Association of anthropometry and weight change with risk of dementia and its major subtypes

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Association of anthropometry and weight change with risk of dementia and its major subtypes: A meta-analysis consisting 2.8 million adults with 57,294 cases of dementia

Crystal ManYing Lee1,2 | Mark Woodward3,4,5 | G. David Batty6,7 | Alexa S. Beiser8,9,10 | Steven Bell11,12,13 | Claudine Berr14,15 | Espen Bjertness16 | John Chalmers4 | Robert Clarke17 | Jean-Francois Dartigues18 | Kendra Davis-Plourde8,10 | Stéphanie Debette19 | Emanuele Di Angelantonio11,12,13 | Catherine Feart20 | Ruth Frikke-Schmidt21,22 | John Gregson23 | Mary N. Haan24 | Linda B. Hassing25 | Kathleen M. Hayden26 | Marieke P. Hoevenaar-Blom27 | Jaakko Kaprio28,29 | Mika Kivimäki6,29 | Georgios Lappas30 | Eric B. Larson31 | Erin S. LeBlanc32 | Anne Lee24 | Li-Yung Lui33 | Eric P. Moll van Charante34 | Toshiharu Ninomiya35 | Liv Tybjærg Nordestgaard21,22 | Tomoyuki Ohara35,36 | Toshiaki Ohkuma4 | Teemu Palviainen28 | Karine Peres20 | Ruth Peters37,38,39 | Nawab Qizilbash23,40 | Edo Richard41,27 | Annika Rosengren30,42 | Sudha Seshadri9,10,43 | Martin Shipley6 | Archana Singh-Manoux44 | Bjorn Heine Strand45,46,47,48 | Willem A. van Gool27 | Eero Vuoksimaa28 | Kristine Yaffe49 | Rachel R. Huxley4,50,51

1School of Psychology and Public Health, La Trobe University, Melbourne, Victoria, Australia
2Boden Institute of Obesity, Nutrition, Exercise & Eating Disorders, University of Sydney, Sydney, New South Wales, Australia
3The George Institute for Global Health, University of Oxford, Oxford, UK
4The George Institute for Global Health, University of New South Wales, Sydney, New South Wales, Australia
5Department of Epidemiology, Johns Hopkins University, Baltimore, Maryland, USA
6Department of Epidemiology and Public Health, University College London, London, UK
7School of Biological & Population Health Sciences, Oregon State University, Corvallis, Oregon, USA
8Department of Biostatistics, Boston University School of Public Health, Boston, Massachusetts, USA
9Department of Neurology, Boston University School of Medicine, Boston, Massachusetts, USA
10Framingham Heart Study, Framingham, Massachusetts, USA
11The National Institute for Health Research Blood and Transplant Unit in Donor Health and Genomics, Strangeways Research Laboratory, University of Cambridge, Cambridge, UK
12UK Medical Research Council/British Heart Foundation Cardiovascular Epidemiology Unit, Department of Public Health and Primary Care, Strangeways Research Laboratory, University of Cambridge, Cambridge, UK
13British Heart Foundation Centre of Excellence, Division of Cardiovascular Medicine, Addenbrooke’s Hospital, Cambridge, UK
14INSERM, U1061, Neuropsychiatry: Epidemiological and Clinical Research, University of Montpellier, Montpellier, France

Abbreviations: BMI, body mass index; CI, confidence intervals; HR, hazard ratio; WC, waist circumference.

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Uncertainty exists regarding the relation of body size and weight change with dementia risk. As populations continue to age and the global obesity epidemic shows no sign of waning, reliable quantification of such associations is important. We examined the relationship of body mass index, waist circumference, and annual percent weight change with risk of dementia and its subtypes by pooling data from 19 prospective cohort studies and four clinical trials using meta-analysis. Compared with body mass index–defined lower-normal weight (18.5-22.4 kg/m²), the risk of all-cause dementia was higher among underweight individuals but lower among those with upper-normal...
1 | INTRODUCTION

Dementia, a disease primarily of aging, affects an estimated 47 million people globally. It is a heterogeneous condition chiefly comprising Alzheimer disease (60-70% of cases) and vascular dementia accounting for about 15% of cases, although the two subtypes frequently co-occur. Aging, family history, and sarcopenia are important risk factors for dementia, and there is growing evidence that vascular risk factors, such as diabetes, may also confer increased risk, particularly for vascular dementia, although findings are inconsistent.

