# Sugar-Sweetened Soda Consumption and Total and Breast Cancer Mortality: The Western New York Exposures and Breast Cancer (WEB) Study



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# ABSTRACT

**Background:** There is growing evidence of an association between sugar-sweetened beverages (SSB) and increased risk of mortality in various populations. However, SSB influence on mortality among patients with breast cancer is unknown.

**Methods:** We assessed the relationship between sugar-sweetened soda and both all-cause and breast cancer mortality among women with incident, invasive breast cancer from the Western New York Exposures and Breast Cancer Study. Breast cancer cases were followed for a median of 18.7 years, with ascertainment of vital status via the National Death Index. Frequency of sugar-sweetened soda consumption was determined via dietary recall using a food frequency questionnaire. Cox proportional hazards, adjusting for relevant variables, were used to estimate HRs and 95% confidence intervals (CI).

**Results:** Of the 927 breast cancer cases, 386 (54.7%) had died by the end of follow-up. Compared with never/rarely sugar-

# Introduction

Breast cancer is the most common incident cancer and the second leading cause of cancer mortality among women in the United States (1). With improved early detection and treatment, the number of breast cancer survivors has grown to more than 3.1 million currently in the United States alone (2). Given the ever-increasing number of patients with breast cancer, understanding the factors related to reduced mortality following a breast cancer diagnosis is critically important. Although dietary changes are potentially modifiable practices that can be implemented by patients with cancer, there is much that remains unknown regarding the relationship, if any, between dietary factors and cancer mortality (3).

One dietary component of potential importance is sugar-sweetened beverages (SSB), which groups together carbonated sugar-sweetened soda, sports or energy drinks, sweetened waters, teas, coffees, and all other beverages containing added caloric sweeteners such as highfructose corn syrup (HFCS), sucrose, glucose, honey, molasses, and even fruit-juice concentrate (4). According to the Global Burden of Disease 2017 Risk Collaborators, worldwide consumption of SSB has sweetened soda drinkers, consumption at  $\geq 5$  times per week was associated with increased risk of both total (HR = 1.62; 95% CI, 1.16–2.26;  $P_{\text{trend}} < 0.01$ ) and breast cancer mortality (HR = 1.85; 95% CI, 1.16–2.94;  $P_{\text{trend}} < 0.01$ ). Risk of mortality was similarly increased among ER-positive, but not ER-negative patients; among women with body mass index above the median, but not below the median; and among premenopausal, but not postmenopausal women for total mortality only.

**Conclusions:** Reported higher frequency of sugar-sweetened soda intake was associated with increased risks of both total and breast cancer mortality among patients with breast cancer.

**Impact:** These results support existing guidelines on reducing consumption of SSB, including for women with a diagnosis of breast cancer.

increased drastically in the past few decades (5). Although the WHO recommends limiting sugar consumption to  $\leq 10\%$  of total energy intake, consumption in the U.S. population generally exceeds that guideline (6). Among the many types of SSB available, sugar-sweetened soda is among the top sources of added sugars and calories in the diet of Americans, but contributes little else nutritionally (7). In fact, in the U.S. population, added sugars contribute about 11% to 17% of daily energy intake, with sugar-sweetened soda contributing about a third of the total added sugar intake (7, 8).

There is considerable evidence of an association of a range of health problems associated with excessive intake of SSB or high sugar diets; these include cardiometabolic diseases, including weight gain, type 2 diabetes, and cardiovascular diseases (CVD; refs. 9, 10). In addition, there is mixed evidence of their association with cancer (11–13), including for breast cancer (14, 15); with some studies indicating increased risk of incident cancer (12, 14), whereas others indicate no or inverse associations (13, 15).

With regard to mortality, SSB consumption has also been identified as one of the primary contributors to increases in total attributable deaths and disability adjusted life years (DALY) globally between 1990 and 2017 (5). Recently, some studies have suggested positive associations between higher intakes of SSB and total mortality in groups suffering from cancers (16, 17) and among general disease-free populations (13, 18-23), but others have found no significant associations (24, 25). There is less evidence regarding all-cancer mortality and cancer-specific mortality in relation to SSB (10, 13, 19, 21, 23, 25). To our knowledge, the Nurses' Health Study (NHS) is the only study that reports on SSB and breast cancer mortality. In the NHS, there was a trend toward increased breast cancer mortality associated with increased SSB intake (HR = 1.09; 95% CI, 1.00–1.18;  $P_{\text{trend}} = 0.02$ ; ref. 20). However, there is no study on this association specifically within a cohort of patients with breast cancer.

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Given the suggestive evidence that SSBs consumption may increase the risk of breast cancer incidence (26–28) and mortality (20), understanding their possible impact on breast cancer prognosis is potentially significant. Sugar-sweetened soda is the most heavily-consumed SSB type in all age groups of the U.S. population and functions as a good proxy for high intakes of added sugars (29); we therefore report here on the association between reported sugar-sweetened soda intake with total and breast cancer mortality in a population-based study of women diagnosed with incident, primary, invasive breast cancer in Western New York.

# **Materials and Methods**

# **Study population**

The Western New York Exposures and Breast Cancer (WEB) Study was a population-based case-control study in Erie and Niagara counties, conducted between 1996 and 2001. For this project, we only used data regarding the breast cancer cases (N = 1,170). A more detailed description of the study design and methods is available elsewhere (30-32). WEB Study breast cancer cases were women ages 35 to 79 years at the time of interview, diagnosed with primary, histologically-confirmed, incident, invasive breast cancer, identified by nurse case finders in all the major hospitals in the study region. Of the eligible cases, 72% agreed to participate. Our analyses are limited to women with complete information on sugar-sweetened soda consumption (n = 28 excluded) and relevant covariates. In addition, to minimize confounding, those who had diabetes at baseline enrolment were excluded (n = 76) because their consumption of sugar-sweetened soft drinks was different from those without diabetes (P < 0.05). Breast cancers diagnosed at stage 0 non-invasive (n = 144) were also excluded. This led to a final sample size of 927 breast cancer cases for our analyses. Written informed consent was obtained from all participants; the study was approved by the institutional review boards of the State University of New York, University at Buffalo, and participating hospitals.

