



Evaluation of a novel biomarker of added sugar intake ($\delta^{13}\text{C}$) compared with self-reported added sugar intake and the Healthy Eating Index-2010 in a community-based, rural US sample

Valisa E Hedrick^{1,*}, Brenda M Davy¹, Grace A Wilburn¹, A Hope Jahren² and Jamie M Zoellner¹

¹Department of Human Nutrition, Foods and Exercise, Virginia Tech, 295 West Campus Drive, 335A Wallace Hall (0430), Blacksburg, VA 24061, USA; ²Department of Geology and Geophysics, University of Hawaii at Manoa, Honolulu, HI, USA

Submitted 29 August 2014; Final revision received 26 January 2015; Accepted 9 March 2015

Abstract

Objective: The $\delta^{13}\text{C}$ value of human blood is an emerging novel biomarker of added sugar (AS) intake for adults. However, no free-living, community-based assessments of comparative validity of this biomarker have been conducted. The purpose of the present investigation was to determine if Healthy Eating Index-2010 (HEI-2010) score, SoFAAS score (HEI-2010 sub-component for solid fat, alcohol and AS), AS and sugar-sweetened beverage (SSB) intakes were associated with $\delta^{13}\text{C}$ value of fingerstick blood in a community-based sample of adults, while controlling for relevant demographics.

Design: A cross-sectional analysis of data obtained from assessments of BMI, dietary intake using 24 h recalls and a fingerstick blood sample was completed. Statistical analyses included descriptive statistics, multiple linear regression and one-way ANOVA.

Setting: Rural Southwest Virginia, USA.

Subjects: Adults (n 216) aged >18 years who consumed at least 837 kJ/d (200 kcal/d) from SSB.

Results: This sample of adult participants with low socio-economic status demonstrated a mean HEI-2010 score of 43.4 (SD 12.2), mean SoFAAS score of 10.2 (SD 5.7), mean AS intake of 93 (SD 65) g/d and mean blood $\delta^{13}\text{C}$ value of -18.88 (SD 0.7) ‰. In four separate regression models, HEI-2010 ($R^2=0.16$), SoFAAS ($R^2=0.19$), AS ($R^2=0.15$) and SSB ($R^2=0.14$) predicted $\delta^{13}\text{C}$ value (all $P\leq 0.001$). Age was also predictive of $\delta^{13}\text{C}$ value, but not sex or race.

Conclusions: These findings suggest that fingerstick $\delta^{13}\text{C}$ value has the potential to be a minimally invasive method for assessing AS and SSB intake and overall dietary quality in community-based settings. Strengths, limitations and future areas of research for using an objective $\delta^{13}\text{C}$ biomarker in diet-related public health studies are discussed.

Keywords
Added sugar
Biomarker
Dietary assessment
Dietary quality

Consumption of added sugars (AS) has been suggested as a major contributor to the development of obesity and related co-morbidities such as diabetes^(1,2). Consequently, as sugar-sweetened beverages (SSB) are the largest source of AS in the diet, excessive consumption may lead to increased energy intake and weight gain⁽³⁾. Intake of SSB and AS represent approximately 7% and 16%, respectively, of total energy intake in US adults^(4,5). Yet, the topic of AS continues to be highly controversial^(6,7). For example, some have argued that 'sugar comprises the single most important cause of the worldwide epidemics of obesity

and diabetes⁽⁸⁾, while others have stated that 'there is no evidential basis for setting a quantitative target for sugar consumption⁽⁹⁾'. A recent review in this area cited the need for methodological advances, specifically instruments to measure dietary intake⁽¹⁰⁾. Thus, objective methods of assessing AS intake are necessary to evaluate the impact of AS and SSB on health status in the US population⁽¹¹⁾, especially in light of recent emphasis on public policies related to AS⁽¹²⁻¹⁴⁾.

Most studies investigating diet and health are limited by their reliance on self-reported measures of habitual dietary



intake, such as 24 h dietary recalls, which can require significant staff resources (i.e. time, personnel, expense) and participant burden^(15–17). These methods are not always practical in large-scale clinical trials and community-based studies; however, dietary biomarkers can overcome some of these limitations^(18–22). It has been demonstrated that rural, low health-literacy populations consume greater amounts of SSB^(23–25) and are more likely to be at risk for health disparities^(26,27); thus low-burden objective biomarkers of dietary intake are needed to assess population health status^(10,18).

The $\delta^{13}\text{C}$ value of human blood, in various mediums, and hair has demonstrated preliminary validity as an AS biomarker for adults in three clinical laboratory-based investigations using self-reported dietary data: capillary fingerstick⁽²⁸⁾; serum, capillary fingerstick and clot⁽²⁹⁾; and serum⁽³⁰⁾; and in one clinical investigation ($n = 5$) which used a feeding study-type design to assess non-fasting plasma glucose $\delta^{13}\text{C}$ values⁽³¹⁾. However, trials of free-living individuals are needed to further assess the utility of the $\delta^{13}\text{C}$ AS biomarker, as only two known studies currently exist in community- or field-based settings, both of which have been conducted in Alaska with a Yup'ik Eskimo population using either red blood cells⁽³²⁾ or hair, plasma and red blood cells⁽³³⁾, and no community-based studies have been conducted within a general US population. High natural concentrations of ^{13}C are found in corn (e.g. high-fructose corn syrup) and in cane plant sugars and their derivatives^(29,34,35). Corn derivatives consist of corn starch, corn syrup, popcorn and corn meal; while molasses, plain cane sugar, brown cane sugar and powdered cane sugar are considered sugar cane derivatives⁽³⁵⁾. US Department of Agriculture data depict an increase in high-fructose corn syrup intake over the past 30 years, while intake of other sugar sources has remained relatively constant⁽³⁶⁾. Moreover, as SSB is a primary dietary source of AS intake, $\delta^{13}\text{C}$ value of blood has shown potential as a biomarker for AS intake due to the high concentration of high-fructose corn syrup in SSB^(28,29,35). Furthermore, it has been established that more research examining the effects of demographic characteristics on biomarker variability is needed for dietary intake biomarkers before generalizability to the overall population can be inferred⁽²⁰⁾. The current biomarker literature has a lack of variability in sex, race and age within the study samples, and consequently limited research is available examining the influence of demographic variance in relationship to biomarkers⁽²⁰⁾. To date, only one investigation has evaluated the independent effects of demographic variance on blood $\delta^{13}\text{C}$ values⁽³⁷⁾.

