

# BMJ Sexual & Reproductive Health

## The relationship between chronic diseases and number of sexual partners: an exploratory analysis

Journal:	<i>BMJ Sexual &amp; Reproductive Health</i>
Manuscript ID	bmjsrh-2019-200352.R3
Article Type:	Original research
Date Submitted by the Author:	08-Dec-2019
Complete List of Authors:	Grabovac, Igor; Medizinische Universitat Wien, Smith, Lee; ARU Yang, Lin; Alberta Health Services Soysal, Pinar Veronese, Nicola Turan ISIK, Ahmet Forwood, Suzanna Jackson, Sarah; UCL,
Keywords:	number of sexual partners, sexual activity, health outcomes, cancer, stroke, self-rated health
Abstract:	<p><b>Background:</b> We investigated sex-specific associations between lifetime number of sexual partners and several health outcomes in a large sample of older adults in England.</p> <p><b>Methods:</b> We used cross-sectional data from 2,537 men and 3,185 women aged <math>\geq 50</math> years participating in the English Longitudinal Study of Ageing. Participants reported the number of sexual partners they had had in their lifetime. Outcomes were self-rated health and self-reported limiting long-standing illness, cancer, coronary heart disease (CHD), and stroke. We used logistic regression to analyse associations between lifetime number of sexual partners and health outcomes, adjusted for relevant sociodemographic and health-related covariates.</p> <p><b>Results:</b> Having had 10 or more lifetime sexual partners was associated with higher odds of reporting a diagnosis of cancer than having had 0-1 sexual partners in men (OR=1.69, 95% CI 1.01-2.83) and women (OR=1.91, 95% CI 1.04-3.51), respectively. Women who had 10 or more lifetime sexual partners also had higher odds of reporting a limiting long-standing illness (OR=1.64, 95% CI 1.15-2.35). No other statistically significant associations were observed.</p> <p><b>Conclusions:</b> A higher lifetime number of sexual partners is associated with increased odds of reported cancer. Longitudinal research is required to establish causality. Understanding the predictive value of lifetime number of sexual partner as a behavioural risk factor may improve clinical assessment of cancer risk in older adults.</p> <p><b>Key words:</b> number of sexual partners; sexual activity; sexual history; health outcomes; self-rated health; cancer; coronary heart disease; stroke.</p>

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60



SCHOLARONE™  
Manuscripts

1  
2  
3 The relationship between chronic diseases and number of sexual partners: an  
4  
5 exploratory analysis  
6  
7

8 Igor Grabovac<sup>1</sup>, Lee Smith<sup>2\*</sup>, Lin Yang<sup>3</sup>, Pinar Soysal<sup>5</sup>, Nicola Veronese<sup>6</sup>, Ahmet Turan Isik<sup>5</sup>, Suzanna  
9 Forwood<sup>7</sup>, Sarah E Jackson<sup>8</sup>  
10  
11

12 <sup>1</sup>Department of Social and Preventive Medicine, Centre for Public Health, Medical University of  
13 Vienna, Vienna, Austria  
14  
15

16 <sup>2</sup>The Cambridge Centre for Sport and Exercise Sciences, Anglia Ruskin University, Cambridge, United  
17 Kingdom  
18  
19

20 <sup>3</sup>Department of Cancer Epidemiology and Prevention Research, Alberta Health Services, Calgary,  
21 Canada  
22  
23

24 <sup>4</sup>Preventive Oncology & Community Health Sciences, Cumming School of Medicine, University of  
25 Calgary, Calgary, Canada  
26  
27

28 <sup>5</sup>Department of Geriatric Medicine, Faculty of Medicine, Bezmialem Vakif University, Istanbul,  
29 Turkey  
30  
31

32 <sup>6</sup>National Research Council, Neuroscience Institute, Aging Branch, Padova, Italy  
33  
34

35 <sup>7</sup>Department of Psychology, Anglia Ruskin University, Cambridge, United Kingdom  
36  
37

38 <sup>8</sup>Department of Behavioural Science and Health, University College London, London, United Kingdom  
39  
40

41  
42 \*Corresponding Author:  
43

44  
45 Dr Lee Smith  
46

47  
48 The Cambridge Centre for Sport and Exercise Sciences, Anglia Ruskin University, Compass House  
49 Cambridge, UK, CB1 1PT; Email: lee.smith@anglia.ac.uk;  
50  
51

52 **Word Count:** 2687  
53  
54  
55  
56  
57  
58  
59  
60

## Abstract

Background: We investigated sex-specific associations between lifetime number of sexual partners and several health outcomes in a large sample of older adults in England.

Methods: We used cross-sectional data from 2,537 men and 3,185 women aged  $\geq 50$  years participating in the English Longitudinal Study of Ageing. Participants reported the number of sexual partners they had had in their lifetime. Outcomes were self-rated health and self-reported limiting long-standing illness, cancer, coronary heart disease (CHD), and stroke. We used logistic regression to analyse associations between lifetime number of sexual partners and health outcomes, adjusted for relevant sociodemographic and health-related covariates.

Results: Having had 10 or more lifetime sexual partners was associated with higher odds of reporting a diagnosis of cancer than having had 0-1 sexual partners in men (OR=1.69, 95% CI 1.01-2.83) and women (OR=1.91, 95% CI 1.04-3.51), respectively. Women who had 10 or more lifetime sexual partners also had higher odds of reporting a limiting long-standing illness (OR=1.64, 95% CI 1.15-2.35). No other statistically significant associations were observed.

Conclusions: A higher lifetime number of sexual partners is associated with increased odds of reported cancer. Longitudinal research is required to establish causality. Understanding the predictive value of lifetime number of sexual partner as a behavioural risk factor may improve clinical assessment of cancer risk in older adults.

**Key words:** number of sexual partners; sexual activity; sexual history; health outcomes; self-rated health; cancer; coronary heart disease; stroke.

