Plasma Adiponectin Levels Correlate Positively with an Increasing Number of Components of Frailty in Male Elders

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Abstract

Objective: Frailty is an important geriatric syndrome. Adiponectin is an important adipokine that regulates energy homeostasis. The aim of this study is to investigate the relationship between plasma adiponectin levels and frailty in elders.

Methods: The demographic data, body weight, metabolic and inflammatory parameters, including plasma glucose, total cholesterol, triglyceride, tumor necrosis factor alpha (TNF-α), c-reactive protein (CRP) and adiponectin levels, were assessed. The frailty score was assessed using the Fried Frailty Index (FFI).

Results: The mean (SD) age of the 168 participants (83 (49.4%) men and 85 (50.6%) women) was 76.86 (6.10) years. Judged by the FFI score, 42 (25%) elders were robust, 92 (54.7%) were pre-frail, and 34 (20.3%) were frail. The mean body mass index was 25.19 (3.42) kg/m². The log-transformed mean (SD) plasma adiponectin (µg/mL) level was 1.00 (0.26). The log-transformed mean plasma adiponectin (µg/mL) levels were 0.93 (0.23) in the robust elders, 1.00 (0.27) in the pre-frail elders, and 1.10 (0.22) in the frail elders, and the differences between these values were statistically significant \( p = 0.012 \). Further analysis showed that plasma adiponectin levels rose progressively with an increasing number of components of frailty in all participants as a whole \( (p \text{ for trend} = 0.024) \) and males \( (p \text{ for trend} = 0.037) \), but not in females \( (p \text{ for trend} = 0.223) \).

Conclusion: Plasma adiponectin levels correlate positively with an increasing number of components of frailty in male elders. The difference between the sexes suggests that certain sex-specific mechanisms may exist to affect the association between adiponectin levels and frailty.

Introduction

Frailty is a geriatric syndrome associated with decreased physiologic reserve, functional decline, and increased vulnerability to stressors, which lead to disability and mortality [1]. The mechanism of frailty is multifactorial. Inadequate nutrition and impairment and dysfunction of the endocrine and immune systems are involved in the development of frailty [2]. Fried’s concept of frailty postulates that frailty is initiated by the accumulation of factors such as lack of physical exercise, inadequate nutrition, injuries, disease, and drugs [3]. These factors may lead to chronic undernutrition, resulting in loss of bone and skeletal muscle mass and an increase in the degree of effort required for a given exercise intensity. Elders are more likely to avoid exercise as they believe that more effort is required. These changes may result in a significant decrease in resting energy metabolism and total energy expenditure [4].

Adiponectin is an important adipokine which possesses insulin sensitizing, anti-atherosclerotic and anti-inflammatory properties [5]. Studies have shown a negative correlation between the circulating levels of adiponectin and obesity, insulin resistance, coronary artery disease, and dyslipidemia [6,7]. High levels of circulating adiponectin have been associated with a reduced risk of cardiovascular disease (CVD) in some population-based studies [8,9]. In addition, adiponectin has a potential role in the central regulation of energy intake and expenditure [10]. Thus, circulating adiponectin may play a potential role in geriatric frailty.

A number of epidemiological studies on adiponectin have been conducted in aged populations. Adiponectin levels were significantly associated with high-density lipoprotein cholesterol (HDL-
C) concentrations in postmenopausal women, which suggested that high adiponectin levels may have a protective effect against atherosclerosis, when the HDL-C concentrations are high [11]. The inverse relationship between regional fat depots and the risk of the metabolic syndrome may be partially mediated by the adiponectin levels and the inflammatory status of middle-aged and older Chinese men and women [12]. Plasma concentrations of adiponectin were also high in middle-aged and older Chinese people with high levels of total physical activity [13]. However, adiponectin levels are known to increase with age [14], and this increase occurs in spite of the increase in visceral fat and insulin resistance that occurs with normal aging [15]. Furthermore, raised adiponectin levels are associated with increased rather than decreased risk of CVD and mortality in older subjects [16]. A study also showed that high levels of adiponectin predict mortality, particularly in patients with prevalent CVD [17]. In addition, a longitudinal study showed that circulating adiponectin levels increase over time in long-lived adults and are associated with greater physical disability and mortality [18]. These conflicting findings suggested a possibility that adiponectin may have different prognostic implications in older subjects [16].