The Healthy Eating Index-2010 (HEI-2010)⁽³⁸⁾ is a dietary measure that evaluates the extent to which an individual's dietary intake conforms to the 2010 Dietary Guidelines for Americans⁽³⁹⁾. The HEI-2010 is composed of twelve dietary components (nine adequacy and three

moderation categories) that combine to provide an overall dietary score and includes total fruit, whole fruit, total vegetables, dark-green vegetables and beans, whole grains, dairy, total protein foods, seafood and plant proteins, fatty acids, refined grains, sodium and empty calories (solid fat, alcohol and AS (SoFAAS)). While the HEI-2010 provides valuable information, it is not always feasible to obtain the scores due to high participant and researcher burden, as extensive dietary intake information is necessary to calculate this score. Therefore, additional research on the use of minimally invasive, objective biomarkers, which could be feasibly used in community or field settings to assess dietary quality, is needed.

The primary aims of the present investigation, conducted in a community-based sample of adults in a rural health-disparate region of Southwest Virginia, are to: (i) determine if HEI-2010 and SoFAAS scores, AS and SSB intakes predict $\delta^{13}\text{C}$ values in fingerstick blood, while controlling for age, sex and race; and (ii) explore the distribution of HEI-2010 scores among tertile levels of $\delta^{13}\text{C}$ values. The hypotheses, tested with four separate regression models, are that HEI-2010 and SoFAAS scores, AS and SSB intakes will each significantly predict $\delta^{13}\text{C}$ values while controlling for age, sex and race. Additionally, mean HEI-2010 and SoFAAS scores will be significantly greater in the low $\delta^{13}\text{C}$ value tertile as compared with the high $\delta^{13}\text{C}$ value tertile.

Experimental methods

Participants and design

The current cross-sectional, comparative validation investigation utilized baseline data from the ongoing clinical trial known as Talking Health⁽⁴⁰⁾ ($n = 224$). Talking Health is a 6-month, community-based, two-arm randomized controlled trial which targets SSB consumption behaviours among low socio-economic status (SES) adult (aged >18 years) residents in rural Southwest Virginia, as compared with a matched-contact control group targeting physical activity behaviours. Participants were recruited from April 2012 to September 2013. To be eligible, participants had to consume at least 837 kJ/d (200 kcal/d) from SSB, as assessed by the BEVQ-15 (an FFQ developed to measure beverage consumption patterns)^(41–44), prior to enrolment. In order to target low-SES adults, recruitment efforts were concentrated on counties that qualified as 'medically underserved areas' by using the benchmark of an Index of Medical Underservice of 62.0 or less⁽⁴⁵⁾. Various recruitment methods were implemented: in-person community outreach efforts in various venues such as day care centres, festivals, community colleges, retail stores, Head Start, health clinics, free clinics, WIC (Special Supplemental Nutrition Program for Women, Infants, and Children) clinics and health departments. Additionally, local extension agents were employed to recruit within the communities, and



other recruitment methods included newspaper advertisements, flyers, email listservs and targeted postcard mailings. Although low-SES adults were targeted, income and education status were not eligibility criteria. The study was conducted according to the guidelines laid down in the Declaration of Helsinki and the Virginia Tech Institutional Review Board approved the study protocol. Participants provided written informed consent prior to enrolment.

Methods

Participants underwent assessments of height, measured in metres without shoes using a portable stadiometer; weight, measured in light clothing without shoes, to the nearest 0.1 kg using a digital scale (model 310GS; Tanita, Tokyo, Japan); calculated BMI; and habitual dietary intake using three 24 h dietary intake recalls^(17,46). The first 24 h food recall was completed in-person and the two remaining food recalls were completed unannounced via telephone; recalls were collected by trained research technicians who were supervised by a doctoral-level registered dietitian. One weekend and two weekdays were recalled to provide a more accurate representation of habitual dietary habits. The dietary intake recalls were analysed using the Nutrition Data System for Research (NDSR) nutritional analysis software (Nutrition Coordinating Center, University of Minnesota, 2011). HEI-2010 scores and AS intake were calculated using dietary intake recall data⁽³⁸⁾. HEI-2010 scores were derived from NDSR output based on guidelines developed by NDSR⁽⁴⁷⁾. The HEI-2010 total and sub-component scores were calculated according to a standardized published protocol, which includes an adjustment for energy intake⁽⁴⁷⁾. Briefly, total fruit, whole fruit, total vegetables, greens and beans, and dairy were converted into one cup equivalents; whole grains, refined grains, total protein foods, and seafood and plant proteins were converted into one ounce equivalents; and sodium was converted into a one gram equivalent. These components were divided by total daily energy intake/1000 to determine serving equivalents per 1000 kcal (4184 kJ), which determined the raw component scores. Fatty acid raw scores were calculated as a ratio of (PUFA+MUFA) to SFA, and the empty calories (SoFAAS) raw score was determined as the percentage of total energy from solid fats, alcohol and added sugars⁽⁴⁷⁾. Standardized scores were calculated from the raw scores based on the HEI-2010 standards for minimum and maximum scores⁽³⁸⁾. HEI-2010 scores range from 0 to 100 (score is a total of all twelve component scores), and SoFAAS scores range from 0 to 20, with higher scores indicating greater adherence to the 2010 Dietary Guidelines for Americans⁽³⁹⁾. HEI-2010 scores can also be divided into three categories based on dietary quality: good (>80), needs improvement (51–80) or poor (<51)⁽³⁸⁾. Participants also completed a multimedia version of the BEVQ-15⁽⁴⁴⁾ to determine SSB consumption and provided demographic information.