1  
2  
3 **Key messages:**  
4  
5  
6

- 7
- 8 • Lifetime numbers of sexual partners is associated with sexually transmitted infection (STI)  
9 acquisition and the associated health risks which may affect health in later life.
  - 10 • Using cross-sectional data from a representative sample of older English adults we found  
11 that higher number of sexual partners is associated with increased risk of reporting a cancer  
12 diagnosis  
13
  - 14 • Sexual history may be a relevant clinical indicator for cancer risk in older patients  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

Confidential: For Review Only

## Background

There is a large body of literature investigating the relationship between sexual activities, including total number of sexual partners, and risk of developing sexually transmitted infections (STIs). To date, most of this research has focused on adolescents and young adults or the cost-effectiveness of preventative strategies<sup>1</sup>. Studies have shown that a greater number of sexual partners is associated with greater risk of contracting STIs in adolescents<sup>2</sup>.

STIs can have long-term consequences for health, including greater risk of specific cancers. Rates of human papillomavirus (HPV) infection in sexually active young females have been consistently reported to range from 19% to 46%<sup>3</sup>. Nearly all cases of cervical cancer can be attributable to HPV infection<sup>4</sup>. Moreover, HPV has been found to be associated with cancers of the mouth, penis and anus<sup>4</sup>, cancers that are most common in older adults<sup>5,6</sup>. Other STIs, such as gonorrhoea infection, have been shown to increase the risk of prostate cancer in black men<sup>7</sup>. The average age for men to be diagnosed with prostate cancer is between 55 and 69 years<sup>8</sup>. Therefore, STIs may have a long lasting negative impact on adults later in life. Investigations into infection-cancer associations have shown that hepatitis B and hepatitis C are associated with a much higher risk of developing liver cancer<sup>9</sup>, a common cancer among older adults with a peak rate between the ages of 85 to 89 years<sup>10</sup>. People living with HIV (PLWHIV) are also more susceptible to several types of cancers<sup>11</sup>. Besides cancer, STIs have also been found to be associated with diseases of the cardiovascular system<sup>12</sup>.

It is plausible that a greater number of lifetime sexual partners in older adults increases the risk of contracting an STI over the lifespan and subsequently increases the risk of developing health complications in later life. Given that STIs often go undiagnosed, number of sexual partners could provide a proxy measure of sexual risk behaviour that is more accurately reported (albeit likely subject to potential underreporting by people with a higher number of partners). Establishing the

1  
2  
3 extent to which number of sexual partners is associated with health problems is important in  
4  
5 gauging the potential utility of this measure as an indicator of risk.  
6  
7

8  
9 Previous studies that have investigated the number of sexual partners and cancer risk have shown  
10  
11 mixed findings. In a sample of black men, those reporting 25 or more sexual partners were found to  
12  
13 be 2.80 (95% CI 1.29, 6.09) times more likely to be diagnosed with cancer compared to men with 5  
14  
15 or fewer partners<sup>13</sup>. Other research has found similar findings in more diverse samples in relation to  
16  
17 anal cancer, prostate cancer, and oral cancer<sup>7 14 15</sup>. In contrast, a study using a Canadian population  
18  
19 found reduced risk of prostate cancer among men with more than 20 sexual partners in the  
20  
21 lifetime<sup>16</sup>. Little research has been carried out examining the role of number of sexual partners as a  
22  
23 behavioural risk factor for wider health outcomes in older adults.  
24  
25

26  
27 To fill these knowledge gaps, the aim of the present paper was to investigate the sex-specific  
28  
29 associations between the number of lifetime sexual partners and several health outcomes in a large  
30  
31 sample of older adults in England. We hypothesised that a greater number of previous sexual  
32  
33 partners would be associated with increased risk of unfavourable health outcomes.  
34  
35  
36  
37  
38  
39  
40

## 41 **Method**

### 42 **Study population**

43  
44  
45 We used cross-sectional data from the English Longitudinal Study of Ageing (ELSA), a population-  
46  
47 representative longitudinal panel study of men and women aged  $\geq 50$  years living in England<sup>17</sup>. The  
48  
49 initial ELSA sample was drawn from households with 1 or more member 50 years or older  
50  
51 responding to the Health Survey for England in 1998, 1999, and 2001. All household members 50  
52  
53 years or older plus partners who were younger than 50 years or had joined the household since the  
54  
55 HSE were invited for interview. Since ELSA began in 2002, data have been collected in biennial waves  
56  
57  
58  
59  
60

1  
2  
3 via computer assisted personal interview and self-completion questionnaires. The present study  
4  
5 uses data from Wave 6 (2012/13) as this is the only wave in which participants have been asked  
6  
7 about their number of sexual partners. The Sexual Relationships and Activities Questionnaire (SRA-  
8  
9 Q) was administered as a self-completion measure and was returned by 7,079 (67%) participants.  
10  
11 We restricted our sample to those who reported their lifetime number of sexual partners and had  
12  
13 complete data on all covariates, leaving a final sample for analysis of 5,722 men and women. All  
14  
15 participants gave fully informed consent to participate in the study, and ethical approval was  
16  
17 obtained from the London Multi-Centre Research Ethics Committee.  
18  
19  
20  
21

## 22 **Measures**

### 23 Exposure: lifetime number of sexual partners

24  
25  
26  
27  
28  
29 Number of sexual partners was assessed as part of the Sexual Relationships and Activities  
30  
31 Questionnaire (SRA-Q)<sup>18</sup>, which participants completed in private and returned in a sealed envelope.  
32  
33 The male and female versions of the SRA-Q are available online at [http://www.elsa-](http://www.elsa-project.ac.uk/study-documentation)  
34  
35 [project.ac.uk/study-documentation](http://www.elsa-project.ac.uk/study-documentation). Participants were asked to indicate the number of sexual  
36  
37 partners (vaginal/oral/anal sex) they had had in their lifetime (0, 1, 2-4, 5-9, 10-19, 20 or more). Due  
38  
39 to low numbers of participants reporting have had 0 or in excess of 20 partners, we combined these  
40  
41 with proximal categories, leaving four groups for analysis: 0-1, 2-4, 5-9 and  $\geq 10$  sexual partners.  
42  
43  
44

### 45 Health outcomes

46  
47  
48  
49 Self-rated health was assessed using a single item: *"Would you say your health is...*  
50  
51 *poor/fair/good/very good/excellent?"* We analysed the proportion of individuals rating their health  
52  
53 as fair/poor, as has been done in previous studies<sup>19 20</sup>. Limiting long-standing illness was self-  
54  
55 reported in response to two questions: (i) *"Do you have any long-standing illness, disability, or*  
56  
57 *infirmity? By long-standing I mean anything that has troubled you over a period of time or that is*  
58  
59 *likely to affect you over a period of time."* If yes, (ii) *"Does this illness or disability limit your activities*  
60