## **Data collection**

# Assessment of consumption of sugar-sweetened soda

We define sugar-sweetened soda as regular carbonated SSB, not including fruit juices, sports drinks, coffees, teas, or flavoured waters. A modified version of the Health Habits and History food frequency questionnaire (FFQ) was used to assess usual dietary intake in the 12 to 24 months prior to diagnosis of breast cancer (33). Participants were asked to recall how often, on average, they consumed sugar-sweetened soda (i.e., not diet soda), using a nine-level categorical frequency scale ranging from (i) rarely/never, (ii) once a month, (iii) two to three times per month, (iv) once a week, (v) twice a week, (vi) three to four times per week, (vii) five to six times per week, (viii) once a day to (ix)  $\geq$ 2 times per day. We collapsed these nine categories into four groups as follows: (i) rarely/never, (ii) ≤ one time per week, (iii) two to four times per week, and (iv)  $\geq$  five times per week. Although, the FFQ queried about serving sizes of sugar-sweetened soda during intake, we do not account for this information in our analyses since more than 40% of the participants were missing portion size information.

#### Assessment of survival

Vital status of breast cancer cases participating in the WEB study was determined from the National Death Index (NDI) through December 31, 2018. We used the International Classification of Diseases, Tenth revision (ICD-10), to systematically identify cause of death. Total mortality was defined as death from any cause, whereas breast cancer mortality was attributed if breast cancer was specifically identified as the underlying cause of death. Participants' contribution to person-time survival was calculated as the number of months of follow up from date of diagnosis until date of death or until the end of follow-up on December 31, 2018, whichever occurred first.

#### Covariate measurement

Recalls from the FFQ were also used to estimate energy intake (daily kilocalories), fruit intake (daily grams), vegetable intake (daily grams), dietary fiber (daily grams), alcohol consumption over the previous 12 to 24 months (ever/never), and other nutrient intakes. Data were also collected by trained interviewers during in-person computerassisted interviews at baseline on demographics (age at diagnosis, race/ ethnicity, education), smoking history (lifetime pack years), reproductive history (age at first birth, age at menarche, menopausal status, age at menopause, parity, ever hormone replacement therapy), and physical activity (metabolic hours per week) obtained from all forms of recreational, work- and chore-related activities performed by participants in the week prior to the interview. All participants who reported a prior physician-diagnosed history of angina, stroke, heart attack (myocardial infarction, transient ischemic heart attack), atrial fibrillation, rheumatoid heart disease, and aortic aneurism were classified as having CVD. History of physician-diagnosed high blood pressure, high blood cholesterol, and high blood glucose were similarly reported by participants. Body mass index (BMI) at baseline was calculated as weight (kg) divided by the square of height  $(m^2)$ . Height and weight were measured by trained interviewers using a standardized protocol (34). Clinical characteristics of tumors from women with breast cancer [cancer stage at diagnosis, estrogen-receptor (ER) status, progesterone-receptor (PR) status] were obtained from baseline medical charts reviewed by trained nurses. During the interview, the women were also asked about the planned or prior treatments for their incident breast cancer including queries regarding surgery, radiation, and chemotherapy.

#### **Statistical analyses**

Descriptive statistics for demographic, personal, and reproductive characteristics of those who died and those who remained alive until the end of the follow-up period were compared. We used T tests for the mean and SD of continuous and normally distributed variables. For non-normal continuous variables, we present the median and interquartile range and made comparisons using the Wilcoxon rank-sum test. Frequencies and percentages are reported for categorical variables and we used the Chi-square test for group comparisons.

Hazard ratios (HR) and the respective 95% confidence intervals (CI) for total and breast cancer mortality were estimated with Cox proportional hazards regression models. We included covariates in the model that altered the unadjusted HRs by 10% or more, as well as other covariates based on a priori knowledge of risk factors for breast cancer survival. The covariates examined included age, race, education, smoking, physical activity, BMI, age at first birth, parity, age at menopause, age at menarche, menopausal status, energy intake, dietary fiber intake, fruit intake, vegetable intake, sodium intake, fat intake, vitamin supplement intake, alcohol consumption, stage at diagnosis, ER status, PR status, radiotherapy, chemotherapy, hormone replacement therapy, high blood pressure, high blood cholesterol, high blood glucose and baseline CVD diagnosis. Our final models were adjusted for age (years), race (white/black), education (years), smoking (life pack years), physical activity (metabolic hours per week), cancer stage (stage I (the reference group), stage II, stages III/IV, stage unknown), BMI (kg/m<sup>2</sup>), alcohol intake (ever/never), energy intake

(Kcal/day), ER status [ER positive (the reference group), ER negative, ER unknown], baseline CVD diagnosis (no/yes), high blood pressure (no/yes), high blood cholesterol (no/yes), vegetable intake (g/day), fruit intake (g/day), and dietary fiber (g/day). The proportional hazards assumption was tested for all models and was found to hold in all analyses. We also assessed whether there was a trend in risk for total and breast cancer mortality across frequencies of sugarsweetened soda consumption using the Wald Chi-square test for trend.

We examined associations stratified by menopausal status (postmenopausal and premenopausal), ER status (ER positive and ER negative) and BMI (< median BMI and ≥ median BMI). Product terms were used to test for the interaction between sugar-sweetened soda intake and menopausal status, ER status, and BMI. In addition, because the probability that terminal cancer and its treatments may affect diet and that diet would be unlikely to affect the survival outcome at later stages, we carried out sensitivity analyses by excluding women who died within 1 year of diagnosis (n = 6). Results remained unchanged and we present the HR from the complete sample. We also performed an analysis excluding participants with reported daily energy intake of less than 500 calories or more than 5,000 calories (N = 2) and those who had missing energy intake information (n = 18). The point estimates were not affected and we present results with the complete sample and after imputing the missing energy intake information. All analyses were conducted using SAS for Windows version 9.4.

### Results

Descriptive characteristics of breast cancer cases from the WEB study included in our analyses are shown in Table 1. Total followup time from the date of diagnosis ranged from 8.9 to 262 months, with a median survival time of 224 months. Of the included 927 women diagnosed with primary, incident, invasive breast cancer, 386 (41.6%) had died by the end of the follow-up period at December 31, 2018. Breast cancer deaths accounted for 39.6% of all deaths (n = 153). Compared with women who survived to the end of the follow-up period, those who died were on average, older, more likely to be postmenopausal, smoked more, were at higher cancer stage at diagnosis, had higher BMI, consumed less fruits and vegetables, were less physically active, were less likely to be alcohol drinkers and were more likely to have had cardiovascular disease, high blood pressure, and high blood cholesterol. Mortality did not differ by age at menarche, age at menopause, hormone therapy use, tumor ER or PR status, age at first childbirth, race, family history of breast cancer, energy intake, dietary fiber intake, or fat intake. Among the participants who died, there was a somewhat higher percentage of women who reported high frequency of sugarsweetened soda consumption compared with the women who remained alive. The frequency of diet soda consumption also differed by survival status (P = 0.03) and by sugar-sweetened soda consumption frequency (P < 0.01).