Although a recent study showed lower fasting levels of adiponectin in frail women, the result was not statistically significant [19]. Thus, the association between plasma adiponectin levels and frailty in the older population warrants more investigation in detail. We aimed to investigate the relationship between plasma adiponectin levels and geriatric frailty in both men and women. We hypothesized that plasma adiponectin levels increase with frailty in elderly.

**Methods**

**Ethics Statement**

The study protocol was approved by the Ethics Committee of the National Taiwan University Hospital (registration number: 200701017R), and written informed consent was obtained from all participants before their inclusion in the study. The items of the consent form include aim, inclusion and exclusion criteria, procedures, harm and benefit, medical care, privacy and right, and withdrawal. All procedures were in accordance with the Helsinki Declaration. In addition, all potential participants who declined to participate or otherwise did not participate were still in care of their family physicians and were not disadvantaged in any other way by not participating in the study.

**Subjects**

From January 2007 to June 2008, the older patients who were aged 65 years and over and were followed up for chronic diseases with their family physicians in a hospital-based program were recruited for a structured comprehensive geriatric assessment if any of the following inclusion criteria was met. These criteria included: (1) functional decline (as measured by new disabilities of activity of daily living or instrumental activity daily living); (2) clinical indication of depression or dementia; (3) mobility impairment; (4) fall in recent one year; (5) weight loss more than 5% per year; (6) multiple comorbidities (≥5 diseases); (7) polypharmacy (≥8 classes of drugs per day); (8) multiple specialty physician visits in recent 6 months (≥3 different specialties with ≥2 visits for each specialty); (9) hospitalization in past year (≥1); (10) frequent emergency room visits in the past year (≥2); and (11) age above 80 years. The exclusion criteria were patients who were bedridden or residing in a nursing home, patients with an expected life expectancy of less than 6 months, and patients with severe hearing or communication disorder.

**Data Collection**

The experienced study nurses collected the data with a structured questionnaire, which included history on demographics, diseases, smoking and drinking habits, current medication, geriatric syndromes, blood pressure level, and body mass index (BMI). The Frailty Index was assessed by modified Fried’s criteria [3]: “weight loss” was defined as self-reported, unintentional weight loss of more than 3 kg (instead of 5 kg, adjusted in proportion to the Chinese body build) or greater than 5% of the body weight in the previous year. “Exhaustion” was indicated if the participants responded with “a moderate amount of the time” or “most of the time” to either of the following 2 statements: “I felt everything I did was an effort” or “I could not get going.” The statements were obtained from the Center for Epidemiological Studies-Depression Scale [20]. “Low physical activity” was defined by sex-specific, low weekly energy expenditure measured using the Taiwan International Physical Activity Questionnaire-Short Form (IPAQ-SF) [21] instead of the Minnesota Leisure Time Physical Activity Questionnaire [22]. “Slow walking speed” based on the time to walk for 5 meters was below certain sex- and height-specific cut-points [3]. “Weakness” was indicated when the maximal grip strength (kilograms) in the dominant hand (3 grips averaged), using a Jamar hand-held dynamometer was lesser than certain sex- and BMI-specific cut-points [3]. The subjects were classified as “robust,” “pre-frail,” or “frail” when 0, 1 or 2, or ≥3 components, respectively, screened positive [3].

