Association of fasting serum insulin and fasting serum glucose levels with cognitive impairment in Chinese nonagenarians/centenarians

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Abstract In this study, we examined the association of fasting serum insulin (FSI) and fasting serum glucose (FSG) with cognitive impairment in the very elderly using a sample of Chinese nonagenarians/centenarians. This study used data from a survey that was conducted in 2005 on all residents aged 90 years or more in a district with 2,311,709 inhabitants. FSG, FSI, and cognitive function were analyzed. The sample included 661 unrelated Chinese individuals (aged 90–108 years; mean, 93.52±3.37 years; 67.17 % women; FSI, 6.27±2.27 mU/mL; FSG levels, 4.46±1.45 mmol/L). The prevalence of cognitive impairment was 61.81 % and that of hypoinsulinemia was 31.92 %. Individuals with hypoinsulinemia showed lower cognitive function scores (14.81±5.79 vs. 15.78±5.24, t=2.160, P=0.031). No differences in cognitive function score between different FSI and FSG groups were significant, and no differences in FSI and FSG between individuals with and without cognitive impairment were statistically significant. Unadjusted multiple logistic regressions showed that hypoinsulinemia, impaired fasting glucose, or diabetes did not change the risk of cognitive impairment significantly. In summary, we found that in elderly subjects, cognitive function appeared associated with FSI, and higher FSI may be associated with enhanced cognitive function.

Keywords Fasting serum insulin · Fasting serum glucose · Nonagenarians/centenarians

Introduction

Type 2 diabetes and its related metabolic disorders including hyperinsulinemia, insulin resistance, and hyperglycemia are important conditions in the elderly (Kesavadev et al. 2003). Recent studies have discovered that the insulin metabolic disorders may be one of the important risk factors for Alzheimer’s disease (AD) (Duron and Hanon 2008; Planel et al. 2007). Some researchers have postulated that independent of the effect of glucose, the level of hyperinsulinemia correlates with cognitive decline in the elderly (van den Berg et al. 2006, 2007). However, other researchers have presented evidence showing no relationship between insulin levels and cognitive dysfunction (Kodl and Seaquist 2008; Park 2001; Kern et al. 2001). Thus, the relationship between fasting serum insulin (FSI) and cognitive dysfunction remains unknown (van den Berg et al. 2006, 2007; Kodl and Seaquist 2008; Park 2001; Kern et al. 2001). Several
additional studies have shown that impaired fasting glucose and type 2 diabetes and their related metabolic disorders correlate with vascular dementia rather than AD (Craft 2009). The association of cognitive function with FSI and fasting serum glucose (FSG) has been the focus of many researchers, but no definitive conclusion has been generated (van den Berg et al. 2006, 2007; Kodl and Seaquist 2008; Park 2001; Kern et al. 2001). However, although some evidence of the correlation exists, no study has examined long-lived subjects. In this study, we provide evidence from long-lived subjects, which may prove useful in clarifying the association of cognitive function with FSI and FSG. We examine a group of Chinese nonagenarians/centenarians.

Subjects and methods

Study subjects

The methods used have been described previously (Zhou et al. 2010; Huang et al. 2009). In brief, a cross-sectional study for age-related diseases was conducted in 870 long-lived subjects (>90 years), which was a part of the Project of Longevity and Aging in Duijiangyan (PLAD) and was based on the Duijiangyan (located in Sichuan Province in southwest China) 2005 census. The PLAD aimed to investigate relationships between the environment, lifestyle, genetics, cognitive function, longevity, and age-related diseases. The basic health of volunteers was examined by trained physicians. Results were entered onto a standard form and included a questionnaire on lifestyles and cognitive function [measured using the 30-item Mini-Mental State Examination (MMSE)]. FSG and FSI were determined using the glucose oxidase method and RIAs, respectively. We excluded those who did not complete the MMSE test or where no information on FSG or FSI was available. Subjects with a previous diagnosis of diabetes (they may have used treatments to influence their FSG, FSI, or insulin resistance (IR)) or receiving diabetes medication were also excluded. Overall, 21 men and 26 women were excluded from the study because they had already died or moved out of the area. Of 262 men and 561 women who were interviewed, 38 men and 124 women did not complete the MMSE test. Those that did not provide FSG or FSI information or those with a previous diagnosis of diabetes did not complete MMSEs. Therefore, the study population ultimately comprised 661 long-lived subjects. Informed consents were obtained from all participants (and their legal proxies). The Research Ethics Committee of Sichuan University approved this study.