To determine $\delta^{13}\text{C}$ values, a fasting blood sample was provided via a routine fingerstick; samples were blotted onto sterilized Whatman spun glass filters (type GF/D), air-dried, then analysed for $\delta^{13}\text{C}$ value using natural abundance stable isotope mass spectrometry (NASIMS)⁽²⁸⁾. Stable isotope values are reported using standard δ notation in units of 'permil' (‰) relative to international standards (Vienna Pee Dee Belemnite). L-Alanine was used as an internal laboratory standard for carbon. The analytical error associated with each sample measurement was less than 0.05 ‰ in every case. The time course of $\delta^{13}\text{C}$ value demonstrated by a specific substrate is dependent upon the turnover rate of the substrate. Hair and nails represent the longest turnover rate (2–6 months depending on area of sampling), red blood cells represent approximately 120 d⁽¹¹⁾ and serum measures intake over a 2- to 3-week period⁽³³⁾. The time course associated with fingerstick blood is 2–3 weeks (due to the serum content); however, $\delta^{13}\text{C}$ value may be diluted as a result of the red blood cell content of fingerstick blood. Davy *et al.* reported a $\delta^{13}\text{C}$ value range of –22.09 to –18.87 ‰ and indicated that higher AS and SSB consumption was associated with higher $\delta^{13}\text{C}$ value (i.e. $\delta^{13}\text{C}$ value closer to 0 corresponded to higher AS intake)⁽²⁸⁾. Therefore, as HEI-2010/SoFAAS scores improve (increase), indicating less AS intake, $\delta^{13}\text{C}$ value should decrease (i.e. become further away from 0), thus creating an inverse relationship; however, when assessing SSB kcal and AS gram intake as related to $\delta^{13}\text{C}$ value, values will demonstrate a direct relationship.

Data analysis

Statistical analyses were performed using the statistical software package IBM SPSS version 21.0 for Windows (2012). Descriptive statistics (means and standard deviations; frequencies) are reported for demographic characteristics. Multiple linear regression models were used to predict $\delta^{13}\text{C}$ values. Variables were entered into a multiple linear regression model using the 'enter' method with two blocks of independent variables. Four separate models predicting $\delta^{13}\text{C}$ values were run using HEI-2010 scores, SoFAAS scores, AS grams and SSB kcal in the first block; and age, sex and race in the second block. Sex and race (Caucasian and other) were dummy coded. One-way ANOVA evaluated differences in HEI-2010 and SoFAAS scores for tertiles of $\delta^{13}\text{C}$ values. Tertiles were created using an equal number of observations within each tertile. Missing data were addressed by using list-wise deletion methods for the multiple linear regression models and case-by-case deletion for ANOVA. The recommended approach for multiple regression analyses ($n \geq 50 + 8m$, where m equals the number of predictor variables) to detect a moderate effect size with 80% power and an α of 0.05 was applied⁽⁴⁸⁾. A *priori* hypothesis included a maximum of four predictor variables per model; therefore a sample size of at least eighty-two participants provided sufficient power.

Results and discussion

Demographics

An initial sample size of 224 was utilized; however, to meet the assumptions of normality, participants with energy intake greater than 2 SD from the mean were excluded (n 8). Thus, a final sample size of 216 was used for the present investigation. Participants were primarily female (83%), Caucasian (94%), with a mean age of 41 (SD 14) years (range 18–81 years) and an annual household income of \$US 24 132 (SD \$US 17 340). Although BMI was widely distributed (mean 32.7 (SD 9.0) kg/m²; range 16.1–71.7 kg/m²), 55% of the sample was considered obese (BMI \geq 30.0 kg/m²). When compared with US census data for this medically underserved region, the participants were representative in terms of age (census mean age = 40.8 years), income level (census mean income = \$US 21 751) and race (census: Caucasian = 93.9%); however, education status for the enrolled participants was slightly higher than in the US census (70% *v.* 58% with an education beyond a high school degree). Additionally, 74% of the sample had an annual income of \$US 35 000 or less, as compared with the state of Virginia (27%) and the USA (33.5%)⁽⁴⁹⁾, indicating a lower SES study population (Table 1).