**Biochemical Assays and Adiponectin Measurement**

Blood samples for the measurement of the adiponectin levels, complete blood count, and biochemical analysis were obtained from the antecubital vein of the subjects after an 8-h fast. For the adiponectin study, the blood was then immediately centrifuged. The plasma samples obtained after centrifugation were then frozen at −70°C until analysis. Total plasma adiponectin levels were determined using a previously described radioimmunoassay method (Linco Research, Inc., St. Charles, MO) [23]. Plasma TNF-α levels were measured by the commercial enzyme-linked immunosorbent assay (ELISA) kits (Assaypro LLC, Saint Charles, Missouri, USA). Plasma CRP levels were measured by the latex agglutination test (Denka Seiken, Gosen, Niigata, Japan).

**Statistical Analyses**

Descriptive statistical data were summarized as frequencies and percentages for categorical variables and means and standard deviations for other continuous variables. An analysis of variance (ANOVA) was used to examine significant differences in the adiponectin levels of the three frailty subgroups. A general linear model was used to test if there is a trend between frailty scores and adiponectin concentrations (log-transformed) after adjusted for age and BMI for both sexes and for all participators. A probability of less than 0.05 (p<0.05) was considered statistically significant. All data were analyzed by using SPSS 14.0 statistical software (SPSS Inc., Chicago, IL).

**Results**

**Plasma Adiponectin Levels in the Study Subjects**

There were 241 patients aged 65 years and over who met the inclusion criteria. Fifty-two patients declined to join the study and 189 patients completed the assessment and frailty screening. Among the 189 participants, 168 had their blood sample collected. Of the 168 elders, 83 (49.4%) were men. Their mean (SD) age was 76.86 (6.10) years. Fifty-one (30.4%) elders had quitted cigarette smoking and only nine (5.4%) smoked cigarettes. The leading factors for the exclusion were patients who were bedridden or residing in a nursing home, patients with an expected life expectancy of less than 6 months, and patients with severe hearing or communication disorder.
comorbidities were hypertension (82.7%), hyperlipidemia (61.3%),
diabetes mellitus (43.5%), and coronary artery disease (31.5%).
Judged by the frailty scoring, 42 (25.0%) elders were robust, 92
were (54.7%) pre-frail, and 34 (20.3%) were frail (Table 1). Their
medication data are summarized in Table 1. There was no
significant difference in demographic variables except education
between male and female subgroups. The mean (SD) BMI was
25.19 (3.42) kg/m². The log-transformed mean (SD) plasma
adiponectin (µg/mL), TNF-α (pg/mL) and CRP (nmol/L) levels
were 1.00 (0.26), 1.51 (0.41), and 1.42 (0.29) respectively (Table 2).
Other laboratory data results are summarized in Table 2. Plasma
total cholesterol (p = 0.001) and uric acid (p = 0.022) concentra-
tions were significantly different between male and female
subgroups.

Plasma Adiponectin Levels in the Three Subgroups with
Different Frailty Levels

Among the three subgroups, participants in the frail subgroup
were significantly older than those in the other two subgroups. The
log-transformed mean (SD) plasma adiponectin (µg/mL) levels in
the robust, pre-frail, and frail subgroups were 0.93 (0.23), 1.00
(0.27), and 1.10 (0.22), respectively. The plasma adiponectin levels
were significantly different among the three subgroups (p = 0.012).

