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Explaining the link between adiposity and colorectal cancer risk in men and postmenopausal women in the UK Biobank: a sequential causal mediation analysis

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Abbreviations used:

CRC: colorectal cancer; CRP: C-reactive protein; HbA1c: hemoglobin-A1c; SHBG: sex hormone binding globulin; NDE: natural direct effect; NIE: natural indirect effect; RR: risk ratio; CI: confidence interval; ICD: International Classification of Disease; IQC: internal quality control; CV: coefficient of variation; WHO: World Health Organization; GMR: geometric mean ratio; OR: odds ratio; TE: total effect; IPW: inverse probability weights; MI: multiple imputation; HPFS: Health Professionals Follow-up Study; SOB-R: soluble leptin receptor; HR: hazard ratio; TyG: triglyceride-glucose index

Novelty and Impact

Mechanisms underlying adiposity-colorectal cancer (CRC) association are not well understood. Here, we used UK Biobank data and performed sequential mediation analysis to quantify mediating effects of C-reactive protein, hemoglobin-A1c, sex-hormone-binding globulin and testosterone in this association in men and postmenopausal women. Our results suggest pathways marked by these obesity-related factors may not

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explain a large proportion of this association and pathways other than those captured here may also be important for understanding the adiposity-CRC link.

Abstract

Mechanisms underlying adiposity-colorectal cancer (CRC) association are incompletely understood. Using UK Biobank data, we investigated the role of C-reactive protein (CRP), hemoglobin-A1c (HbA1c), and (jointly) sex hormone-binding globulin (SHBG) and testosterone, in explaining this association. Total effect of obesity vs normal-weight (based on waist circumference, body mass index, waist-hip ratio) on CRC risk was decomposed into natural direct (NDE) and indirect (NIE) effects using sequential mediation analysis. After a median follow-up of 7.1 years, 2070 incident CRC cases (men=1280; postmenopausal women=790) were recorded. For men, the adjusted risk ratio (RR) for waist circumference (≥ 102 vs ≤ 94 cm) was 1.37 (95% confidence interval (CI), 1.19-1.58). The RRs^{NIE} were 1.08 (95%CI, 1.01-1.16) through all biomarkers, 1.06 (95%CI, 1.01-1.11) through pathways influenced by CRP, 0.99 (95%CI, 0.97-1.01) through HbA1c beyond [the potential influence of] CRP, and 1.03 (95%CI, 0.99-1.08) through SHBG and testosterone combined beyond CRP and HbA1c. The RR^{NDE} was 1.26 (95%CI, 1.09-1.47). For women, the RR for waist circumference (≥ 88 vs ≤ 80 cm) was 1.27 (95%CI, 1.07-1.50). The RRs^{NIE} were 1.08 (95%CI, 0.94-1.22) through all biomarkers, 1.08 (95%CI, 0.99-1.17) through CRP, 1.00 (95%CI, 0.98-1.02) through HbA1c beyond CRP, and 1.00 (95%CI, 0.92-1.09) through SHBG and testosterone combined beyond CRP and HbA1c. The RR^{NDE} was 1.18 (95%CI, 0.96-1.45). For men and women, pathways influenced by CRP explained a small proportion of the adiposity-CRC association. Testosterone and SHBG also explained a small proportion of this association in men. These results suggest that pathways marked by these obesity-related factors may not explain a large proportion of the adiposity-CRC association.

Introduction

Globally, colorectal cancer (CRC) is the third most commonly diagnosed cancer among men and women ¹. There is a wide disparity in CRC incidence across the world regions. High-income countries have three times higher incidence rate compared with low- and middle-income countries ¹, while the latter, especially countries undergoing significant developments, have faced an increased burden of CRC more recently ². These patterns highlight the role that some of the consequences of a 'Westernized' lifestyle may have in developing CRC ¹.

Excess body fatness, a hallmark of 'Westernized' lifestyle, is an established risk factor for CRC, as well as for 12 other cancers ³. Based on a recent meta-analysis, the relative risk (RR) of CRC for the highest vs lowest category of waist circumference (a measure of central adiposity) was 1.48 (95% confidence interval (CI), 1.30-1.67) for men and 1.44 (95%CI, 1.30-1.60) for women ⁴. Similar associations were reported for BMI (obese vs normal), with a more evident difference in the strength of association for men (RR 1.47; 95%CI, 1.36-1.58) compared with women (RR 1.15; 95%CI, 1.08-1.23) ⁴.

Overweight and obesity already make a substantial contribution to the burden of CRC. In 2012, 13% (90% confidence interval (CI), 11%-14%) of colon and 6% (90%CI, 5%-7%) of rectal cancer cases diagnosed in men and 8% (90%CI, 7%-8%) of colon and 4% (90%CI, 3%-4%) of rectal cancer cases diagnosed in women were attributable to overweight and obesity (BMI ≥ 25 kg/m²) ⁵. Globally, the prevalence of adults living with obesity (BMI >30 kg/m²) has increased from ~3% in 1975 to ~11% in 2014 in men and from ~6% in 1975 to ~15% in 2014 in women ⁶. If these trends continue, by 2025, 18% of men and 21% of women will have obesity ⁷. It can, therefore, be expected for obesity to make considerably larger contributions to CRC burden in the future.

Adiposity-induced chronic inflammation, dysregulation of insulin signaling and glucose homeostasis, and sex-steroid hormones production are three interconnected pathways hypothesized to mediate the adiposity–CRC association⁸. Inflammation might exert its influence on CRC risk directly (through its angiogenic, mitogenic, and antiapoptotic effects)⁸ and through contributions to the development of insulin resistance and dysregulated insulin signaling⁹ and sex-steroid hormones production pathways^{10, 11}. Insulin resistance may increase cancer risk through hyperglycemia and exposure to elevated levels of insulin which has mitogenic and antiapoptotic effects⁸ and its influences on insulin-like growth factor-1^{12, 13}, and sex-steroid hormones levels^{10, 11, 14}. The potential role of sex-steroid hormones in colorectal carcinogenesis is not well-understood, but the higher incidence of CRC in men, a stronger adiposity–CRC association in men⁸, differences in the associations between adiposity and sex-steroid hormone levels in men and women (inverse association with testosterone levels in men, positive association in women)¹⁵, and the inverse association between hormone therapy and CRC in women¹⁶ are consistent with a role of sex-steroid hormones in the etiology of CRC⁸. Overall, however, the evidence for associations between sex-steroid hormones and CRC risk in men and women is limited and inconclusive⁸.

We used data from the UK Biobank to investigate the extent to which biomarkers of these three pathways mediate the effect of adiposity (measured by waist circumference, BMI, and waist-hip ratio) on CRC risk in men and postmenopausal women. A sequential causal mediation analysis approach was used to quantify the mediating effects through the three pathways without assuming they acted independently¹⁷. The assessed biomarkers were high sensitivity C-reactive protein (CRP, a non-specific biomarker of systemic inflammation), glycated hemoglobin (HbA1c, a biomarker of glycemic control and a correlate of insulin resistance¹⁸), and sex-hormone-binding globulin (SHBG) and testosterone (treated jointly to capture some aspects of altered sex-steroid hormone levels in the obese state).