Dietary intake and $\delta^{13}\text{C}$ value

The overall completion rate for the dietary intake recalls was 89% (75% had three complete days, 17.5% had two and 7.5% had only one complete day). Participants had a

Table 1 Characteristics of the participants: adults aged >18 years (n 216), rural Southwest Virginia, USA, April 2012–September 2013

Characteristic	n	%
Sex		
Male	36	17
Female	180	83
Race/ethnicity		
White	203	94
African American	8	3.5
Other	1	0.5
More than one race	4	2
BMI category		
Underweight (\leq 18.4 kg/m ²)	4	2
Normal weight (18.5–24.9 kg/m ²)	42	19
Overweight (25.0–29.9 kg/m ²)	51	24
Obese (\geq 30.0 kg/m ²)	119	55
Obese class 1 (30.0–34.9 kg/m ²)	38	17.5
Obese class 2 (35.0–39.9 kg/m ²)	36	17
Obese class 3 (\geq 40.0 kg/m ²)	45	20.5
Education level		
<High school	22	10
High school graduate	43	20
Some college	83	38.5
College graduate	68	31.5
Annual household income (\$US)		
<14 999	86	40
15 000–34 999	74	34
35 000–54 999	27	12.5
\geq 55 000	29	13.5

mean daily intake of 1094 (SD 828) ml (37 (SD 28) fl oz) and 1782 (SD 1443) kJ (426 (SD 345) kcal) from SSB, 7255 (SD 2499) kJ (1734 (SD 597) kcal) from all foods and beverages, and 93 (SD 65) g of AS. Total HEI-2010 score ranged from 14.4 to 81.1 with a mean of 43.4 (SD 12.2); SoFAAS score ranged from 0 to 20 with a mean of 10.2 (SD 5.7). In comparison, the mean total HEI-2010 score for the general US population was considered slightly healthier (49.9 (SD 0.5)); however, the mean SoFAAS score was comparable (9.4 (SD 0.2)) to that in the present sample⁽⁵⁰⁾. The majority of this sample's dietary intake quality was considered 'poor' (75%), with 24.5% 'needing improvement', and only 0.5% were in the 'good' category.

Fingerstick $\delta^{13}\text{C}$ value ranged from -21.05 to -17.00 ‰, with a mean value of -18.88 (SD 0.7)‰. This mean $\delta^{13}\text{C}$ value suggests a higher AS and SSB intake than observed in a prior clinical laboratory-based investigation, which included individuals with a lower habitual AS (66 (SD 5) g/d) and SSB (561 (SD 105) kJ/d, 134 (SD 25) kcal/d) intake and a mean fingerstick $\delta^{13}\text{C}$ value of -19.94 (SD 0.1)‰⁽²⁸⁾.

Multiple linear regression analysis for variables predicting $\delta^{13}\text{C}$ value

In step 1, HEI-2010, SoFAAS, AS grams and SSB kcal were all significant predictors (all $P \leq 0.001$) of $\delta^{13}\text{C}$ value (Table 2). With the addition of the demographic factors (age, sex and race), the models remained significant with R^2 values increasing by about twofold. The β weights remained relatively consistent and significant for HEI-2010, SoFAAS, AS grams and SSB kcal (all $P \leq 0.001$). Age was also a significant predictor of $\delta^{13}\text{C}$ value in all four models (HEI-2010, SoFAAS, AS grams and SSB kcal; all $P \leq 0.001$); however, sex and race were not predictive of $\delta^{13}\text{C}$ value, with the exception of sex for HEI-2010 SoFAAS ($P = 0.04$). The significant contribution of age when predicting $\delta^{13}\text{C}$ value may be explained by the fact that different age groups consume varying amounts of SSB and AS (typically older adults consume less SSB and AS than younger adults)^(5,51). While controlling for demographic factors, for every 1 SD increase in HEI-2010 and SoFAAS scores, there is a decrease of 0.23 and 0.28 SD in $\delta^{13}\text{C}$ values. That is, as HEI-210 and SoFAAS scores increase (indicating greater adherence to the Dietary Guidelines for Americans), $\delta^{13}\text{C}$ values significantly decrease (indicating a lower level of AS consumption). Similarly, for every 1 SD increase in consumption of AS grams and SSB kcal, there is an increase of 0.21 and 0.19 SD in $\delta^{13}\text{C}$ value, respectively.

Prior biomarker studies have been shown to predict dietary intake with wide variance, ranging from 14% to 99%⁽²⁰⁾, and a wide range of correlations have been reported in validity studies of dietary biomarkers (0.03–0.73)⁽¹⁷⁾. Even so, acceptable correlations for this area of research should range from 0.5 to 0.7⁽¹⁷⁾. In the present study, although significant, the 14–19% variance explained by HEI-2010 and SoFAAS scores, AS grams and SSB kcal when predicting $\delta^{13}\text{C}$ value is on the low end of the spectrum. Importantly, there are

**Table 2** Summary of multiple linear regression analysis for variables predicting $\delta^{13}\text{C}$ value among adults aged >18 years (n 216), rural Southwest Virginia, USA, April 2012–September 2013

Model	Step 1		Step 2	
	Model R^2	β	Model R^2	β
Model 1			0.16***	
Step 1				
HEI-2010 total score†	0.09***	-0.30***		-0.23***
Step 2				
Age				-0.26***
Sex‡				-0.09
Race§				0.04
Model 2			0.19***	
Step 1				
HEI-2010 SoFAAS score	0.10***	-0.31***		-0.28***
Step 2				
Age				-0.27***
Sex				-0.13*
Race				0.05
Model 3			0.15***	
Step 1				
AS grams¶	0.06***	0.25***		0.21**
Step 2				
Age				-0.28***
Sex				-0.10
Race				0.06
Model 4			0.14***	
Step 1				
SSB kcal††	0.07***	0.26***		0.19**
Step 2				
Age				-0.26***
Sex				-0.09
Race				0.04

* $P \leq 0.05$, ** $P \leq 0.01$, *** $P \leq 0.001$.

†Healthy Eating Index-2010 (HEI-2010) score ranges from 0 to 100, higher scores indicate greater adherence to the 2010 Dietary Guidelines for Americans.

‡Sex is dummy-coded.

