BMJ Open

Heavier smoking may lead to a relative increase in waist circumference: evidence for a causal relationship from a Mendelian randomisation meta-analysis. The CARTA consortium


ABSTRACT

Objectives: To investigate, using a Mendelian randomisation approach, whether heavier smoking is associated with a range of regional adiposity phenotypes, in particular those related to abdominal adiposity.

Design: Mendelian randomisation meta-analyses using a genetic variant (rs16969968/rs1051730 in the CHRNA5-CHRNA3-CHRNB4 gene region) as a proxy for smoking heaviness, of the associations of smoking heaviness with a range of adiposity phenotypes.

Participants: 148 731 current, former and never-smokers of European ancestry aged ≥16 years from 29 studies in the consortium for Causal Analysis Research in Tobacco and Alcohol (CARTA).

Primary outcome measures: Waist and hip circumferences, and waist-hip ratio.

Results: The data included up to 66 809 never-smokers, 43 009 former smokers and 38 913 current daily cigarette smokers. Among current smokers, for each extra minor allele, the geometric mean was lower for waist circumference by −0.40% (95% CI −0.57% to −0.22%), with effects on hip circumference, waist-hip ratio and body mass index (BMI) being −0.31% (95% CI −0.42% to −0.19%), −0.08% (−0.19% to 0.03%) and −0.74% (−0.96% to −0.51%), respectively. In contrast, among never-smokers, these effects were higher by 0.23% (0.09% to 0.36%), 0.17% (0.08% to 0.26%), 0.07% (−0.01% to 0.15%) and 0.35% (0.18% to 0.52%), respectively. When adjusting the three central adiposity measures for BMI, the effects among current smokers changed direction and were higher by 0.14% (0.05% to 0.22%) for waist circumference, 0.02% (−0.05% to 0.08%) for hip circumference and 0.10% (0.02% to 0.19%) for waist-hip ratio, for each extra minor allele.

Conclusions: This is a very large Mendelian randomisation study of the relationship between smoking and several anthropometric phenotypes relating to regional adiposity. Data included never, former and current smokers from a very wide spectrum of ages among 29 studies. By using a genetic variant associated with smoking heaviness as a proxy for smoking heaviness, bias from confounding is minimised and findings are not affected by reverse causality. Data for direct measures of fat, such as fat mass, and the biomarker leptin were available for only about one fifth of the participants whose height, waist and hip were measured. Participants were exclusively of self-reported European ancestry, and were mostly recruited in European countries.

Strengths and limitations of this study

- This is a very large Mendelian randomisation study of the relationship between smoking and several anthropometric phenotypes relating to regional adiposity.
- Data included never, former and current smokers from a very wide spectrum of ages among 29 studies.
- By using a genetic variant associated with smoking heaviness as a proxy for smoking heaviness, bias from confounding is minimised and findings are not affected by reverse causality.
- Data for direct measures of fat, such as fat mass, and the biomarker leptin were available for only about one fifth of the participants whose height, waist and hip were measured. Participants were exclusively of self-reported European ancestry, and were mostly recruited in European countries.

CrossMark

For numbered affiliations see end of article.

Correspondence to Professor Richard W Morris; richard.morris@bristol.ac.uk

To view the table of contents for this issue, visit http://dx.doi.org/10.1136/bmjopen-2015-008808
INTRODUCTION
Tobacco is the single most important cause of preventable death globally: one in two young people taking up lifelong cigarette smoking will die of causes related to it. Enormous efforts have gone into developing interventions for smoking cessation. Spontaneous cessation rates are low due to the high proportion of smokers who are dependent on nicotine, and effective treatments are still not widely available. One barrier to smoking cessation is the fear of weight gain. In a study of almost 2000 smokers in the USA, recruited into a trial of bupropion and/or nicotine inhalers to promote cessation, 50% of female and 26% of male smokers reported that gaining weight discouraged them from trying to quit,\(^2\) while among adults in Finland, daily smokers were found to report more weight concerns than former smokers or occasional smokers.\(^3\)

A genetic variant in the chromosome 15 \textit{CHRNA5}-\textit{CHRN}\textit{A3-CHRN}\textit{B4} gene region (rs16969968) codes for a functional amino acid change D398N in the nicotinic receptor \(\alpha\) 5 subunit. The SNP rs16969968, which is in perfect linkage disequilibrium with SNP rs1051730 in European populations, is associated with smoking quantity among smokers.\(^4\) The minor allele of this variant is associated with an average increase in smoking amount of one cigarette per day in smokers and increases in cotinine (a metabolite of nicotine) levels.\(^5\)\(^6\) It has also been found that the variant was associated with a lower mean body mass index (BMI),\(^7\)\(^8\)\(^9\) thus adding evidence that heavier smoking leads to lower BMI. The latter study also noted lower waist and hip circumferences among smokers with the variant.\(^5\) However, prior observational evidence suggests that waist circumference and waist-hip ratio may be higher in smokers than in non-smokers. This indicates by lower hip circumferences in smokers.\(^10\) It has also been observed that smoking in adolescence predicts abdominal obesity in adulthood.\(^11\) Moreover, heavy smokers exhibit greater central adiposity than light smokers, based on an analysis of middle-aged smokers of European ancestry.\(^12\) These studies suggest that smoking leads to a central fat accumulation at the expense of peripheral fat loss, particularly in women.\(^13\) In addition, there are also suggestions that smoking may lead to loss of muscle mass as indicated by lower hip circumferences in smokers. This is of high public health relevance in view of the reportedly greater impact of increased central adiposity both on mortality\(^14\)\(^15\) and on the development of diabetes, especially among women.\(^16\)\(^17\) and since smoking is associated with an increased risk of type 2 diabetes.\(^18\)

We previously used Mendelian randomisation methods to investigate the effect of smoking quantity on BMI.\(^7\)\(^9\) This method exploits Mendel’s laws concerning the random assortment of alleles at the time of gamete formation so that individuals are allocated at random to having 0, 1 or 2 alleles in the rs1051730/rs16969968 genotype. The effect of this genotype on smoking quantity among smokers has been demonstrated,\(^9\) and thus the inverse relationship between allele count and BMI is not subject to effects of confounding and reverse causality. Using a substantial pool of studies in the consortium for Causal Analysis Research in Tobacco and Alcohol (CARTA), we have extended our use of Mendelian randomisation methods to examine the effect of smoking quantity on a range of adiposity phenotypes. We test the hypotheses that (1) phenotypes representing central adiposity are affected by smoking quantity differentially from other phenotypes, and (2) these effects are more marked among women than among men.