Methods

The UK Biobank is a prospective cohort of 229,134 men and 273,402 women recruited between 2006 and 2010 following an invitation letter sent to about 9.2 million individuals (~5.5% response rate) ¹⁹. Invitees were registered with the UK National Health Service, were mostly 40-69 years old, and lived within approximately 25 miles of one of 22 assessment centers in England, Wales, and Scotland. Participants gave informed consent, and the North West Multi-Centre Research Ethics committee, the National Information Governance Board for Health and Social Care in England and Wales, and the Community Health Index Advisory Group in Scotland approved the study. In 2004, the Ethics and Governance Council was established to ensure adherence to the Ethics and Governance Framework (<http://www.ukbiobank.ac.uk/ethics/>) of the study. This research was conducted using the UK Biobank Resource under application number 25897.

The current study used data from all participants, with the outcome defined as CRC diagnosis between one year after baseline (i.e. recruitment) and end of follow-up (see *Follow-up and outcome assessment*).

Data collection

At baseline, information on demographics, early life and lifestyle exposures, health history and medication use were collected using a self-administered touchscreen questionnaire available at <http://www.ukbiobank.ac.uk/key-documents/>. Weight, height, waist and hip circumference were measured following prespecified protocols ²⁰, and blood samples were collected, centrifuged, and stored at -80°C or liquid nitrogen within ~24 hours after venipuncture ²¹.

Follow-up and outcome assessment

Incident cancer cases and cancer cases recorded first in death certificates were identified through linkage to national cancer and death registries. Complete follow-up was available through 31 March 2016 for England

and Wales and 31 October 2015 for Scotland. The 10th Revision of the International Classification of Disease (ICD-10) was used to define CRC cases (C18, C19, and C20 for malignant neoplasms of colon, rectosigmoid junction, and rectum respectively).

Selection of participants eligible for the study

Participants were ineligible for this study if they had withdrawn consent; had cancer diagnosed before baseline; had less than one-year follow-up; were diagnosed with CRC within the first year after baseline; were diagnosed with cancer of the appendix (ICD-10 C18.1; <1%); died (<2%) or were lost to follow-up (<0.5%) before the end of follow-up; did not have body size measurements collected or had BMI<18.5 kg/m² at baseline; had unknown diabetes status, had diabetes or reported taking diabetes medication at baseline (Figure 1a-1b). The last exclusion criterium was imposed because diabetes and diabetes medication use may change the link between adiposity and insulin signaling²². Additionally, women who, at baseline, were premenopausal or ≤55 years old with unknown menopause status, who reported taking hormone therapy, or hormonal contraceptives were ineligible (Figure 1b). We did not include premenopausal women in this analysis because the mechanistic pathways, especially the sex-steroid hormones pathway, might be different in pre- and postmenopausal women. Instead of multiply imputing missing confounder data (the approach used to handle missing biomarker data), we also excluded participants with missing data for any of the selected confounders (see *Confounder selection*) because they were a relatively small proportion (≤10%) of all the participants.

Biomarkers and laboratory methods

Descriptions of laboratory methods and protocols for measuring the biomarkers have been published^{23, 24}. We did not include estradiol in our analyses because most participants had poor quality data²⁴.

Measurements were performed in blood samples (red blood cells for HbA1c, serum for all other biomarkers) collected at baseline (2006-2010) as part of the UK Biobank Biomarker Project. Immuno-turbidimetric (Beckman Coulter AU5800) was used to measure CRP (with average within-laboratory coefficient of variation (CV) of 2.3% for low and 1.7% for high internal quality control (IQC) samples); high-performance liquid chromatography (Bio-Rad Variant II Turbo) to measure HbA1c (CV 2.1% for low, 1.5% for high IQC samples); and chemiluminescent immunoassay (Beckman Coulter DXI 800) to measure SHBG (CV 5.7% for low, 5.2% for high IQC samples), and testosterone (CV 8.3% for low, 4.2% for high IQC samples)^{23, 24}. An algorithm was developed to ensure each laboratory batch included a random selection of participants to reduce the likelihood of any systematic differences in assay results between participants²⁴. When applicable, the data we received had also been corrected for aliquot-dilution effect (SHBG, testosterone) and date of the assay (HbA1c, SHBG, testosterone)²¹. Free testosterone was calculated from measured testosterone, SHBG, and albumin²⁵. For a subgroup of participants, biomarker measurements were additionally performed in blood samples collected at repeat assessment visit (2012-2013). Intraclass correlation coefficients (ICC) were calculated using the two measurements to assess the reproducibility of the measures (see Statistical analysis).

Statistical analysis

Waist circumference, BMI, and waist-hip ratio were used as measures of adiposity in separate analyses. For BMI, we used the World Health Organization (WHO) recommended cut-off values to compare participants with obesity vs normal weight (≥ 30 vs < 25 kg/m²). For waist circumference, we also used the WHO suggested cut-off values to compare participants defined as “at substantially increased risk of metabolic complications” (referred to as with obesity hereafter) with normal (> 102 vs ≤ 94 cm for men, > 88 vs ≤ 80 cm for women)²⁶. For waist-hip ratio, we used sex-specific quartiles to define with obesity (4th quartile > 0.98

for men, >0.86 for women) vs normal (1st quartile ≤0.89 for men, ≤0.77 for women) because the WHO does not have a suggested cut-off value for normal ²⁶. We treated CRC diagnosis as a binary outcome because methods for handling multiple mediators and survival outcome are not fully developed yet ²⁷.

To calculate the ICCs for biomarker measurements, linear mixed-effects modelling was used to obtain estimates of the between- and within-person variances.

Confounder Selection: The following variables were identified *a priori* and included as potential confounders (i.e. common causes for exposure-outcome, mediator-outcome, or exposure-mediator associations ²⁸): age at baseline (fitted as splines), educational attainment, Townsend deprivation index, first-degree relative history of CRC, physical activity, alcohol intake, smoking status, total red or processed meat intake, regular aspirin or ibuprofen use; additionally for women number of live births, ever use of oral contraceptive pill, and ever use of hormone therapy.

Adiposity-biomarker associations: For each biomarker, the geometric mean ratio (GMR; 95% CI) was estimated in relation to adiposity measures (comparing with obesity vs normal) using linear regression models fitted to the log-transformed biomarker variable adjusting for potential confounders.

