; Manczak, E.M. Adipocytokine correlates of childhood and adolescent mental health: A systematic review. Dev. Psychobiol. 2023, 65, e22379. [CrossRef] [PubMed]
- 23. Comeras, L.B.; Herzog, H.; Tasan, R.O. Neuropeptides at the crossroad of fear and hunger: A special focus on neuropeptide Y. *Ann. N. Y. Acad. Sci.* **2019**, 1455, 59–80. [CrossRef] [PubMed]
- 24. Beccuti, G.; Pannain, S. Sleep and obesity. Curr. Opin. Clin. Nutr. Metab. Care 2011, 14, 402–412. [CrossRef] [PubMed]
- 25. Mota, G.R.D.; Zanesco, A. Leptin, ghrelin, and physical exercise. Arq. Bras. Endocrinol. Metabol. 2007, 51, 25–33. [CrossRef]
- 26. Kraemer, R.R.; Chu, H.; Castracane, V.D. Leptin and exercise. Exp. Biol. Med. 2002, 227, 701–708. [CrossRef] [PubMed]
- Voss, S.C.; Nikolovski, Z.; Bourdon, P.C.; Alsayrafi, M.; Schumacher, Y.O. The effect of cumulative endurance exercise on leptin and adiponectin and their role as markers to monitor training load. *Biol. Sport* 2016, 33, 23–28. [PubMed]
- Cicchella, A.; Stefanelli, C.; Jürimäe, T.; Saar, M.; Purge, P. Moderate physical activity correlates with elevated leptin in physically active 10–12-year-old boys with normal BMI. *Percept. Mot. Skills* 2013, 117, 358–366. [CrossRef] [PubMed]
- Gullu, E.; Gullu, A.; Düzova, H.; Ozgor, B.İ.L.G.E.; Kilinç, E.; Akçinar, F. The relationship between serum leptin and VO2_{max} levels in pre-puberty swimmer girls: Effect of acute exercise. *Prog. Nutr.* 2020, 22, 177–184.
- van Niekerk, E.; Botha-Le Roux, S.; Mels, C.M.; Swanepoel, M.; Delles, C.; Welsh, P.; Kruger, R. Twenty-four-hour ambulatory, but not clinic blood pressure associates with leptin in young adults with overweight or obesity: The African-PREDICT study. *Hypertens. Res.* 2024, 47, 478–486. [CrossRef]
- 31. Sigit, F.S.; Trompet, S.; Tahapary, D.L.; Sartono, E.; van Dijk, K.W.; Yazdanbakhsh, M.; de Mutsert, R. The associations of leptin and adiponectin with the metabolic syndrome in an Indonesian and a Dutch population. *Nutr. Metab. Cardiovasc. Dis.* **2021**, *31*, 2426–2435. [CrossRef] [PubMed]
- Baratta, R.; Amato, S.; Degano, C.; Farina, M.G.; Patanè, G.; Vigneri, R.; Frittitta, L. Adiponectin relationship with lipid metabolism is independent of body fat mass: Evidence from both cross-sectional and intervention studies. *J. Clin. Endocrinol. Metab.* 2004, 89, 2665–2671. [CrossRef] [PubMed]
- 33. Morrison, C.D. Leptin signaling in brain: A link between nutrition and cognition? *Biochim. Biophys. Acta Mol. Basis Dis.* 2009, 1792, 401–408. [CrossRef] [PubMed]
- Li, G.; Feng, D.; Qu, X.; Fu, J.; Wang, Y.; Li, L.; Gao, S. Role of adipokines FGF21, leptin and adiponectin in self-concept of youths with obesity. *Eur. Neuropsychopharmacol.* 2018, 28, 892–902. [CrossRef] [PubMed]
- 35. Pan, W.; Kastin, A.J. Leptin: A biomarker for sleep disorders? Sleep Med. Rev. 2014, 18, 283–290. [CrossRef] [PubMed]
- 36. Wu, Y.T.; Huang, W.Y.; Kor, C.T.; Liu, K.H.; Chen, T.Y.; Lin, P.T.; Wu, H.M. Relationships between depression and anxiety symptoms and adipocyte-derived proteins in postmenopausal women. *PLoS ONE* **2021**, *16*, e0248314. [CrossRef] [PubMed]