METHODS
Study populations
We used data on individuals (\(\geq 16\) years) of self-reported European ancestry from 29 studies from the CARTA consortium (http://www.bris.ac.uk/expsych/research/brain/targ/research/collaborations/carta/): the 1958 Birth Cohort (1958BC), the Avon Longitudinal Study of Parents and Children (ALSPAC, including both mothers and children), the British Regional Heart Study (BRHS), the British Women’s Heart and Health Study (BWHHS), the Caerphilly Prospective Study (CaPS), the Christchurch Health and Development Study (CHDS), CoLaus, the Danish Monica study (Dan-MONICA), the Exeter Family Study of Child Health (EFSOCH), the English Longitudinal Study of Ageing (ELSA), the National FINRISK studies, GEMINAKAR, GS:SFHS (Generation Scotland: Scottish Family Health Study), the Genomics of Overweight Young Adults (GOYA) females, GOYA males, the Helsinki Birth Cohort Study (HBCS), Health2006, Health2008, the Nord-Trøndelag Health Study (HUNT), Inter99, MIDSPAN, the Northern Finland Birth Cohorts (NFBC 1966 and NFBC 1986), the National Health and Nutrition Examination Survey (NHANES), the MRC National Survey of Health & Development (NSHD), the Netherlands Twin Register (NTR), the PROspective Study of Pravastatin in the Elderly at Risk (PROSPER) and Whitehall II. All studies received ethics approval from the local research ethics committees. Further details of these studies are provided in online supplementary material.

Genotype
Within each study, individuals were genotyped for one of two single nucleotide polymorphisms (SNPs) in the \textit{CHRNA5-A3-B4} nicotinic receptor subunit gene cluster, either rs16969968 or rs1051730. These SNPs are in perfect linkage disequilibrium with each other in Europeans (\(R^2=1.00\) in HapMap 3, http://hapmap.ncbi.nlm.nih.gov/) and therefore represent the same genetic signal. Where studies had data available for both SNPs, we used the SNP that was genotyped in the largest number of individuals. Details of genotyping methods within each study are provided in online supplementary material.

Adiposity measures
Direct physical measurements included weight, height, waist and hip circumferences, arm circumference,
triceps skinfold and subscapular skinfold thickness. Fat mass and fat-free mass were available from bioimpedance measurements, while leptin and adiponectin were the two biochemical markers related to fat mass.

**BMI** (weight/height^2^) and waist-hip ratio (waist/hip) were calculated.

Waist circumference and waist-hip ratio were taken as key measures of central adiposity, while BMI acted as a non-specific measure of adiposity for purposes of adjustment in regression analysis.

**Smoking status**

Smoking status was self-reported (either by questionnaire or interview) at the same time as regional adiposity measures for all studies, with the exception of 1958 BC (see online supplementary material). Individuals were classified as current, former, ever (ie, current and former combined) or never cigarette smokers. Where information on pipe and cigar smoking was available, individuals reporting being current or former smokers of pipes or cigars but not cigarettes were excluded from all analyses.

For studies with adolescent populations (ALSPAC children and NFBC 1986), analyses were restricted to current daily smokers who reported smoking at least one cigarette per day (current smokers) and individuals who had never tried smoking (never-smokers).

**Statistical analysis**

Analyses were conducted within each contributing study using Stata (Stata Corp, College Station, Texas, USA) and R (R Foundation for Statistical Computing, Vienna, Austria. http://www.R-project.org) software, following the same analysis plan. Analyses were restricted to individuals with full data on smoking status and rs1051730/rs16969968 genotype, and having data on at least one of the regional adiposity phenotypes.

Within each study, genotype frequencies were tested for deviation from the Hardy Weinberg Equilibrium (HWE) using a χ^2^ test. Mendelian randomisation analyses of the association between rs1051730/rs16969968 and each regional adiposity phenotype were performed using linear regression, stratified by smoking status (never, former and current) and sex, and adjusted for age. Apart from height, natural logarithmic transforms were taken of every anthropometric phenotype. An additive genetic model was assumed on log values, so that each effect size could be exponentiated to represent the percentage increase per minor (risk) allele. These analyses were presented separately for each smoking status category. All phenotypic measures were further adjusted for log(BMI) (apart from weight, height and BMI itself), thus assessing the effect of the particular adiposity measure after adjusting for this global weight measure. Log(weight) was adjusted for height instead of log(BMI). Since adjustment for ratio variables in anthropometric studies has been criticised,^{19} we further adjusted waist circumference for log(weight) and height. Finally, we repeated analysis of waist circumference adjusted for BMI restricted to participants with BMI under 30 kg/m^2^; 95% CIs have been quoted for all effect sizes.

Meta-analysis was also carried out of the relationship between reported daily cigarette consumption and rs1051730/rs16969968 genotype, among current smokers.

Although analyses were carried out separately for males and females, the estimates were combined where no evidence for separate sex effects was seen. For NHANES, which has a survey design, Taylor series linearisation was implemented to estimate variances. For studies including related family members, appropriate methods were used to adjust SEs: in GEMINAKAR, twin pair identity was included as a cluster variable in the model; in MIDSPAN, linear mixed effects regression models fitted using restricted maximum likelihood were used to account for related individuals, while in NTR, only unrelated individuals were included. ALSPAC mothers and children were analysed as separate samples; as there are related individuals across these samples, sensitivity analyses were performed excluding each of these studies in turn.

Results from individual studies were meta-analysed in Stata (V13) using the ‘metan’ command from Stata. Where there was evidence of heterogeneity between studies (I^2^ >50%), it was planned that both fixed and random effects analyses would be performed: however, as this never occurred, results for fixed effects analysis only are shown. Meta-regression analysis, using the ‘metareg’ command from Stata, was used to examine whether SNP effects varied by smoking status or by sex, or by a smoking by sex combination.

**RESULTS**

**Descriptive statistics**

The maximum sample size available, with genotype recorded, was 148 751 for weight, height and BMI over 29 studies. The data on individuals with weight, height, smoking status and genotype recorded included 66 809 never-smokers, 43 009 former smokers and 38 913 current smokers. Waist circumference was available in 28 studies (n=142 381), and hip circumference and waist-hip ratio in 25 studies (n=139 667). Measures of fat mass and fat-free mass were provided by 10 studies (n=28 231), arm circumference by nine studies (n=72 536), and skinfolds by five studies (n=7758). Finally, leptin and adiponectin were measured in nine studies (n=23 630 and 19 191, respectively). Overall, 47% of the combined study population was male. The median age within the contributing studies ranged from 16–74 years. Descriptive statistics for each of the study populations are found in the supplementary material (see online supplementary table S1).

Minor allele frequency for rs1051730/rs16969968 ranged between 0.31 and 0.36. There was no strong evidence for deviation from the Hardy-Weinberg
Equilibrium in any of the studies (p values all ≥0.09, see online supplementary table S2).