§Race is dummy-coded into 'Caucasian' and 'other'.

||SoFAAS, the empty calorie component of the HEI-2010, is comprised of solid fats, added sugars and alcohol intake. Score ranges from 0 to 20, higher scores indicate greater adherence to the 2010 Dietary Guidelines for Americans (i.e. higher score = less sugar intake).

¶AS, added sugar intake in grams.

††SSB, sugar-sweetened beverage intake in kilocalories.

known limitations to self-reported dietary intake methods (i.e. underestimating intake⁽¹⁷⁾, especially socially undesirable foods such as sugar-rich foods⁽⁵²⁾), which may partially explain this lower variance when using self-reported dietary intake to predict a biomarker. However, given that there is limited available research on biomarkers which assess AS intake and/or overall dietary quality, falling within a similar correlation range as other dietary biomarker studies indicates promise and the need for further investigations utilizing $\delta^{13}\text{C}$ analysis. To fully understand the potential of using the minimally invasive fingerstick $\delta^{13}\text{C}$ value to predict dietary intake in large-scale community trials, $\delta^{13}\text{C}$ validation studies using controlled feeding methods as the comparison are also needed. With improved accuracy in the comparison method, it is plausible to suggest that the amount of variance explained by similar $\delta^{13}\text{C}$ models would increase.

Differences in HEI-2010 and SoFAAS scores for tertiles of $\delta^{13}\text{C}$ value

ANOVA were also used to further explore potential differences in mean HEI-2010 and SoFAAS scores across $\delta^{13}\text{C}$ value tertiles to determine if $\delta^{13}\text{C}$ value was able to distinguish between different levels of dietary quality. HEI-2010 scores were significantly different for upper and lower $\delta^{13}\text{C}$ value tertiles (39.6 (SD 10) and 48.1 (SD 14), respectively; absolute difference = 8.5, $P \leq 0.0001$) and middle and lower tertiles (42.4 (SD 11) and 48.1 (SD 14), respectively; absolute difference = 5.8, $P \leq 0.01$). SoFAAS scores were significantly different for upper and lower $\delta^{13}\text{C}$ value tertiles (7.7 (SD 6) and 12.0 (SD 5), respectively; absolute difference = 4.3, $P \leq 0.0001$) and upper and middle tertiles (7.7 (SD 6) and 10.7 (SD 5), respectively; absolute difference = 3.0, $P \leq 0.01$; Fig. 1(a) and (b)). In line with the stated hypothesis that $\delta^{13}\text{C}$ value is associated with AS and SSB, it was expected that differences in HEI-2010 scores would be slightly less robust than SoFAAS scores across $\delta^{13}\text{C}$ value tertiles, in that HEI-2010 scores are comprised of twelve dietary components while SoFAAS scores are more representative of AS and SSB consumption. Even though SoFAAS scores are a part of total HEI-2010 scores, the mean difference in upper and lower $\delta^{13}\text{C}$ value tertiles for HEI-2010 scores was double the difference for SoFAAS scores (8.5 *v.* 4.3), indicating an effect of AS intake on the diet beyond the SoFAAS component. Although additional research is needed in this area, these data suggest that $\delta^{13}\text{C}$ value may be predictive of overall dietary quality. While dietary biomarkers have been previously utilized to develop and validate HEI scores⁽⁵³⁾, no investigations have assessed the potential of $\delta^{13}\text{C}$ to serve as an indicator of overall dietary quality.

Strengths/limitations

The present investigation is the first to report $\delta^{13}\text{C}$ fingerstick values in a free-living community-based sample within a general US population, and the first examination of associations of $\delta^{13}\text{C}$ fingerstick blood values with overall dietary quality in adults while controlling for relevant demographic factors. One limitation of the study is lack of variability in race/ethnicity and sex, as well as small sample sizes for non-Caucasian (n 13, 6%) and male participants (n 36, 17%). However, the targeted Southwest Virginia region is 95% Caucasian and thus our sample is representative of the study region^(40,49). Furthermore, given that eligibility criteria included an average intake of at least 837 kJ (200 kcal) from SSB per day, overall AS intake was high. Nevertheless, HEI-2010 scores had an adequate range from 14.4 to 81.1. Given these limitations, examining fingerstick blood $\delta^{13}\text{C}$ values among more ethnic/racially diverse participants and among those with lower AS consumption should be a future research priority. An additional limitation is the reliance on self-reported dietary intake data, which is subject to reporting error and