**Table 1.** Demographic data of study participants.

| Variables               | Overall (n = 168) | Male (n = 83) | Female (n = 85) | t(p)/x²(p) |
|-------------------------|------------------|--------------|----------------|------------|
| Age                     | 76.86±6.10       | 77.71±5.87   | 76.04±6.24     | 1.79 (.075) |
| Education               |                  |              |                |            |
| Less than 6 years       | 86 (51.2%)       | 31 (37.3%)   | 55 (64.7%)     | 17.99 (.000) |
| Junior High             | 23 (13.7%)       | 10 (12.0%)   | 13 (15.3%)     |            |
| Senior High             | 24 (14.3%)       | 16 (19.3%)   | 8 (9.4%)       |            |
| College or more         | 35 (20.8%)       | 26 (31.3%)   | 9 (10.6%)      |            |
| Smoking status          |                  |              |                |            |
| Never                   | 108 (64.3%)      | 25 (30.1%)   | 83 (97.6%)     | N/A        |
| Quitted                 | 51 (30.4%)       | 50 (60.2%)   | 1 (1.2%)       |            |
| Smoking                 | 9 (5.4%)         | 8 (9.6%)     | 1 (1.2%)       |            |
| Comorbidity             |                  |              |                |            |
| Hypertension            | 139 (82.7%)      | 66 (79.5%)   | 73 (85.9%)     | 1.19 (.275) |
| Hyperlipidemia          | 103 (61.3%)      | 47 (56.6%)   | 56 (65.9%)     | 1.52 (.218) |
| Diabetes mellitus       | 73 (43.5%)       | 34 (41.0%)   | 39 (45.9%)     | 0.41 (.520) |
| Coronary artery disease | 53 (31.5%)       | 27 (32.5%)   | 26 (30.6%)     | 0.07 (.787) |
| Stroke                  | 45 (26.8%)       | 25 (30.1%)   | 20 (23.5%)     | 0.93 (.335) |
| Medication              |                  |              |                |            |
| Aspirin                 | 74 (44.0%)       | 39 (47.0%)   | 35 (41.2%)     | 0.57 (.448) |
| β-blockers              | 41 (24.4%)       | 21 (25.3%)   | 20 (23.5%)     | 0.07 (.789) |
| Calcium channel blockers| 80 (47.6%)       | 38 (45.8%)   | 42 (49.4%)     | 0.22 (.638) |
| ACEIs or ARBs           | 94 (56.0%)       | 46 (53.4%)   | 48 (56.5%)     | 0.02 (.891) |
| Metformin               | 44 (26.2%)       | 18 (21.7%)   | 26 (30.6%)     | 1.72 (.190) |
| Sulfonylureas           | 53 (31.5%)       | 22 (26.5%)   | 31 (36.5%)     | 1.93 (.165) |
| Thiazolidinediones      | 13 (7.7%)        | 6 (7.2%)     | 7 (8.2%)       | 0.06 (.807) |
| Acarbose                | 5 (3.0%)         | 3 (3.6%)     | 2 (2.4%)       | N/A        |
| Repaglinide             | 4 (2.4%)         | 1 (1.2%)     | 3 (3.5%)       | N/A        |
| Statin                  | 56 (33.3%)       | 22 (26.5%)   | 34 (40.0%)     | 3.44 (.064) |
| Frailty Score (Level)   |                  |              |                |            |
| 0 (Robust)              | 42 (25.0%)       | 22 (26.5%)   | 20 (23.5%)     | N/A        |
| 1 (Pre-frail)           | 53 (31.5%)       | 30 (36.1%)   | 23 (27.1%)     |            |
| 2 (Pre-frail)           | 39 (23.2%)       | 17 (20.5%)   | 22 (25.9%)     |            |
| 3 (Frail)               | 24 (14.3%)       | 9 (10.8%)    | 15 (17.6%)     |            |
| 4 (Frail)               | 9 (5.4%)         | 5 (6.0%)     | 4 (4.7%)       |            |
| 5 (Frail)               | 1 (0.6%)         | 0 (0.0%)     | 1 (1.2%)       |            |

Note: a: n (%) for categorical data, mean (SD) for continuous data.
N/A: Chi-square not available due to having cells with less than 5 cases.
No variable was adjusted for those models in this table.
ACEIs: angiotensin-converting enzyme inhibitors; ARBs: angiotensin II receptor blockers.
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The results of the post hoc tests showed that the people in the frail subgroup had higher plasma adiponectin levels than those in the robust subgroup. We further analyzed the effect of sex on the relationship between frailty severity and plasma adiponectin level. In both subgroups, plasma adiponectin levels were not associated with frailty severity. With regard to comorbidity and medication, a relationship between frailty severity and plasma adiponectin level. However, both studies found no significant relationship between adiponectin levels correlate positively with an increasing number of components of frailty in male elders. Our findings support the link between high adiponectin levels and mortality in the general older population [16,18].