Data collection and measurements

Assessment of cognitive function

The methods used to assess cognitive function have been reported previously (Zhou et al. 2010; Huang et al. 2009). Cognitive function was measured using the 30-item MMSE, which is a global test that includes components that test orientation, attention, calculation, language, and recall. To decrease methodological errors and assure method reliability, the administrator of the test was provided with professional training comprising: (1) reviewing the MMSE procedure and grading system, as outlined in a short booklet and video; (2) observing a geriatrician conduct the MMSE on residents who were not part of the study; and (3) receiving supervision while conducting the MMSE on residents who were not part of the study. Individuals who gave consent to participate in the study were tested using the MMSE. The individuals were categorized as follows: possible dementia (scores between 0 and 18), mild cognitive impairment (scores between 19 and 24), and normal (scores between 25 and 30) (Zhou et al. 2010; Huang et al. 2009; Tombaugh and McIntyre 1992).

The MMSE relies heavily on visual and auditory abilities, in particular at advanced ages (Zhou et al. 2010; Huang et al. 2009), and there was a high prevalence of visual or hearing impairment among the nonagenarians and centenarians. Twenty-eight men and 72 women did not complete the MMSE test due to visual or hearing impairment. To address this issue, these subjects were excluded at data analysis. In the oldest-old group, only 35 subjects had scores higher than 24 (30 in men and 5 in women); therefore, we merged data from subjects with mild cognitive impairment and normal cognitive function into an “impossible” dementia group when the data were analyzed.

Measurement of FSG and FSI

Blood samples for measurement of FSG and FSI were obtained from all subjects at 08:00 following an overnight fast. FSG was measured using the glucose oxidase method (glucose B-test, Wako Pure Chemical, Osaka, Japan) (Kouzuma et al. 2002). FSI levels were determined using the RIA technique. Sera used to measure serum insulin concentrations by RIA were stored at
−80 °C (Bowsher et al. 1999; Singhal et al. 2005). FSG and FSI were determined using standard laboratory techniques (performed by a technician in the biochemistry laboratory of Sichuan University). Abnormal FSG and FSI levels were defined based on criteria provided by the China Diabetes Association as follows: normal FSG, <6.00 mmol/L; IFG, 6.00–7.00 mmol/L; diabetes, >7.00 mmol/L; hypo-FSI, <5.00 mU/mL; normal FSI, 5–15.00 mU/mL; and hyper-FSI, >15.00 mU/mL (Song et al. 2011; Xue et al. 2010), respectively.

Assessment of covariates

The methods used have been described previously (Zhou et al. 2010; Huang et al. 2009). The baseline examination obtained information on age (years), gender (male/female), body mass index (BMI), serum uric acid (SUA), and serum lipid/lipoprotein levels. Blood pressure in the right arm (sitting or recumbent position) was measured twice to the nearest 2 mmHg using a standard mercury sphygmomanometer (Korotkoff phases I and V) by trained nurses or physicians. Serum lipid/lipoprotein levels (including serum triglyceride (TG), total cholesterol (TC), high density lipoprotein (HDL) cholesterol, and low density lipoprotein (LDL) cholesterol) and SUA levels were determined using standard laboratory techniques (performed by a technician in the biochemistry laboratory of Sichuan University).