**Mendelian randomisation analysis**

Table 1 shows the per-allele increases in each phenotype within each smoking status category. As previously shown,^9^ the increase in BMI was positive in never-smokers: +0.35% (95% CI 0.18% to 0.52%; p=6.38×10^{-5}), non-significant in former smokers: −0.14% (95% CI −0.34% to +0.07%; p=0.19) and significantly inverse in current smokers: −0.74% (95% CI −0.96% to −0.51%; p=2.10^{-10}). The full results for each contributing study are shown in online supplementary figure S1.

The waist circumference was higher per minor allele in never-smokers: +0.23% (95% CI 0.09% to 0.36%; p=0.0012), non-significantly related in former smokers: −0.07% (95% CI −0.24% to 0.09%; p=0.37) and lower in current smokers: −0.40% (95% CI −0.57 to −0.22 p=1.69×10^{-5}); differences among smoking groups were highly significant (p=3.89×10^{-7}; see online supplementary figure S2). The per-allele effect on waist circumference in current smokers was about half the magnitude of that seen for BMI. After adjustment for log(BMI), the minor allele of rs1051730-rs16969968 was not associated with waist circumference in either never-smokers: +0.01% (95% CI −0.06 to 0.08; p=0.72) or former smokers +0.06% (95% CI −0.02% to 0.15%; p=0.15). However, in current smokers, the minor allele was associated with a 0.14% (95% CI 0.05% to 0.22%; p=0.003) higher waist circumference after adjustment for log (BMI). Very similar results were seen in all three smoking status categories after waist was adjusted for log(BMI). The mean difference in daily cigarette consumption was 0.77 among current smokers (95% CI 0.67 to 0.88, I^2=17%).

**Discussion**

This meta-analysis of 29 studies comprising almost 150 000 participants with key adiposity phenotypes has demonstrated, first, that a variant associated with increased cigarette consumption was associated not only with lower BMI among current smokers, consistent with earlier findings,^7^ but also with lower waist and hip circumferences. Second, the inverse association of the variant with lower waist circumference among current smokers appeared more marked among women (p=0.067), but this effect was no longer apparent after adjusting for BMI (p=0.51).

For several other phenotypes, per-allele decreases were observed in current smokers that exceeded those seen either in former or never-smokers (see online supplementary table S4). However, there was only statistical evidence for decreases among current smokers for arm circumference (p=8.4×10^{-5}) and leptin (p=0.025), while the difference between smoking groups was only significant for arm circumference (p=3.29×10^{-7}). Both effects became non-significant after adjustment for log(BMI). Fat mass and fat-free mass, after adjustment by height, showed differences in effects by smoking group. These effects were more due to per-allele increases seen among never-smokers than decreases among current smokers.

Meta-regression analyses showed no clear evidence for associations between genotype and each adiposity phenotype being modified by sex: p values exceeded 0.1 for all phenotypes, adjusted or unadjusted, apart from hip circumference. The per-allele decreases in hip circumference among current smokers appeared more marked among women (p=0.067), but this effect was no longer apparent after adjusting for BMI (p=0.51).

The mean difference in daily cigarette consumption was 0.77 among current smokers (95% CI 0.67 to 0.88, I^2=17%).
Table 1  Per allele percentage increases in measures of regional adiposity (BMI, weigh, waist circumference, hip circumference, waist-hip ratio) among never, ex and current smokers, before and after adjustment for BMI

<table>
<thead>
<tr>
<th>BMI (kg/m²)</th>
<th>Adjusted for age</th>
<th>Adjusted for age and BMI</th>
<th>p For interaction*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Never-smokers</td>
<td>Former smokers</td>
<td>Current smokers</td>
</tr>
<tr>
<td>% increase</td>
<td></td>
<td></td>
<td>p For interaction*</td>
</tr>
<tr>
<td>0.35</td>
<td>−0.14</td>
<td>−0.74</td>
<td>−</td>
</tr>
<tr>
<td>95% CI</td>
<td>(0.18 to 0.52)</td>
<td>(−0.34 to 0.07)</td>
<td>(−0.96 to −0.51)</td>
</tr>
<tr>
<td>p</td>
<td>6.38×10⁻⁵</td>
<td>0.19</td>
<td>2.00×10⁻¹⁰</td>
</tr>
<tr>
<td>N</td>
<td>66 809</td>
<td>43 009</td>
<td>38 912</td>
</tr>
<tr>
<td>I²</td>
<td>14%</td>
<td>0%</td>
<td>0%</td>
</tr>
</tbody>
</table>

| % increase  | 0.23             | −0.07                    | −0.40              |
| 95% CI      | (0.09 to 0.36)   | (−0.24 to 0.09)          | (−0.57 to −0.22)   |
| p           | 0.0012           | 0.37                     | 1.69×10⁻⁵          |
| N           | 64 265           | 40 756                   | 37 360             |
| I²           | 14%              | 0%                       | 10%                |

| % increase  | 0.17             | −0.07                    | −0.31              |
| 95% CI      | (0.08 to 0.26)   | (−0.17 to 0.04)          | (−0.42 to −0.19)   |
| p           | 2.95×10⁻⁴        | 0.23                     | 2.55×10⁻⁷          |
| N           | 62 323           | 40 512                   | 36 833             |
| I²           | 7%               | 0%                       | 16%                |

| % increase  | 0.07             | −0.08                    | −0.01              |
| 95% CI      | (−0.01 to 0.15)  | (−0.10 to 0.10)          | (−0.19 to 0.03)    |
| p           | 0.087            | 0.97                     | 0.14               |
| N           | 62 322           | 40 512                   | 36 833             |
| I²           | 21%              | 9%                       | 15%                |

*Interaction assessed by assessing heterogeneity between effect estimates according to smoking status, with a fixed effects model.

BMI, body mass index.
we have already noted among never-smokers an unexpected positive association of the gene variant with BMI; the current analysis demonstrates this same association with waist and hip circumferences. This occurred in the opposite direction to the inverse association of various adiposity measures with the gene variant seen in current smokers (before adjustment for BMI).

The analysis consisted of never, former and current smokers from a very wide spectrum of ages among the 29 studies. The sample size was very large for the primary phenotypes considered here. Participants were exclusively of self-reported European ancestry, and were mostly recruited in European countries. Data for direct measures of fat, such as fat mass, and the biomarker leptin were available for only about one-fifth of the participants whose weight, height, waist and hip were measured. Effects according to genotype for these phenotypes showed broadly similar results for the three smoking categories to those seen for BMI.