A recent study on 73 community-dwelling women, aged 84–95 years without a diagnosis of diabetes, showed fasting levels of adiponectin were lower in frail subjects, but not significant statistically [19]. There are two important differences in the study subjects between their study and ours. First, their study population did not have diabetes whereas 43% of our study subjects did. Second, their study population was limited in women only. However, both studies found no significant relationship between plasma adiponectin levels and frailty in females.

Geriatric frailty, which is usually concurrent with sarcopenia and aging, is associated with increased inflammatory activity and is reflected by increased levels of circulating TNF-α, interleukin (IL)-6, cytokine antagonists, and acute-phase proteins. Epidemiologic studies suggest that chronic low-grade inflammation during aging promotes an atherogenic profile and is related to age-associated disorders, such as Alzheimer’s disease, atherosclerosis, type 2 diabetes and enhances mortality risk. The development of frailty is characterized by the dysregulated production of proinflammatory adiponectin levels correlate positively with an increasing number of components of frailty in male elders. Our findings support the link between high adiponectin levels and mortality in the general older population [16,18].

| Laboratory tests | Overall | Male | Female | t(p) |
|------------------|---------|------|--------|------|
| RBC (M/μL) | 4.38 (0.58) | 4.49 (0.65) | 4.27 (0.47) | 2.42 (0.017) |
| Hb (g/dL) | 12.98 (1.60) | 13.37 (1.75) | 12.58 (1.34) | 3.05 (0.003) |
| Platelet (K/μL) | 223.01 (71.77) | 205.92 (67.09) | 240.11 (72.66) | -2.97 (0.003) |
| WBC (K/μL) | 6.54 (1.55) | 6.51 (1.47) | 6.56 (1.64) | -0.20 (0.843) |
| Albumin (g/dL) | 4.56 (0.33) | 4.54 (0.42) | 4.57 (0.23) | -0.39 (0.697) |
| Glucose AC (mmol/L) | 5.00 (1.92) | 6.00 (1.97) | 6.60 (1.89) | -0.65 (0.519) |
| Total-Cholesterol (mmol/L) | 4.89 (0.88) | 4.64 (0.80) | 5.11 (0.90) | -3.41 (0.001) |
| Triglyceride (mmol/L) | 1.69 (1.01) | 1.56 (0.75) | 1.81 (1.20) | -1.55 (0.122) |
| AST (Ukat/L) | 0.41 (0.20) | 0.43 (0.25) | 0.39 (0.14) | 1.01 (0.312) |
| ALT (Ukat/L) | 0.37 (0.24) | 0.39 (0.28) | 0.34 (0.19) | 1.45 (0.148) |
| BUN (mmol/L) | 0.75 (0.34) | 0.83 (0.40) | 0.67 (0.23) | 2.95 (0.003) |
| Creatinine (μmol/L) | 110.12 (48.31) | 129.77 (58.77) | 91.23 (23.47) | 5.29 (0.000) |
| MDRD-simplify-GFR (mL/min/1.73 m²) | 57.88 (16.37) | 56.58 (17.55) | 59.12 (15.16) | -0.96 (0.338) |
| Log(TNF-α) (pg/mL) | 1.51 (0.41) | 1.50 (0.41) | 1.52 (0.40) | -0.30 (0.762) |
| Log (CRP (nmol/L)) | 1.42 (0.29) | 1.44 (0.29) | 1.41 (0.30) | -0.50 (0.619) |
| Log (Adiponectin (μg/mL)) | 1.00 (0.26) | 0.97 (0.22) | 1.03 (0.28) | -1.30 (0.195) |

MDRD: Modification of diet in renal disease.

MDRD-simplify-GFR (mL/min/1.73 m²) = 186 × [(CRE)⁻¹.154 × (age)⁻⁰.²₀⁰] / [(CRE)⁻¹.154 × (age)⁻⁰.²₀⁰] if male).

MDRD-simplify-GFR (mL/min/1.73 m²) = 186 × [(CRE)⁻¹.154 × (age)⁻⁰.²₀⁰] × 0.742 (if female).