Excess body weight, typically defined as having a high body mass index (BMI), has been causally linked to a large number of chronic conditions, particularly vascular disease. While the relationship between BMI and dementia has been the subject of several large-scale epidemiological studies, the findings have been inconsistent: some studies have reported that only individuals at the extreme ends of the body size spectrum (ie, underweight and obese) experience an increased dementia risk, while others have documented positive and even inverse associations. Measures of central obesity, such as waist circumference (WC), have been argued to be more informative measures of obesity-related risk compared with BMI. Currently, little is known about the association between central obesity and dementia risk.

As nonvascular and vascular dementia have different pathophysiological mechanisms, any association with body size may similarly differ according to endpoint. Distinguishing between possible dementia subtypes in any analysis with measures of body size may, therefore, prove informative in explaining some of the observed heterogeneity. Further, whether sex differences exist, or if the association between body size and dementia risk differs in middle and later life remains unclear.

In this meta-analysis, we examined the relationships of body size with all-cause dementia and possible vascular and nonvascular dementia by sex and baseline age in participants free of dementia who had their body size assessed at baseline and were later followed up on dementia status. Using repeat measures, where available, we explored the association between standardized annual weight change during follow-up with subsequent dementia risk.

2 | METHODS

The investigators of studies that were either identified from previous systematic reviews and meta-analyses or who were known to...
Dementia Pooling Project collaborators, were contacted and asked to contribute results for their studies (Figure S1). Seven of the 14 studies from the Dementia Pooling Project contributed data that were included in the analyses. Additional relevant data from 28 studies were identified from five previous systematic reviews and meta-analyses of the association. Fourteen studies that had not contributed to previous overviews were identified through a PUBMED search by one investigator (CMYL). This was limited to human subjects and the period up to 1 September 2017, based on the search strategy: [(body mass index OR body weight OR obesity OR waist circumference) AND dementia] OR (clinical trial AND incident dementia). Studies were deemed eligible for inclusion if BMI or WC were collected at baseline and dementia status was available at follow-up. Subsequently, 42 invitations were issued inviting study investigators to collaborate in this pooling study. Of these, 10 studies contributed results. Collaborators provided information on a further eight studies that had not been identified by the literature search, or which had not been included in previous overviews, as the data were unpublished; six of these studies provided data. A total of 23 studies responded (n = 2 790 753).

BMI was calculated as weight in kilogrammes divided by height in metres squared. Directly measured (n = 2 760 602; 98.9%) and self-report (n = 30 151; 1.1%) height and weight were used in the calculation. WC was measured either at the midpoint between rib and iliac crest (n = 217 051; 29.9%), narrowest waist (n = 9768; 1.3%), or narrowest waist or umbilicus (n = 486 275; 67.0%). Annual percent weight change was calculated as follows:

\[
\frac{100 \times (\text{last body weight measure prior to dementia follow up} - \text{baseline body weight measure})}{\text{baseline body weight measure} - \text{date at last body weight measure} - \text{date at baseline body weight measure}} \times \frac{365.25}{365.25}
\]

WC was measured either at the midpoint between rib and iliac crest (n = 217 051; 29.9%), narrowest waist (n = 9768; 1.3%), or narrowest waist or umbilicus (n = 486 275; 67.0%). Annual percent weight change was calculated as follows:

Five BMI categories were distinguished: underweight: less than 18.5 kg/m²; lower-normal weight: 18.5-22.4 kg/m²; upper-normal weight: 22.5-24.9 kg/m²; overweight: 25.0-29.9 kg/m²; and obese: greater than or equal to 30.0 kg/m². Dementia endpoints were defined by investigators of each individual study. Ascertained of dementia was by medical examination in 12 studies (Table 1). These studies classified dementia based solely on the Diagnostic and Statistical Manual of Method Disorders criteria or in combination with the National Institute of Neurological and Communication Disorders and Stroke and the Alzheimer’s Disease and Related Disorders Association criteria for Alzheimer disease or the Association Internationale pour la Recherche et l’Enseignement en Neurosciences criteria for vascular dementia.

Other methods used in dementia ascertainment included health records (two studies), death records (five studies), and death and health or hospitalization records (four studies). Ten of these studies that used other methods to ascertain dementia classified the disorder based on the International Classification of Diseases and one study used Read Codes to classify dementia (Table 1).