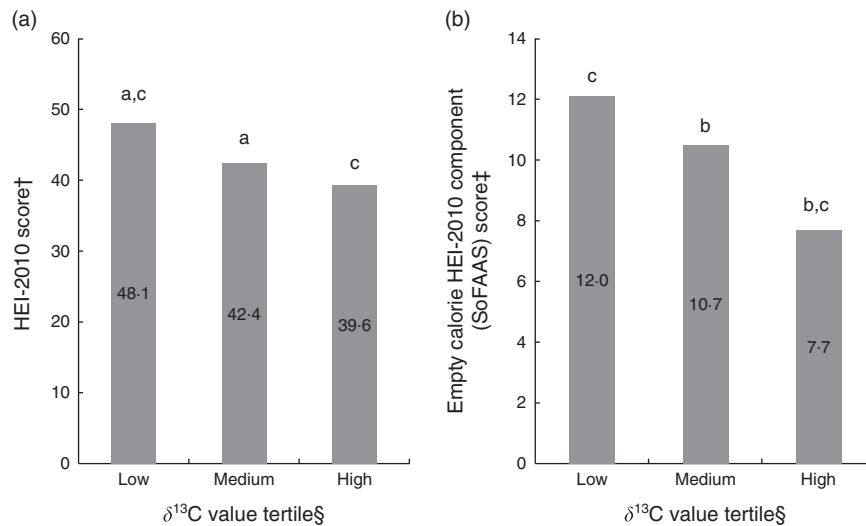


Fig. 1 (a) Mean Healthy Eating Index-2010 (HEI-2010) score and (b) mean empty calorie HEI-2010 component (SoFAAS) score by tertile level of $\delta^{13}\text{C}$ value among adults aged >18 years (n 216), rural Southwest Virginia, USA, April 2012–September 2013. †HEI-2010 score ranges from 0 to 100, higher scores indicate greater adherence to the 2010 Dietary Guidelines for Americans. ‡SoFAAS (solid fats, alcohol and added sugars) score ranges from 0 to 20, higher scores indicate greater adherence to the 2010 Dietary Guidelines for Americans (i.e. higher score = less sugar intake). § $\delta^{13}\text{C}$ value tertiles determined by low (mean -19.60 (SD 0.46) ‰), medium (mean -18.90 (SD 0.15) ‰) and high (mean -18.13 (SD 0.40) ‰) $\delta^{13}\text{C}$ value. ^aLow and medium $\delta^{13}\text{C}$ value tertiles significantly different (HEI-2010; $P \leq 0.01$). ^bMedium and high $\delta^{13}\text{C}$ value tertiles significantly different (SoFAAS; $P \leq 0.01$). ^cLow and high $\delta^{13}\text{C}$ value tertiles significantly different (HEI-2010 and SoFAAS; $P \leq 0.0001$)

participant bias^(17,52). However, gold-standard dietary recall methodology and state-of-the-art nutritional analysis (NDSR) software were used in efforts to mitigate this limitation. Other dietary items (corn, beet sugar, honey, animal protein) are known to have a similar $\delta^{13}\text{C}$ value to AS and SSB⁽¹¹⁾. Several investigations have preliminarily assessed the potential confounds from these items by utilizing a dual-isotope model to explain AS intake using $\delta^{13}\text{C}$ as a predictor and $\delta^{15}\text{N}$ as a covariate, which may increase the biomarker's sensitivity for AS intake^(32,54); however, further research is warranted. Two preliminary validation studies have reported that fasting plasma glucose $\delta^{13}\text{C}$ is not associated with AS or SSB^(31,33); however, postprandial plasma glucose levels demonstrated strong positive correlations with recent AS intake⁽³¹⁾. Collectively, these findings suggest that carbon present in fasting blood glucose reflects acute (e.g. recent meal) and not usual AS consumption, possibly because fasting glucose is primarily derived from non-carbohydrate sources via gluconeogenesis^(11,31,33). However, when our group compared the variances of $\delta^{13}\text{C}$ values between a fasting whole-blood fingerstick sample^(40,55) (n 202; mean -19.19 (SD 0.87) ‰) and a non-fasting whole-blood fingerstick sample⁽²⁸⁾ (n 60; mean -19.94 (SD 0.76) ‰) from two separate cross-sectional studies in adults, no significant differences in the variance of the two groups were found ($F = 1.31$, $P = 0.22$); thus demonstrating the robustness of whole-blood fingerstick $\delta^{13}\text{C}$ value against acute dietary intake. Additional research is needed to determine the optimal substrate and analytical process to assess AS intake using $\delta^{13}\text{C}$ values and the time frame reflected by various approaches.

Conclusions

The present results suggest that the $\delta^{13}\text{C}$ value of fingerstick blood may be useful as an objective indicator of AS intake and overall dietary quality. It may be able to identify individuals at a low *v.* high risk for poor dietary quality, which might be especially useful in large-scale public health studies. Additionally, the present investigation helps to address gaps in the biomarker literature by examining the effects of age, sex and race on the variance of $\delta^{13}\text{C}$ value. However, given the relatively low amount of variance explained by the prediction models, additional studies are needed, with an emphasis on controlled feeding studies. Future directions include assessing changes in associations of $\delta^{13}\text{C}$ value and HEI-2010 over time, determining the ability of the $\delta^{13}\text{C}$ biomarker to reflect long-term SSB and AS consumption habits and assessing the sensitivity to change of the $\delta^{13}\text{C}$ biomarker, comparing various $\delta^{13}\text{C}$ tissue substrates, as well as the inclusion of more ethnic/racially diverse participants and those with lower AS consumption. Also, given the influence of age, age should be controlled for in future research pertaining to $\delta^{13}\text{C}$ value and dietary quality. These preliminary findings contribute to research aimed at constructing a ' $\delta^{13}\text{C}$ fingerstick blood level' guideline that could eventually be used in research and in clinical or community settings to objectively evaluate AS intake, based on the expansion and validation of the association between $\delta^{13}\text{C}$ in human blood and AS intake. In conclusion, since HEI-2010 scores are resource-intensive to calculate and are based upon a subjective measure of dietary intake, fingerstick $\delta^{13}\text{C}$ values



show preliminary promise for use as a minimally invasive, objective measure of AS intake, and an indicator of overall dietary quality and adherence to the Dietary Guidelines for Americans, in community settings.

Acknowledgements

Financial support: This work was supported by the National Institutes of Health (J.M.Z. Principal Investigator, grant number 1R01CA154364-01A1) and the American Heart Association (V.E.H. Principal Investigator, grant number 13POST16560008). The National Institutes of Health and the American Heart Association had no role in the design, analysis or writing of this article. **Conflict of interest:** None. **Authorship:** V.E.H., B.M.D., G.A.W., A.H.J. and J.M.Z. designed the research, wrote the paper and had primary responsibility for final content; V.E.H. and G.A.W. conducted the research; V.E.H. and G.A.W. analysed the data; V.E.H. performed the statistical analysis; A.H.J. provided essential reagents and materials. All authors read and approved the final manuscript. **Ethics of human subject participation:** This study was conducted according to the guidelines laid down in the Declaration of Helsinki and the Virginia Tech Institutional Review Board approved the study protocol. Participants provided written informed consent prior to enrolment.