Compared with those who rarely or never drank sugar-sweetened soda, reported consumption at  $\geq$ 5 times per week was associated with increased breast cancer mortality (HR = 1.85; 95% CI, 1.16–2.94;  $P_{\rm trend} < 0.01$ ) and total mortality (HR = 1.62; 95% CI, 1.16–2.26;  $P_{\rm trend} < 0.01$ ; **Table 2**). As part of our sensitivity analyses, we ran additional models that included treatment for the breast cancer such as chemotherapy and radiation therapy. The results were similar (total mortality: HR = 1.49; 95% CI, 1.16–2.08;  $P_{\rm trend} = 0.02$ ; breast cancer mortality: HR = 1.78; 95% CI, 1.11–2.85;  $P_{\rm trend} < 0.01$ ). Similarly, including diet soda consumption as a covariate in the models did not

change our measures of associations (total mortality: HR = 1.46; 95% CI, 1.05–2.04;  $P_{trend} = 0.02$ ; breast cancer mortality: HR = 1.65; 95% CI, 1.01–2.68;  $P_{trend} = 0.03$ ).

Tests for interactions were not significant for BMI (P = 0.65), menopausal status (P = 0.13), or ER status (P = 0.89). In analyses stratified by menopausal status (Table 3), associations were stronger among the pre- than the postmenopausal women for breast cancer mortality but not total mortality. In analyses stratified by ER status, women with ER-positive breast cancer had higher risk of both total (HR = 1.65; 95% CI, 1.10–2.50;  $P_{\text{trend}} = 0.02$ ) and breast cancer mortality (HR = 2.41; 95% CI, 1.31–4.46;  $P_{\text{trend}} < 0.01$ ) with increasing frequency of sugar-sweetened soda consumption, whereas among those who were ER-negative, associations for total mortality and breast cancer mortality included the null. In BMI-stratified analyses, among women with BMI above the median, higher sugar-sweetened soda intake was associated with increased risks of total mortality (HR = 1.87; 95% CI, 1.21–2.89;  $P_{\text{trend}} < 0.01$ ) and breast cancer mortality (HR = 2.02; 95% CI, 1.05–3.90;  $P_{\text{trend}} = 0.03$ ). The associations were not significant for women with BMI below the median.

### Discussion

In this study of women with primary, incident, histologicallyconfirmed, invasive breast cancer, we found a dose-response trend of increasing risk for both total mortality and breast cancer mortality with higher consumption frequency of sugar-sweetened, non-diet soda  $(P_{\text{trend}} < 0.01 \text{ for both})$ , after adjusting for age, race, education, smoking, physical activity, cancer stage, BMI, menopausal status, alcohol intake, energy intake, ER status, baseline CVD, high blood pressure, high blood cholesterol, vegetable intake, fruit intake, dietary fiber. With higher frequency of sugar-sweetened soda consumption, there was increased risk of total and breast cancer mortality among women with ER positive ( $P_{\text{trend}} < 0.01$ ), but not ER-negative tumors ( $P_{\rm trend} > 0.05),$  with BMI above the median ( $P_{\rm trend} < 0.05),$  but not below the median BMI ( $P_{\rm trend} > 0.05$ ) and for premenopausal women  $(P_{\text{trend}} < 0.05)$ ; for postmenopausal women, there was increased risk for breast cancer mortality ( $P_{\text{trend}} = 0.03$ ), but not total mortality  $(P_{\rm trend} = 0.07).$ 

To our knowledge, this is the first study specifically among patients with breast cancer regarding the association between sugar-sweetened soda and mortality. Among patients with upper aerodigestive tract cancer (16) and colon cancer (17), SSB consumption has been found to be positively associated with increased risk of all-cause mortality. Among cohorts initially free from any cancer, such as European Prospective Investigation into Cancer and Nutrition (EPIC; ref. 23), Health Professionals Follow-up Study (HPFS; ref. 20), NHS (20), Vitamins and Lifestyle Study (VLS; ref. 19), and the Singapore Chinese Health Study (SCHS; ref. 25), the association of SSB with all-cause or cancer-specific mortality have been less consistent. For instance, VLS, HPFS, and NHS all reported significantly higher risk of all-cause and cancer mortality, whereas EPIC reported higher risk for all-cause mortality, but not for cancer mortality and SCHS reported higher risk for cancer mortality, but not for all-cause mortality. Of these five studies, only NHS reported on breast cancer-specific mortality. They found increased risk of breast cancer death with increasing SSB intake. Findings likely differ primarily because of the consumption patterns in these different populations or based on the definition of SSB in each study and the SSB parameterization used in the statistical analyses.

Sugar-sweetened soda, as part of the SSB group, are among the leading sources of added sugars in the American diet, irrespective of age (7, 8, 29, 35). High intake of sugar-sweetened soda, in the quantities

Table 1. Sugar-sweetened soda and breast	cancer prognosis - WEB st	udy: descriptive characteristic	s by mortality status ( $N = 927$ ).