Statistical analyses

All statistical analyses for this study were performed using the SPSS for Windows software package, version 11.5 (SPSS Inc., Chicago, IL, USA). Comparisons were made between baseline characteristics of the different FSI groups (out of all participants, only 5 showed hyperinsulinemia and 211 showed hypoinsulinemia; the five individuals with hyperinsulinemia were merged into the normal insulin group) and between individuals with and without cognitive impairment using unpaired Student’s t tests for continuous variables or Pearson Chi squared or Fisher’s exact tests (where an expected cell count was <5) for categorical variables. Baseline characteristics between the different FSG groups were also compared using analysis of variance for continuous variables. Multiple logistic regression was used to estimate the odds ratio and 95 % confidence interval of hypoinsulinemia or abnormal FSG as a function of

Table 1 Baseline demographics based on fasting serum glucose (FSG; n=661)

| Demographic                        | All (n=661) | Fasting serum glucose | Diabetes (n=41) | \(\chi^2\) or \(F\) | \(P\) value |
|------------------------------------|-------------|-----------------------|-----------------|-----------------|-------------|
|                                    | Normal (n=587) | Impaired fasting glucose (n=33) |                  |                 |             |
| Age (years)                        | 93.52±3.37 | 95.56±3.34 | 92.79±3.52 | 93.51±3.62 | 0.818 | 0.442 |
| Gender (male/female)               | 217/444 | 187/400 | 15/18 | 15/26 | 2.964 | 0.227 |
| MMSE score                         | 15.48±5.44 | 15.46±5.45 | 16.63±5.15 | 14.90±5.66 | 0.939 | 0.392 |
| Cognitive impairment (yes/no)      | 409/252 | 366/221 | 16/17 | 15/26 | 1.599 | 0.420 |
| BMI (kg/m²)                        | 19.05±3.64 | 19.11±3.67 | 18.40±3.40 | 18.57±3.46 | 0.956 | 0.385 |
| DBP (mmHg)                         | 72.59±12.06 | 72.58±12.03 | 70.52±12.48 | 74.46±12.10 | 0.984 | 0.374 |
| SBP (mmHg)                         | 139.71±23.31 | 140.21±22.78 | 140.36±28.58 | 135.61±18.91 | 0.780 | 0.459 |
| TG (mmol/L)                        | 1.22±0.64 | 1.23±0.64 | 0.99±0.32 | 1.27±0.74 | 2.264 | 0.105 |
| TC (mmol/L)*                       | 4.15±0.85 | 4.18±0.85 | 3.94±0.82 | 3.89±0.12 | 3.170 | 0.043 |
| HDL (mmol/L)                       | 1.58±0.59 | 1.59±0.61 | 1.56±0.43 | 1.40±0.35 | 1.327 | 0.222 |
| LDL (mmol/L)                       | 2.28±0.97 | 2.30±0.97 | 2.09±0.57 | 2.12±0.53 | 2.862 | 0.105 |
| SUA (μmol/L)**                     | 318.72±87.01 | 314.84±81.53 | 338.91±95.80 | 357.62±134.16 | 5.642 | 0.004 |
| FSI (mU/L)                         | 6.27±2.27 | 6.24±2.24 | 6.46±2.58 | 6.49±2.56 | 0.359 | 0.669 |

Baseline demographics were compared between different SUA groups
SUA serum uric acid, BMI body mass index, FSG fasting blood glucose, HDL high density lipoprotein, LDL low density lipoprotein, TC total cholesterol, TG triglyceride, SBP systolic blood pressure, DBP diastolic blood pressure
*\(P<0.05\); ** \(P<0.01\) vs. low SUA group using \(\chi^2\) or Fisher’s exact tests (where an expected cell count was <5) for categorical variables or unpaired Student’s t tests for continuous variables
increased risk for increased cognitive impairment. We adjusted age, gender, and educational levels as general covariates in model 1. In view of the fact that general or central adiposity, elevated blood pressure, dyslipidemia, hyperglycemia, and hyperuricacidemia may be components of metabolic syndrome and that they may be involved in the association of hypoinsulinemia or abnormal FSG with cognitive impairment, we adjusted BMI, serum lipid/lipoprotein, blood pressure, and SUA in model 2. We adjusted all of these covariates in model 3. A $P$ value of $<0.05$ was considered to be statistically significant, and all $P$ values were two sided.