1  
2  
3 *in any way?"* Affirmation of a long-standing illness and the reporting of any form of limitation  
4  
5 classified the participant as having a limiting long-standing illness. Information about doctor-  
6  
7 diagnosed cancer, CHD and stroke (ever in their lifetime) was self-reported.  
8  
9

### 10 11 Covariates

12  
13  
14 All potential confounders were selected *a priori* based on previous literature reporting links between  
15  
16 these variables and our exposure and outcomes of interest. Demographic information collected  
17  
18 included age, ethnicity (white vs. non-white) and partnership status (married/cohabiting,  
19  
20 separated/divorced, widowed, or single/never married). Socio-economic status (SES) was based on  
21  
22 household non-pension wealth (which has been identified as particularly relevant to health  
23  
24 outcomes in this age group<sup>21</sup>, categorised into quintiles across all wave 6 ELSA participants. We also  
25  
26 included data on several health-related variables. Participants reported their current smoking status  
27  
28 (smoker or non-smoker) and frequency of alcohol intake, categorised as never/rarely (never – once  
29  
30 or twice a year), regularly (once every couple of months – twice a week), or frequently (3 days a  
31  
32 week – almost every day). Physical activity was assessed with three items that asked participants  
33  
34 how often they took part in vigorous, moderate and low-intensity activities (more than once a week,  
35  
36 once a week, 1-3 times a month, hardly ever/never)<sup>22</sup>, and further categorised into three groups, as  
37  
38 previously described<sup>23</sup>: inactive (no moderate/vigorous activity on a weekly basis); moderate activity  
39  
40 at least once a week; and vigorous activity at least once a week. Depressive symptoms were  
41  
42 assessed using the 8-item Centre of Epidemiological Studies Depression (CES-D) scale, a scale highly  
43  
44 validated for use in older adults<sup>24</sup>. These variables have been shown to be linked with number of  
45  
46 sexual partners and differences in perceived health and the diagnoses of interest here<sup>25-33</sup>.  
47  
48  
49  
50  
51  
52

### 53 **Patient and Public Involvement**

54  
55  
56  
57 There was no involvement of patients or the public in the design of any aspect of the present study.  
58  
59  
60

## Statistical analysis

Analyses were performed using IBM SPSS Statistics 22 on complete cases. Data were weighted to correct for sampling probabilities and for differential non-response and to calibrate back to the 2011 National Census population distributions for age and sex. The weights accounted for the differential probability of being included in wave 6 of ELSA and for non-response to the SRA-Q. Details can be found

at [https://www.ucl.ac.uk/drupal/site\\_iehc/sites/iehc/files/5050\\_elsa\\_w6\\_technical\\_report\\_v1.pdf](https://www.ucl.ac.uk/drupal/site_iehc/sites/iehc/files/5050_elsa_w6_technical_report_v1.pdf)<sup>34</sup>.

Associations between lifetime number of sexual partners and covariates were assessed using one-way analyses of variance (ANOVAs) for continuous variables and chi-square tests for categorical variables. We used logistic regression to analyse associations between lifetime number of sexual partners and fair/poor self-rated health, limiting long-standing illness, cancer, CHD and stroke. All models were adjusted for age, partnership status, ethnicity, wealth, smoking status, alcohol intake, physical activity, and depressive symptoms. Separate analyses were carried out on men and women. For each outcome, we report the odds ratio (OR) and 95% confidence interval (CI) for those who reported having had 2-4, 5-9 and  $\geq 10$  sexual partners in their lifetime, relative to those who had had 0-1 sexual partners. To check whether our categorisation of the number of sexual partners (grouping together the 0 and 1 responses and 10-19 and  $\geq 20$  responses) influenced the results, we ran sensitivity analyses with distinct groups for those reporting 0, 1, 10-19 and  $\geq 20$  sexual partners.

## Results

There were 2,537 men and 3,185 women in the sample. The mean age of participants was 64.25 (SD 9.75) years in men and 65.34 (SD 10.06) years in women. The majority were married or cohabiting (73.6% of men, 60.6% of women), white (93.7% of men, 96.0% of women) non-smokers (85.5% of men, 86.9% of women) who drank alcohol regularly or frequently (84.0% of men, 69.9% of women) and were moderately or vigorously active at least once a week (80.2% of men, 74.8% of women). Among men, 28.5% reported having had 0-1 sexual partners in their lifetime, 29.0% had had between 2 and 4 partners, 20.2% had had between 5 and 9 partners, and 22.2% had had 10 or more partners. Among women, the respective figures were 40.8% (0-1), 35.5% (2-4), 15.8% (5-9) and 7.8% ( $\geq 10$ ).

Sample characteristics in relation to number of sexual partners are summarised in Table 1. In both men and women, a higher number of sexual partners was associated with younger age, being unmarried, and being in either the highest or lowest quintile of wealth. Those with a higher number of sexual partners were also more likely to report smoking, frequent alcohol intake and engaging in vigorous physical activity on a weekly basis. There was an association between higher number of sexual partners and white ethnicity in women but not in men, and an association between higher number of sexual partners and a greater number of depressive symptoms in men but not in women.

1  
23 *Table 1 Associations between lifetime number of sexual partners and covariates in men and women*