Covariates	Total	Alive	Dead	Pa
N (%)	927 (100)	541 (58.4)	386 (41.6)	
White ( <i>n</i> , %)	856 (92.3)	503 (93.0)	353 (91.5)	0.39
Age (years) <sup>b,c</sup>	57.7 (11.3)	54.5 (9.6)	62.4 (11.9)	<0.01
Education (years) <sup>b,c</sup>	13.5 (2.6)	13.9 (2.5)	13.0 (2.6)	<0.01
Lifetime smoking (life pack years) <sup>d,e</sup>	0.1 (18.3)	0.2 (14.2)	2.1 (23.5)	<0.01
Physical activity (METh/week) <sup>d</sup>	235.5 (14.8)	236.8 (16.0)	234.5 (14.3)	<0.01
BMI (kg/m <sup>2</sup> ) <sup>b,c</sup>	28.4 (6.4)	27.8 (6.2)	29.3 (6.5)	<0.01
Age at first childbirth (n, %)				0.97
Nulliparous	160 (17.3)	91 (16.8)	69 (17.9)	
<24 years old	272 (29.3)	161 (29.8)	111 (28.8)	
24 to 27 years old	266 (28.7)	155 (28.7)	111 (28.8)	
>27 years old	229 (24.7)	134 (24.8)	95 (24.6)	
Post-menopause $(n, \%)^{c}$	653 (70.4)	342 (63.2)	311 (80.6)	<0.01
Energy intake (kcal/d) <sup>b,f</sup>	1,446 (683)	1,532 (571.5)	1,564.4 (697.3)	0.45
Dietary fiber (g/day) <sup>b,f</sup>	11.4 (6.8)	12.3 (5.6)	11.7 (5.5)	0.12
Vegetable intake (g/day) <sup>d,f</sup>	56.8 (57.5)	59.2 (56.7)	52.7 (57.0)	0.04
Fruit intake (g/day) <sup>d,f</sup>	94 (148.8)	100.1 (157.4)	88.2 (114.0)	0.02
Saturated fat (g/day) <sup>d,f</sup>	20.7 (14.5)	20.7 (13.7)	20.6 (14.9)	0.26
Polyunsaturated fatty acids (g/day) <sup>d,f</sup>	8.9 (7.1)	8.8 (7.1)	9.2 (7.1)	0.18
Ever alcohol consumption $(n, \%)^{f}$	470 (50.7)	297 (54.9)	173 (44.8)	< 0.01
Stage at diagnosis $(n, \%)^{c}$				<0.01
Stage I	444 (47.9)	295 (54.5)	149 (38.6)	
Stage II	284 (30.6)	155 (28.7)	129 (33.4)	
Stage III/IV	56 (6.0)	14 (2.6)	42 (10.9)	
Unknown	143 (15.4)	77 (14.2)	66 (17.1)	
ER status (n, %)				0.69
Negative	259 (27.9)	153 (28.3)	106 (27.5)	
Positive	615 (66.3)	360 (66.5)	255 (66.0)	
Unknown	53 (5.7)	28 (5.2)	25 (6.5)	
PR status (n, %)				0.28
Negative	326 (37.8)	183 (36.3)	143 (39.9)	
Positive	536 (62.2)	321 (63.7)	215 (60.1)	
Unknown	65 (7.0)	37 (7.0)	28 (7)	
Radiation therapy $(n, \%)$	488 (53.3)	305 (57.1)	183 (48.0)	<0.01
Chemotherapy (n, %)	400 (44.2)	236 (44.3)	164 (44.0)	0.93
Hormone replacement therapy $(n, \%)^{c}$	381 (41.1)	221 (40.9)	160 (41.5)	0.85
Family history of breast cancer $(n, \%)^c$	181 (19.5)	113 (20.9)	68 (17.6)	0.45
Prior benign breast disease $(n, \%)^c$	325 (35.6)	205 (38.6)	120 (31.4)	0.03
High blood pressure $(n, \%)^{c}$	301 (32.5)	138 (25.5)	163 (42.2)	<0.01
High blood cholesterol $(n, \%)^{c}$	314 (33.9)	164 (30.3)	150 (38.9)	<0.01
High blood glucose $(n, \%)^c$	21 (2.3)	14 (2.6)	7 (1.8)	0.44
Baseline CVD $(n, \%)^{c}$	212 (22.9)	100 (18.5)	112 (29.0)	<0.01
Sugar sweetened soda (n, %) <sup>f</sup>	212 (22.3)	100 (10.5)	112 (2010)	0.07
Never/rarely	480 (51.8)	290 (53.6)	190 (48.2)	0.07
≤ once per week	201 (21.7)	123 (22.7)	78 (20.2)	
2-4 times per week	133 (14.4)	74 (13.7)	59 (15.3)	
≥5 times per week	113 (12.2)	54 (10.0)	59 (15.3)	
Underlying causes of death $(n, \%)$	110 (12.2)	J- (10.0)	55 (15.5)	
All cancer mortality			211 (54.7)	
Breast cancer mortality			153 (39.6)	
CVD mortality			76 (19.7)	
Other causes			99 (25.7)	
Other Causes			JJ (ZJ./)	

<sup>a</sup>P value displayed compares patients with breast cancer who were alive and those who died; comparison of means by t test, of medians by Kruskal-Wallis test, and of frequencies by chi-square test. <sup>b</sup>Mean (SD).

<sup>c</sup>At baseline interview at the time of breast cancer diagnosis.

<sup>d</sup>Median (interquartile range).

<sup>e</sup>Lifetime considered from any smoking start age.

<sup>f</sup>Measured as dietary recall via FFQ in the 12 to 24 months prior to breast cancer diagnosis.

Table 2. Association between s	ugar-sweetened so	da and mortal	ity amon	g WEB study b	reast cance	er cases (N =	= 927).

Frequency of consumption	Rarely/never (n = 480)	≤1 time per week ( <i>n</i> = 201)	2–4 times per week ( <i>n</i> = 133)	≥5 times per week ( <i>n</i> = 113)	<b>P</b> <sub>trend</sub>
Total mortality ( $N = 386$ )	190	78	59	59	
Age-adjusted	Ref.	1.04 (0.80-1.35)	1.41 (1.05–1.89)	2.01 (1.49-2.70)	<0.01
Multivariable-adjusted <sup>a</sup>	Ref.	1.00 (0.76-1.31)	1.26 (0.92-1.72)	1.62 (1.16-2.26)	<0.01
Breast cancer mortality ( $N = 153$ )	63	27	29	34	
Age-adjusted	Ref.	1.02 (0.65-1.61)	1.71 (1.10-2.66)	2.38 (1.56-3.64)	<0.01
Multivariable-adjusted <sup>a</sup>	Ref.	0.93 (0.58-1.47)	1.49 (0.93–2.39)	1.85 (1.16–2.94)	<0.01

<sup>a</sup>Multivariable-adjusted: age (years), race (white/black), education (years), smoking (life pack years), physical activity (metabolic hours/week), cancer stage [stage I (the reference group), stage II, stage III/IV, stage unknown], BMI (kg/m<sup>2</sup>), menopausal status (premenopausal/postmenopausal), alcohol intake (ever/never), energy intake (Kcal/day), ER status [ER positive (the reference group), ER negative, ER unknown], baseline CVD (no/yes), high blood pressure (no/yes), high blood cholesterol (no/yes), vegetable intake (g/day), fruit intake (g/day), dietary fiber (g/day).

Table 3. Association between sugar-sweetened soda and mortality stratified by menopausal status, ER status, and BMI.