Adiposity-CRC & Biomarker-CRC associations: Odds ratios (OR; 95% CI) for the adiposity-CRC and biomarker-CRC associations were estimated using logistic regression models adjusted for confounders. For each biomarker-CRC model, we additionally included waist circumference and other biomarkers that might have confounded the association (i.e. CRP for HbA1c-CRC model; and CRP and HbA1c for SHBG-CRC and testosterone-CRC models, selected based on the hypothesized causal sequence – see **Figure 2**). The linearity of the biomarker-CRC associations was investigated in models with biomarker variables fitted as restricted cubic splines (2 degrees of freedom).

Sequential mediation analysis: Using a causal mediation analysis approach ¹⁷, the total adiposity-CRC association (total effect Risk Ratio (RR)^{TE}) was decomposed into a natural indirect effect (RR^{NIE}; effect explained by all the included biomarkers, i.e. CRP, HbA1c, SHBG, and testosterone) and a natural direct effect (RR^{NDE}; effect not explained by any of the included biomarkers) (see ²⁸ for a formal definition of NDE and NIE). Assuming that CRP levels might have also influenced HbA1c or biomarker of sex-steroid hormone pathway ^{9, 11}, and HbA1c might have influenced biomarker of sex-steroid hormone pathway ^{11, 14}, but not vice versa, we took a sequential approach to further decompose the estimated NIE into the following NIEs: i) through pathways influenced by CRP; ii) through HbA1c excluding the possible influence of CRP on HbA1c; and iii) through SHBG and testosterone combined excluding the influences of CRP or HbA1c.

Figure 2 provides a schematic explanation of these effects and the sequential process used to estimate them

¹⁷. The mediation analysis involved calculating inverse probability weights (IPWs) based on the predicted probabilities of being obese from logistic regression models for exposure conditional on confounders ¹⁷.

Then, under different counterfactual scenarios, potential outcomes weighted by the IPWs (to take confounding into account) were estimated based on logistic regression models for the outcome conditional on exposure, confounders, with and without the biomarkers ¹⁷. Potential outcomes were expressed as risks, using the following formulae $risk = \frac{1}{(1 + \exp(-\log(odds)))}$ ²⁹. To accommodate non-linear biomarker-CRC associations, relevant biomarker variables were included as restricted cubic splines. Outcome models used to estimate each effect are presented in **Figure 2**.

Missing data: Missing biomarker data were handled using multiple imputation (MI) based on chained equations with 20 iterations for men and 35 for women (18% of men and 35% of women had missing values for at least one biomarker (**Table 1**)). For each adiposity measure, MI was performed with separate

imputation models for obese and non-obese individuals³⁰. The imputation models included all variables in the mediation analyses and the biomarker-biomarker and biomarker-CRC interaction terms (i.e. treating these interactions as 'just another variable' with missing values)³⁰. Then, the obesity-biomarker interaction terms were generated from the multiply imputed datasets³⁰.

Mediation analysis was performed for each imputed dataset and standard errors for TE, NDE, and NIEs were estimated using 1000 bootstrap samples. Rubin's rules were then applied to calculate the final MI estimates of effects and 95% CIs by pooling the estimated effects across the imputed datasets and deriving the standard errors from the within- and between-imputation variances³¹.

Additional Analyses: The following additional mediation analyses were performed: i) for the outcome models, including an interaction term between the adiposity measure and each biomarker, as well as between CRP-HbA1c, CRP-SHBG, CRP-testosterone, HbA1C-SHBG, HbA1C-testosterone, and SHBG-testosterone; ii) using calculated free testosterone as the biomarker of sex-steroid hormone pathway (instead of testosterone and SHBG combined); iii) excluding rectal cancer cases and limiting analyses to colon cancer; and iv) excluding colorectal cancer cases diagnosed within the first two years of follow-up and non-cases with follow-up ≤ 2 years.

All analyses were performed using Stata version 15³².

Data availability

UK Biobank is an open-access resource. Bonafide researchers can apply to use the UK Biobank dataset by registering and applying at <http://ukbiobank.ac.uk/register-apply/>.

Results

Overall, 179,936 men and 136,578 women met the eligibility criteria for this analysis (**Figure 1**). Men with waist circumference >102 vs ≤ 94 cm had higher median red or processed meat intake, and a higher proportion had lower physical activity levels, were former smokers, and reported regular aspirin or ibuprofen use. Similar differences were observed for women with waist circumference >88 vs ≤ 80 (**Table 1**; see **Supplementary Tables 1 and 2** for distributions of baseline characteristics by BMI and waist-hip ratio categories respectively). During a median follow-up time of 7.1 years, 1,280 and 790 CRC cases were diagnosed in men and postmenopausal women, respectively.

The ICCs (95% CI) for the biomarkers ranged from 0.56 (95%CI, 0.55-0.57) to 0.86 (95%CI, 0.86-0.87) for CRP and SHBG respectively in men and 0.67 (95%CI, 0.66-0.68) to 0.81 (95%CI, 0.80-0.82) for CRP and SHBG respectively in women (**Supplementary Table 3**).

Adiposity-biomarker associations: For men with waist circumference >102 vs ≤ 94 cm, an inverse association was observed for SHBG (GMR 0.79; 95%CI, 0.79-0.79) and testosterone (GMR 0.83; 95%CI, 0.82-0.83); a positive association for CRP (GMR 2.00; 95%CI, 1.97-2.02); and a weak positive association for HbA1c (GMR 1.04; 95%CI, 1.04-1.05). For women with waist circumference >88 vs ≤ 80 cm, an inverse association was observed for SHBG (GMR 0.63; 95%CI, 0.63-0.63); a positive association for CRP (GMR 2.64; 95%CI, 2.60-2.67); and weak positive associations for testosterone (GMR 1.08; 95%CI, 1.07-1.09) and HbA1c (GMR 1.04; 95%CI, 1.04-1.04) (**Table 2**). The associations were similar for BMI and waist-hip ratio in both men and women. (See **Supplementary Table 4** for complete case analysis).

Adiposity-CRC & Biomarker-CRC associations: The adjusted OR for CRC risk for men with waist circumference >102 vs ≤ 94 cm was 1.38 (95% CI, 1.20-1.59). Similar associations were observed for BMI and waist-hip ratio. A positive association was observed for CRP after adjusting for confounders and waist circumference (OR per doubling concentration 1.07; 95% CI, 1.02-1.11), and there was suggestion for an

inverse association for SHBG (OR 0.92; 95% CI, 0.82-1.03) and testosterone (OR 0.92; 95% CI, 0.81-1.05) after additionally adjusting for CRP, and HbA1c. There was no evidence for an association between HbA1c and CRC risk (OR additionally adjusted for CRP 0.82; 95% CI, 0.58-1.14) (**Table 3** multiple imputation analysis; **Supplementary Table 5** complete case analysis). No strong evidence for departure from linearity for adiposity-CRC or biomarker-CRC associations was observed (**Table 3**).