### 2.1 Data analysis

Sex-specific hazard ratios (HRs) and 95% confidence intervals (CIs) were obtained for all-cause dementia in relation to (a) each of five BMI categories, with lower-normal weight as the referent group; (b) each fifth of WC, with the first fifth as the referent group; and (c) each of three annual percent weight change categories (greater than or equal to 0.5% annual weight loss, less than 0.5% annual weight change, and greater than or equal to 0.5% annual weight gain), with less than 0.5% annual weight change as the referent group. Following a pre-specified common analytic protocol, effect estimates were adjusted for age (model 1); age, smoking, and education or socioeconomic status (model 2; which was the primary model—used in the reporting of outcomes herein); and age, smoking, education or socioeconomic status, diabetes, systolic blood pressure, total cholesterol, blood pressure-lowering medication, cholesterol-lowering medication, and glucose-lowering medication where available (model 3). For stud-
| Study                                      | Country     | Baseline Year | Baseline Age (years) | N (% female) | Dementia Cases | Dementia Ascertainment | Dementia Criteria | Anthropometric Measurement (waist protocol) |
|--------------------------------------------|-------------|---------------|----------------------|--------------|----------------|------------------------|------------------|---------------------------------------------|
| Adult Changes in Thought study\(^{16}\)    | USA         | 1994-1996     | ≥65                  | 4343 (58.3)  | 1096           | Medical examination    | DSM-IV, NINCDS-ADRDA | Measured (narrowest waist)                  |
| Action in Diabetes and Vascular Disease Preteraz and Diaclonic MR Controlled Evaluation trial\(^{17}\) | International | 2001-2003     | 55-88               | 11 136 (42.5) | 109             | Medical examination    | DSM-IV            | Measured (midpoint between rib and iliac crest) |
| Aging Multidisciplinary Investigation (AMI) cohort\(^{18}\) | France      | 2007          | ≥65                  | 563 (38.7)   | 65             | Medical examination    | DSM-III-R, NINCDS-ADRDA, NINDS-AIREN | Measured (midpoint between rib and iliac crest) |
| The Copenhagen City Heart Study\(^{19}\)   | Denmark     | 1976-1978     | ≥20                  | 9037 (55.7)  | 969            | Health record          | ICD-8, ICD-10     | Measured (umbilicus)                        |
| Cache County Memory Study\(^{20}\)        | USA         | 1995          | ≥65                  | 3185 (57.4)  | 507            | Medical examination    | DSM-III-R, NINCDS-ADRDA, NINDS-AIREN | Self-report                   |
| The Copenhagen General Population Study\(^{21}\) | Denmark     | 2003          | ≥20                  | 10 4506 (55.2) | 1906          | Health record          | ICD-8, ICD-10     | Measured (midpoint between rib and iliac crest) |
| Clinical Practice Research Datalink\(^{9}\) | UK          | 1992-2007     | ≥40                  | 19 9819 (54.8) | 45507         | Health and death record | Read Codes       | Measured                                    |
| Finnish Twin Cohort\(^{22}\)             | Finland     | 1975          | ≥18                  | 25 814 (51.1) | 960            | Death record           | ICD-8, ICD-9, ICD-10 | Self-report                   |
| Framingham Heart Study\(^{23}\)          | USA         | 1992-1996, 1998-2001 | ≥60                  | 2232 (56.0)  | 289            | Medical examination    | DSM-IV, NINCDS-ADRDA, NINDS-AIREN | Measured (umbilicus)                        |
| General Post Office Study\(^{24}\)       | UK          | 1966-1967     | 35-70                | 1385 (37.0)  | 18             | Death record           | ICD-8, ICD-9, ICD-10 | Measured                                    |
| Hisayama Study\(^{25}\)                  | Japan       | 1988          | ≥60                  | 1192 (58.3)  | 350            | Medical examination    | DSM-III-R, NINCDS-ADRDA, NINDS-AIREN | Measured (umbilicus)                        |
| Health Survey for England and Scottish Health Survey\(^{26}\) | UK          | 1995-2008     | 16-99                | 90 685 (54.7) | 524            | Death record           | ICD-9, ICD-10     | Measured (midpoint between rib and iliac crest) |
| Hypertension in the Very Elderly Trial\(^{27}\) | International | 2000          | ≥80                  | 3337 (60.4)  | 263            | Medical examination    | DSM-IV            | Measured                                    |
| Norwegian Counties Study\(^{28}\)        | Norway      | 1974-1978     | 35-50                | 40 978 (50.6) | 1173           | Death record           | ICD-9, ICD-10     | Measured                                    |
| Origins of Variance in the Old-Old\(^{29}\) | Sweden      | 1963          | 45-65                | 1152 (69.0)  | 312            | Health record and interview | DSM-III-R, NINCDS-ADRDA, NINDS-AIREN | Self-report                   |
| Prevention of Dementia by Intensive Vascular Care trial\(^{30}\) | The Netherlands | 2006-2009    | 70-78                | 3526 (54.4)  | 233            | Medical examination    | DSM-IV            | Measured (midpoint between rib and iliac crest) |