	Men					Women				
	0-1 (n=785) <sup>1</sup>	2-4 (n=779)	5-9 (n=487)	≥10 (n=486)	<i>p</i>	0-1 (n=1285)	2-4 (n=1178)	5-9 (n=483)	≥10 (n=239)	<i>p</i>
Age (mean [SD] years)	68.54 (9.87)	65.53 (9.34)	62.21 (8.46)	60.71 (7.97)	<.001	69.39 (9.82)	64.36 (8.90)	60.52 (7.31)	59.88 (7.48)	<.001
Partner status										
Married/cohabiting	84.4	78.6	70.6	55.7	<.001	66.6	58.6	56.3	46.5	<.001
Separated/divorced	1.8	7.3	17.0	25.0	-	4.2	20.3	28.8	32.6	-
Widowed	8.3	8.2	3.8	3.0	-	24.2	16.7	8.0	5.7	-
Single/never married	5.4	6.0	8.6	16.3	-	5.0	4.4	6.9	15.2	-
Ethnicity										
White	93.5	93.2	93.0	95.3	.325	94.3	96.6	97.6	98.7	.001
Non-white	6.5	6.8	7.0	4.7	-	5.7	3.4	2.4	1.3	-
Wealth quintile										
1 (poorest)	12.8	13.0	16.7	29.5	<.001	17.8	20.6	20.6	26.1	.004
2	23.0	18.3	17.4	15.7	-	20.6	21.1	18.9	21.7	-
3	22.2	23.4	17.8	13.5	-	22.1	22.1	16.6	17.9	-
4	22.2	24.6	23.8	17.2	-	21.8	18.5	21.7	13.0	-
5 (richest)	19.8	20.6	24.4	24.0	-	17.8	17.8	22.2	21.3	-
Smoking status										
Non-smoker	90.5	90.4	81.7	76.2	<.001	92.1	85.9	82.4	73.0	<.001
Smoker	9.5	9.6	18.3	23.8	-	7.9	14.1	17.6	27.0	-
Alcohol intake <sup>1</sup>										
Never/rarely	20.5	16.7	12.5	12.3	<.001	37.1	28.6	17.4	25.1	<.001
Regularly	44.2	42.3	40.6	39.5	-	41.9	44.9	48.3	38.5	-
Frequently	35.3	40.9	47.0	48.2	-	21.0	26.5	34.3	36.4	-
Physical activity										

59  
60

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

Inactive	22.2	17.3	15.4	24.0	<.001		32.1	21.2	19.2	19.6	<.001
Moderately active at least once a week	48.2	43.9	44.0	35.9	-		46.7	51.2	44.6	50.4	-
Vigorously active at least once a week	29.6	38.7	40.7	40.1	-		21.3	27.6	36.2	30.0	-
Depressive symptoms (0-8) (mean [SD])	1.06 (1.76)	0.92 (1.62)	0.99 (1.72)	1.45 (2.05)	<.001		1.50 (1.91)	1.56 (2.04)	1.53 (1.99)	1.60 (2.05)	.796

Unweighted sample sizes.  
SD = standard deviation.  
Never/rarely = never – once or twice a year; regularly = once every couple of months – twice a week; frequently = 3 days a week – almost every day.  
Notes: Values are percentages unless otherwise stated. All figures are weighted for sampling probabilities and differential non-response.

Associations between lifetime number of sexual partners and health outcomes are presented in Table 2. Among women, there was a statistically significant association between number of sexual partners and risk of limiting long-standing illness. Relative to women who reported having had 0-1 sexual partners, the odds of reporting a limiting long-standing illness were 64% higher for those who had had between 5 and 9 sexual partners in their lifetime (OR=1.64, 95% confidence interval [CI] 1.15-2.05,  $p=0.003$ ), and 64% higher for those who had had 10 or more partners (OR=1.64, 95% CI 1.15-2.35,  $p=0.007$ ). There was also a statistically significant association between number of sexual partners and cancer risk in both women and men. In women, differences between those reporting 2-4 and 5-9 sexual partners and those reporting 0-1 sexual partners were not statistically significant ( $p>0.13$ ), but those who had had 10 or more sexual partners in their lifetime had 91% higher odds of reporting a diagnosis of cancer than those who had had 0-1 sexual partners (OR=1.91, 95% CI 1.04-3.51,  $p=0.038$ ). In men, odds of cancer were increased by 57% among those reporting 2-4 lifetime sexual partners (OR=1.57, 95% CI 1.02-2.42,  $p=0.039$ ) and by 69% among those reporting  $\geq 10$  sexual

1  
2  
3 partners (OR=1.69, 95% CI 1.01-2.83,  $p=0.047$ ), relative to those who had had 0-1 partners. The  
4  
5 difference between men reporting 5-7 sexual partners and those reporting 0-1 partners did not  
6  
7 reach statistical significance, although the effect size was in the same region as that for the group  
8  
9 reporting 2-4 sexual partners (OR=1.50, 95% CI 0.89-2.51,  $p=0.127$ ). Lifetime number of sexual  
10  
11 partners was not statistically significantly associated with self-rated health, coronary heart disease  
12  
13 or stroke in either sex, or with limiting long-standing illness in men.  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

Confidential: For Review Only

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46

Table 2 Associations between lifetime number of sexual partners and health outcomes in men and women

	n <sup>1</sup>	% (SE) <sup>2</sup>				2-4		5-9		≥10	
		0-1	2-4	5-9	≥10	OR [95% CI] <sup>3</sup>	p	OR [95% CI]	p	OR [95% CI]	p
<b>Men</b>											
Fair/poor self-rated health	2535	24.0 (1.3)	23.5 (1.2)	25.7 (1.5)	28.8 (1.5)	0.91 [0.69-1.20]	.481	1.00 [0.72-1.38]	.975	1.26 [0.91-1.74]	.173
Limiting long-standing illness	2537	29.7 (1.4)	28.1 (1.3)	33.4 (1.6)	32.5 (1.6)	0.84 [0.65-1.09]	.184	1.17 [0.87-1.57]	.289	1.09 [0.80-1.47]	.598
Cancer	2537	4.1 (0.8)	7.0 (0.8)	6.5 (1.0)	6.9 (1.0)	1.57 [1.02-2.42]	.039	1.50 [0.89-2.51]	.127	1.69 [1.01-2.83]	.047
Coronary heart disease	2537	11.0 (1.0)	10.8 (1.0)	10.2 (1.2)	9.9 (1.2)	1.03 [0.74-1.43]	.876	0.90 [0.60-1.37]	.634	0.89 [0.58-1.65]	.568
Stroke	2537	4.6 (0.7)	2.7 (0.6)	4.0 (0.8)	4.5 (0.8)	0.62 [0.36-1.07]	.084	0.86 [0.45-1.63]	.634	1.01 [0.54-1.92]	.967
<b>Women</b>											
Fair/poor self-rated health	3185	24.5 (1.1)	25.3 (1.1)	27.4 (1.7)	24.7 (2.4)	1.12 [0.89-1.42]	.344	1.26 [0.91-1.74]	.161	0.94 [0.62-1.44]	.790
Limiting long-standing illness	3184	32.9 (1.2)	34.9 (1.3)	39.9 (1.9)	41.8 (2.7)	1.17 [0.95-1.45]	.137	1.64 [1.15-2.05]	.003	1.64 [1.15-2.35]	.007
Cancer	3185	4.8 (0.7)	6.1 (0.7)	6.1 (1.0)	8.9 (1.5)	1.35 [0.91-1.99]	.135	1.21 [0.71-2.08]	.482	1.91 [1.04-3.51]	.038
Coronary heart disease	3185	6.2 (0.7)	7.5 (0.7)	7.1 (1.1)	5.0 (1.5)	1.28 [0.90-1.82]	.178	1.06 [0.59-1.91]	.854	0.38 [0.13-1.12]	.080
Stroke	3185	3.0 (0.5)	3.4 (0.5)	4.5 (0.8)	3.5 (1.1)	1.19 [0.73-1.95]	.483	1.52 [0.76-3.06]	.241	1.18 [0.44-3.16]	.740