For women with waist circumference >88 vs ≤ 80 cm, the adjusted OR for CRC was 1.28 (95% CI, 1.08-1.51). A weaker association was observed for BMI (≥ 30 vs < 25 kg/m², OR 1.11; 95% CI, 0.91-1.36) and a stronger association for waist-hip ratio (>0.89 vs ≤ 0.77 , OR 1.33; 95% CI, 1.07-1.65). The point estimates and 95% CI were suggestive of weak-to-null associations between biomarkers and CRC (adjusted OR per doubling CRP 1.03; 95% CI, 0.98-1.09; HbA1C 1.06; 95% CI, 0.66-1.71; SHBG 1.03, 95% CI, 0.90-1.18; testosterone 1.00; 95% CI, 0.88-1.13) (**Table 3** multiple imputation analysis; **Supplementary Table 5** complete case analysis). There was evidence for a non-linear SHBG-CRC association for women.

Therefore, in the models for mediation analysis for women, SHBG was included as restricted cubic splines.

Sequential mediation analysis: For men with waist circumference >102 vs ≤ 94 cm, the RR^{NIE} was 1.08 (95% CI, 1.01-1.16) through all the biomarkers, 1.06 (95% CI, 1.01-1.11) through CRP and pathways influenced by CRP, 0.99 (95% CI, 0.97-1.01) through HbA1c excluding the influence of CRP on HbA1c levels, and 1.03 (95% CI, 0.99-1.08) through SHBG and testosterone combined excluding the influences of CRP and HbA1c on SHBG and testosterone levels. The RR^{NDE} not through any of the included biomarkers was 1.26 (95% CI, 1.09-1.47). Similar patterns of mediation were observed for BMI and waist-hip ratio (**Table 4**, see **Supplementary Table 6** for complete case analysis).

For women, the RR^{NIE} was 1.08 (95% CI, 0.95-1.22) through all the biomarkers, 1.08 (95% CI, 0.99-1.17) through pathways influenced by CRP, 1.00 (95% CI, 0.98-1.02) through HbA1c excluding the influence of

CRP, and 1.00 (95% CI, 0.92-1.09) through SHBG and testosterone combined excluding the influences of CRP and HbA1c. The RR^{NDE} not through any of the included biomarkers was 1.18 (95% CI, 0.96-1.45). Similar mediation effects were observed for waist-hip ratio. We did not attempt to perform mediation analysis for BMI, because no total effect of BMI on CRC was observed in the multiple imputation (**Table 4**) or complete case analyses (**Supplementary Table 6**).

Additional analyses

Patterns of mediation were similar in analyses that included all possible adiposity-mediator and mediator-mediator interaction terms (**Supplementary Table 7**), when calculated free testosterone was used as the biomarker of sex-steroid hormone pathway (**Supplementary Table 8**), in analyses limited to colon cancer cases (**Supplementary Table 9**), and in analyses limited to colorectal cancer cases diagnosed after two years of follow-up and non-cases with more than two years of follow-up (**Supplementary Table 10**).

Potential influence of missing data

We had missing biomarker data for 18% of men and 35% of women. Baseline characteristics were comparable for men with and without missing biomarker data. A slightly larger proportion of women with missing biomarker data had waist circumference ≤ 80 cm (41% vs 39%) or BMI < 25 kg/m² (40% vs 37%) compared with women with complete biomarker data (**Supplementary Table 11**).

For all analyses, results from complete case analyses are presented (**Supplementary Table 4 to 11**).

Multiple imputation and complete case analyses yielded similar results but the 95% CIs from multiple imputation analyses were slightly narrower.

Discussion

For men and postmenopausal women, the effect of adiposity on CRC not explained by the included biomarkers was larger than the estimated indirect effect through all biomarkers. Pathways originating with CRP explained a small proportion of the adiposity-CRC association. In men, a small part of the adiposity-CRC association was explained by adiposity-induced reduced levels of SHBG and testosterone.

To our knowledge, this is one of the first and largest studies that has attempted to quantify the mediating effects of multiple pathways on the adiposity-CRC association, while allowing for correlation between mediators and interrelation between pathways¹⁷. The causal mediation analysis approach was able to handle non-linear mediator-outcome associations and exposure-mediator or mediator-mediator interactions¹⁷.

These are important advantages over other statistical methods that have traditionally been used to assess mediation²⁸. The path-specific indirect effects were estimated using a sequential approach. An unverifiable assumption required by this approach is that the prespecified causal ordering of the pathways is true¹⁷. For this analysis, based on current evidence, we assumed that inflammatory and insulin signaling pathways preceded and possibly influenced SHBG and testosterone levels, while inflammatory pathway preceded dysregulated insulin signaling and glucose homeostasis^{9, 14}.

We made use of all the relevant biomarkers in the UK Biobank, but measurements were available for only a limited number of biomarkers, which was perhaps the most important limitation of this study. The observed mediating role for the pathways, or lack thereof, would have depended on how well the available biomarkers captured the biological characteristics of each pathway. For all participants, we used measurements for the biomarkers at one time point only. The differences in temporal variability and measurement quality of the biomarkers may have introduced varying degrees of information bias into our estimated associations and influenced the strength of the observed indirect effects. Overall, however, the biomarker measurements were of high quality (all CVs <9%)^{21, 23, 24} and, uniquely, were available in nearly all cohort participants.

Additionally, for all biomarkers, the ICCs calculated for the subgroup of participants who had measurements

available for two time points (approximately four years apart) were >0.55 , suggesting that single measurements provided a reasonable estimate of longer-term exposures. In the UK Biobank, height, weight, waist and hip circumference were measured following a published protocol to reduce measurement error²⁰. In addition to BMI, we used waist circumference and waist-hip ratio as measures of adiposity, which may perform better in identifying individuals with obesity³³. Our study did not provide evidence for an association between BMI and colorectal cancer risk in women, which was similar to a previous analysis of the UK Biobank data (hazard ratio (HR) for highest vs lowest quintile 1.11; 95%CI, 0.89-1.38)³⁴, but inconsistent with the results of a recent meta-analysis of studies investigating the association between BMI and colorectal cancer (summary RR for women per 5 kg/m² 1.05; 95%CI, 1.02-1.08)³⁵. It is unclear why we found no association between BMI and colorectal cancer for women. One possibility is that a longer duration of follow-up and a larger number of cases may be required to detect the expected weaker (compared with men) BMI and colorectal cancer association. In support of this supposition, in the European Prospective Investigation into Cancer and Nutrition (EPIC) study after an average of 6.1 years of follow-up (n=1570 cases), no association was found between BMI and colorectal cancer for women³⁶. However, in a more recent EPIC study that included data after an average of 14.9 years of follow-up (n=6291 cases), the expected positive association between BMI and colorectal cancer for women was observed³⁷.”

The sequential mediation analysis relied on assumptions of no unmeasured confounding of exposure-outcome, mediator-outcome, exposure-mediator, and mediator-mediator relations¹⁷. We had data on several potential confounders and took those into account in our analyses, but due to the observational nature of this study, residual confounding due to unmeasured or mismeasured confounders cannot be ruled out. For the adiposity-biomarker associations, this study was cross-sectional and, thus, prone to reverse causation. We

excluded CRC cases diagnosed within the first year of follow-up to reduce the possibility of outcome having influenced adiposity or biomarker measurements.