(Continues)
## Table 1 (Continued)

| Study | Country | Baseline Year | Baseline Age (years) | N (% female) | Dementia Cases | Dementia Ascertainment | Dementia Criteria | Anthropometric Measurement (waist protocol) |
|-------|---------|---------------|----------------------|--------------|----------------|------------------------|-----------------|---------------------------------------------|
| Primary Prevention Study<sup>31</sup> | Sweden | 1970-1973 | 45-55 | 7394 (0) | 788 | Death and hospitalization records | ICD-8, ICD-9, ICD-10 | Measured |
| The Perindopril Protection Against Recurrent Stroke Study<sup>32</sup> | International | 1995-1997 | 26-91 | 5865 (29.7) | 380 | Medical examination | DSM-IV | Measured |
| Study of Osteoporotic Fractures<sup>33</sup> | USA | 1986-1988 | $\geq$ 65 | 1019 (100) | 232 | Medical examination | DSM-IV | Measured (narrowest waist) |
| Three City Study<sup>34</sup> | France | 1999-2000 | $\geq$ 65 | 6721 (61.4) | 832 | Medical examination | DSM-IV | Measured (midpoint between rib and iliac crest) |
| UK Biobank<sup>35</sup> | UK | 2006-2010 | 39-74 | 48 627 (54.6) | 344 | Death and hospitalization records | ICD | Measured (narrowest waist or umbilicus) |
| Whitehall I Study<sup>36</sup> | UK | 1967-1969 | 40-69 | 17 167 (0) | 288 | Death record | ICD-8, ICD-9, ICD-10 | Measured |
| Whitehall II Study<sup>37</sup> | UK | 1985-1988 | 35-55 | 5050 (28.3) | 149 | Death and hospitalization records | ICD-10 | Measured (narrowest waist) |

Abbreviations: DSM-III, Diagnostic and Statistical Manual of Mental Disorders third edition criteria; DSM-III-R, Diagnostic and Statistical Manual of Mental Disorders third edition revised criteria; DSM-IV, Diagnostic and Statistical Manual of Mental Disorders fourth edition criteria; ICD-8, International Classification of Diseases eighth revision; ICD-9, International Classification of Diseases ninth revision; ICD-10, International Classification of Diseases tenth revision; and Related Health Problems; NINCDS-ADRDA, National Institute of Neurological and Communication Disorders and Stroke and the Alzheimer's Disease and Related Disorders Association criteria; NINDS-AIREN, National Institute of Neurological Disorders and Stroke and Association Internationale pour la Recherche et l'Enseignement en Neurosciences criteria;
RESULTS

Results from 23 studies comprised 2,790,753 participants (54% women) in whom 57,294 cases of all-cause dementia were accumulated during a mean of 9.6 years of disease surveillance. Of these cases, 6,792 were classed as nonvascular dementia, and 1214 were recorded as vascular dementia. Of the studies included, information from 14 cohort studies (n = 728,959; 26.1%) were previously unpublished, and four were from clinical trial populations (n = 23,864; 0.9%). Most studies were from Europe (n = 2,758,444; 98.8%), with one study contributing more than 70% of participants (79% cases of all-cause dementia). Study characteristics are shown in Tables S1 and S2. In brief, mean age at baseline ranged from 36 to 83 years in men and 37 to 84 years in women. Mean duration of follow-up varied from 4 to 38 years, with an overall median of 9 years. Mean BMIs at study baseline were 21.9 to 28.2 kg/m² in men and 22.4 to 28.8 kg/m² in women; for WC, the corresponding results were 96.9 to 100.7 and 80.7 to 97.5 cm.