Frequency of consumption	Rarely/never (n = 480)	≤1 time per week ( <i>n</i> = 201)	2-4 times per week ( <i>n</i> = 133)	≥5 times per week ( <i>n</i> = 113)	<b>P</b> trend
Premenopause $(N = 274)^{a}$					
Total deaths ( $N = 75$ )	21	20	16	18	
Total mortality (HR, 95% CI)	Ref	1.76 (0.94-3.31)	1.96 (0.96-4.00)	3.21 (1.53-6.73)	<0.01
Breast cancer deaths ( $N = 56$ )	18	13	12	13	
Breast cancer mortality (HR, 95% CI)	Ref	1.33 (0.63-2.79)	1.55 (0.68-3.50)	2.64 (1.14-6.15)	0.02
Postmenopause $(N = 653)^{a}$					
Total deaths ( $N = 311$ )	169	58	43	41	
Total mortality (HR, 95% CI)	Ref	0.83 (0.61-1.14)	1.08 (0.76-1.55)	1.42 (0.97-2.08)	0.07
Breast cancer deaths ( $N = 97$ )	45	14	17	21	
Breast cancer mortality (HR, 95% CI)	Ref	0.71 (0.38-1.32)	1.48 (0.81-2.69)	1.81 (1.02-3.23)	0.03
ER-positive $(N = 615)^{b}$					
Total deaths ( $N = 255$ )	137	48	32	38	
Total mortality (HR, 95% CI)	ref	0.83 (0.59-1.16)	1.16 (0.77-1.75)	1.65 (1.10-2.50)	0.02
Breast cancer deaths ( $N = 91$ )	41	15	15	20	
Breast cancer mortality (HR, 95% CI)	Ref	0.90 (0.49-1.66)	1.51 (0.80-2.85)	2.41 (1.31-4.46)	<0.01
ER-negative $(N = 259)^{b}$					
Total deaths ( $N = 106$ )	43	26	22	15	
Total mortality (HR, 95% CI)	Ref	1.41 (0.84-2.35)	1.35 (0.75-2.46)	0.94 (0.46-1.89)	0.95
Breast cancer deaths ( $N = 57$ )	21	12	13	11	
Breast cancer mortality (HR, 95% CI)	Ref	1.01 (0.47-2.15)	1.50 (0.69-3.24)	1.14 (0.50-2.64)	0.65
BMI < Median $(N = 463)^{c}$					
Total deaths ( $N = 167$ )	81	40	24	22	
Total mortality (HR, 95% CI)	Ref	0.92 (0.62-1.36)	0.77 (0.47-1.27)	1.27 (0.75-2.15)	0.52
Breast cancer deaths ( $N = 74$ )	30	16	13	15	
Breast cancer mortality (HR, 95% CI)	Ref	0.90 (0.48-1.68)	1.04 (0.51-2.14)	1.75 (0.84-3.64)	0.12
$BMI \ge Median (N = 464)^c$					
Total deaths ( $N = 219$ )	109	38	35	37	
Total mortality (HR, 95% CI)	Ref	1.04 (0.71-1.52)	1.70 (1.13-2.55)	1.87 (1.21-2.89)	<0.01
Breast cancer deaths ( $N = 79$ )	33	11	16	19	
Breast cancer mortality (HR, 95% CI)	Ref	0.95 (0.47-1.90)	1.91 (1.00-3.66)	2.02 (1.05-3.90)	0.03

<sup>a</sup>Interaction between sugar-sweetened soda consumption and menopausal status prior to stratification: P = 0.13.

All models were adjusted for age (years), race (white/black), education (years), smoking (life pack years), physical activity (metabolic hours/week), cancer stage [stage I (the reference group), stage II, stage III/IV, stage unknown], BMI (kg/m<sup>2</sup>), alcohol intake (ever/never), energy intake (Kcal/day), ER status [ER positive (the reference group), ER negative, ER unknown], baseline CVD (no/yes), high blood pressure (no/yes), high blood cholesterol (no/yes), vegetable intake (g/day), fruit intake (g/day), dietary fiber (g/day).

<sup>b</sup>Interaction between sugar-sweetened soda consumption and ER status prior to stratification: P = 0.89.

All models were adjusted for age (years), race (white/black), education (years), smoking (life pack years), physical activity (metabolic hours/week), cancer stage [stage I (the reference group), stage II, stage III/IV, stage unknown], BMI (kg/m<sup>2</sup>), menopausal status (premenopausal/postmenopausal), alcohol intake (ever/ never), energy intake (Kcal/day), baseline CVD (no/yes), high blood pressure (no/yes), high blood cholesterol (no/yes), vegetable intake (g/day), fruit intake (g/ day), dietary fiber (g/day).

<sup>c</sup>Interaction between sugar-sweetened soda consumption and BMI prior to stratification: P = 0.65.

All models were adjusted for age (years), race (white/black), education (years), smoking (life pack years), physical activity (metabolic hours/week), cancer stage [stage I (the reference group), stage II, stage III/IV, stage unknown], BMI (kg/m<sup>2</sup>), menopausal status (premenopausal/postmenopausal), alcohol intake (ever/ never), energy intake (kcal/day), ER status [ER positive (the reference group), ER negative, ER unknown], baseline CVD (no/yes), high blood pressure (no/yes), high blood cholesterol (no/yes), vegetable intake (g/day), fruit intake (g/day), dietary fiber (g/day).

consumed by many in the United States, has been associated with weight gain and high adiposity (36–38), both of which are well-established risk factors for cancers, particularly postmenopausal breast cancer (39, 40) and for reduced survival after breast cancer diagnosis (41–43). BMI was associated with both sugar-sweetened soda consumption frequency (P = 0.04) and with mortality status (P < 0.01) in our WEB population. In analyses stratified by BMI, those with BMI  $\geq$  median ( $\sim 27 \text{ kg/m}^2$ ) had higher risks for both total and breast cancer mortality, whereas the HR were much smaller and did not reach statistical significance for those with BMI less than the median. Although this difference in the mortality risks between BMI groups may be indicative of its role as a potential effect measure modifier in the association between sugar-sweetened soda consumption and mortality, there was no significant interaction.

Postmenopausal women had significantly higher BMI compared with premenopausal women (P < 0.01). We found increased risk of breast cancer mortality ( $P_{\rm trend} = 0.03$ ) with higher frequency of sugar-sweetened soda consumption among postmenopausal women with breast cancer, even with adjustment for BMI. However, we did not find an association with total mortality in this older group. Although we also found significantly higher risks of total and breast cancer mortality among premenopausal women, the number of women in this group in the WEB study are small and CIs for the HR were wide. We know of no other study on this association between sugar-sweetened soda and breast cancer mortality with data stratified by menopausal status.