No variable was adjusted for those models in this table.

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Table 2. Results of the physical examination and laboratory tests of the 168 elders.
High Plasma Adiponectin Levels in Frail Elders

(a) Log (Adiponectin levels)

(b) Log (Adiponectin levels)

(c) Log (Adiponectin levels)
cytokines [24]. Therefore, frailty is a critical condition in the elderly. It is noteworthy that plasma adiponectin levels are high in frail elders, which has anti-inflammatory, insulin sensitization, and anti-atherogenic properties [25]. However, plasma TNF-α and CPR levels were not elevated in the frail elders in our study. Thus, elevation of plasma adiponectin levels in the frail elders is not due to compensatory upregulation. A recent study has shown that age-related increases in adiponectin and inflammatory markers are unrelated [26], which supports our finding.

Several studies have also shown that plasma adiponectin levels are high and indicate a poor prognosis in some critical conditions such as severe CVDs [16,17]. Accumulation of adiponectin in patients with diseases such as heart failure or chronic kidney disease may reflect the emaciation and malnutrition that characterize these diseases, and therefore, such an accumulation is a marker of poor prognosis [27]. However, the nutritional status, including albumin levels, hemoglobin levels, and BMI, was similar among the three subgroups with different levels of frailty severity in this study. Thus, malnutrition is not the only factor affecting plasma adiponectin levels in elders.

Aging is associated with weight loss and loss of skeletal muscle mass and strength (sarcopenia) [28], which are prominent predictors of mortality in the elderly population [29]. Body weight is an important factor in the regulation of adiponectin expression. Plasma adiponectin levels decreased in obese patients [6] and increased after weight reduction [30]. It has been suggested that high adiponectin levels in elderly persons may be a consequence of weight loss and sarcopenia. Even though the weight and BMI values were not different among the different subgroups in this study, the sample size in this study is not enough to exclude the effect of weight.

Age-related decline in renal function might cause a reduction in renal adiponectin clearance that may contribute to the increase in adiponectin levels in the elderly [24]. However, renal function, which was estimated using modification of diet in renal disease (MDRD)-simplify-GFR, was similar among the different subgroups in this study.

Table 3. Comparisons among elders with different frailty levels.

| Frailty     | Robust | Pre-frail | Frail | Statistics* | F(p)/x²(p) |
|------------|--------|-----------|-------|-------------|------------|
| Age (years) | 74.69±6.68 | 77.04±5.97 | 79.06±4.86 | 5.14 (.007) |
| Sex (female) | 20 (47.6%) | 45 (48.9%) | 20 (58.8%) | 1.17 (.556) |
| Body mass index (kg/m²) | 25.04±3.32 | 25.52±3.55 | 24.46±3.15 | 1.26 (.287) |
| MDRD-simplify-GFR (mL/min/1.73 m²) | 62.43±12.56 | 56.68±18.23 | 55.62±14.81 | 2.04 (.143) |
| Log (TNF-α (pg/mL)) | 1.55±0.38 | 1.50±0.41 | 1.47±0.43 | 0.36 (.699) |
| Log (CRP (nmol/L)) | 1.42±0.30 | 1.42±0.28 | 1.43±0.32 | 0.03 (.974) |
| Log (Adiponectin (μg/mL)): | | | | |
| Overall | 0.93±0.23 | 1.00±0.27 | 1.10±0.22 | 4.51 (.012) |
| Male | 0.89±0.21 | 0.99±0.23 | 1.06±0.18 | 2.66 (.076) |
| Female | 0.96±0.25 | 1.01±0.31 | 1.13±0.25 | 1.92 (.154) |
| Diseases | | | | |
| Hypertension | 38 (90.5%) | 75 (81.5%) | 26 (76.5%) | 2.79 (.248) |
| Diabetes mellitus | 10 (23.8%) | 44 (47.8%) | 19 (55.9%) | 9.45 (.009) |
| Hyperlipidemia | 32 (76.2%) | 57 (62.0%) | 14 (41.2%) | 9.75 (.008) |
| Stroke | 5 (11.9%) | 25 (27.2%) | 15 (44.1%) | 9.96 (.007) |
| CAD | 13 (31.0%) | 33 (35.9%) | 7 (20.6%) | 2.69 (.260) |
| Medication | | | | |
| Aspirin | 14 (33.3%) | 39 (42.4%) | 21 (61.8%) | 6.39 (.041) |
| β-blockers | 12 (28.6%) | 21 (22.8%) | 8 (23.5%) | 0.53 (.766) |
| CCBs | 20 (47.6%) | 47 (51.1%) | 13 (38.2%) | 1.64 (.440) |
| ACEIs or ARBs | 24 (57.1%) | 48 (52.2%) | 22 (64.7%) | 1.61 (.446) |
| Metformin | 7 (16.7%) | 24 (26.1%) | 13 (38.2%) | 4.52 (.104) |
| Sulfonylureas | 9 (21.4%) | 31 (33.7%) | 13 (38.2%) | 2.89 (.235) |
| Statins | 22 (52.4%) | 29 (31.5%) | 5 (14.7%) | 12.30 (.002) |