Results

Baseline characteristics of genotype frequencies of the IR and cognitive impairment

The sample group included 661 unrelated Chinese (aged 90–108 years; mean, 93.52±3.37 years; 67.3 % women; FSI, 6.27±2.27 mU/mL; FSG, 4.46±1.45 mmol/L; MMSE score, 15.49±5.44). In this sample, 409 individuals (61.8 %) were cognitively impaired.

To demonstrate the association between demographics or clinical characteristics and FSG, the means (or percentages) of various demographic and clinical characteristics grouped by FSG are shown in Table 1. Subgroups with impaired fasting glucose and diabetes showed significantly higher SUA ($F^2=5.642$, $P=0.004$) and TC ($F^2=3.170$, $P=0.043$) levels. There were no significant differences between the different subgroups in other covariates (age, gender, FSI, serum lipid/lipoprotein levels, blood pressure, and BMI; see Table 1).

The means (or percentages) of various demographic or clinical characteristics in the hypoinsulinemia and normal insulin groups are shown in Table 2 to demonstrate the association between demographics or clinical characteristics and FSI. Fewer males showed hypoinsulinemia compared with females ($\chi^2=5.280$, $P=0.029$). Individuals with hypoinsulinemia showed significantly lower SUA ($t=4.486$, $P<0.0001$), TC ($t=5.376$, $P<0.0001$), HDL ($t=3.321$, $P=0.010$), and LDL ($t=2.353$, $P=0.027$) and higher diastolic blood pressure ($t=1.964$, $P=0.049$) levels. There were no significant differences between the different subgroups in other covariates (age, TG, SBP, and BMI; see Table 2).

Table 2 Baseline demographics based on fasting serum insulin (FSI; $n=661$)

| Demographic              | All ($n=661$) | Fasting serum insulin | Hypoinsulinemia ($<5$ mU/mL; $n=211$) | Normal insulin ($>5$ mU/mL; $n=450$) | $\chi^2$ or $t$ | $P$  |
|--------------------------|--------------|-----------------------|---------------------------------------|-------------------------------------|-----------------|------|
| Age (years)              | 93.52±3.37   | 93.40±3.16            | 93.57±3.47                            | 0.598                               | 0.550           |
| Gender (male/female)*    | 217/444      | 58/153                | 158/291                               | 5.280                               | 0.029           |
| MMSE scores*             | 15.48±5.44   | 14.81±5.79            | 15.78±5.24                            | 2.160                               | 0.031           |
| Cognitive impairment (yes/no) | 249/252    | 139/72                | 268/177                               | 2.960                               | 0.228           |
| BMI (kg/m²)              | 19.05±3.64   | 19.17±3.54            | 18.99±3.69                            | 0.573                               | 0.567           |
| DBP (mmHg)*              | 72.59±12.06  | 73.94±11.91           | 71.96±12.14                           | 1.964                               | 0.049           |
| SBP (mmHg)               | 139.71±23.31 | 141.12±21.94          | 139.38±23.38                          | 0.913                               | 0.932           |
| TG (mmol/L)              | 1.22±0.64    | 1.28±0.69             | 1.19±0.61                             | 1.598                               | 0.111           |
| TC (mmol/L)**            | 4.15±0.85    | 3.99±0.85             | 4.22±0.84                             | 5.376                               | <0.0001         |
| HDL (mmol/L)*            | 1.58±0.59    | 1.50±0.41             | 1.61±0.65                             | 3.321                               | 0.010           |
| LDL (mmol/L)*            | 2.28±0.97    | 2.16±0.60             | 2.33±1.10                             | 2.353                               | 0.027           |
| SUA (μmol/L)**           | 318.72±87.01 | 296.75±79.90          | 328.97±88.37                          | 4.486                               | <0.0001         |
| FSG (mmol/L)             | 4.47±1.45    | 4.49±1.49             | 4.47±1.44                             | 0.180                               | 0.858           |