Mendelian randomisation has proved a powerful tool for eliciting causal associations between phenotypic measures. In the present analysis, Mendel’s laws concerning random assignment of genotype should produce an unconfounded comparison between the genotype influencing smoking consumption and the outcomes of interest, namely anthropometric phenotypes. Furthermore, since this random assignment occurs at the very outset of life, the associations between genotype and anthropometric measures cannot be due to reverse causality. If the genotype only influences smoking consumption, and not the initiation of smoking, then the relationship between genotype and anthropometric outcomes would only be expected among smokers.

In fact, while the variant was associated with lower waist and hip circumferences among current smokers, it was associated with greater waist and hip circumferences among never-smokers. This suggests that the true effect among current smokers may be even greater than estimated. When we adjusted waist circumference for BMI, there was no association with the gene variant among never-smokers. The relative proportions of ever-smokers and never-smokers were not clearly associated with genotype in the CARTA consortium, as reported elsewhere. The reversal of the association between waist circumference and allele count from negative to positive among current smokers after adjustment for BMI may be consistent with alternative explanations. First, heavy smokers may have less muscle mass; however, no association between allele count and fat-free mass could be detected in our analysis among smokers. Second, the test for interaction for smoking status and allele count on waist circumference after adjustment was of weak statistical significance. Third, the adjustment of one measure of adiposity with another with which it is highly correlated may have caused a spurious association. We repeated our analysis for participants with BMI under 30 only, where the correlation was more modest, and obtained similar results, albeit with reduced evidence for an effect.

Stratification of our analyses by smoking status could, in theory, introduce bias by conditioning on a collider (rs1051730/rs16969968). This variant shows some evidence for association with smoking cessation (current vs former smoking). While this is a possibility, no effect modifications of this variant with potential confounders by smoking status were demonstrated among 56 625 participants in the HUNT study.

Cross-sectional observational data from Switzerland has demonstrated that waist and hip circumferences were more strongly related to the number of cigarettes smoked per day than was BMI, while in Scotland being a smoker was associated with greater central adiposity among women. In a Finnish longitudinal twin cohort study, smoking in adolescence predicted abdominal obesity in adulthood. Observational data are, however, prone to confounding and reverse causality, and the present study adds some evidence that the associations reported are likely to be causal.

Some observational studies have noted that low fat-free mass and bone mineral density were more common among smokers. The present analysis has not substantiated the association with fat-free mass, although our sample size was much more limited for this phenotype.

Our findings resonate with observational studies which have shown associations between smoking and risk of diabetes, especially as analysis of the British Women’s Heart and Health Study showed that abdominal adiposity was a stronger predictor of diabetes than BMI. Waist circumference and waist-to-hip ratio were strongly associated, independently of BMI, with the risk of death among 359 387 participants from nine countries in the European Prospective Investigation into Cancer and Nutrition. Therefore, the health hazards of smoking could well be enhanced or partly mediated through increasing abdominal adiposity. In addition, the desire of many smokers to use smoking as a means of weight control might be counterproductive if a loss of weight is accompanied by a relative increase in waist circumference: this possibility could be used in counselling people seeking to quit smoking.

People who quit smoking appear to be at increased risk of acquiring diabetes in the short term but this was not explained by weight gain in a Japanese population. This study took place almost exclusively among white European participants, and replication of the findings among other ethnic populations would be of great value. This is especially urgent on a global scale since smoking levels are increasing among several non-white ethnic groups, and this is seen to be partly responsible for increases in coronary heart disease mortality in Beijing, China, in Syria and in Tunisia among women. In addition, increases in average waist circumference have been observed even when average BMI levels have remained constant, and metabolic disorders, especially diabetes, have increased in prevalence. It is thus possible that increased CHD mortality will be partly fuelled by increasing smoking levels.
Mendelian randomisation studies have more potential than traditional observational epidemiological studies to establish causality for specific exposures, and they should now be used to investigate other impacts of smoking, in particular on pathways leading to type 2 diabetes, as well as on type 2 diabetes itself. The findings of this study could now be further tested by assembling data from randomised trials of smoking cessation, where postintervention data on measures of central adiposity are available. If confirmed, a tendency for smokers to acquire an ‘apple shape’ due to increasing central adiposity might provide a novel health promotion message to encourage smoking cessation, and appropriate new interventions should then be designed and evaluated as part of overall tobacco control policies in society.