References

- Elliott SS, Keim NL, Stern JS *et al.* (2002) Fructose, weight gain, and the insulin resistance syndrome. *Am J Clin Nutr* **76**, 911–922.
- Yang Q, Zhang Z, Gregg EW *et al.* (2014) Added sugar intake and cardiovascular diseases mortality among US adults. *JAMA Intern Med* **174**, 516–524.
- Johnson R, Appel L, Brands M *et al.* (2009) Dietary sugars intake and cardiovascular health: a scientific statement from the American Heart Association. *Circulation* **120**, 1011–1020.
- Kit BK, Fakhouri TH, Park S *et al.* (2013) Trends in sugar-sweetened beverage consumption among youth and adults in the United States: 1999–2010. *Am J Clin Nutr* **98**, 180–188.
- Marriott BP, Olsho L, Hadden L *et al.* (2010) Intake of added sugars and selected nutrients in the United States, National Health and Nutrition Examination Survey (NHANES) 2003–2006. *Crit Rev Food Sci Nutr* **50**, 228–258.
- Sievenpiper JL & de Souza RJ (2013) Are sugar-sweetened beverages the whole story? *Am J Clin Nutr* **98**, 261–263.
- Willett WC & Ludwig DS (2013) Science souring on sugar. *BMJ* **346**, e8077.
- Ludwig DS (2013) Examining the health effects of fructose. *JAMA* **310**, 33–34.
- Cottrell RC (2013) Science souring on sugar. *BMJ* **346**, e8077.
- Althuis MD & Weed DL (2013) Evidence mapping: methodologic foundations and application to intervention and observational research on sugar-sweetened beverages and health outcomes. *Am J Clin Nutr* **98**, 755–768.
- Jahren AH, Bostic JN & Davy BM (2014) The potential for a carbon stable isotope biomarker of dietary sugar intake. *J Anal At Spectrom* **29**, 795–816.
- Farley T, Just DR & Wansink B (2012) Clinical decisions. Regulation of sugar-sweetened beverages. *N Engl J Med* **367**, 1464–1466.
- Pomeranz JL (2012) The bittersweet truth about sugar labeling regulations: they are achievable and overdue. *Am J Public Health* **102**, e14–e20.
- Sturm R, Powell LM, Chriqui JF *et al.* (2010) Soda taxes, soft drink consumption, and children's body mass index. *Health Aff (Millwood)* **29**, 1052–1058.
- Marshall TA, Eichenberger Gilmore JM, Broffitt B *et al.* (2003) Relative validation of a beverage frequency questionnaire in children ages 6 months through 5 years using 3-day food and beverage diaries. *J Am Diet Assoc* **103**, 714–720.
- Thomson CA, Giuliano A, Rock CL *et al.* (2003) Measuring dietary change in a diet intervention trial: comparing food frequency questionnaire and dietary recalls. *Am J Epidemiol* **157**, 754–762.
- Willett W & Lenart E (1998) Reproducibility and validity of food-frequency questionnaires. In *Nutritional Epidemiology*, 2nd ed., pp. 101–147 [WC Willett, editor]. New York: Oxford University Press.
- Institute of Medicine of the National Academies (2007) *Dietary Reference Intakes: Research Synthesis Workshop Summary*. Washington, DC: The National Academies Press.
- Hardin DS (2009) Validating dietary intake with biochemical markers. *J Am Diet Assoc* **109**, 1698–1699.
- Hedrick VE, Dietrich AM, Estabrooks PA *et al.* (2012) Dietary biomarkers: advances, limitations and future directions. *Nutr J* **11**, 109.
- McCabe-Sellers B (2010) Advancing the art and science of dietary assessment through technology. *J Am Diet Assoc* **110**, 52–54.
- Thompson FE & Subar AF (2008) Dietary assessment methodology. In *Nutrition in the Prevention and Treatment of Disease*, 2nd ed., pp. 3–39 [A Coulstan and C Boushey, editors]. Oxford: Elsevier, Inc.
- Han E & Powell LM (2013) Consumption patterns of sugar-sweetened beverages in the United States. *J Acad Nutr Diet* **113**, 43–53.
- Thompson FE, McNeel TS, Dowling EC *et al.* (2009) Interrelationships of added sugars intake, socioeconomic status, and race/ethnicity in adults in the United States: National Health Interview Survey, 2005. *J Am Diet Assoc* **109**, 1376–1383.
- Ogden C, Kit BK, Carroll M *et al.* (2011) *Consumption of Sugar Drinks in the United States, 2005–2008*. NCHS Data Brief no. 11. Hyattsville, MD: National Center for Health Statistics.
- Zahnd WE, Scaife SL & Francis ML (2009) Health literacy skills in rural and urban populations. *Am J Health Behav* **33**, 550–557.
- Paasche-Orlow MK, Parker RM, Gazmararian JA *et al.* (2005) The prevalence of limited health literacy. *J Gen Intern Med* **20**, 175–184.
- Davy BM, Jahren AH, Hedrick VE *et al.* (2011) Association of $\delta^{13}\text{C}$ in fingerstick blood with added-sugar and sugar-sweetened beverage intake. *J Am Diet Assoc* **111**, 874–878.
- Kraft RA, Jahren AH & Saudek CD (2008) Clinical-scale investigation of stable isotopes in human blood: $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ from 406 patients at the Johns Hopkins Medical Institutions. *Rapid Commun Mass Spectrom* **22**, 3683–3692.
- Fakhouri TH, Jahren AH, Appel LJ *et al.* (2014) Serum carbon isotope values change in adults in response to changes in sugar-sweetened beverage intake. *J Nutr* **144**, 902–905.
- Cook CM, Alvig AL, Liu YQ *et al.* (2010) The natural ^{13}C abundance of plasma glucose is a useful biomarker of recent dietary caloric sweetener intake. *J Nutr* **140**, 333–337.
- Nash SH, Kristal AR, Bersamin A *et al.* (2013) Carbon and nitrogen stable isotope ratios predict intake of sweeteners in a Yup'ik study population. *J Nutr* **143**, 161–165.