Baseline demographics were compared between different FSG groups

SU4 Serum uric acid, BMI body mass index, FSG fasting blood glucose, HDL high density lipoprotein, LDL low density lipoprotein, TC total cholesterol, TG triglyceride, SBP systolic blood pressure, DBP diastolic blood pressure

* $P<0.05$; ** $P<0.01$ vs. low SUA group using $\chi^2$ or Fisher’s exact tests (where an expected cell count was <5) for categorical variables and unpaired Student’s $t$ tests for continuous variables
The means (or percentages) of various demographic or clinical characteristics grouped by cognitive impairment are shown in Table 3 to demonstrate the association between demographics or clinical characteristics and cognitive impairment. Fewer males showed cognitive impairment compared with females ($\chi^2=56.16, P<0.0001$). Individuals with higher educational levels showed less cognitive impairment ($\chi^2=95.32, P<0.0001$). Individuals with cognitive impairment showed higher TC ($t=2.098, P=0.036$). There were no significant differences between the different subgroups in other covariates (see Table 3).

Association of cognitive impairment with FSG

Among the different FSG groups, there was no significant difference in MMSE score ($F=0.939, P=0.392$) or in prevalence of cognitive impairment ($F=0.939, P=0.392$; see Table 1). Furthermore, there was no significant difference in FSGs between those with or without cognitive impairment ($t=0.147, P=0.883$; see Tables 3 and 4).

Association of cognitive impairment with FSI

In the different FSI groups, individuals with a normal insulin level showed a significantly higher MMSE score compared with those in the hypoinsulinemia groups ($t=2.160, P=0.031$) and also showed a non-significant increased prevalence of cognitive impairment ($\chi^2=2.960, P=0.228$; see Table 2). There was also no significant difference in FSI between those with and without cognitive impairment ($t=1.609, P=0.108$; see Table 3). Hypoinsulinemia nonsignificantly increased the risk of cognitive impairment (see Table 5).

### Discussion

This study examined the association between FSG or FSI and cognitive impairment among Chinese nonagenarians/centenarians. Using cross-sectional observations, nonagenarians/centenarians from one community who had no previous diagnosis of diabetes

### Table 3 Baseline demographics based on cognitive function (n=661)

| Demographic          | All (n=661) | Non-cognitively impaired (n=252) | Cognitive impairment (n=409) | $\chi^2$ or $t$ | $P$ value |
|----------------------|------------|----------------------------------|----------------------------|----------------|-----------|
| Age (years)          | 93.52±3.37 | 93.73±3.40                       | 93.17±3.28*                | 2.058          | 0.040     |
| Gender (male/female) | 216/444    | 127/125                          | 90/319**                   | 56.16          | <0.0001   |
| FSG (mmol/L)         | 4.46±1.45  | 4.45±1.52                        | 4.47±1.40                  | 0.147          | 0.883     |
| FSI                  | 6.27±2.27  | 6.45±2.39                        | 6.16±2.18                  | 1.609          | 0.108     |
| DBP (mmHg)           | 72.59±12.06| 72.08±12.08                      | 72.91±12.05                | 0.864          | 0.388     |
| SBP (mmHg)           | 139.71±23.31| 139.71±23.31                    | 140.07±22.63               | 0.199          | 0.862     |
| TG (mmol/L)          | 1.22±0.64  | 1.19±0.54                        | 1.24±0.70                  | 1.074          | 0.283     |
| TC (mmol/L)          | 4.15±0.85  | 4.06±0.86                        | 4.20±0.84*                 | 2.098          | 0.036     |
| HDL (mmol/L)         | 1.58±0.59  | 1.59±0.66                        | 1.57±0.54                  | 0.759          | 0.222     |
| LDL (mmol/L)         | 2.28±0.97  | 2.27±0.59                        | 2.28±1.17                  | 0.240          | 0.810     |
| SUA (μmol/L)         | 318.72±87.01| 321.17±92.73                    | 316.88±82.53               | 0.623          | 0.534     |
| BMI                  | 19.05±3.64 | 18.93±3.25                       | 19.11±3.85                 | 0.629          | 0.530     |

Baseline demographics were compared between groups with differing cognitive function scores. A $P$ value of $<0.05$ was considered statistically significant.