Author affiliations
1School of Social and Community Medicine, University of Bristol, Bristol, UK
2Department of Primary Care and Population Health, UCL, London, UK
3MRC Integrative Epidemiology Unit (IEU) at the University of Bristol, University of Bristol, Bristol, UK
4UK Centre for Tobacco and Alcohol Studies and School of Experimental Psychology, University of Bristol, Bristol, UK
5Department of Public Health, Faculty of Medicine, Norwegian University of Science and Technology, Trondheim, Norway
6Forensic Department, Research Centre Brøset, St Olav’s University Hospital Trondheim, Trondheim, Norway
7Department of Endocrinology, St. Olav’s Hospital, Trondheim University Hospital, Trondheim, Norway
8Department of Laboratory Medicine, Children’s and Women’s Health, The Faculty of Medicine, Norwegian University of Science and Technology, Trondheim, Norway
9Medical Genetics Section, Centre for Genomic and Experimental Medicine, Institute of Genetics and Molecular Medicine, University of Edinburgh, Edinburgh, UK
10Centre for Cognitive Ageing and Cognitive Epidemiology, University of Edinburgh, Edinburgh, UK
11Queensland Brain Institute, University of Queensland, Brisbane, Queensland, Australia
12Institute for Social and Economic Research, University of Essex, Colchester, UK
13Department of Public Health, Hjelt Institute, University of Helsinki, Helsinki, Finland
14Institute of Public Health and Clinical Nutrition, University of Eastern Finland, Kuopio, Finland
15National Institute for Health and Welfare, Helsinki, Finland
16Institute of Public Health and Clinical Nutrition, University of Eastern Finland, Kuopio, Finland
17Institute of Health Sciences, University of Oulu, Oulu, Finland
18Biocenter Oulu, University of Oulu, Oulu, Finland
19Centre for Environmental and Preventive Medicine, Wolfson Institute of Preventive Medicine, Barts and the London School of Medicine and Dentistry, Queen Mary University of London, London, UK
20Population, Policy and Practice, UCL Institute of Child Health, University College London, UK
21Department of Gerontology and Geriatrics, Leiden University Medical Center, Leiden, The Netherlands
22Netherlands Consortium of Healthy Ageing, Leiden, The Netherlands
23Research Centre for Prevention and Health, the Capital Region of Denmark, Denmark
24Novo Nordisk Foundation Center for Basic Metabolic Research, Section of Metabolic Genetics, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark
25Steno Diabetes Center, Gentofte, Denmark
26COPSAC, Copenhagen Prospective Studies on Asthma in Childhood, Herlev and Gentofte Hospital, University of Copenhagen, Copenhagen, Denmark
27Department of Biological Psychology, Netherlands Twin Register, VU University, Amsterdam, The Netherlands
28Faculty of Epidemiology and Population Health, London School of Hygiene & Tropical Medicine, London, UK
29Institute of Behavioural Sciences, University of Helsinki, Helsinki, Finland
30Folkhalsåren Research Centre, Helsinki, Finland
31Wellcome Trust Sanger Institute, Wellcome Trust Genome Campus, Cambridge, UK
32Institute for Molecular Medicine Finland (FIMM), University of Helsinki, Finland
33The Medical and Population Genomics Program, The Broad Institute of MIT and Harvard, Cambridge, Massachusetts, USA
34Robertson Centre for Biostatistics, University of Glasgow, Glasgow, UK
35Institute of Cardiovascular and Medical Sciences, University of Glasgow, Glasgow, UK
36MRC Unit for Lifelong Health and Ageing at UCL, London, UK
37Department of Environmental Medicine, Institute of Public Health, University of Southern Denmark, Odense, Denmark
38European Centre for Environment and Human Health, University of Exeter Medical School, Exeter, UK
39Genetics of Complex Traits, University of Exeter Medical School, Exeter, UK
40Department of Psychological Medicine, University of Otago, Christchurch, New Zealand
41Department of Pathology, University of Otago, Christchurch, New Zealand
42Institute for Clinical Research, University of Southern Denmark, Odense, Denmark
43Department of Epidemiology, Biostatistics and Biodemography, Institute of Public Health, University of Southern Denmark, Denmark
44University of Glasgow, Glasgow, UK
45Department of General Practice and Primary Health Care, University of Helsinki, Helsinki, Finland
46Unit of General Practice, Helsinki University Central Hospital, Helsinki, Finland
47Vasa Central Hospital, Vasa, Finland
48Population Health Research Institute, St George’s University of London, London, UK
49Department of Clinical Experimental Research, Glostrup University Hospital, Glostrup, Denmark.
50Department of Clinical Medicine, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark
51Department of Cardiology, Leiden University Medical Center, Leiden, The Netherlands
52Durrer Center for Cardiogenetic Research, Amsterdam, The Netherlands
53Interuniversity Cardiology Institute of the Netherlands, Utrecht, The Netherlands
54Centre for Population Health Research, School of Health Sciences and Sansom Institute of Health Research, University of South Australia, Adelaide, Australia
55South Australian Health and Medical Research Institute, Adelaide, South Australia, Australia
56Institute of Health and Welfare, Helsinki, Finland
57Institute of Internal Medicine, Lausanne University Hospital, Lausanne, Switzerland
58Institute of Health Sciences, University of Oulu, Oulu, Finland
59Biocenter Oulu, University of Oulu, Oulu, Finland
60Population, Policy and Practice, UCL Institute of Child Health, University College London, UK
61Department of Gerontology and Geriatrics, Leiden University Medical Center, Leiden, The Netherlands
62Division of Population Health Sciences, University of Dundee, Ninewells Hospital and Medical School, Dundee, UK
63Medical Research Council Human Genetics Unit, Institute of Genetics and Molecular Medicine, University of Edinburgh, Edinburgh, UK
64Institute of Preventive Medicine, Bispebjerg and Frederikberg Hospitals, The Capital Region, Copenhagen, Denmark
65Faculty of Medicine, BHF Glasgow Cardiovascular Research Centre, Glasgow, UK


Open Access
Acknowledgements 1958BC: Statistical analyses were funded by the Academy of Finland (Project 24300796 and SALVE/PREV/MDSVYN). DNA collection was funded by MRC grant G0009834 and cell-line creation by Wellcome Trust grant 068545/2/0. This research used resources provided by the Type 1 Diabetes Genetics Consortium, a collaborative clinical study sponsored by the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK), National Institute of Allergy and Infectious Diseases, National Human Genome Research Institute, National Institute of Child Health and Human Development, and Juvenile Diabetes Research Foundation International (JDRF) and supported by U01 DK082418. This study makes use of data generated by the Wellcome Trust Case-Control Consortium. A full list of investigators who contributed to generation of the data is available from the Wellcome Trust Case-Control Consortium website. Funding for the project was provided by the Wellcome Trust under the award 076113. The work was supported by the Department of Health Policy Research Programme through the Public Health Research Consortium (PHRC). The views expressed in the publication are those of the authors and not necessarily those of the Department of Health. Information about the wider programme of the PHRC is available from http://phrc.ishtm.ac.uk. Great Ormond Street Hospital/University College London, Institute of Child Health receives a proportion of funding from the Department of Health’s National Institute for Health Research (NIHR) (‘Biomedical Research Centres’ funding). ALSPAC: We are extremely grateful to all the families who took part in this study, the midwives for their help in recruiting them, and the whole ALSPAC team, which includes interviewers, computer and laboratory technicians, clerical workers, research scientists, volunteers, managers, receptionists and nurses. The UK Medical Research Council and the Wellcome Trust (Grant ref: 092731) and the University of Bristol provide core support for ALSPAC. Ethics approval for the study was granted for the BWHHS from the London Multi-Centre Research Ethics Committee. Each participant has given written informed consent.

BWHHS: The British Women’s Heart and Health Study has been supported by funding from the British Heart Foundation, Cancer Research UK, Economic and Social Research Council, Medical Research Council, and the National Institute for Health Research, under the auspices of the UK Clinical Research Collaboration, is gratefully acknowledged. This work was supported by the Wellcome Trust (grant number 086884) and the Medical Research Council (grant numbers MR/J01351X/1, G0800612, G0802736, MC_UU_12013/1, MC_UU_12013/6).

BRHS: The British Regional Heart Study is a British Heart Foundation (BHF) Research Group. The BRHS has local (from each of the districts in which the study was based) and multicentre ethical committee approvals. BWHHS: The British Women’s Heart and Health Study has been supported by funding from the British Heart Foundation (BHF) (grant PG/09/022) and the UK Department of Health Policy Research Programme (England) (grant 0090049). The BWHHS HumanCVD data were funded by the BHF (PG/07/131/24254). We thank all BWHHS participants, the general practitioners and their staff who have supported data collection since the study inception. Ethics approval was granted for the BWHHS from the London Multi-Centre Research Ethics Committee and 23 Local Research Ethics Committees.