CAD: coronary artery disease; CCBs: calcium channel blockers; ACEIs: angiotensin-converting enzyme inhibitors; ARBs: angiotensin II receptor blockers.

[Post hoc: Frail:Robust for AP and log (AP)].

No variable was adjusted for those models in this table.

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Thus, sex is an important factor not only affecting plasma adiponectin levels [31], but also their relations to frailty severity. The finding suggests that adiponectin may have different regulation in frail male and female elders. This may not relate to the adiponectin gene that explains 6.7% of the phenotypic variation because the gene is unlikely to cause sex differences in adiponectin levels [32]. A potential explanation could be sex hormone levels. Sex-hormone modulation of adiponectin production may provide protection from adiponectin elevation with frailty in the female elders.

In addition to sex, nutritional status and body weight, other factors affecting plasma adiponectin levels include menopausal status, and medications such as β-blockers and angiotensin II receptor blockers [33,34]. However, the age of the study subjects was greater than 65 years, and therefore, all the female participants were menopausal. Furthermore, the use of β-blockers and angiotensin II receptor blockers was not different among the three subgroups.

Factors such as malnutrition, weight loss, comorbid diseases, and medication use did not explain the high levels of plasma adiponectin in frail elders. Geriatric frailty, which causes disturbances in energy homeostasis, is associated with sarcopenic obesity and physical inactivity [35]. Thus, the elevation of plasma adiponectin levels in the frail elders may be due to age-related homeostatic dysregulation [18]. Further investigation is warranted to define the underlying mechanisms.

The small sample size is a limitation in this study, which limits the number of covariates that can be added to regression models and is underpowered in detecting a significant interaction by sex. A larger-scale study to confirm the results obtained in the small-scale pilot study is necessary. On the other hand, circulating adiponectin has three isoforms: high-, medium-, and low-molecular-weight adiponectin [36]. Previous studies have shown that high-molecular-weight adiponectin exerts a protective role as an anti-diabetic and anti-atherogenic hormone [36,37]. However, a recent study showed that low-molecular-weight adiponectin has a protective role in aging [38]. Therefore, further studies on the adiponectin isoforms in elders are necessary to elucidate the role of adiponectin in geriatric frailty.

In conclusion, plasma adiponectin levels correlate positively with an increasing number of components of frailty especially in male elders. The difference between the sexes suggests that certain sex-specific mechanisms may exist to affect the association between adiponectin levels and frailty. Further study is required to elucidate the underlying mechanisms.

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Author Contributions
Conceived and designed the experiments: JST LMC Ching-Yu Chen. Performed the experiments: JST KCH Chin-Ying Chen CIC Ching-Yu Chen. Analyzed the data: CHW SCC. Contributed reagents/materials/analysis tools: CHW SCC. Wrote the paper: JST LMC Ching-Yu Chen.

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