Another mechanism by which sugar-sweetened soda may promote carcinogenesis and potentially affect prognosis after breast cancer diagnosis, is through glycemic response (44). With the large quantities of sucrose and fructose that make up sugar-sweetened soda, these beverages have the highest glycemic load in comparisons to other foods or drinks (45, 46). Their contribution to high concentrations of glucose and insulin may lead to hyperinsulinemia, impaired glucose tolerance, and higher circulating insulin-like growth factor (IGF) levels (47-49); all of which have been associated with higher risk of breast cancer through enhanced tumor development and tumor cell migration (28, 50-56). Hyperglycemia after sugar-sweetened soda consumption also induces oxidative stress (57, 58). Most sugar-sweetened soda contain large amounts of fructose from the sweetening agent, HFCS, which can produce advanced glycation end-products, found to contribute to the development and progression of cancers in nonhuman studies (28, 59). In a prospective cohort, patients with breast cancer in the United States with higher prediagnostic blood glucose were found to have lower overall survival (60). In another cohort of nonmetastatic patients with breast cancer, all-cause mortality was associated with high fasting blood glucose level. In the Health, Eating, Activity and Lifestyle (HEAL) Study of Breast Cancer Prognosis cohort, there was a nonsignificant trend toward higher all-cause and breast cancer mortality risks associated with high glycemic index and glycemic loads (61).

When the association between mortality and sugar-sweetened soda was stratified by ER-status among the WEB breast cancer cases, there were higher risks for both total mortality and breast cancer mortality for women with ER positive, but not ER negative tumors. To our knowledge, there are no other studies that have examined this association in a cohort of patients with breast cancer stratified by ER status. Additional research, accounting for adequate statistical power, should be performed to investigate the interaction between SSB intake and ER status in the association with mortality among women with breast cancer.

These results need to be considered in the context of the strengths and limitations of the study. As an observational study, we made every effort to control for other possible variables that might affect the associations under investigation, but residual confounding cannot be entirely ruled out. Another limitation of these findings is the single measure of diet used as exposure measure; participants were asked to recall their usual dietary intake in the 12 to 24 months prior to the breast cancer diagnosis. The effect of diet on breast cancer survival is likely to be a complex combination of prediagnostic and postdiagnostic consumption (62, 63); we did not have dietary data following diagnosis or during follow-up. Women in the WEB study may have changed their diet following their breast cancer diagnosis, potentially with changes toward a healthier diet and lifestyle, perhaps including reduced sugar-sweetened soda intake (62). If this change did occur, it would likely have attenuated the observed associations.

Another limitation in the examination of sugar-sweetened soda is the potential underestimation of total sugar intake which is considered the primary dietary culprit in SSB for poor survival post breast cancer diagnosis. Because of the nature of the questionnaire, we could not isolate individual types of sugar-containing beverages, such as juices, teas, coffees, sports drinks, or flavored waters. However, since sugar-sweetened soda makes up the majority of all SSB consumption (29), our reported associations are most likely underestimates of the true associations. It is also important to note that high frequency of sugar-sweetened soda consumption is not an isolated feature of the diet, but is usually part of an unhealthy diet pattern. In our analyses, we attempted to adjust for overall diet quality, by adjusting for fruit, vegetable, and dietary fiber. Results remained similar to models not including diet quality variables. In addition, because sugar-sweetened nondiet soda is perceived as an unhealthy choice, the information provided by the WEB participants may not be completely accurate due to social desirability bias. We would not expect the bias to be differential between those who remained alive and those who died because diet information was collected prior to the outcome occurrence. All these sources of error would likely result in an underestimation of the actual intake of sugar-sweetened soda and an underestimation of the true association between sugar-sweetened soda and mortality in this population.

This study also has multiple strengths. We used data from a largescale population-based series of incident, histologically-confirmed breast cancer cases. In addition, where other studies have focused on the general disease-free population, we specifically focused on a population of patients with breast cancer and survivors. Survival was ascertained prospectively from the time of enrolment, using the NDI, which has been widely utilized and validated for mortality studies (64). Furthermore, we had a relatively long follow-up time that is adequate for survival studies (median: 224; range: 8.88–262 months). Finally, comparisons of descriptive characteristics between women who died with those still alive at the end of the follow-up period identified expected risk factors associated with mortality, suggesting that our sample is representative of the general population of patients with breast cancer.

In summary, our study contributes to the current growing evidence of mortality risk associated with the consumption of SSB. Our findings suggest that higher frequency of consumption of sugar-sweetened soda is associated with greater risk of both total mortality and breast cancer mortality among women diagnosed with incident invasive breast cancer. Overall, even though we cannot infer causality from this study, our findings coupled with the nutritional knowledge that sugarsweetened soda is a nutrient-poor beverage option, having a high contribution of unnecessary added sugars to the diet, support recommendations for their reduced consumption, even among patients with breast cancer and survivors.

#### **Authors' Disclosures**

M. Trevisan reports grants from SUNY Buffalo during the conduct of the study. J.L. Freudenheim reports grants from NIH during the conduct of the study. No disclosures were reported by the other authors.

### **Authors' Contributions**

N. Koyratty: Formal analysis, writing-original draft, writing-review and editing. S.E. McCann: Conceptualization, writing-review and editing. A.E. Millen: Writingreview and editing. J. Nie: Data curation, software. M. Trevisan: Writing-review and

#### References

- CDC. Breast cancer statistics. 2019; Available at: https://www.cdc.gov/cancer/ breast/statistics/index.htm. [Accessed May 28, 2019].
- ACS. How common is breast cancer? 2018; Available at: https://www.cancer.org/ cancer/breast-cancer/about/how-common-is-breast-cancer.html. [Accessed April 20, 2018].
- World Cancer Research Fund/American Institute for Cancer Research. Diet, nutrition, physical activity and cancer: a global perspective. Continuous Update Project Expert Report 2018.
- U.S. Department of Health and Human Services and U.S. Department of Agriculture. 2015–2020 dietary guidelines for Americans. 8th edition. December 2015. Available at: https://health.gov/sites/default/files/2019-09/2015-2020\_ Dietary\_Guidelines.pdf.
- Stanaway JD, Afshin A, Gakidou E, Lim SS, Abate D, Abate KH, et al. Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks for 195 countries and territories, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. Lancet 2018;392:1923–94.
- 6. WHO. Guideline: sugars intake for adults and children. Geneva; 2015.
- Huth PJ, Fulgoni VL, Keast DR, Park K, Auestad N. Major food sources of calories, added sugars, and saturated fat and their contribution to essential nutrient intakes in the U.S. diet: data from the National Health and Nutrition Examination Survey (2003–2006). Nutr J 2013;12:116.
- Drewnowski A, Rehm CD. Consumption of added sugars among US children and adults by food purchase location and food source. Am J Clin Nutr 2014;100: 901–7.
- Keller A, O'Reilly EJ, Malik V, Buring JE, Andersen I, Steffen L, et al. Substitution of sugar-sweetened beverages for other beverages and the risk of developing coronary heart disease: results from the Harvard Pooling Project of Diet and Coronary Disease. Prev Med 2020;131:105970.
- Malik VS, Pan A, Willett WC, Hu FB. Sugar-sweetened beverages and weight gain in children and adults: a systematic review and meta-analysis. Am J Clin Nutr 2013;98:1084–102.
- Genkinger JM, Li R, Spiegelman D, Anderson KE, Albanes D, Bergkvist L, et al. Coffee, tea, and sugar-sweetened carbonated soft drink intake and pancreatic cancer risk: a pooled analysis of 14 cohort studies. Cancer Epidemiol Biomarkers Prev 2012;21:305–18.
- Inoue-Choi M, Robien K, Mariani A, Cerhan JR, Anderson KE. Sugar-sweetened beverage intake and the risk of type I and type II endometrial cancer among postmenopausal women. Cancer Epidemiol Biomarkers Prev 2013;22:2384–94.
- Tasevska N, Jiao L, Cross AJ, Kipnis V, Subar AF, Hollenbeck A, et al. Sugars in diet and risk of cancer in the NIH-AARP Diet and Health Study. Int J Cancer 2012;130:159–69.
- Hodge AM, Bassett JK, Milne RL, English DR, Giles GG. Consumption of sugarsweetened and artificially sweetened soft drinks and risk of obesity-related cancers. Public Health Nutr 2018;21:1618–26.
- Lu S, Huang X, Yu H, Yang J, Han R, Su J, et al. Dietary patterns and risk of breast cancer in Chinese women: a population-based case-control study. Lancet North Am Ed 2016;388:S61.
- Miles FL, Chang SC, Morgenstern H, Tashkin D, Rao JY, Cozen W, et al. Association of sugary beverages with survival among patients with cancers of the upper aerodigestive tract. Cancer Causes Control 2016;27:1293–300.