Caps: The Caerrphilly Prospective Study was conducted by the former MRC Epidemiology Unit (South Wales). The Caerphilly archive is now maintained by the School of Social and Community Medicine in Bristol University. We thank the Health and Social Care Information Centre (HSCIC) for helping us maintain long-term follow-up with the cohort. We thank all the men who have given their time to be participants in CaPS. Ethics approval was obtained from the South Glamorgan Area Health Authority, the Gwent REC and the South Wales Research Ethics Committee. CDHS: The Christchurch Health and Development Study has been supported by funding from the Health Research Council of New Zealand, the National Child Health Research Foundation (Cure Kids), the Canterbury Medical Research Foundation, the New Zealand Lottery Grants Board, the University of Otago, the Carney Centre for Pharmacogenomics, the James Hume Bequest Fund, US NIH grant MH077874, and NIDA grant “A developmental model of gene-environment interplay in SUDs.” (R01DA024413) 2007–2012. All phases of the study have received ethics approval from the regional Health and Disability Ethics Committee and all forms of data collection have been subject to the signed consent of research participants.

Colaus: The Colaus/PsyCoLaus study was supported by four grants of the Swiss National Science Foundation (#105993, 118308, 119468 and 122661), two unrestricted grants from GlaxoSmithKline as well as by the Faculty of Biology and Medicine of the University of Lausanne. Colaus and PsyCoLaus were approved by the Institutional Ethics Committee of the University of Lausanne.

Dan-MONICA: The Dan-MONICA10 was sponsored by The Danish Heart Foundation; the Danish Medical Research Council; The Danish Hospital Foundation of Medical Research, region of Copenhagen, the Faroe Islands and Greenland; The Danish Health Insurance Foundation; The Foundation of E. & M. Wedel-Wedellsborg; Landsforeningen til Bekampelse af Kredsløbssygdomme; The Augustinus Foundation; The Becket Foundation; and The Foundation of senior registrar J. & L. Boserup. All participants gave written consent and the study was conducted in accordance with the Second Helsinki Declaration and approved by the Ethics Committee for Copenhagen County. EFSoCH: The Exeter Family Study of Childhood Health (EFSoCH) was supported by the South West NHS Research and Development, Exeter NHS Research and Development, the Darlington Trust, and the Peninsula National Institute of Health Research (NIHR) Clinical Research Facility at the University of Exeter. The opinions given in this paper do not necessarily represent those of NIHR, the NHS or the Department of Health. Ethics approval was given by the North and East Devon Local Research Ethics Committee. ELSA: ELSA is funded by the National Institute on Aging in the US (R01 AG017644/R01AG176440651) and by a consortium of UK Government departments (including: Department for Communities and Local Government, Department for Transport, Department for Work and Pensions, Department of Health, HM Revenue and Customs and Office for National Statistics). ELSA has been approved by the National Research Ethics Service and all participants have given informed consent. Finrisk: This study was supported by the Academy of Finland, Center of Excellence in Complex Disease Genetics (grant numbers 213506, 129680), the Academy of Finland (grant numbers 139635, 129494, 136895, 263836 and 141054), the Sigrid Juselius Foundation and ENGAGE—European Network for Genetic and Genomic Epidemiology, FP7-HEALTH-F4-2007, grant agreement number 201413. The 2002 and 2007 FINRISK surveys have been approved by the Coordinating Ethics Committee of the Helsinki University Hospital District. Each participant has given written informed consent.

Geminakar: The GEMINAKAR study was supported by grants from the Medical Research Fund, the Danish Diabetes Association, the NOVO Foundation and the Danish Heart Foundation. The study was approved by the relevant Danish Ethics Committee (baseline, S-VF-19970271) and Danish Data Protection Board (baseline, 1999-1200-441). All participants provided written informed consent.

Generation Scotland: Generation Scotland has received core funding from the Chief Scientists Office of the Scottish Government Health Directorates CED/16/6 and the Scottish Funding Council HR03006. We are grateful to all the families who took part, the general practitioners and the Scottish School of Primary Care for their help in recruiting them, and the whole Generation Scotland team, which includes interviewers, computer and laboratory technicians, clerical workers, research scientists, volunteers, managers, receptionists, healthcare assistants and nurses. Genotyping of the GS:SFHS samples was carried out by the Genetics Core Laboratory at the Wellcome Trust Clinical Research Facility, Edinburgh, Scotland and was funded by the UK Medical Research Council (MRC). Ethics approval for the study was given by the NHS Tayside committee on research ethics (reference 05/S1401/9).

Goya: The GÖYA study was conducted as part of the activities of the Danish Obesity Research Centre (DanORC, http://www.danorc.dk) and The MRC centre for Causal Analyses in Translational Epidemiology (MRC CAITE). The genotyping for GÖYA was funded by the Wellcome Trust (WT 084762). GÖYA is a nested study within The Danish National Birth Cohort which was established with major funding from the Danish National Research Foundation. Additional support for this cohort has been obtained from the Pharmacy Foundation, the Egmont Foundation, The March of Dimes Birth Defects Foundation, the Augustinus Foundation and the Health Foundation. TSA was supported by the Gene Diet Interactions in Obesity (GENDINOB, http://www.gendinob.dk) postdoctoral fellowship grant. LP is funded by an MRC Population Health Scientist Fellowship (MR/J012165/1). The study was approved by the regional scientific ethics committee and by the Danish Data Protection Board.