Using causal mediation analysis in a nested case-control study within the Health Professionals Follow-up Study (HPFS), it was observed that a large proportion of the BMI-CRC association (OR per 3.6 kg/m² 1.40; 95%CI, 1.14-1.73) was explained by all inflammatory (CRP, interleukin-6, tumour necrosis factor receptor 2, and macrophage inhibitory cytokine-1) and metabolic (adiponectin, C-peptide, and soluble leptin receptor (sOB-R)) biomarkers (OR^{NIE} 1.26; 95%CI, 0.97-1.52)³⁸. The authors repeated the mediation analysis separately for the two biomarker groups, assuming that the pathways were independent. Although the HPFS included more inflammatory biomarkers compared with ours, similar to our results, the observed indirect effect through inflammation was small (OR^{NIE} 1.05; 95%CI, 0.96-1.14)³⁸. Also, in a nested case-control study within the European Investigation into Cancer and Nutrition, adjusting for CRP only changed the RR for waist circumference-CRC association from 1.33 (95%CI, 1.26-1.41) per 10 cm increment to 1.31 (95%CI, 1.24-1.39) in men and from 1.20 (95%CI, 1.14-1.25) to 1.19 (95%CI, 1.14-1.25) in women³⁹. Additional studies with a wider range of inflammatory biomarkers are needed to further explore the mediating role of inflammation in adiposity-CRC association.

The HPFS nested case-control study found a larger OR^{NIE} for the metabolic biomarkers (OR^{NIE} 1.24; 95%CI, 0.92-1.55), which was mainly driven by adiponectin and sOB-R³⁸. When the causal mediation analysis was repeated for individual biomarkers, weak to no mediating effect was observed for C-peptide (a biomarker for insulin signaling; OR^{NIE} 1.05; 95%CI, 0.91-1.18)³⁸. In a cohort study with 4,032 colon and 2,430 rectal cancer cases, the total BMI-colon (HR per 5 kg/m² 1.14; 95%CI, 1.10-1.19) and -rectal cancer (HR 1.09; 95%CI, 1.03-1.15) associations were weakly mediated by triglyceride-glucose index (TyG; a proxy for insulin resistance) (HR^{NIE} 1.03; 95%CI, 1.01-1.04 for colon and HR^{NIE} 1.03; 95%CI, 1.01-1.05 for rectal cancer)⁴⁰. The strength of the mediating effect was similar in men and women, and after excluding

participants who did not have diabetes ⁴⁰. In our study, we did not observe any mediating effect through HbA1c. The observed indirect effect through TyG in the study by Fritz *et al.* may have been overestimated because the analysis did not allow for potential confounding by the preceding inflammatory pathway ⁴⁰. In our study, the estimated indirect effect through HbA1c excluded the possible influence of CRP on HbA1c, thus removed the confounding effect of CRP ¹⁷. This issue, however, is unlikely to fully explain the null indirect effect for HbA1c we observed. In our study of men and postmenopausal women who at baseline did not have diabetes and did not report taking diabetes medication, there was no evidence for an association between HbA1c and CRC, regardless of adjustment for CRP. Similarly, a recent Mendelian randomization analysis found no evidence of an effect between genetically-predicted HbA1c concentration and colorectal cancer risk (OR per genetically determined 1 standard deviation (SD) increment 1.02; 95% CI, 0.85 – 1.22) ⁴¹. Fritz *et al.* observed an association between TyG and colon (HR per 1 SD 1.07; 95% CI, 1.03-1.10) and rectal (HR 1.09; 95% CI, 1.04-1.14) cancers ⁴⁰. Although HbA1c is an established long-term biomarker of glucose and insulin metabolism ⁴², it may not be the ideal biomarker of dysregulated insulin signaling on its own.

We are not aware of other studies that have investigated the mediating role of sex-steroid hormones in the adiposity-CRC association. We were also limited in exploring the role of this pathway fully because we did not have measures for estrogens. Our results provided evidence for a small role of adiposity-induced reduced levels of SHBG and testosterone in explaining the increased CRC risk in men with obesity. This observation is in line with the existing evidence on the potential protective role of androgens on CRC development ⁴³. It may also partly explain the stronger adiposity-CRC association observed in men because as found in ours and previous studies, adiposity is associated with reduced testosterone levels in men, but not in women ⁴³.

In summary, increased CRP and reduced SHBG and testosterone levels explained a small proportion of the effect of adiposity on CRC in men. Increased CRP also had a small mediating effect on the association in postmenopausal women, comparable to the mediating effect observed in men. We did not observe any mediating effect for HbA1c. A large proportion of the effect of adiposity on CRC was not explained by the included biomarkers. These results suggest that pathways other than those captured here may also be important for understanding the mechanisms linking adiposity and CRC and for identifying targets for CRC prevention in individuals with obesity. Future studies with a wider range of biomarkers reflecting inflammatory status, impaired insulin signaling and hyperinsulinemia, and altered sex-steroid hormone levels in the obese state are needed to perform a comprehensive evaluation of the role of these pathways in explaining the effect of adiposity on CRC link. Proper causal mediation analysis approaches would allow such investigation, while appropriately accounting for correlations between biomarkers.

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Tables

Table 1 - Baseline Characteristics of men and women eligible for the study by categories of **waist circumference**