| All-cause dementia | BMI (kg/m²) | N (cases) | HR (95% CI) | I² |
|--------------------|-------------|-----------|-------------|----|
|                    | <18.5       | 44408 (2510) | 1.26 (1.20, 1.31) | 0% |
|                    | 18.5-22.4   | 431624 (12110) | 1.00 | |
|                    | 22.5-24.9   | 583832 (13402) | 0.85 (0.83, 0.88) | 0% |
|                    | 25.0-29.9   | 1067974 (20573) | 0.86 (0.81, 0.92) | 65.5% |
|                    | ≥30.0       | 661475 (8675) | 0.91 (0.83, 1.01) | 78.4% |

| Non-vascular dementia | BMI (kg/m²) | N (cases) | HR (95% CI) | I² |
|-----------------------|-------------|-----------|-------------|----|
|                       | <18.5       | 1601 (127) | 1.28 (1.06, 1.54) | 0% |
|                       | 18.5-22.4   | 27102 (1224) | 1.00 | |
|                       | 22.5-24.9   | 38389 (1649) | 0.89 (0.83, 0.97) | 0% |
|                       | 25.0-29.9   | 61814 (2787) | 0.89 (0.81, 0.98) | 29.1% |
|                       | ≥30.0       | 24073 (1005) | 0.91 (0.78, 1.05) | 49.5% |

| Vascular dementia | BMI (kg/m²) | N (cases) | HR (95% CI) | I² |
|-------------------|-------------|-----------|-------------|----|
|                   | <18.5       | 1443 (34) | 1.76 (1.20, 2.58) | 1.4% |
|                   | 18.5-22.4   | 26624 (212) | 1.00 | |
|                   | 22.5-24.9   | 37490 (294) | 1.00 (0.82, 1.23) | 13.0% |
|                   | 25.0-29.9   | 60046 (490) | 1.03 (0.84, 1.25) | 13.8% |
|                   | ≥30.0       | 23316 (184) | 1.48 (1.10, 2.00) | 38.2% |

FIGURE 1 Association between body mass index (BMI) and incident fatal and nonfatal dementia and its major subtypes. Hazard ratios (HRs) and 95% confidence intervals (CIs) adjusted for age, smoking, and education or socio-economic status.
A non-linear association between BMI with all-cause dementia was observed: compared with the referent group (BMI: 18.5-22.4 kg/m²) those who were underweight had a one-quarter greater risk of dementia (HR: 1.26, 95% CI, 1.20-1.31; Figure 1). This association remained similar after excluding the first 3, 5, or 10 years (median 10 years) of follow-up in a subset of studies (1.35 [1.24-1.46]; Figure S2a; Table S3). Relative to the referent group, individuals with upper-normal BMI, overweight or obesity had a 10% to 15% lower dementia risk (Figure 1). Similar results were obtained after adjustment for age (Figures S3-S6). Additional adjustment for cardiometabolic risk factors did not materially alter the relationship (model 3; Table S4). Neither were results significantly different in a range of sensitivity analyses (Table S5; Figure S7). Further, the associations were comparable between studies with baseline mean age younger than 60 and 60 years and older (Figure S8) and before and after exclusion of the first 3, 5, or 10 years (median 10 years; Figure S2a; Table S3).

As with all-cause dementia, the association between BMI and nonvascular dementia risk was non-linear. Relative to lower-normal BMI, individuals categorized as underweight were at approximately one-quarter increased dementia risk (1.28 [1.06-1.54]; Figure 1; Table S4). Adjustment for cardiometabolic risk factors did not materially affect the association (Table S4). The findings were similarly robust when restricting the analysis to studies that used measured height and weight. Findings were also consistent between clinical trials and observational studies and between studies with baseline mean age younger 60 and 60 years and older (Figures S8 and S9). There was no evidence of a sex difference in the association (Figure 2). Relative to the referent group, individuals with upper-normal BMI, overweight, or obesity had 10% lower dementia risk (Figure 1). After excluding the first few years of follow-up (median 7.5 years), the increased risk of nonvascular dementia in the underweight category remained, but the association was no longer significant in the overweight and obese categories (Figure 2b; Table S3).

Both individuals with underweight or obesity were at increased vascular dementia risk. Compared with the referent group, those who were underweight had an approximate 80% greater dementia risk, and for those in the obese category, the risk was approximately 50% higher (Figure 1). The excess risk among the obese group was largely mediated by cardiometabolic risk factors (HR for model 3: 1.26 [0.87-1.84]; Table S4). Sensitivity analyses indicated findings compatible with the main result (Figure S9). There was no evidence of a sex difference in the association between BMI and vascular dementia risk (Figure 2). The association was broadly similar in studies with baseline mean age younger than 60 and 60 years and older (Figure S8). Excluding the first few years of follow-up (median 7.5 years) did not materially influence the relationship (Figure S2c; Table S3).