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- Fuchs MA, Sato K, Niedzwiecki D, Ye X, Saltz LB, Mayer RJ, et al. Sugarsweetened beverage intake and cancer recurrence and survival in CALGB 89803 (alliance). PLoS One 2014;9:e99816.
- Anderson JJ, Gray SR, Welsh P, Mackay DF, Celis-Morales CA, Lyall DM, et al. The associations of sugar-sweetened, artificially sweetened and naturally sweet juices with all-cause mortality in 198,285 UK Biobank participants: a prospective cohort study. BMC Med 2020;18:97.
- Barrington WE, White E. Mortality outcomes associated with intake of fast-food items and sugar-sweetened drinks among older adults in the Vitamins and Lifestyle (VITAL) study. Public Health Nutr 2016;19:3319–26.
- Malik Vasanti S, Li Y, Pan A, De Koning L, Schernhammer E, Willett WC, et al. Long-term consumption of sugar-sweetened and artificially sweetened beverages and risk of mortality in US adults. Circulation 2019;139:2113–25.
- Ramne S, Alves Dias J, Gonzalez-Padilla E, Olsson K, Lindahl B, Engstrom G, et al. Association between added sugar intake and mortality is nonlinear and dependent on sugar source in 2 Swedish population-based prospective cohorts. Am J Clin Nutr 2019;109:411–23.
- Collin LJ, Judd S, Safford M, Vaccarino V, Welsh JA. Association of sugary beverage consumption with mortality risk in US adults: a secondary analysis of data from the REGARDS Study. JAMA Netw Open 2019;2:e193121.
- Mullee A, Romaguera D, Pearson-Stuttard J, Viallon V, Stepian M, Freisling H, et al. Association between soft drink consumption and mortality in 10 European countries. JAMA Intern Med 2019;179:1479–90.
- Paganini-Hill A, Kawas CH, Corrada MM. Non-alcoholic beverage and caffeine consumption and mortality: the Leisure World Cohort Study. Prev Med 2007;44: 305–10.
- Odegaard AO, Koh WP, Yuan JM, Pereira MA. Beverage habits and mortality in Chinese adults. J Nutr 2015;145:595–604.
- Chazelas E, Srour B, Desmetz E, Kesse-Guyot E, Julia C, Deschamps V, et al. Sugary drink consumption and risk of cancer: results from NutriNet-Sante prospective cohort. BMJ 2019;366:12408.
- Makarem N, Nicholson JM, Bandera EV, McKeown NM, Parekh N. Consumption of whole grains and cereal fiber in relation to cancer risk: a systematic review of longitudinal studies. Nutr Rev 2016;74:353–73.
- Jiang Y, Pan Y, Rhea PR, Tan L, Gagea M, Cohen L, et al. A sucrose-enriched diet promotes tumorigenesis in mammary gland in part through the 12-lipoxygenase pathway. Cancer Res 2016;76:24–9.
- Han E, Powell LM. Consumption patterns of sugar-sweetened beverages in the United States. J Acad Nutr Diet 2013;113:43–53.
- McCann SE, Muti P, Vito D, Edge SB, Trevisan M, Freudenheim JL. Dietary lignan intakes and risk of pre- and postmenopausal breast cancer. Int J Cancer 2004;111:440–3.
- Freudenheim JL, Marshall JR, Graham S, Laughlin R, Vena JE, Bandera E, et al. Exposure to breastmilk in infancy and the risk of breast cancer. Epidemiology 1994;5:324–31.
- McCann SE, Sempos C, Freudenheim JL, Muti P, Russell M, Nochajski TH, et al. Alcoholic beverage preference and characteristics of drinkers and nondrinkers in western New York (United States). Nutr Metab Cardiovasc Dis 2003;13:2–11.
- Block G, Hartman AM, Dresser CM, Carroll MD, Gannon J, Gardner L. A databased approach to diet questionnaire design and testing. Am J Epidemiol 1986; 124:453–69.