- 37. Bouillon-Minois, J.B.; Trousselard, M.; Thivel, D.; Benson, A.C.; Schmidt, J.; Moustafa, F.; Dutheil, F. Leptin as a biomarker of stress: A systematic review and meta-analysis. *Nutrients* **2021**, *13*, 3350. [CrossRef] [PubMed]
- 38. Overvad, K.I.M.; Jønneland, A.T.; Haraldsdóttir, J.; Ewertz, M.; Jensen, O.M. Development of a semiquantitative food frequency questionnaire to assess food, energy and nutrient intake in Denmark. *Int. J. Epidemiol.* **1991**, *20*, 900–905. [CrossRef] [PubMed]
- Craig, C.L.; Marshall, A.L.; Sjöström, M. International physical activity questionnaire: 12-country reliability and validity. *Med. Sci.* Sports Exerc. 2003, 35, 1381–1395. [CrossRef]
- Guidelines for the Prevention and Control of Overweight and Obesity in Chinese Adults (Trial). Issued by the Department of Disease Control and Prevention of the National Health and Family Planned parenthood Commission. 2003. Available online: https://pubmed.ncbi.nlm.nih.gov/15807475/ (accessed on 21 May 2024).
- Norton, K. Standards for Anthropometry Assessment. In *Kinanthropometry and Exercise Physiology*; Routledge: New York, NY, USA, 2018; pp. 68–137.
- 42. ACSM's Guidelines for Exercise Testing and Prescription, 11th ed.; Lippincott Williams & Wilkins: Philadelphia, PA, USA, 2021.
- 43. Tan, S.; Wang, J.; Cao, L. Exercise training at the intensity of maximal fat oxidation in obese boys. *Appl. Physiol. Nutr. Metab.* **2016**, 41, 49–54. [CrossRef]
- 44. Wang, Y.; Guo, X.; Liu, L.; Xie, M.; Lam, W.K. Effects of Tai-chi and running exercises on cardiorespiratory fitness and biomarkers in sedentary middle-aged males: A 24-week supervised training study. *Biology* **2022**, *11*, 375. [CrossRef] [PubMed]
- Lins, L.; Carvalho, F.M. SF-36 total score as a single measure of health-related quality of life: Scoping review. SAGE Open Med. 2016, 4, 2050312116671725. [CrossRef] [PubMed]
- Alomari, M.A.; Keewan, E.F.; Qhatan, R.; Amer, A.; Khabour, O.F.; Maayah, M.F.; Hurtig-Wennlöf, A. Blood pressure and circulatory relationships with physical activity level in young normotensive individuals: IPAQ validity and reliability considerations. *Clin. Exp. Hypertens.* 2011, 33, 345–353. [CrossRef] [PubMed]
- 47. Thompson, E. Hamilton rating scale for anxiety (HAM-A). Occup. Med. 2015, 65, 601. [CrossRef] [PubMed]
- Leung, C.M.; Wing, Y.K.; Kwong, P.K.; Shum, A.L.K. Validation of the Chinese-Cantonese version of the Hospital Anxiety and Depression Scale and comparison with the Hamilton Rating Scale of Depression. *Acta Psychiatr. Scand.* 1999, 100, 456–461. [CrossRef] [PubMed]

- Meng, R.; Li, J.; Wang, Z.; Zhang, D.; Liu, B.; Luo, Y.; Yu, C. The Chinese version of the Perceived Stress Questionnaire: Development and validation amongst medical students and workers. *Health Qual. Life Outcomes* 2022, 18, 70. [CrossRef] [PubMed]
- 50. Sepehry, A.A. Self-rating depression scale (SDS). In *Encyclopedia of Quality of Life and Well-Being Research;* Springer International Publishing: Cham, Switzerland, 2024; pp. 6269–6276.