A non-linear association was observed between WC and all-cause dementia among the 13 studies (725 522 individuals; 54.5% women; 7057 cases) that contributed to the analysis. Compared with individuals in the lowest fifth of WC, individuals with larger WC had 10% to 22% lower all-cause dementia risk (Figure 3). Similar results were obtained when adjusted for age or after adjustment for cardiometabolic risk factors (Figures S10-S13; Table S6). The estimates tended to be larger for studies that used death records to ascertain dementia status rather than those that used medical examination (Figure S14). Data from clinical trials produced similar results to those from nontrial populations (Figure S15). The estimate of effect was more pronounced for studies with baseline mean age younger than 60 than 60 years and older especially at higher WC categories (Figure S16).

Pooling data from ten studies (5319 nonvascular dementia cases) indicated that, compared with the lowest fifth, all WC categories were associated with lower nonvascular dementia risk (Figure 3; Table S6).

![Figure 2](https://example.com/figure2.png)

**Figure 2** Associations between body mass index (BMI) and incident fatal and nonfatal dementia subtypes by sex. Hazard ratios (HRs) and 95% confidence intervals (CIs) adjusted for age, smoking, and education or socio-economic status.
Adjustment for cardiometabolic risk factors did not alter the association (Table S6). Data from clinical trials produced a similar pattern (Figure S15), and there was little evidence of a sex difference in the association (Figure 4).

For vascular dementia (808 cases), compared with the referent category, individuals in the highest fifth of WC had roughly one-quarter higher risk (1.23 [0.97-1.55]) that was attenuated after further adjustment for cardiometabolic risk factors (Figure 3; Table S6). No sex difference was evident (Figure 4). Data from clinical trial participants contained too few cases to draw meaningful conclusions (Figure S15). Only the highest WC category in studies with baseline mean age 60 years and older was associated with increased vascular dementia risk (Figure S16).

In the analysis of weight change (111 620 individuals; 44.8% female; 5626 dementia cases), compared with individuals who maintained a relatively stable weight during follow-up, individuals with greater than or equal to 0.5% annual weight loss had approximately one-third greater all-cause dementia risk (1.32 [1.18-1.47]; Table 2). In contrast, greater than or equal to 0.5% annual weight gain was not associated with dementia risk (1.00 [0.89-1.12]; Table 2). The results remained unchanged for models 1 and 3, and when studies were stratified by baseline mean age (Figures S17-S19; Table 2). For

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**FIGURE 3** Association between waist circumference and incident fatal and nonfatal dementia and its major subtypes. Hazard ratios (HRs) and 95% confidence intervals (CIs) adjusted for age, smoking, and education or socio-economic status.
greater than or equal to 0.5% annual weight loss, the estimate of effect was more pronounced for studies that ascertained dementia by medical examination than studies that used death records (Figure S20).

As with all-cause dementia, greater than or equal to 0.5% annual weight loss was associated with higher nonvascular dementia risk (1.41 [1.19-1.67]), whereas greater than or equal to 0.5% annual weight gain was not associated with risk (0.99 [0.83-1.19]; Table 2). For vascular dementia, opposing trends were observed: a trend towards increased risk was observed in those with greater than or equal to 0.5% annual weight gain (1.21 [0.98-1.49]) but only in those with baseline mean age younger than 60 years (Figure S19). Weight

<FIGURE 4> Associations between waist circumference and incident fatal and nonfatal dementia subtypes by sex. Hazard ratios (HRs) and 95% confidence intervals (CIs) adjusted for age, smoking, and education or socio-economic status

<TABLE 2> Adjusted hazard ratios (HR) with 95% confidence intervals (CI) of dementia by annual percent weight change