HBCS: The Helsinki Birth Cohort Study has been supported by grants from the Academy of Finland, the Finnish Diabetes Research Society, Samfunnet Folkhälsan, Novo Nordisk Foundation, Finska Läkarealliansen, Signe and Ane Gunnarsson Foundation, University of Helsinki,
Ministry of Education, Ahokas Foundation and Emil Aaltonen Foundation. The research plan of the HBCS was approved by the Institutional Review Board of the National Public Health Institute and all participants have signed informed consent forms. Health2006/Health2008/Inter99: LLNH was supported by the Health Insurance Foundation (grant No. 2010 B 131). The studies have been approved by the Ethical Committee of Copenahgen. HUNT: Nord-Trondelag Health Study (The HUNT Study) is a collaboration between the HUNT Research Centre (Faculty of Medicine, Norwegian University of Science and Technology NTNU), the Nord-Trondelag County Council and the Norwegian Institute of Public Health. Use of data in this study was approved by the Regional Committee for Medical Research Ethics (Reference no. 2013/1127/ REK midt). Midsnap: The Midsnap Family Study was funded as part of the NIH Research and Development Cardiovascular Research Programme. Ethics approval was obtained from the Argyll and Clyde Health Board Local Research Ethics. NFBC: NFBC1966 and NFBC1966 received financial support from the Academy of Finland (project grants 104781, 120315, 129269, 1114194, 24300796, 141042 Center of Excellence in Complex Disease Genetics and SALVE), University Hospital Oulu, Biocenter, University of Oulu, Finland (75617), NHLBI grant 5R01HL087679–02 through the STAMPEED program (1R1LM080328–01), NIH/NIMH (5R01MH67307–02), the European Commission (EURO-BLCS, Framework 5 award QLG1-CT-2000–01643), ENGAGE project and grant agreement HEALTH-F4-2007–20143, EU FP7 EuroHeartAgeing–277849, the Medical Research Council, UK (G0505539, G0600705, G1002319, PrevMetSyn/SALVE) and the MRC, Centenary Early Career Award. The DNA extractions, sample quality controls, biobank upkeep and aliquoting was performed in the National Public Health Institute, Biomedicalc Helsinki, Finland and supported financially by the Academy of Finland and Biocentrum Helsinki. We thank the late Professor Paula Rautakallio (launch of NFBCs) and Ms Outi Tornwall and Ms Minttu Jussila (DNA biobanking). The authors would like to acknowledge the contribution of the late Academician of Science Leena Peltonen. The University of Oulu Ethics Committee and the Ethical Committee of Northern Ostrobothnia Hospital District have approved the study. Participants provided written informed consent. NHANES: The National Health and Nutrition Examination Survey (NHANES) (http://www.cdc.gov/nchs/nhanes.htm) is a program of health surveys run by the National Center for Health Statistics, part of the Centers for Disease Control and Prevention in the USA. Data collection for NHANES was approved by the NCHS Research Ethics Review Board. Analysis of de-identified data from the survey is exempt from the federal regulations for the protection of human research participants. Analysis of restricted data through the NCHS Research Data Center is also approved by the NCHS ERB. The findings and conclusions in this paper are those of the author(s) and do not necessarily represent the views of the Research Data Center, the National Center for Health Statistics or the Centers for Disease Control and Prevention. NSHD: We are very grateful to the members of this birth cohort for their continuing interest and participation in the study. We would like to acknowledge the Sallow group at University College London, who performed the DNA extractions. This work was funded by the Medical Research Council [MC_UU_12019/1]. Ethics approval was given by the Central Manchester Research Ethics Committee. NTR: This study was supported by the European Research Council (ERC Starting Grant 284167 PI Vink), Netherlands Organization for Scientific Research (NWO: MagW/ZonMW grants 904-61-090, 985-10-002, 904-61-193, 480-04-004, 400-05-717, Addiction-31600081 Middelgroot-911–09–032, Spinozepremie 56–464– 14192), BBRMI-NL (Biobanking and Biomolecular Resources Research Infrastructure), VU University’s Institutes for Health and Care Research and Neuroscience Campus Amsterdam. The NTR study was approved by the Central Ethics Committee on Research Involving Human Subjects of the VU University Medical Center, Amsterdam (IRB number IRB-2991 under Federaalwide Assurance 3703: IRB/Institute code 03–180), and all subjects provided written informed consent. PROSPER: The Prospective Study of Pravastatin in the Elderly at Risk (PROSPER) trial was supported by an investigator initiated grant from Bristol-Myers Squibb, USA. The study was conducted, analysed and reported independently of the company. The GWAS project PHASE has received funding from the European Union’s Seventh Framework Programme (FP7/2007–2013) under grant agreement HEALTH-F2–2009–223004. A part of the genotyping was funded by The Netherlands Consortium for Healthy Ageing (NGI: 05060810). JW is an established clinical investigator of The Netherlands Heart Foundation (2001 D 032). PROSPER was approved by the Argyll and Clyde Local Research Ethics Committee, the Glasgow Royal Infirmary Local Research Ethics Committee, Greater Glasgow Primary Care and Mental Health Research Ethics Committee, Lanarkshire Health Board Local Research Ethics Committee, Dumfries and Galloway Health Board Local Research Ethics Committee, Forth Valley Health Board Local Research Ethics Committee, METC board of Leiden University Medical Center and the Clinical Research Ethics Committee of The Cork Teaching Hospitals, and all participants gave written informed consent.

Whitehall II: The Whitehall II study has been supported by grants from the Medical Research Council (K013531), British Heart Foundation; Health and Safety Executive; Department of Health; National Heart Lung and Blood Institute (NHLBI: HL36310) and National Institute on Aging (AG13196), US, NIH; Agency for Health Care Policy Research (HS06156), and the John D and Catherine T MacArthur Foundation Research Networks on Successful Midlife Development and Socio-economic Status and Health. MeKu is partially supported by the Economic and Social Research Council International Centre for Life Course Studies in Society and Health (RES-596–28–0001). MK is partially supported by the Medical Research Council and the Economic and Social Research Council. Ethics approval for the Whitehall II study was obtained from the University College London Medical School committee on the ethics of human research. Informed consent was gained from every participant.

Collaborators
Allan Linneberg.

Contributors
RWM, AET, TIAS, MRN and NS conceived the study and contributed to the writing of the manuscript. RWM conducted the final analyses. All other authors conducted individual study analyses and contributed to the writing of the manuscript.

Funding
This research received no specific grant from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests
All authors have completed the Unified Competing Interest form at http://www.icmje.org/coi_disclosure.pdf (available on request from the corresponding author) and declare: support for the submitted work (LP, AW, DK: Medical Research Council, LIK, DMF: New Zealand Health Research Council, MAK: Jim and Mary Carney Charitable Trust, New Zealand Health Research Council, BHS: Scottish Government Chief Scientist Office) and financial relationships with any organisations that might have an interest in the submitted work in the previous three years (MF: grants from Swiss National Science Foundation and from GlaxoSmithKline, during the conduct of the study, TK: consulted for Pfizer in 2011–2015 on nicotine dependence, JK: grants from Academy of Finland, during the conduct of the study and personal fees from Pfizer, outside the submitted work). There are no other relationships or activities that could appear to have influenced the submitted work.

Ethics approval
The manuscript describes approval given for each of the 29 studies.

Provenance and peer review
Not commissioned; externally peer reviewed.

Data sharing statement
1958BC: This study makes use of data generated by the Wellcome Trust Case-Control Consortium. A full list of investigators who contributed to generation of the data is available from the Wellcome Trust Case-Control Consortium website. The 1958 birth cohort data can be accessed via the UK Data Service (http://ukdataservice.ac.uk/). ALSAPC: Data used for this submission will be made available on request to the ALSAPC executive committee (alsapc@bristol.ac.uk). The ALSAPC data management plan (available here:http://www.bristol.ac.uk/alsapc/researchers/data-access/) describes in detail the policy regarding data sharing, which is through a system of managed open access. BRHS: We welcome proposals for collaborative projects and data sharing (http://www.ucl.ac.uk/pcph/data-access/) describes in detail the policy regarding data sharing, which is through a system of managed open access.