- Cappelleri, J.C.; Bushmakin, A.G.; Gerber, R.A.; Leidy, N.K.; Sexton, C.C.; Lowe, M.R.; Karlsson, J. Psychometric analysis of the Three-Factor Eating Questionnaire-R21: Results from a large diverse sample of obese and non-obese participants. *Int. J. Obes.* 2009, 33, 611–620. [CrossRef] [PubMed]
- 52. Zhang, T.; Hao, Y.; Yuan, H.; Bu, P. Validity and reliability of Chinese version of Three-Factor Eating Questionnaire R21 for assessment of sample of college students. *Chin. Nurs. Res.* **2016**, *30*, 4137–4141.
- 53. Shi, C.; Guo, Y.; Ma, H.; Zhang, M. Psychometric validation of the 14-item perceived stress scale in Chinese medical residents. *Curr. Psychol.* **2019**, *38*, 1428–1434. [CrossRef]
- 54. Guo, S.; Sun, W.; Liu, C.; Wu, S. Structural validity of the Pittsburgh sleep quality index in Chinese undergraduate students. *Front. Psychol.* **2016**, *7*, 1126. [CrossRef]
- 55. Diamond, F.B., Jr.; Cuthbertson, D.; Hanna, S.; Eichler, D. Correlates of adiponectin and the leptin/adiponectin ratio in obese and non-obese children. *J. Pediatr. Endocrinol. Metab.* **2004**, *17*, 1069–1076. [CrossRef]
- 56. Salimi, S.; Farajian-Mashhadi, F.; Naghavi, A.; Mokhtari, M.; Shahrakipour, M.; Saravani, M.; Yaghmaei, M. Different profile of serum leptin between early onset and late onset preeclampsia. *Dis. Markers* **2014**, *2014*, *6*28476. [CrossRef] [PubMed]
- 57. Pardo, I.M.; Geloneze, B.; Tambascia, M.A.; Barros-Filho, A.A. Hyperadiponectinemia in newborns: Relationship with leptin levels and birth weight. *Obes. Res.* 2004, *12*, 521–524. [CrossRef] [PubMed]
- Paduszyńska, A.; Sakowicz, A.; Banach, M.; Maciejewski, M.; Dąbrowa, M.; Bielecka-Dąbrowa, A. Cardioprotective properties of leptin in patients with excessive body mass. *Ir. J. Med. Sci.* 2020, 189, 1259–1265. [CrossRef] [PubMed]
- 59. Wang, H.; Necheles, J.; Birne, J.S.; Li, Z.; Xing, H.; Tang, G.; Christoffel, K.K.; Brickman, W.J.; Zimmerman, D.; Wang, X. Association of adipokines with blood pressure in rural Chinese adolescents. *J. Hum. Hypertens.* **2012**, *26*, 493–501. [CrossRef] [PubMed]
- 60. Izadi, V.; Larijani, B.; Azadbakht, L. Is coffee and green tea consumption related to serum levels of adiponectin and leptin? *Int. J. Prev. Med.* **2018**, *9*, 106. [PubMed]
- 61. Enomoto, N.; Oyama, Y.; Yasui, H.; Karayama, M.; Hozumi, H.; Suzuki, Y.; Kono, M.; Furuhashi, K.; Fujisawa, T.; Inui, N.; et al. Analysis of serum adiponectin and leptin in patients with acute exacerbation of idiopathic pulmonary fibrosis. *Sci. Rep.* **2019**, *9*, 10484. [CrossRef] [PubMed]
- 62. Winnicki, M.; Somers, V.K.; Accurso, V.; Phillips, B.G.; Puato, M.; Palatini, P.; Pauletto, P. Fish-rich diet, leptin, and body mass. *Circulation* **2022**, *106*, 289–291. [CrossRef]
- 63. Olmedillas, H.; Sanchis-Moysi, J.; Fuentes, T.; Guadalupe-Grau, A.; Ponce-González, J.G.; Morales-Alamo, D.; Santana, A.; Dorado, C.; Calbet, J.A.; Guerra, B. Muscle hypertrophy and increased expression of leptin receptors in the musculus triceps brachii of the dominant arm in professional tennis players. *Eur. J. Appl. Physiol.* **2010**, *108*, 749–758. [CrossRef]