| Dementia Type         | ≥0.5% Weight Loss per Year | Reference | ≥0.5% Weight Gain per Year |
|-----------------------|-----------------------------|-----------|----------------------------|
| All-cause dementia, 15 studies |                             |           |                            |
| N (cases)             | 23 590 (1668)               | 44 425 (2254) | 43 605 (1704)               |
| Model 2<sup>a</sup>  | HR (95% CI) 1.32 (1.18-1.47)| 1.00      | 1.00 (0.89-1.12)            |
| i² (%)                | 46.2                       |           |                            |
| Model 3<sup>b</sup>  | HR (95% CI) 1.28 (1.15-1.42)| 1.00      | 0.99 (0.89-1.11)            |
| i² (%)                | 41.1                       |           |                            |
| Nonvascular dementia, 10 studies |                             |           |                            |
| N (cases)             | 8106 (898)                  | 10 929 (932) | 9926 (652)                  |
| Model 2<sup>a</sup>  | HR (95% CI) 1.41 (1.19-1.67)| 1.00      | 0.99 (0.83-1.19)            |
| i² (%)                | 51.2                       |           |                            |
| Model 3<sup>b</sup>  | HR (95% CI) 1.36 (1.15-1.61)| 1.00      | 0.98 (0.81, 1.17)           |
| i² (%)                | 49.0                       |           |                            |
| Vascular dementia, 7 studies |                             |           |                            |
| N (cases)             | 6000 (151)                  | 9034 (225) | 8275 (181)                  |
| Model 2<sup>a</sup>  | HR (95% CI) 1.11 (0.88-1.39)| 1.00      | 1.21 (0.98-1.49)            |
| i² (%)                | 0                          |           |                            |
| Model 3<sup>b</sup>  | HR (95% CI) 1.09 (0.87-1.36)| 1.00      | 1.13 (0.92-1.40)            |
| i² (%)                | 0                          |           |                            |

<sup>a</sup>Hazard ratio adjusted for age, smoking, education/socio-economic status.

<sup>b</sup>Hazard ratios adjusted for age, smoking, education/socio-economic status, diabetes, systolic blood pressure, total cholesterol, blood pressure lowering medication, cholesterol lowering medication, and glucose lowering medication.
loss was not associated with increased vascular dementia risk (Table 2).

4 | DISCUSSION

This is the largest study to characterize the relationship between measures of body size and weight change with dementia outcomes. We demonstrated that the relationship between body size, weight change, and subsequent risk varies by dementia subtype. When considering all-cause dementia risk, and hence its major subtype of nonvascular dementia, there was no evidence that excess body weight (measured by either BMI or WC) conferred higher risk. Rather, levels of BMI greater than or equal to 22.5 kg/m² (and higher WCs) were associated with a slightly lower dementia risk in later life. Conversely, and in agreement with previous findings, individuals who were categorized as underweight had higher all-cause dementia risk compared with those with lower-normal BMI. For vascular dementia, only the highest levels of BMI and WC were associated with increased risk relative to lower-normal BMI and the lowest fifth of WC.

The main novel finding, however, relates to how weight change appears to influence dementia risk in later life. Relative to weight maintenance, weight loss over follow-up was associated with approximately 30% and 40% increased risk of all-cause dementia and nonvascular dementia, respectively. The association between weight loss and nonvascular dementia may reflect the subclinical expression of prodromal dementia (ie, where an individual has early cognitive impairment but remains functionally independent), or in epidemiological terms, reverse causality. In support of this argument, weight loss has been observed in the preclinical stage of autosomal dominant Alzheimer disease, suggesting that decreasing BMI could be a consequence, rather than a risk factor, of dementia.

An additional, albeit more speculative, explanation may be pathophysiological such as weight loss-induced cortical thinning, as cerebral atrophy is a characteristic of dementia. In a cohort of healthy elderly individuals, faster cognitive decline and accelerated atrophy rate were observed in those with relative weight loss greater than or equal to 5% (equivalent to greater than or equal to 1.2% annual loss) compared with those with relative weight loss less than 5% (equivalent to less than 1.2% annual loss). Similarly, a Norwegian study that assessed percent change in BMI in midlife reported that while greater than or equal to 5% loss (equivalent to approximately greater than or equal to 0.6% annual loss) was associated with increased risk of dementia-related mortality, a gain of greater than or equal to 20% (equivalent to approximately greater than or equal to 2.2% annual gain) was associated with reduced risk.

Conversely, for vascular dementia, weight gain was associated with a modest 20% increased risk but only in those aged younger than 60 years at study baseline. This is consistent not only with what we know about weight gain being a risk factor for other vascular conditions such as coronary heart disease but also with the diminution of the strength in the association between vascular risk factors such as diabetes and blood pressure with vascular risk at older ages. Moreover, data from animal studies have indicated that weight gain is associated with increased vascular dementia risk.

For all-cause dementia, and its major subtype, that of nonvascular dementia, there was no evidence that carrying excess body weight (either in general or more centrally) conferred increased risk. Rather, individuals with a BMI of greater than or equal to 22.5 kg/m² (and higher WCs) had a slightly lower risk of dementia in later life. Conversely, and in agreement with some previous findings, individuals who were categorized as underweight had a one-quarter greater risk of developing all-cause dementia compared with those with lower-normal BMI. When we attempted to exclude those individuals who may have had undetected signs of cognitive impairment at study baseline by excluding the first few years of follow-up, the relationship remained.