SUA serum uric acid, BMI body mass index, FSG fasting serum glucose, FSI fasting serum insulin, HDL high density lipoprotein, LDL low density lipoprotein, TC total cholesterol, TG triglyceride

* $P<0.05$; ** $P<0.01$ vs. non-cognitively impaired using $\chi^2$ or Fisher’s exact tests (where an expected cell count was $<5$) for categorical variables and unpaired Student’s $t$ tests for continuous variables.
showed a high prevalence of cognitive impairment and hypoinsulinemia. The prevalence of impaired fasting glucose was 4.99%, and that of diabetes was 6.20%. This study showed that among Chinese nonagenarians/centenarians, cognitive function correlated with FSI and not FSG. Higher insulin levels may be a preventative factor for cognitive impairment.

Insulin sensitivity declines and insulin resistance increases with aging (Lin et al. 2011; Frisardi et al. 2010; Barzilai et al. 2012; Penninx et al. 2009). In this study, the prevalence of hypoinsulinemia was almost a third higher (34.53%) than in adults in general including those aged over 60 years. However, in this sample where a high prevalence of hypoinsulinemia existed, only 12.11% of subjects showed abnormally high FSGs. If a high prevalence of insulin resistance exists, those with hypoinsulinemia are at risk for hyperglycosemia; thus, the prevalence of insulin resistance in this sample is low.

In general, it is thought that insulin resistance and hyperinsulinemia increase with aging (Lin et al. 2011; Frisardi et al. 2010; Barzilai et al. 2012; Penninx et al. 2009), that in the elderly, obesity- and insulin-resistant individuals are more long lived, and that mortality in the elderly could lead to the retention of obesity- and insulin-resistant individuals (Lamming et al. 2012; Blanchard 2005; Andres 1980). However, our results in this study of long-lived subjects provide contradictory evidence. To the best of our knowledge, this study is the only study to focus on FSI and FSG levels in nonagenarians/centenarians. These data indicate that the effectiveness and production of insulin in long-lived subjects differs from adults in general. The mechanisms underlying these differences in long-lived subjects compared with the elderly in general should be examined in future studies.

In this study, the prevalence of both hypoinsulinemia and cognitive impairment was high, FSI significantly and positively correlated with cognitive score, and subjects with hypoinsulinemia showed lower cognitive function scores than those with normal insulin levels. Together, these data show that insulin may play a key role in cognitive function (Zhong et al. 2012; Shemesh et al. 2012). Hypoinsulinemia may underlie cognitive impairment in long-lived subjects. Previous studies provided evidence of the role insulin played in cognitive processes; specifically, in hippocampal function, training on a spatial memory task increased hippocampal insulin receptor expression and hypothalamic insulin-regulated memory processes (Costello et al. 2012; McNay and Recknagel 2011). Insulin receptors are also found in the amygdala and other brain regions involved in cognitive processes (Huang et al. 2010). Our study

| Table 4 Odds risk (OR) of abnormal fasting serum glucose (FSG) on the prevalence of cognitive impairment in long-lived subjects |
|---------------------------------|---------------------------------|-----------------|
| Normal (n=587) | Impaired fasting glucose (n=33) | Diabetes (n=41) |
| Number of cases | Unadjusted OR | 1 (reference) | 1.047 (0.543–2.019) | 1.631 (0.642–4.146) |
| | Model 1, adjusted OR | 1 (reference) | 0.906 (0.450–1.827) | 1.125 (0.410–1.084) |
| | Model 2, adjusted OR | 1 (reference) | 1.009 (0.508–2.007) | 1.416 (0.543–3.694) |
| | Model 3, adjusted OR | 1 (reference) | 0.948 (0.639–1.405) | 1.129 (0.365–3.496) |