	Men			Women		
Waist circumference, cm	≤94 N=83,940	>94 - ≤102 N=51,228	>102 N=44,768	≤80 N=54,339	>80 - ≤88 N=35,820	>88 N=46,419
Colorectal cancer affected - yes, n (%)	478 (1)	402 (1)	400 (1)	288 (1)	174 (0)	328 (1)
Age at recruitment, median (25th - 75th percentiles)	56.0 (48.0-62.0)	58.0 (50.0-63.0)	59.0 (51.0-64.0)	60.0 (56.0-64.0)	61.0 (57.0-65.0)	61.0 (57.0-65.0)
Age at cancer diagnosis or the end of follow up, median (25th - 75th percentiles)	63.3 (55.4-69.4)	65.2 (57.5-70.5)	65.7 (58.5-70.6)	67.4 (63.2-71.0)	68.2 (64.3-71.8)	68.2 (64.4-71.8)
Follow-up time - years, median (25th - 75th percentiles)	7.1 (6.5-7.7)	7.2 (6.4-7.8)	7.1 (6.4-7.7)	7.1 (6.5-7.7)	7.1 (6.4-7.7)	7.1 (6.4-7.7)
Highest educational attainment, n (%)						
None of the below	10,106 (12)	8,040 (16)	8,674 (19)	9,026 (17)	7,609 (21)	11,596 (25)
CSEs/O-levels/GCSEs or equivalent	20,321 (24)	11,739 (23)	10,056 (22)	14,853 (27)	9,648 (27)	11,801 (25)
NVQ/HND/HNC/A-levels or equivalent	16,959 (20)	10,607 (21)	9,456 (21)	6,667 (12)	4,461 (12)	6,277 (14)
Other professional qualifications. (e.g. nurse/teacher)	24,048 (29)	14,462 (28)	11,878 (27)	17,327 (32)	10,530 (29)	12,928 (28)
College/university degree	12,506 (15)	6,380 (12)	4,704 (11)	6,466 (12)	3,572 (10)	3,817 (8)
Townsend deprivation index at recruitment - quintiles, n (%)						
1 (least degree of deprivation)	18,372 (22)	11,232 (22)	8,861 (20)	12,886 (24)	7,751 (22)	8,605 (19)
2	17,559 (21)	10,944 (21)	8,758 (20)	12,218 (22)	7,824 (22)	9,144 (20)
3	16,739 (20)	10,443 (20)	8,943 (20)	11,392 (21)	7,736 (22)	9,369 (20)
4	16,157 (19)	9,772 (19)	9,184 (21)	10,152 (19)	7,005 (20)	9,669 (21)
5 (highest degree of deprivation)	15,113 (18)	8,837 (17)	9,022 (20)	7,691 (14)	5,504 (15)	9,632 (21)
First degree relative history of colorectal cancer - yes, n (%)	8,923 (11)	6,041 (12)	5,441 (12)	6,316 (12)	4,383 (12)	5,680 (12)
Physical activity - MET hr/wk, n (%)						
<10	13,575 (16)	10,941 (21)	12,768 (29)	9,968 (18)	7,964 (22)	14,117 (30)
10-<20	13,279 (16)	9,006 (18)	8,094 (18)	9,513 (18)	6,678 (19)	8,771 (19)
20-<40	20,937 (25)	12,163 (24)	9,805 (22)	13,531 (25)	8,840 (25)	10,257 (22)
40-<60	12,273 (15)	6,546 (13)	4,739 (11)	7,739 (14)	4,530 (13)	5,088 (11)
60+	23,876 (28)	12,572 (25)	9,362 (21)	13,588 (25)	7,808 (22)	8,186 (18)
Alcohol intake, n (%)						
never	4,431 (5)	2,522 (5)	2,537 (6)	4,340 (8)	2,884 (8)	5,091 (11)
Current occasionally to						
1-3 times/month	12,412 (15)	7,200 (14)	7,668 (17)	12,576 (23)	9,328 (26)	14,923 (32)
1-2 times/week	21,613 (26)	13,007 (25)	11,956 (27)	13,647 (25)	9,239 (26)	11,637 (25)
3-4 times/week	23,358 (28)	14,520 (28)	11,335 (25)	12,790 (24)	7,724 (22)	8,101 (17)
daily or almost daily	22,126 (26)	13,979 (27)	11,272 (25)	10,986 (20)	6,645 (19)	6,667 (14)
Smoking status, n (%)						
Never	47,086 (56)	24,891 (49)	19,373 (43)	33,719 (62)	20,990 (59)	25,575 (55)
Former	26,615 (32)	20,485 (40)	20,250 (45)	16,693 (31)	12,151 (34)	17,221 (37)
Current	10,239 (12)	5,852 (11)	5,145 (11)	3,927 (7)	2,679 (7)	3,623 (8)

Total red and processed meat intake - times/week, median (25th - 75th percentiles)	3.5 (2.5-5.0)	4.0 (2.5-5.0)	4.5 (2.5-5.5)	2.5 (2.0-4.0)	2.5 (2.0-4.5)	3.0 (2.0-4.5)
Regular aspirin or ibuprofen use - yes, n (%)	18,700 (22)	14,187 (28)	14,657 (33)	10,656 (20)	8,427 (24)	12,954 (28)
Number of live births, median (25th - 75th percentiles)	-	-	-	2.0 (1.0-2.0)	2.0 (1.0-3.0)	2.0 (1.0-3.0)

Table 1 - Baseline Characteristics of men and women eligible for the study by categories of **waist circumference**

Men				Women		
Ever use of oral contraceptive pill - yes, n (%)	-	-	-	43,179 (79)	28,021 (78)	35,539 (77)
Ever use of hormone therapy - yes, n (%)	-	-	-	24,233 (45)	17,395 (49)	22,551 (49)
Adiposity measures						
Waist Circumference - cm, median (25th - 75th percentiles)	88.0 (84.0-92.0)	98.0 (96.0-100.0)	108.0 (105.0-114.0)	75.0 (71.0-78.0)	84.0 (82.0-86.0)	96.0 (92.0-102.0)
Body Mass Index - kg/m ² , n (%)						
≥18.5-<25	44,047 (52)	3,134 (6)	93 (0)	41,836 (77)	9,267 (26)	1,213 (3)
≥25-<30	38,951 (46)	39,700 (77)	12,529 (28)	12,243 (23)	23,690 (66)	17,688 (38)
≥30	942 (1)	8,394 (16)	32,146 (72)	260 (0)	2,863 (8)	27,518 (59)
Body Mass Index - kg/m ² , median (25th - 75th percentiles)	24.9 (23.3-26.3)	27.9 (26.6-29.3)	31.6 (29.7-34.0)	23.3 (21.8-24.8)	26.4 (24.9-28.0)	30.9 (28.5-34.0)
Waist-Hip Ratio, n (%)						
≤0.89 M; ≤0.77 W	44,766 (53)	3,681 (7)	409 (1)	27,045 (50)	3,140 (9)	874 (2)
>0.89 - ≤0.98 M; >0.77 - ≤0.86 W	37,589 (45)	37,444 (73)	16,277 (36)	26,254 (48)	25,958 (72)	18,284 (39)
0.98 M; >0.86 W	1,585 (2)	10,103 (20)	28,082 (63)	1,040 (2)	6,722 (19)	27,261 (59)
Waist-Hip ratio, median (25th - 75th percentiles)	0.9 (0.9-0.9)	0.9 (0.9-1.0)	1.0 (1.0-1.0)	0.8 (0.7-0.8)	0.8 (0.8-0.9)	0.9 (0.8-0.9)
Biomarkers, median (25th - 75th percentiles)						
C-reactive protein - mg/L	0.9 (0.5-1.7)	1.3 (0.8-2.4)	2.0 (1.1-3.6)	0.8 (0.4-1.6)	1.4 (0.8-2.6)	2.5 (1.4-4.7)
C-reactive protein missing value - yes, n (%)	4,369 (5)	2,633 (5)	2,601 (6)	3,053 (6)	2,049 (6)	2,943 (6)
Glycated hemoglobin-nmol/mol	34.4 (32.1-36.6)	35.0 (32.6-37.4)	36.0 (33.4-38.6)	35.2 (33.2-37.3)	35.7 (33.6-37.9)	36.6 (34.3-39.1)
Glycated hemoglobin-missing value - yes, n (%)	4,731 (6)	2,844 (6)	2,643 (6)	3,284 (6)	2,277 (6)	3,134 (7)
Sex Hormone Binding Globulin - nmol/L	40.8 (31.3-52.4)	35.2 (27.0-45.3)	32.9 (24.9-42.6)	67.5 (52.4-85.7)	53.1 (40.3-68.8)	42.1 (31.3-55.9)
Sex Hormone Binding Globulin missing value - yes, n (%)	11,043 (13)	6,543 (13)	5,843 (13)	8,083 (15)	5,256 (15)	6,986 (15)
Testosterone - pmol/L	12.7 (10.5-15.2)	11.4 (9.4-13.7)	10.4 (8.5-12.6)	0.9 (0.7-1.3)	1.0 (0.7-1.3)	1.0 (0.7-1.4)
Testosterone missing value - yes, n (%)	4,882 (6)	2,897 (6)	2,895 (6)	13,080 (24)	8,261 (23)	10,116 (22)
Missing for any of the biomarkers - yes, n (%)	15,223 (18)	9,074 (18)	8,130 (18)	18,919 (35)	12,077 (34)	15,121 (33)