<sup>1</sup>Unweighted sample sizes.

<sup>2</sup>Percentage (with standard error) reporting health problem.

<sup>3</sup>Adjusted odds ratios (OR) and 95% confidence intervals (CI) for reporting health problem relative to the group who had had 0-1 sexual partners.

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46

Notes: All percentages and odds ratios are adjusted for age, partnership status, ethnicity, wealth, smoking status, alcohol intake, physical activity and depressive symptoms, and weighted for sampling probabilities and differential nonresponse.

Confidential: For Review Only



1  
2  
3 Sensitivity analyses in which those reporting 0, 1, 10-19 and  $\geq 20$  lifetime sexual partners were  
4 analysed separately revealed no notable differences in the pattern of results (Supplementary Table  
5  
6  
7 1).

## 11 12 13 14 **Discussion**

### 15 16 17 **Summary of findings**

18  
19  
20  
21 These results provide some evidence that the number of lifetime sexual partners is associated with  
22 adverse health outcomes in a sample of older adults in England. In both men and women, a higher  
23 number of sexual partners was associated with increased risk of cancer. In women, there was also a  
24 statistically significant positive association between number of sexual partners and risk of limiting  
25 long-standing illness. We observed no statistically significant association between number of lifetime  
26 sexual partners and self-rated health, CHD or stroke in either sex, or with limiting long-standing  
27 illness in men.  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37

### 38 **Comparison with previous studies**

39  
40  
41 These findings provide some support for the aforementioned hypothesis that a greater number of  
42 previous sexual partners is likely to increase the risk of adverse health outcomes in older adults.  
43 Perhaps most importantly, the present findings show that a greater number of lifetime sexual  
44 partners is associated with increased risk of cancer in older adults. Our findings, using a more  
45 proximal outcome, are in line with a large body of literature that suggests that specific STIs may lead  
46 to several cancers, such as HPV and cervical, oral, penile and anal cancers<sup>4 5 6</sup>, hepatitis C and B and  
47 liver cancer<sup>9</sup> as well as gonorrhoea infection and prostate cancer<sup>7</sup>. This finding supports previous  
48 research<sup>13-15</sup> in suggesting that lifetime number of sexual partners can aid in the identification of  
49 those older adults who are at a higher potential risk of cancer. The limited number of specific cancer  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

1  
2  
3 cases precluded further analysis by cancer type. We speculate the heightened risk of cancer might  
4  
5 be driven by those types known to be associated with STIs. Further studies using large sample to  
6  
7 elucidate such associations is important in evaluating the potential utility of health practitioners  
8  
9 screening older adults for number of lifetime sexual partners when considering risk of specific  
10  
11 cancers.  
12

13  
14  
15 The finding that number of lifetime sexual partners is associated with limiting long-standing illness in  
16  
17 women and not men should be noted. This gender difference is interesting, but an explanation is  
18  
19 elusive, especially when men have a greater number of lifetime sexual partners than women, as  
20  
21 shown in this study, and women are more likely to seek medical screening for STIs<sup>35</sup> and are thus less  
22  
23 likely to experience negative long-term health complications. It is possible that the limiting long-  
24  
25 standing illness was attained early in life among these women, resulting in a lower number of sexual  
26  
27 partner during their lifetime. Further research is required to identify mechanisms that explain this  
28  
29 observed association and the divergent pattern between men and women.  
30  
31  
32

33  
34 While some previous studies have suggested that STIs are associated with diseases of the  
35  
36 cardiovascular system<sup>12</sup>, the present results provide no evidence that the number of lifetime sexual  
37  
38 partners is associated with risk of CHD or stroke. It may be that only specific STIs, such as HIV and  
39  
40 hepatitis C, are associated with cardiovascular health<sup>12</sup>. It is possible that these infections are  
41  
42 contracted less often than the multiple STIs that can lead to cancer and thus a precise measure of  
43  
44 these exposures is needed to observe a statistically significant association. Moreover, the biological  
45  
46 processes between STIs and cardiovascular disease may be weaker than those driving cancer.  
47  
48 Further research is required to ascertain the causal mechanisms driving such associations.  
49  
50  
51

### 52 53 **Implications and directions for further research**

54  
55  
56 Our findings indicate a potential utility of lifetime number of sexual partners as a behavioural factor  
57  
58 for cancer risk assessment. In our analysed sample, divergent lifestyle profiles have been observed in  
59  
60

1  
2  
3 relation to lifetime number of sexual partners. Those with a greater number of sexual partners were  
4  
5 more likely to smoke cigarettes and drink alcohol frequently; behaviours known to be associated  
6  
7 with cancer risk<sup>36 37</sup>. It is possible that the number of sexual partners one has had captures a  
8  
9 combination of likelihood of exposure to STIs and lifestyle profile.  
10

11  
12  
13 Further research is required to replicate and confirm our findings. We tested a number of models  
14  
15 and it is possible that the association between number of sexual partners and cancer was a chance  
16  
17 finding. If the same associations were observed, it would be interesting to explore the extent to  
18  
19 which the associations are moderated by key sociodemographic (e.g. socioeconomic position),  
20  
21 health-related (e.g. physical activity) and sex-related (e.g. sexual orientation) variables. One could  
22  
23 also explore the predictive value of lifetime number of sexual partners as a simple question (e.g.  
24  
25 please indicate the number of sexual partners (vaginal/oral/anal sex) you have had in your lifetime  
26  
27 [0, 1, 2-4, 5-9, 10-19, 20 or more]) embedded in routine clinical assessment for cancer risk.  
28  
29  
30