33. Nash SH, Kristal AR, Hopkins SE *et al.* (2014) Stable isotope models of sugar intake using hair, red blood cells, and plasma, but not fasting plasma glucose, predict sugar intake in a Yup'ik study population. *J Nutr* **144**, 75–80.
34. Jahren AH & Kraft RA (2008) Carbon and nitrogen stable isotopes in fast food: signatures of corn and confinement. *Proc Natl Acad Sci U S A* **105**, 17855–17860.
35. Jahren AH, Saudek C, Yeung EH *et al.* (2006) An isotopic method for quantifying sweeteners derived from corn and sugar cane. *Am J Clin Nutr* **84**, 1380–1384.
36. US Department of Agriculture, Economic Research Service (2010) US per capita food availability: caloric sweeteners by subgroup. <http://www.ers.usda.gov/Data/FoodConsumption/app/reports/displayCommodities> (accessed July 2010).
37. Yeung EH, Saudek CD, Jahren AH *et al.* (2010) Evaluation of a novel isotope biomarker for dietary consumption of sweets. *Am J Epidemiol* **172**, 1045–1052.
38. Guenther PM, Casavale KO, Reedy J *et al.* (2013) Update of the healthy eating index: HEI-2010. *J Acad Nutr Diet* **113**, 569–580.
39. US Department of Health and Human Services & US Department of Agriculture (2010) *Dietary Guidelines for Americans*, 7th ed. Washington, DC: US Government Printing Office.
40. Zoellner J, Chen Y, Davy B *et al.* (2014) Talking Health, a pragmatic randomized-controlled health literacy trial targeting sugar-sweetened beverage consumption among adults: rationale, design & methods. *Contemp Clin Trials* **37**, 43–57.
41. Hedrick VE, Comber DL, Estabrooks PA *et al.* (2010) The beverage intake questionnaire: determining initial validity and reliability. *J Am Diet Assoc* **110**, 1227–1232.
42. Hedrick VE, Comber DL, Ferguson KE *et al.* (2013) A rapid beverage intake questionnaire can detect changes in beverage intake. *Eat Behav* **14**, 90–94.
43. Hedrick VE, Savla J, Comber DL *et al.* (2012) Development of a brief questionnaire to assess habitual beverage intake (BEVQ-15): sugar-sweetened beverages and total beverage energy intake. *J Acad Nutr Diet* **112**, 840–849.
44. Riebl SK, Paone AC, Hedrick VE *et al.* (2013) The comparative validity of interactive multimedia questionnaires to paper-administered questionnaires for beverage intake and physical activity: pilot study. *JMIR Res Protoc* **2**, e40.
45. US Department of Health and Human Services, Health Resources and Services Administration (1995) Medically Underserved Areas/Populations: Guidelines for MUA and MUP Designation. <http://www.hrsa.gov/shortage/mua/> (accessed October 2014).
46. Monsen E (2003) *Research: Successful Approaches*, 2nd ed. Chicago, IL: American Dietetic Association.
47. University of Minnesota, Nutrition Data System for Research (2013) *Guide to creating variables needed to calculate scores for each component of the Health Eating Index-2010 (HEI-2010)*. Minneapolis, MN: Regents of the University of Minnesota.
48. Green S (1991) How many subjects does it take to do a regression analysis. *Multivariate Behav Res* **26**, 499–510.
49. US Census Bureau (2010) American FactFinder fact sheet: Lee County, Giles County, Pulaski County, Washington County, Grayson County, Wise County, VA, Virginia, United States. <http://factfinder2.census.gov> (accessed March 2014).
50. Guenther PM, Kirkpatrick SI, Reedy J *et al.* (2014) The Healthy Eating Index-2010 is a valid and reliable measure of diet quality according to the 2010 Dietary Guidelines for Americans. *J Nutr* **144**, 399–407.
51. Popkin BM (2010) Patterns of beverage use across the lifecycle. *Physiol Behav* **100**, 4–9.
52. Kuhnle GG (2012) Nutritional biomarkers for objective dietary assessment. *J Sci Food Agric* **92**, 1145–1149.
53. Weinstein SJ, Vogt TM & Gorrion SA (2004) Healthy Eating Index scores are associated with blood nutrient concentrations in the third National Health And Nutrition Examination Survey. *J Am Diet Assoc* **104**, 576–584.
54. Hedrick V, Zoellner J, Jahren AH *et al.* (2015) A dual-carbon-and-nitrogen stable isotope ratio model is not superior to a single-carbon stable isotope ratio model for predicting added sugar intake in Southwest Virginian adults. *J Nutr* (In the Press).
55. Hedrick V, Davy B, Wilburn G *et al.* (2013) Evaluation of a validated clinical biomarker of added sugar intake compared to the Healthy Eating Index and dietary records in a community-based, rural sample. Poster abstract presented at *The Obesity Society Annual Meeting 'Obesity Week 2013'*, Atlanta, GA, 11–16 November 2013; www.obesityweek.com.