CSE Certificate of Secondary Education; GCSE General Certificate of Secondary Education; NVQ National Vocational Education; HND Higher National Diploma; HNC Higher National Certificate; MET Metabolic Equivalent of Task; W women; M men

Baseline Characteristics of men and women eligible for the study by categories of body mass index and waist-hip ratio are presented in Supplementary Table 1 and Supplementary Table 2 respectively.

Table 2 - Association between adiposity measures and biomarkers; **multiple imputation analyses** excluding women in the middle category of each adiposity measure

Ratio of geometric means (95% confidence interval)							
Biomarker	Waist circumference		Body Mass Index		Waist-Hip Ratio		
	>102 vs. ≤94 M >88 vs. ≤80 F		≥30 vs. <25 kg/m ²		>0.98 vs. ≤89 M >0.86 vs. ≤77 F		
	N=128,708		N=88,756		N=88,626		
Men	CRP	2.00 (1.97 to 2.02)	2.31 (2.28 to 2.34)	2.10 (2.07 to 2.13)			
	HbA1C	1.04 (1.04 to 1.05)	1.05 (1.05 to 1.05)	1.05 (1.05 to 1.05)			
	SHBG	0.79 (0.79 to 0.79)	0.72 (0.72 to 0.73)	0.77 (0.76 to 0.77)			
	Testosterone	0.83 (0.82 to 0.83)	0.80 (0.80 to 0.80)	0.83 (0.83 to 0.84)			
	N=100,758		N=82,957		N=66,082		
Women	CRP	2.64 (2.60 to 2.67)	3.19 (3.15 to 3.24)	2.19 (2.16 to 2.23)			
	HbA1C	1.04 (1.04 to 1.04)	1.04 (1.04 to 1.04)	1.05 (1.05 to 1.05)			
	SHBG	0.63 (0.63 to 0.63)	0.61 (0.60 to 0.61)	0.63 (0.62 to 0.63)			
	Testosterone	1.08 (1.07 to 1.09)	1.12 (1.11 to 1.13)	1.04 (1.03 to 1.04)			

Results from complete case analysis are provided in Supplementary Table 5.

Abbreviations: CRP C-reactive protein; HbA1c glycated hemoglobin; SHBG sex-hormone-binding globulin

Missing biomarker data were multiply imputed (complete case analysis results are presented in Supplementary Table 4).

Models were adjusted for age at recruitment, education, Townsend deprivation index at recruitment, first-degree relative history of colorectal cancer, physical activity, alcohol intake, smoking status, red and processed meat intake, and regular use of aspirin or ibuprofen. Models for women were additionally adjusted for the number of live births, ever use of oral contraceptive pill, and ever use of hormone therapy.

Table 3 - Estimated odds ratios and 95% confidence intervals for the association between adiposity measures (exposure) or biomarkers (mediators) and colorectal cancer (outcome) in men and women; **multiple imputation analysis**

Men				Women			
Waist Circumference, cm		>102 vs <94		>88 vs <80			
OR (95% CI)		1.38 (1.20 to 1.59)		1.28 (1.08 to 1.51)			
Body Mass Index, kg/m²		≥30 vs <25		≥30 vs <25			
OR (95% CI)		1.37 (1.16 to 1.62)		1.11 (0.91 to 1.36)			
Waist-Hip ratio		>0.98 vs <0.89		>0.86 vs <0.77			
OR (95% CI)		1.42 (1.19 to 1.70)		1.33 (1.07 to 1.65)			
Biomarker		Per doubling concentration	P value for evidence against linearity		Per doubling concentration	P value for evidence against linearity	
CRP							
Model 1		1.07 (1.02 to 1.11)	0.02		1.03 (0.98 to 1.09)	0.75	
HbA1C							
Model 1		0.85 (0.61 to 1.19)			1.08 (0.67 to 1.75)		
Model 2 (additionally adjusted for CRP)		0.82 (0.58 to 1.14)	0.15		1.06 (0.66 to 1.71)	0.75	
SHBG							
Model 1		0.92 (0.82 to 1.02)			1.01 (0.89 to 1.15)		
Model 2 (additionally adjusted for CRP and HbA1c)		0.92 (0.82 to 1.03)	0.20		1.03 (0.90 to 1.18)	0.00	
Testosterone							
Model 1		0.91 (0.80 to 1.04)			1.00 (0.88 to 1.13)		
Model 2 (additionally adjusted for CRP and HbA1c)		0.92 (0.81 to 1.05)	0.56		1.00 (0.88 to 1.13)	0.19	

Results from complete case analysis are provided in Supplementary Table 6.

Abbreviations: CRP C-reactive protein; HbA1c glycated hemoglobin; SHBG sex-hormone-binding globulin

Missing biomarker data were multiply imputed (complete case analysis results are presented in Supplementary Table 5).

Models were adjusted for age at recruitment, education, Townsend deprivation index at recruitment, first-degree relative history of colorectal cancer, physical activity, alcohol intake, smoking status, red and processed meat intake, and regular use of aspirin or ibuprofen. Models for women were additionally adjusted for the number of live births, ever use of oral contraceptive pill, and ever use of hormone therapy. Model 1 for biomarkers were additionally adjusted for waist circumference.