### 31 32 **Strengths and limitations**

33  
34  
35 Strengths of this study include the large sample and statistical adjustment for a number of important  
36  
37 covariates. Moreover, the older age of the sample corresponds to the time of life when our  
38  
39 outcomes of interest tend to become more prevalent. However, the findings from the present study  
40  
41 must be interpreted in light of its limitations. First, all data were self-reported, which introduces a  
42  
43 number of potential biases. For example, the measure of number of sexual partners may have been  
44  
45 subject to social desirability bias, although this item was asked in a paper-based questionnaire  
46  
47 returned by post rather than in the face-to-face interview, to minimise participant embarrassment  
48  
49 and encourage honest responses. Diagnoses of cancer, CHD and stroke may not have accurately  
50  
51 been recalled, although previous studies have shown high agreement between self-reported cancer  
52  
53 diagnoses and medical record validation in population-based samples<sup>38-40</sup>. There is also the  
54  
55 possibility that self-reports may lack accuracy in older participants with memory problems. Second,  
56  
57 the number of sexual partners was assessed on a categorical response scale which asked participants  
58  
59  
60

1  
2  
3 to select the range within which their number of sexual partners fell. This meant we did not have the  
4  
5 precise number of sexual partners for most participants (i.e. all those who had had more than one  
6  
7 sexual partner) and as such it was not possible to model this exposure as a continuous variable,  
8  
9 which would have provided more easily interpretable results regarding the increase in risk of cancer  
10  
11 associated with each additional sexual partner. Third, no data were available on partner gender, for  
12  
13 example men having sex with men, which may be linked with higher risk of contracting certain STIs.  
14  
15 Fourth, the small number of cancer diagnoses meant we lacked statistical power to analyse this  
16  
17 outcome broken down by cancer type. It is likely that the association between number of sexual  
18  
19 partners and cancer would be stronger for those known to be associated with STIs. Fifth, there was a  
20  
21 substantial amount of missing data due to non-response to the survey assessing sexual relationships,  
22  
23 so these findings cannot be presumed to generalise to the entire population of older adults in  
24  
25 England. Further research is required to replicate our analysis and examine whether the same  
26  
27 associations are observed in different populations. Moreover, if those participants who had had  
28  
29 fewer (or indeed, more) partners were more likely to live to the time of survey then associations  
30  
31 between number of sexual partners and health outcomes may have been underestimated. There  
32  
33 may be a potential survival bias in the present analyses since the reference group (0-1) was  
34  
35 statistically significantly older than people in the >10 group. In other words, it is possible that cancer  
36  
37 rates were even higher in the group with more sexually partners, but these people had already died.  
38  
39 Finally, the analyses were cross-sectional and as such, it is not possible to determine causality, or  
40  
41 even whether the number of sexual partners pre-dated any health diagnoses. It is possible that some  
42  
43 people may react to diagnosis of a chronic condition by 'living life to the full', resulting in a rise in  
44  
45 their number of lifetime sexual partners. Further research using a prospective design could provide  
46  
47 further insight.  
48  
49  
50  
51  
52  
53  
54  
55

## 56 **Conclusions**

57  
58  
59  
60

1  
2  
3 In this large sample of older adults in England, we found that a greater number of previous sexual  
4 partners was associated with increased odds of cancer in men and women, and increased odds of  
5 limiting long-standing illness in women only. Enquiring about the number of sexual partners a  
6 patient has had may be a simple and cost-effective complement to existing cancer screening  
7 programmes in identifying those at risk of certain cancers, although further work is required first in  
8 order to replicate our findings and establish whether a causal relationship exists.  
9  
10  
11  
12  
13  
14  
15  
16  
17

### 18 **Author Contribution Statement**

19  
20 IG, LS, and SEJ conceived the idea, carried out the analyses, interpreted the results, and drafted the  
21 manuscript. All authors provided critical revisions and approved the final manuscript before  
22 submission.  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60

## References

1. Korenromp EL, Wi T, Resch S, et al. Costing of National STI Program Implementation for the Global STI Control Strategy for the Health Sector, 2016-2021. *PLoS One* 2017;12(1):e0170773. doi: 10.1371/journal.pone.0170773 [published Online First: 2017/01/28]
2. Stergachis A, Scholes D, Heidrich FE, et al. Selective screening for Chlamydia trachomatis infection in a primary care population of women. *Am J Epidemiol* 1993;138(3):143-53. doi: 10.1093/oxfordjournals.aje.a116840 [published Online First: 1993/08/01]
3. Moscicki AB, Hills N, Shiboski S, et al. Risks for incident human papillomavirus infection and low-grade squamous intraepithelial lesion development in young females. *JAMA* 2001;285(23):2995-3002. doi: 10.1001/jama.285.23.2995 [published Online First: 2001/06/30]
4. de Martel C, Plummer M, Vignat J, et al. Worldwide burden of cancer attributable to HPV by site, country and HPV type. *Int J Cancer* 2017;141(4):664-70. doi: 10.1002/ijc.30716 [published Online First: 2017/04/04]
5. Arya M, Li R, Pegler K, et al. Long-term trends in incidence, survival and mortality of primary penile cancer in England. *Cancer Causes Control* 2013;24(12):2169-76. doi: 10.1007/s10552-013-0293-y [published Online First: 2013/10/09]
6. Wilkinson JR, Morris EJ, Downing A, et al. The rising incidence of anal cancer in England 1990-2010: a population-based study. *Colorectal Dis* 2014;16(7):O234-9. doi: 10.1111/codi.12553 [published Online First: 2014/01/15]
7. Sarma AV, McLaughlin JC, Wallner LP, et al. Sexual behavior, sexually transmitted diseases and prostatitis: the risk of prostate cancer in black men. *J Urol* 2006;176(3):1108-13. doi: 10.1016/j.juro.2006.04.075 [published Online First: 2006/08/08]
8. Hayes JH, Barry MJ. Screening for prostate cancer with the prostate-specific antigen test: a review of current evidence. *JAMA* 2014;311(11):1143-9. doi: 10.1001/jama.2014.2085 [published Online First: 2014/03/20]
9. Perz JF, Armstrong GL, Farrington LA, et al. The contributions of hepatitis B virus and hepatitis C virus infections to cirrhosis and primary liver cancer worldwide. *J Hepatol* 2006;45(4):529-38. doi: 10.1016/j.jhep.2006.05.013 [published Online First: 2006/08/02]
10. UK; CR. Liver cancer incidence statistics 2018 [Available from: <https://www.cancerresearchuk.org/health-professional/cancer-statistics/statistics-by-cancer-type/liver-cancer/incidence> accessed 31.08. 2018.
11. IeDEA AI-dCPWGf, EuroCoord Ci. Comparison of Kaposi Sarcoma Risk in Human Immunodeficiency Virus-Positive Adults Across 5 Continents: A Multiregional Multicohort Study. *Clin Infect Dis* 2017;65(8):1316-26. doi: 10.1093/cid/cix480 [published Online First: 2017/05/23]
12. Freiberg MS, Chang CC, Skanderson M, et al. The risk of incident coronary heart disease among veterans with and without HIV and hepatitis C. *Circ Cardiovasc Qual Outcomes* 2011;4(4):425-32. doi: 10.1161/CIRCOUTCOMES.110.957415 [published Online First: 2011/06/30]
13. Daling JR, Madeleine MM, Johnson LG, et al. Human papillomavirus, smoking, and sexual practices in the etiology of anal cancer. *Cancer* 2004;101(2):270-80. doi: 10.1002/cncr.20365 [published Online First: 2004/07/09]
14. Dennis LK, Dawson DV. Meta-analysis of measures of sexual activity and prostate cancer. *Epidemiology* 2002;13(1):72-9. doi: 10.1097/00001648-200201000-00012 [published Online First: 2002/01/24]