Registered: All authors have completed the Unified Competing Interest form at http://www.icmje.org/coi_disclosure.pdf (available on request from the corresponding author) and declare: support for the submitted work (LP, AW, DK: Medical Research Council, LIK, DMF: New Zealand Health Research Council, MAK: Jim and Mary Carney Charitable Trust, New Zealand Health Research Council, BHS: Scottish Government Chief Scientist Office) and financial relationships with any organisations that might have an interest in the submitted work in the previous three years (MF: grants from Swiss National Science Foundation and from GlaxoSmithKline, during the conduct of the study, TK: consulted for Pfizer in 2011–2015 on nicotine dependence, JK: grants from Academy of Finland, during the conduct of the study and personal fees from Pfizer, outside the submitted work). There are no other relationships or activities that could appear to have influenced the submitted work.

Ethics approval
The manuscript describes approval given for each of the 29 studies.

Provenance and peer review
Not commissioned; externally peer reviewed.

Data sharing statement
1958BC: This study makes use of data generated by the Wellcome Trust Case-Control Consortium. A full list of investigators who contributed to generation of the data is available from the Wellcome Trust Case-Control Consortium website. The 1958 birth cohort data can be accessed via the UK Data Service (http://ukdataservice.ac.uk/). ALSAPC: Data used for this submission will be made available on request to the ALSAPC executive committee (alsapc@bristol.ac.uk). The ALSAPC data management plan (available here:http://www.bristol.ac.uk/alsapc/researchers/data-access/) describes in detail the policy regarding data sharing, which is through a system of managed open access. BRHS: We welcome proposals for collaborative projects and data sharing (http://www.ucl.ac.uk/pcph/data-access/) describes in detail the policy regarding data sharing, which is through a system of managed open access.
No data should be passed on to any third party unless they were specified in the original application. CaPS: Data used for the Caerphilly Prospective Study (CaPS) was made available by the CaPS access committee (Chair: Professor Kay Tee Khaw). More information about its managed access procedure is available on the study website (http://www.bris.ac.uk/social-community-medicine/people/project/1392). CHDS: Data contributed for this submission are available on request from the CHDS (John horwood@otago.ac.nz). Colauς/PsyCoLaus: Data from the CoLaus/PsyCoLaus study can be requested according to the procedure described on the CoLaus website (http://www.colaus.ch/en/cls_home/cls_prf_home/cls-research-3. htm). ELSA: ELSA data are made available through the ESDS website (http://www.elsa-project.eu/availableData). FINRISK: Data used for this submission will be made available on request to the FINRISK Management Group, according to the given ethical guidelines and Finnish legislation. Generation Scotland: Data are available on request (access@generationscotland.org). GOYA females: An anonymised copy of the data used for this submission will be made available on request to the GOYA analysts after permission has been given by the DNBC executive committee (www.dncb.dk). HBCS: Data used for this submission will be made available on request to the HBCS executive committee (johan.eriksson@helsinki.fi). Health2006/Health2008/Inter99: Data used for this submission can be made available on request to the Research Centre for Prevention and Health (http://www.regionh.dk/kfcs/Menu/). Please contact LLNH (lise.lotte.nystrup.husemoen@regionh.dk) or AL (allan.linneberg@regionh.dk). HUNT: Data used from the HUNT Study for this submission will be made available on request to the HUNT Data Access Committee (hunt@medisin.ntnu.no). The HUNT data access information (http://www.ntnu.edu/hunt/data) describes in detail the policy regarding data availability. NFBC: Data used for this submission can be made available on request to Tuija Ylitalo (tuija.ylitalo@oulu.fi), Minna Mannikko (minna.amnikko@oulu.fi) or M-RJ (m.jarvelin@imperial.ac.uk). NHANES: NHANES data can be accessed here: (http://www.cdc.gov/nchs/nhanes.htm). The genotype used in this analysis is a restricted variable. Applications for access to these data must be made through the Research Data Center: (http://www.cdc.gov/research/). NASHO: The NASHO data are made available to researchers who submit data requests (tom.chia.swiftinfo@ucl.ac.uk). More information is available in the full policy documents (http://www.nshd.mrc.ac.uk/data.aspx). Managed access is in place for this study to ensure that use of the data is within the bounds of consent given previously by participants, and to safeguard any potential threat to anonymity since the participants are all born in the same week. NTR: Data used for this submission will be made available on request to the NTR committee (ntr@ipps.vu.nl). Whitehall: Data from the Whitehall II study are made publicly available as described in the Whitehall II data sharing policy (http://www.ucl.ac.uk/whitehallII/datab sharinging). TIAS, MRM and NS are joint senior authors.

Open Access This is an Open Access article distributed in accordance with the terms of the Creative Commons Attribution (CC BY 4.0) license, which permits others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for the terms of the Creative Commons Attribution (CC BY 4.0) license, which allows others to distribute, remix, adapt and build upon this work, for

REFERENCES

Heavier smoking may lead to a relative increase in waist circumference: evidence for a causal relationship from a Mendelian randomisation meta-analysis. The CARTA consortium


BMJ Open 2015 5:
doi: 10.1136/bmjopen-2015-008808

Updated information and services can be found at:
http://bmjopen.bmj.com/content/5/8/e008808

These include:

Supplementary Material
Supplementary material can be found at:
http://bmjopen.bmj.com/content/suppl/2015/08/11/bmjopen-2015-008808.DC1.html
http://bmjopen.bmj.com/content/suppl/2015/08/12/bmjopen-2015-008808.DC2.html

References
This article cites 30 articles, 10 of which you can access for free at:
http://bmjopen.bmj.com/content/5/8/e008808#BIBL

Open Access
This is an Open Access article distributed in accordance with the terms of the Creative Commons Attribution (CC BY 4.0) license, which permits others to distribute, remix, adapt and build upon this work, for commercial use, provided the original work is properly cited. See:
http://creativecommons.org/licenses/by/4.0/

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/
Correction

Morris RW, Taylor AE, Fluharty ME, et al. Heavier smoking may lead to a relative increase in waist circumference: evidence for a causal relationship from a Mendelian randomisation meta-analysis. The CARTA consortium. *BMJ Open* 2015;5:e008808. The author name Tarun Veer Singh Ahluwalia should be spelt Tarunveer Singh Ahluwalia, and the abbreviation is Ahluwalia TS. Also, the surname of Maiken Elvestad Gabrielsen is ‘Gabrielsen’ only so should be abbreviated to Gabrielsen ME as opposed to Elvestad Gabrielsen M.

*BMJ Open* 2015;5:e008808. doi:10.1136/bmjopen-2015-008808corr1

CrossMark