To date, evidence from Mendelian randomization studies has provided little support of a relationship between low levels of BMI with future risk of Alzheimer disease, implying that reverse causality or residual confounding may be driving the observed effect in observational studies. However, as Mendelian randomization studies do not use methods that are suited to capturing non-linear associations, the potential for other explanations, other than reverse causality, to explain the observed association remains.

For vascular dementia, the relationship with body size was similar to other vascular conditions with both underweight and obesity conferring increased risk. Underweight individuals had 75% greater vascular dementia risk, which was unaltered by excluding the first 5 years of data. Individuals with obesity had 50% greater risk, possibly due to the adverse effects of high levels of BMI on other vascular risk factors as the relationship was significantly attenuated after adjustment for vascular risk factors. An increased vascular dementia risk was observed for WC but only among those with the highest levels of central obesity.

Recent studies have suggested that the association between BMI and dementia risk is dependent on the age when BMI was assessed. We investigated the association separately for midlife and late life by stratifying studies by their baseline mean age. Aside from the effect of weight gain on vascular dementia risk, our results indicated that there was no evidence of any difference between the two age groups, although the crude dichotomization of age precludes us from definitely concluding that there is no evidence of an age effect in the association between body size and dementia risk. Similarly, this study found no evidence to suggest that the associations reported herein differed between women and men. In addition to its large sample size and number of dementia cases, key strengths of the study included the ability to look at the effect of a measure of central obesity and the influence of weight change on the association between body size and dementia outcomes. BMI and WC were divided into five categories to allow the study of the relationship with dementia in more detail. The lowest fifth of WC was chosen as referent as four studies already included women with abdominal obesity in this group. Nevertheless, using the second or third fifth as referent would not have changed the relationship between WC and dementia. Limitations included the between-study differences in
impaired at study baseline. In our analysis of weight change, we attempted to address the potential for reverse causality by excluding the first 3, 5, or 10 years of follow-up in order to fully negate the effects of inadvertently including individuals with early cognitive impairment at study baseline. In our analysis of weight change, we were unable to distinguish between intentional (eg, because of a diagnosis of hypertension or diabetes in midlife) versus unintentional weight loss (ie, because of pre-existing disease), which may have diluted the observed associations. Studies that contributed to the weight change analysis also varied in the length of time between first and last weight measurement. We standardized weight change across studies by calculating annual percent weight change; however, this method assumes a consistent weight change over time, which may not be valid. In addition, we attempted to distinguish between vascular dementia and nonvascular dementia separately, which may not be valid. In addition, we attempted to distinguish between vascular dementia and nonvascular dementia separately, but the dichotomization is problematic as the two subtypes frequently co-occur. Finally, as body size (both underweight and overweight) is related to a wide range of chronic illnesses, and dementia is a disease of aging, individuals may have died before they had the opportunity to develop dementia. We were unable to apply competing risks methodology in the current analysis, which may have resulted in an underestimation of the relationship between body size and dementia risk. It potentially could also explain why above-normal levels of BMI and WC were not associated with increased all-cause dementia risk.

Excess body weight was not associated with risk of all-cause dementia and its major subtype of nonvascular dementia, whereas it was positively associated with risk of vascular dementia. Underweight was related to increased risk of all-cause dementia and both its subtypes, possibly due to reverse causality. Weight loss in midlife to late life was associated with an increased risk of developing dementia and nonvascular dementia. Future studies should focus on examining the basis between weight loss and increased nonvascular dementia risk to determine if it has a pathophysiological basis or is due to limitations in the epidemiology. Given the known adverse effects of excess body weight on a wide range of health outcomes, from a public health perspective, maintaining a healthy body weight and minimizing weight fluctuation in adult life should continue to be widely promoted.

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CONFLICT OF INTEREST

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outside the submitted work; NQ reported other from pharmaceutical industry outside the submitted work; EV reported grants from The Academy of Finland during the conduct of the study; KY serves on DSMBs for Takeda and Eli Lilly outside the submitted work and is a member of the Beeson Scholars in Aging Scientific Advisory Board and of the Senate of the German Center for Neurodegenerative Diseases.

ORCID
Crystal ManYing Lee https://orcid.org/0000-0001-6613-5491

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SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section at the end of this article.