15. Schwartz SM, Daling JR, Doody DR, et al. Oral cancer risk in relation to sexual history and evidence of human papillomavirus infection. *J Natl Cancer Inst* 1998;90(21):1626-36. doi: 10.1093/jnci/90.21.1626 [published Online First: 1998/11/12]
16. Spence AR, Rousseau MC, Parent ME. Sexual partners, sexually transmitted infections, and prostate cancer risk. *Cancer Epidemiol* 2014;38(6):700-7. doi: 10.1016/j.canep.2014.09.005 [published Online First: 2014/10/04]
17. Steptoe A, Breeze E, Banks J, et al. Cohort profile: the English longitudinal study of ageing. *International journal of epidemiology* 2013;42(6):1640-8. doi: 10.1093/ije/dys168 [published Online First: 2012/11/13]
18. Lee DM, Nazroo J, O'Connor DB, et al. Sexual Health and Well-being Among Older Men and Women in England: Findings from the English Longitudinal Study of Ageing. *Arch Sex Behav* 2016;45(1):133-44. doi: 10.1007/s10508-014-0465-1 [published Online First: 2015/01/28]
19. DeSalvo KB, Bloser N, Reynolds K, et al. Mortality prediction with a single general self-rated health question. A meta-analysis. *J Gen Intern Med* 2006;21(3):267-75. doi: 10.1111/j.1525-1497.2005.00291.x [published Online First: 2005/12/13]
20. Steptoe A, Jackson SE. The Life Skills of Older Americans: Association with Economic, Psychological, Social, and Health Outcomes. *Sci Rep* 2018;8(1):9669. doi: 10.1038/s41598-018-27909-w [published Online First: 2018/07/07]
21. Banks J, Karlsen S, Oldfield Z. Socio-economic position. In: Marmot A, ed. *Health, Wealth and Lifestyle of the Older Population in England*. London: Institute for Fiscal Studies 2003.
22. Demakakos P, Hamer M, Stamatakis E, et al. Low-intensity physical activity is associated with reduced risk of incident type 2 diabetes in older adults: evidence from the English Longitudinal Study of Ageing. *Diabetologia* 2010;53(9):1877-85. doi: 10.1007/s00125-010-1785-x [published Online First: 2010/05/25]
23. Hamer M, Molloy GJ, de Oliveira C, et al. Leisure time physical activity, risk of depressive symptoms, and inflammatory mediators: the English Longitudinal Study of Ageing. *Psychoneuroendocrinology* 2009;34(7):1050-5. doi: 10.1016/j.psyneuen.2009.02.004 [published Online First: 2009/03/11]
24. Steffick DE. Documentation of affective functioning measures in the Health and Retirement Study. HRS Documentation Report., 2000.
25. Anand P, Kunnumakara AB, Sundaram C, et al. Cancer is a Preventable Disease that Requires Major Lifestyle Changes. *Pharmaceutical research* 2008;25(9):2097-116. doi: 10.1007/s11095-008-9661-9
26. Chiuve SE, Rexrode KM, Spiegelman D, et al. Primary prevention of stroke by healthy lifestyle. *Circulation* 2008;118(9):947-54. doi: 10.1161/CIRCULATIONAHA.108.781062 [published Online First: 2008/08/12]
27. Franks P, Gold MR, Fiscella K. Sociodemographics, self-rated health, and mortality in the US. *Soc Sci Med* 2003;56(12):2505-14. doi: 10.1016/s0277-9536(02)00281-2 [published Online First: 2003/05/14]
28. Jackson SE, Yang L, Veronese N, et al. Sociodemographic and behavioural correlates of lifetime number of sexual partners: findings from the English Longitudinal Study of Ageing. *BMJ Sexual & Reproductive Health* 2019;45(2):138-46. doi: 10.1136/bmjsexrh-2018-200230
29. Mackenbach JP, Stirbu I, Roskam A-JR, et al. Socioeconomic Inequalities in Health in 22 European Countries. *New England Journal of Medicine* 2008;358(23):2468-81. doi: 10.1056/NEJMsa0707519
30. Poole L, Steptoe A. Depressive symptoms predict incident chronic disease burden 10 years later: Findings from the English Longitudinal Study of Ageing (ELSA). *Journal of Psychosomatic Research* 2018;113:30-36. doi: <https://doi.org/10.1016/j.jpsychores.2018.07.009>
31. Robards J, Evandrou M, Falkingham J, et al. Marital status, health and mortality. *Maturitas* 2012;73(4):295-99. doi: <https://doi.org/10.1016/j.maturitas.2012.08.007>