Unadjusted a Wald χ² test with df=1 was used (multiple logistic regression was used to adjust for covariates), Model 1 adjustment made using the general covariates age, gender, and educational level, Model 2 adjustment made using components of metabolic syndrome (serum lipid/lipoprotein, arterial blood pressure, body mass index, and serum uric acid), Model 3 adjustment made using all covariates

| Table 5 Odds risk (OR) of hypoinsulinemia on the prevalence of cognitive impairment in long-lived subjects |
|---------------------------------|---------------------------------|
| Hypoinsulinemia | Normal insulin |
| N=211 | N=450 |
| Number of cases | Unadjusted OR | 1 (reference) | 0.777 (0.552–1.093) |
| | Model 1, adjusted OR | 1 (reference) | 0.907 (0.627–1.313) |
| | Model 2, adjusted OR | 1 (reference) | 0.832 (0.578–1.198) |
| | Model 3, adjusted OR | 1 (reference) | 0.948 (0.639–1.405) |

OR odds ratio, Unadjusted a Wald χ² test with df=1 was used (multiple logistic regression was used to adjust for covariates), Model 1 adjustment made using the general covariates age, gender, and educational levels, Model 2 adjustment made using components of metabolic syndrome (serum lipid/lipoprotein, arterial blood pressure, body mass index, and serum uric acid), Model 3 adjustment made using all covariates
provides important evidence from long-lived subjects in a clinic trial setting; this was the interesting finding of this study.

In our sample, the cognitive score was very low; 41 of 661 subjects showed a normal score, the mean score was only 15.48, and the prevalence of serious cognitive impairment was 61.8 % (almost two thirds). It has been confirmed that individuals with cognitive impairment show greater than double mortality rate than those without cognitive impairment (Cano et al. 2012; Sachs et al. 2011). Insulin is an important hormone with essential physiologic functions, and the high prevalence of hypoinsulinemia may be a manifestation of the deterioration in pancreatic island function with age. Individuals with hypoinsulinemia may also show a high rate of mortality (Langer et al. 1986). Therefore, due to this increased rate of mortality, at advanced ages, only those with both normal cognitive function and normal insulin levels remain. This theory is conducive with the finding that hypoinsulinemia correlated with cognitive impairment. However, the high mortality rate could underlie the phenomenon that more individuals with cognitive impairment show hypoinsulinemia but cannot explain the fact that FSI was significantly and positively correlated with cognitive scores. Thus, insulin must play a role in cognitive function.

Our study had some limitations that must be considered. First, 870 subjects aged 90 years or older volunteered for the PLAD Study. Among these 870 volunteers, only 661 had nonmissing data for the two main variables involved in this current analysis. Second, due to the cross-sectional nature of this study, the subjects might change their diets and the conditions related to the association of FSG and FSI with cognitive impairment. However, all of the nonagenarians/centenarians had lived for long periods in the country in southwest China (an underdeveloped area in China), and in that area, the diet was typically vegetarian. The lifestyles and food habits of the nonagenarians/centenarians were relatively consistent and similar. Their diets and conditions were similar; thus, it was impossible that diet and conditions affected the results of this study. Third, due to the cross-sectional nature of this study, we could not conduct analyses of causes. Fourth, because this study is part of the PLAD study, a survival bias may exist. However, this issue is inherent in a study of individuals of this age group. Finally, there was a difference in the prevalence of cognitive impairment between genders. Gender may influence the association of FSG or FSI with cognitive impairment.

Data analysis based on gender may remove this influence. However, this analysis could reduce the sample size thereby preventing the generation of a conclusion.

In summary, we observed that among long-lived subjects, cognitive function appeared to correlate with FSI, and higher FSI was associated with enhanced cognitive function. Insulin and its role in cognitive function must underline this finding.

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