Table 4- Estimated natural direct and indirect effects using sequential mediation analysis for the association between adiposity measures and colorectal cancer for men and women, analyses exclude participants categorized as overweight and do not include any interaction terms between adiposity measures and biomarkers (exposure-mediator interactions) or biomarkers (mediator-mediator interactions) - **multiple imputation analysis**

		Risk Ratio (95% CI)	
Effect		Men	Women
Waist Circumference >102 vs. ≤94 cm M >88 vs. ≤80 cm F	No. Colorectal Cancer affected/ No. unaffected	878/127,830	616/100,142
	Total effect	1.37 (1.19 to 1.58)	1.27 (1.07 to 1.50)
	Natural indirect effect through all the mediators	1.08 (1.01 to 1.16)	1.08 (0.95 to 1.22)
	Natural indirect effect through CRP	1.06 (1.01 to 1.11)	1.08 (0.99 to 1.17)
	Natural indirect effect through HbA1C, excluding the influence of CRP	0.99 (0.97 to 1.01)	1.00 (0.98 to 1.02)
	Natural indirect effect through SHBG and testosterone, excluding the influence of CRP and HbA1c	1.03 (0.99 to 1.08)	1.00 (0.92 to 1.09)
	Natural direct effect not through any of the mediators	1.26 (1.09 to 1.47)	1.18 (0.96 to 1.45)
Body Mass Index ≥30 vs. <25 kg/m ²	No. Colorectal Cancer affected/ No. unaffected	614/88,142	451/82,506
	Total effect	1.36 (1.15 to 1.60)	1.12 (0.92 to 1.38)
	Natural indirect effect through all the mediators	1.11 (1.00 to 1.22)	- - -
	Natural indirect effect through CRP	1.06 (0.99 to 1.13)	- - -
	Natural indirect effect through HbA1C, excluding the influence of CRP	0.99 (0.96 to 1.02)	- - -
	Natural indirect effect through SHBG and testosterone, excluding the influence of CRP and HbA1c	1.05 (0.99 to 1.12)	- - -
	Natural direct effect not through any of the mediators	1.23 (1.02 to 1.48)	- - -
Waist-Hip Ratio >0.98 vs. <0.89 M >0.86 vs. ≤0.77 F	No. Colorectal Cancer affected/ No. unaffected	600/88,026	378/65,704
	Total effect	1.31 (1.09 to 1.58)	1.33 (1.06 to 1.66)
	Natural indirect effect through all the mediators	1.12 (1.03 to 1.23)	1.10 (0.96 to 1.26)
	Natural indirect effect through CRP	1.07 (1.00 to 1.14)	1.09 (1.00 to 1.19)
	Natural indirect effect through HbA1C, excluding the influence of CRP	0.99 (0.96 to 1.02)	1.01 (0.97 to 1.05)
	Natural indirect effect through SHBG and testosterone, excluding the influence of CRP and HbA1c	1.06 (1.01 to 1.11)	1.00 (0.91 to 1.11)
	Natural direct effect not through any of the mediators	1.17 (0.95 to 1.44)	1.20 (0.92 to 1.57)

Results from complete case analysis are provided in Supplementary Table 7.

Abbreviations: CRP C-reactive protein; HbA1c glycated hemoglobin; SHBG sex-hormone-binding globulin

Models were adjusted for age at recruitment, education, Townsend deprivation index at recruitment, first-degree relative history of colorectal cancer, physical activity, alcohol intake, smoking status, red and processed meat intake, and regular use of aspirin or ibuprofen. Models for women were additionally adjusted for the number of live births, ever use of oral contraceptives, and ever use of hormone therapy.

Figure Legends

Figure 1a- Flow diagram demonstrating the selection process of men in the UK Biobank for the present study

Figure 1b- Flow diagram demonstrating the selection process of women in the UK Biobank for the present study

Figure 2 - Schematic interpretation of the estimated total, natural direct, and natural indirect effects

In this study, C-reactive protein (CRP) was used as a general marker for inflammation, HbA1C as a marker for glycemic control and a correlate of insulin resistance, and sex-hormone-binding globulin (SHBG) and testosterone treated jointly as makers of the sex-steroid hormone pathway. Waist circumference, BMI, and waist-hip ratio were used as proxy measures for adiposity in separate sets of analyses. In each provided outcome model needed to estimate the effects, y is the outcome (colorectal cancer incidence), a is the exposure (each adiposity measure comparing obese vs normal), c confounders, $m1$ CRP, $m2$ HbA1c, $m3a$ SHBG, and $m3b$ testosterone.

Figure 2 – Schematic interpretation of the estimated total, natural direct, and natural indirect effects

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The total effect of adiposity and CRC

This effect captures the adiposity-CRC association through the included biomarkers (the indirect effect) and not through the included biomarkers (referred to as the direct effect).

Required models for the outcome

The natural direct effect (NDE) not through the included biomarkers

This effect captures the adiposity-CRC association not through any of the included biomarkers (the solid arrow).

logistic $y \sim x + c$

Required models for the outcome

The natural indirect effect through all included biomarkers

This effect captures the adiposity-CRC association through all the included biomarkers (the solid arrows).

logistic $y \sim x + m1 + m2 + m3a + m3b + c$ & logistic $y \sim x + c$

Required models for the outcome

The natural indirect effect through pathways originating with CRP

This effect captures the effect of adiposity-induced changes in CRP levels on CRC risk directly (1), or indirectly through influencing HbA1C (2) or SHBG and testosterone (3) levels.

logistic $y \sim x + m1 + m2 + m3a + m3b + c$ & logistic $y \sim x + c$

Required models for the outcome

The natural indirect effect through HbA1C beyond the influence of C-reactive protein

This effect (C) is estimated as the difference between the indirect effect through CRP and HbA1C (A) and the indirect effect through CRP (B). It thus excludes the possible influence of CRP on HbA1C levels and captures the effect of adiposity-induced changes in HbA1C on CRC risk directly (1), or indirectly through influencing SHBG and testosterone levels (2)

logistic $y \sim x + m1 + m2 + c$ & logistic $y \sim x + c$

Required models for the outcome

The natural indirect effect through sex-hormone-binding globulin and testosterone beyond the influences of C-reactive protein and HbA1C

This effect (C) is estimated as the difference between the indirect effect through CRP, HbA1C, and SHBG and testosterone (A) and the indirect effect through CRP and HbA1C (B). It thus excludes the possible influence of CRP or HbA1C on SHBG and testosterone levels and captures the effect adiposity-induced change in SHBG and testosterone have on CRC risk (1).

logistic $y \sim x + m1 + m2 + m3a + m3b + c$ & logistic $y \sim x + c$

Required models for the outcome

logistic $y \sim x + m1 + m2 + m3a + m3b + c$ & logistic $y \sim x + c$

logistic $y \sim x + m1 + m2 + c$ & logistic $y \sim x + c$

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Mechanisms underlying the association between adiposity and colorectal cancer (CRC) are not well understood. Here, using UK Biobank data and sequential mediation analysis, the authors quantified the effects of C-reactive protein, hemoglobin-A1c, sex-hormone-binding globulin (SHBG), and testosterone on the adiposity-CRC association in men and postmenopausal women. Analyses show that, in both men and women, increased CRP had small mediating effects on adiposity and CRC. In men, a small proportion of the effect was further explained by reduced SHBG and testosterone levels. The results suggest that pathways other than those captured in the study are important for understanding the adiposity-CRC link.

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