- Adelantado-Renau, M.; Jiménez-Pavón, D.; Beltran-Valls, M.R.; Ponce-González, J.G.; Chiva-Bartoll, Ó.; Moliner-Urdiales, D. Fitness and academic performance in adolescents. The mediating role of leptin: DADOS study. *Eur. J. Pediatr.* 2018, 177, 1555–1563. [CrossRef]
- 65. Muñoz, J.; Gower, B.A. Relationship between serum leptin concentration and low-density muscle in postmenopausal women. *J. Clin. Endocrinol. Metab.* **2003**, *88*, 1157–1161. [CrossRef] [PubMed]
- 66. Gysel, T.; Tonoli, C.; Pardaens, S.; Cambier, D.; Kaufman, J.M.; Zmierczak, H.G.; Goemaere, S.; Lapauw, B.; Calders, P. Lower insulin sensitivity is related to lower relative muscle cross-sectional area, lower muscle density and lower handgrip force in young and middle aged non-diabetic men. *J. Musculoskelet. Neuronal Interact.* 2016, 16, 302–309. [PubMed]
- 67. Perry, H.M.; Miller, D.K.; Patrick, P.; Morley, J.E. Testosterone and leptin in older African-American men: Relationship to age, strength, function, and season. *Metabolism* **2000**, *49*, 1085–1091. [CrossRef] [PubMed]
- Díez, J.J.; Iglesias, P. The role of the novel adipocyte-derived hormone adiponectin in human disease. *Eur. J. Endocrinol.* 2003, 148, 293–300. [CrossRef] [PubMed]
- 69. Pénicaud, L.; Meillon, S.; Brondel, L. Leptin and the central control of feeding behavior. *Biochimie* 2012, 94, 2069–2074. [CrossRef] [PubMed]
- 70. Hebebrand, J.; Denecke, S.; Antel, J. The role of leptin in rodent and human sleep: A transdiagnostic approach with a particular focus on anorexia nervosa. *Neurosci. Biobehav. Rev.* **2023**, *149*, 105164. [CrossRef] [PubMed]
- Dahmen, N.; Engel, A.; Helfrich, J.; Manderscheid, N.; Löbig, M.; Forst, T.; Pfützner, A.; Tonn, P. Peripheral leptin levels in narcoleptic patients. *Diabetes Technol. Ther.* 2007, *9*, 348–353. [CrossRef] [PubMed]
- Milaneschi, Y.; Simmons, W.K.; van Rossum, E.F.; Penninx, B.W. Depression and obesity: Evidence of shared biological mechanisms. *Mol. Psychiatry* 2019, 24, 8–33. [CrossRef]
- 73. Nieman, D.C.; Stear, S.J.; Castell, L.M.; Burke, L.M. A-Z of nutritional supplements: Dietary supplements, sports nutrition foods and ergogenic aids for health and performance: Part 24. *Br. J. Sports Med.* **2011**, *45*, 1005–1007.
- 74. Soares Bonato, R.C.; Abel Mapengo, M.A.; de Azevedo-Silva, L.J.; Janson, G.; de Carvalho Sales-Peres, S.H. Tooth movement, orofacial pain, and leptin, interleukin-1β, and tumor necrosis factor-α levels in obese adolescents. *Angle Orthod.* 2022, 92, 95–100. [CrossRef]

- 75. Peterlin, B.L.; Sacco, S.; Bernecker, C.; Scher, A.I. Adipokines and migraine: A systematic review. *Headache* **2016**, *56*, 622–644. [CrossRef] [PubMed]
- 76. Tanaka, Y.; Kita, S.; Nishizawa, H.; Fukuda, S.; Fujishima, Y.; Obata, Y.; Shimomura, I. Adiponectin promotes muscle regeneration through binding to T-cadherin. *Sci. Rep.* **2019**, *9*, 16. [CrossRef] [PubMed]

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