- 1  
2  
3 32. Smith GD, Chaturvedi N, Harding S, et al. Ethnic inequalities in health: A review of UK  
4 epidemiological evidence. *Critical Public Health* 2000;10(4):375-408. doi:  
5 10.1080/09581590010005331  
6  
7 33. Van der Kooy K, van Hout H, Marwijk H, et al. Depression and the risk for cardiovascular diseases:  
8 systematic review and meta analysis. *International Journal of Geriatric Psychiatry*  
9 2007;22(7):613-26. doi: 10.1002/gps.1723  
10  
11 34. Bridges S, Hussey D, Blake M. The dynamics of aging: The 2012 English Longitudinal Study of  
12 Ageing (Wave 6) - Technical Report. UK: NatCen Social Research, 2015.  
13  
14 35. Trienekens SC, van den Broek IV, Donker GA, et al. Consultations for sexually transmitted  
15 infections in the general practice in the Netherlands: an opportunity to improve STI/HIV  
16 testing. *BMJ Open* 2013;3(12):e003687. doi: 10.1136/bmjopen-2013-003687 [published  
17 Online First: 2014/01/02]  
18  
19 36. Agudo A, Bonet C, Travier N, et al. Impact of cigarette smoking on cancer risk in the European  
20 prospective investigation into cancer and nutrition study. *J Clin Oncol* 2012;30(36):4550-7.  
21 doi: 10.1200/JCO.2011.41.0183 [published Online First: 2012/11/22]  
22  
23 37. Bagnardi V, Blangiardo M, La Vecchia C, et al. A meta-analysis of alcohol drinking and cancer risk.  
24 *Br J Cancer* 2001;85(11):1700-5. doi: 10.1054/bjoc.2001.2140 [published Online First:  
25 2001/12/18]  
26  
27 38. Bergmann MM, Byers T, Freedman DS, et al. Validity of self-reported diagnoses leading to  
28 hospitalization: a comparison of self-reports with hospital records in a prospective study of  
29 American adults. *Am J Epidemiol* 1998;147(10):969-77. doi:  
30 10.1093/oxfordjournals.aje.a009387 [published Online First: 1998/05/22]  
31  
32 39. Bush TL, Miller SR, Golden AL, et al. Self-report and medical record report agreement of selected  
33 medical conditions in the elderly. *Am J Public Health* 1989;79(11):1554-6. doi:  
34 10.2105/ajph.79.11.1554 [published Online First: 1989/11/01]  
35  
36 40. Simpson CF, Boyd CM, Carlson MC, et al. Agreement between self-report of disease diagnoses  
37 and medical record validation in disabled older women: factors that modify agreement. *J Am*  
38 *Geriatr Soc* 2004;52(1):123-7. doi: 10.1111/j.1532-5415.2004.52021.x [published Online  
39 First: 2003/12/23]  
40  
41  
42  
43  
44  
45  
46  
47  
48  
49  
50  
51  
52  
53  
54  
55  
56  
57  
58  
59  
60



Supplementary Table 1 Sensitivity analyses

	0		2-4		5-9		10-19		≥20	
	OR [95% CI] <sup>1</sup>	p	OR [95% CI]	p	OR [95% CI]	p	OR [95% CI]	p	OR [95% CI]	p
<b>Men</b>										
Fair/poor self-rated health	0.52 [0.25-1.07]	.077	0.86 [0.25-1.07]	.289	0.94 [0.67-1.31]	.714	1.36 [0.92-2.02]	.123	1.01 [0.67-1.53]	.970
Limiting long-standing illness	0.76 [0.39-1.48]	.414	0.82 [0.64-1.07]	.142	1.15 [0.85-1.55]	.369	1.03 [0.71-1.49]	.883	1.09 [0.75-1.59]	.664
Cancer	1.39 [0.46-4.18]	.562	1.62 [1.04-2.52]	.033	1.54 [0.91-2.61]	.110	1.83 [0.99-3.39]	.054	1.64 [0.85-3.17]	.139
Coronary heart disease	0.68 [0.31-1.52]	.347	0.99 [0.71-1.38]	.954	0.87 [0.57-1.33]	.524	0.58 [0.33-1.03]	.063	1.15 [0.69-1.91]	.587
Stroke	1.00 [0.30-3.33]	.995	0.62 [0.36-1.07]	.088	0.85 [0.45-1.63]	.634	0.92 [0.41-2.03]	.830	1.13 [0.51-2.53]	.760
<b>Women</b>										
Fair/poor self-rated health	1.76 [1.01-3.06]	.048	1.17 [0.92-1.49]	.201	1.32 [0.95-1.83]	.101	1.19 [0.71-1.99]	.503	0.78 [0.41-1.46]	.435
Limiting long-standing illness	1.33 [0.77-2.29]	.304	1.19 [0.96-1.48]	.104	1.57 [1.18-2.09]	.002	1.89 [1.23-2.92]	.004	1.41 [0.83-2.39]	.204
Cancer	1.81 [0.75-4.37]	.188	1.42 [0.95-2.12]	.088	1.28 [0.74-2.21]	.377	2.06 [0.99-4.27]	.052	2.03 [0.84-4.89]	.115
Coronary heart disease	1.31 [0.63-2.74]	.474	1.31 [0.91-1.89]	.142	1.09 [0.60-1.98]	.772	0.33 [0.08-1.43]	.137	0.50 [0.11-2.30]	.372
Stroke	2.52 [0.97-6.54]	.058	1.29 [0.78-2.14]	.317	1.65 [0.81-3.37]	.165	2.03 [0.71-5.78]	.185	0.36 [0.03-4.64]	.431
<sup>1</sup> Adjusted odds ratios (OR) and 95% confidence intervals (CI) for reporting health problem relative to the group who had had 1 sexual partner. Notes: All percentages and odds ratios are adjusted for age, partnership status, ethnicity, wealth, smoking status, alcohol intake, physical activity and depressive symptoms, and weighted for sampling probabilities and differential nonresponse.										

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42  
43  
44  
45  
46

Confidential: For Review Only