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- Barba M, McCann SE, Nie J, Vito D, Stranges S, Fuhrman B, et al. Perinatal exposures and breast cancer risk in the Western New York Exposures and Breast Cancer (WEB) Study. Cancer Causes Control 2006;17:395–401.
- CDC. Get the facts: sugar-sweetened beverages and consumption; 2017. Available at: https://www.cdc.gov/nutrition/data-statistics/sugar-sweetenedbeverages-intake.html. [Accessed August 2018].
- Schulze MB, Manson JE, Ludwig DS, Colditz GA, Stampfer MJ, Willett WC, et al. Sugar-sweetened beverages, weight gain, and incidence of type 2 diabetes in young and middle-aged women. JAMA 2004;292:927–34.
- Apovian CM. Sugar-sweetened soft drinks, obesity, and type 2 diabetes. JAMA 2004;292:978–9.
- Vartanian LR, Schwartz MB, Brownell KD. Effects of soft drink consumption on nutrition and health: a systematic review and meta-analysis. Am J Public Health 2007;97:667–75.
- Huang Z, Hankinson SE, Colditz GA, Stampfer MJ, Hunter DJ, Manson JE, et al. Dual effects of weight and weight gain on breast cancer risk. JAMA 1997;278: 1407–11.
- Mulholland HG, Murray LJ, Cardwell CR, Cantwell MM. Dietary glycaemic index, glycaemic load and breast cancer risk: a systematic review and metaanalysis. Br J Cancer 2008;99:1170–5.
- Cecchini RS, Swain SM, Costantino JP, Rastogi P, Jeong JH, Anderson SJ, et al. Body mass index at diagnosis and breast cancer survival prognosis in clinical trial populations from NRG Oncology/NSABP B-30, B-31, B-34, and B-38. Cancer Epidemiol Biomarkers Prev 2016;25:51–59.
- 42. Guo Q, Burgess S, Turman C, Bolla MK, Wang Q, Lush M, et al. Body mass index and breast cancer survival: a Mendelian randomization analysis. Int J Epidemiol 2017;46:1814–22.
- Nelson SH, Marinac CR, Patterson RE, Nechuta SJ, Flatt SW, Caan BJ, et al. Impact of very low physical activity, BMI, and comorbidities on mortality among breast cancer survivors. Breast Cancer Res Treat 2016; 155:551–7.
- 44. McCann SE, McCann WE, Hong CC, Marshall JR, Edge SB, Trevisan M, et al. Dietary patterns related to glycemic index and load and risk of premenopausal and postmenopausal breast cancer in the Western New York Exposure and Breast Cancer Study. Am J Clin Nutr 2007;86:465–71.
- Bray GA, Nielsen SJ, Popkin BM. Consumption of high-fructose corn syrup in beverages may play a role in the epidemic of obesity. Am J Clin Nutr 2004;79: 537–43.
- Brand-Miller JC, Stockmann K, Atkinson F, Petocz P, Denyer G. Glycemic index, postprandial glycemia, and the shape of the curve in healthy subjects: analysis of a database of more than 1000 foods. Am J Clin Nutr 2009;89:97–105.
- Jenkins DJ, Jenkins AL, Wolever TM, Vuksan V, Rao AV, Thompson LU, et al. Low glycemic index: lente carbohydrates and physiological effects of altered food frequency. Am J Clin Nutr 1994;59:7068–98.
- Liu S, Manson JE, Stampfer MJ, Holmes MD, Hu FB, Hankinson SE, et al. Dietary glycemic load assessed by food-frequency questionnaire in relation to plasma high-density-lipoprotein cholesterol and fasting plasma triacylglycerols in postmenopausal women. Am J Clin Nutr 2001;73:560–6.
- Ludwig DS. The glycemic index: Physiological mechanisms relating to obesity, diabetes, and cardiovascular disease. JAMA 2002;287:2414–23.

- Lawlor DA, Smith GD, Ebrahim S. Hyperinsulinaemia and increased risk of breast cancer: findings from the British Women's Heart and Health Study. Cancer Causes Control 2004;15:267–75.
- Kaaks R, Lundin E, Rinaldi S, Manjer J, Biessy C, Soderberg S, et al. Prospective study of IGF-I, IGF-binding proteins, and breast cancer risk, in northern and southern Sweden. Cancer Causes Control 2002;13:307–16.
- Del Giudice ME, Fantus IG, Ezzat S, McKeown-Eyssen G, Page D, Goodwin PJ. Insulin and related factors in premenopausal breast cancer risk. Breast Cancer Res Treat 1998;47:111–20.
- Giles GG, Simpson JA, English DR, Hodge AM, Gertig DM, MacInnis RJ, et al. Dietary carbohydrate, fibre, glycaemic index, glycaemic load and the risk of postmenopausal breast cancer. Int J Cancer 2006;118:1843–7.
- 54. Samani AA, Yakar S, LeRoith D, Brodt P. The role of the IGF system in cancer growth and metastasis: overview and recent insights. Endocr Rev 2007;28:20–47.
- Sieri S, Pala V, Brighenti F, Agnoli C, Grioni S, Berrino F, et al. High glycemic diet and breast cancer occurrence in the Italian EPIC cohort. Nutr Metab Cardiovasc Dis 2013;23:628–34.
- Romieu I, Ferrari P, Rinaldi S, Slimani N, Jenab M, Olsen A, et al. Dietary glycemic index and glycemic load and breast cancer risk in the European Prospective Investigation into Cancer and Nutrition (EPIC). Am J Clin Nutr 2012;96:345–55.
- Title LM, Cummings PM, Giddens K, Nassar BA. Oral glucose loading acutely attenuates endothelium-dependent vasodilation in healthy adults without diabetes: an effect prevented by vitamins C and E. J Am Coll Cardiol 2000;36:2185–91.
- Loader J, Meziat C, Watts R, Lorenzen C, Sigaudo-Roussel D, Stewart S, et al. Effects of sugar-sweetened beverage consumption on microvascular and macrovascular function in a healthy population. Arterioscler Thromb Vasc Biol 2017; 37:1250–60.
- Peeters K, Van Leemputte F, Fischer B, Bonini BM, Quezada H, Tsytlonok M, et al. Fructose-1,6-bisphosphate couples glycolytic flux to activation of Ras. Nat Commun 2017;8:922.
- 60. Monzavi-Karbassi B, Gentry R, Kaur V, Siegel ER, Jousheghany F, Medarametla S, et al. Pre-diagnosis blood glucose and prognosis in women with breast cancer. Cancer Metab 2016;4:7.
- Belle FN, Kampman E, McTiernan A, Bernstein L, Baumgartner K, Baumgartner R, et al. Dietary fiber, carbohydrates, glycemic index, and glycemic load in relation to breast cancer prognosis in the HEAL cohort. Cancer Epidemiol Biomarkers Prev 2011;20:890–9.
- Terranova CO, Protani MM, Reeves MM. Overall dietary intake and prognosis after breast cancer: a systematic review. Nutr Cancer 2018;70:153–63.
- 63. George SM, Ballard-Barbash R, Shikany JM, Caan BJ, Freudenheim JL, Kroenke CH, et al. Better postdiagnosis diet quality is associated with reduced risk of death among postmenopausal women with invasive breast cancer in the Women's Health Initiative. Cancer Epidemiol Biomarkers Prev 2014;23:575–83.
- Johnson CJ, Weir HK, Fink AK, German RR, Finch JL, Rycroft RK, et al. The impact of National Death Index linkages on population-based cancer survival rates in the United States. Cancer Epidemiol 2013;37:20–8.