Research in the early 1990s established that increases in prenatal intakes of folate or folic acid early in pregnancy can significantly reduce the risk of neural tube defects (NTDs) in newborns. NTDs are one of the most common birth defects in this country and affect approximately 2500 live births and 1500 terminated pregnancies each year. In an effort to reduce the number of NTDs, the US Food and Drug Administration (FDA) in 1998 implemented the most significant fortification policy since the 1940s, and mandated that all enriched grain products be fortified with folic acid. This policy was expected to add approximately 100 μg of folic acid per day to the diet of the average American. In addition, the percentage of women of childbearing age who consumed at least 400 μg/day of the vitamin was expected to increase from 29% to 50%.

In addition to reducing the risk of NTDs, folic acid fortification has the potential to decrease risks of heart disease and colon cancer. However, excessive intake may mask anemia that results from a vitamin B12 deficiency, which could delay diagnosis and treatment in elderly persons. Masking can facilitate the progression of irreversible neurological manifestations of the deficiency, such as numbness, balance problems, weakness, and paralysis. The FDA currently recommends that adults do not consume more than 1000 μg of folate per day, an amount defined as a “tolerable upper intake level.”

Studies have estimated the effects of the FDA fortification policy at a population level, and showed that the average age-specific total folate intake has increased by approximately 100 μg/day for both adult men and women from the early 1990s to the end of the decade. However, these analyses did not correct for the fact that the dietary intake data represents what the respondents ate in a single day but is not necessarily representative of an average day of the respondents’ dietary intake. Without such correction, it is not possible to accurately estimate the number of women who reach the FDA’s 400 μg/day threshold. In addition, no study has quantified national and population-based intakes by age, gender, and racial/ethnic subgroups. Our analysis provides national, population-based estimates of folate consumption levels by age, gender, and racial/ethnic subgroups and accounts for food and supplement intake, corrected for measurement error because of within-person variation.

**METHODS**

**Data**

We analyzed food and dietary supplement data from 2 periods of the National Health and Nutrition Examination Surveys (NHANES III, 1988–1994; NHANES 1999–2000). NHANES are surveys conducted by the National Center for Health Statistics of the Centers for Disease Control and Prevention and are designed to monitor trends in risk behaviors, environmental exposures, diet, nutrition, and health. Data are collected from personal interviews and physical health exams. Nutrient intake data are based primarily on one 24-hour dietary recall measure from the interview component. Data on supplement use are collected during the physical examination component and entail detailed information on specific types and amounts of supplement use over the previous month. Nutrient intake values are calculated by coding the survey data with the US Department of Agriculture’s Survey Nutrient Database, which incorporates folate values and other nutrients that have been added by fortification.

We used data from NHANES III (1988–1994) to represent prefortification folate intake levels and the 1999–2000 data release of the current NHANES for postfortification estimates. We assessed folate intake levels for men and women aged 15 years and older, to incorporate women of childbearing age as defined by the Centers for Disease Control and Prevention (aged 15–44 years) who may be at risk for a NTD-affected pregnancy.
as well as adults who may be at risk for colon cancer, coronary heart disease, and B_12 deficiency. Age was categorized into 3 groups: 15–44 years, 45–64 years, and 65 years and older. Only subjects who had complete data for both food folate and supplemental folic acid intake levels were included in our analysis, which excluded 19% of subjects for NHANES III and 16% of subjects for NHANES 1999–2000. We included non-Hispanic Whites—hereafter “Whites”—(N=11,332, NHANES III; 2062, NHANES 1999–2000) and non-Hispanic Blacks—hereafter “Blacks”—(N=49,084, NHANES III; 1114, NHANES 1999–2000) as well as Mexican Americans (N=554, NHANES III; 1655, NHANES 1999–2000); other racial/ethnic groups were of insufficient sample size. All analyses used sampling weights to adjust for survey design.

Analysis Overview

We estimated population distributions of total folate consumption (from food and supplements) by grouping intake levels into categories of 100 µg/day. For each age, gender, and race/ethnicity, we estimated intake frequencies and medians, the percentage of the population taking folic acid-containing supplements, and the proportions of the population taking folate at levels above 200 µg/day, 400 µg/day, and 1000 µg/day. All analyses were performed using SAS version 9.1.3 (SAS Institute Inc, Cary, NC) unless otherwise noted.

Dietary Data

Because a 24-hour recall measure for nutrient intake does not represent an individual’s average long-term daily intake, population distribution estimates of dietary folate intake (folate consumption from all food sources, including fortified foods) will be distorted by measurement error. The result is an exaggerated standard deviation caused by overestimating the number of observations in the distribution tails. Such error causes particular problems when estimating the proportions of a population above or below certain thresholds, as was done in our analysis.

We adjusted for measurement error using a subsample of NHANES III subjects who had provided 2 separate 24-hour recalls. There were 1271 observations (7.5%) in the subsample who had complete food folate data for both 1-day measures. Second 24-hour recalls were provided at least 1 week after initial measures for over 96% of the sample, and at least 5 days after initial measure for over 99% of the sample.

To correct for measurement error, we log-transformed all food folate data and conducted visual tests of normality (histograms, straight normal probability plots, and comparable means and medians) to verify the normality assumption. We used results from 1-way analyses of variance to estimate the within-subject and between-subject variances for the log-transformed 2-measure sample. We calculated the “true” between-subject variance by assuming that the ratio of within-subject to between-subject variances is the same for the subjects with only 1 day of data as it is for those with 2 days of data, and to thereby remove the effects of the day-to-day variation in intake. This “true” between-person variance was then used to obtain the corrected food folate intake frequency distributions.

We used gender-, age-, and race-specific within-subject variances for the 2-measure, between-subject variance estimates among Whites and Blacks, and applied gender- and age-specific variances from combined racial/ethnic groups (Whites, Blacks, and Mexican Americans) to the Mexican American subgroup because of small sample sizes of Mexican Americans with two 24 recalls. Because of restricted availability of the 2-measure data for NHANES 1999–2000, we used the same within-person variability estimates from NHANES III to correct for measurement error in the postfortification sample.

Total Folate Data

Participants were asked about their supplement use over 1 month, and therefore, the supplemental folate intake is not subject to measurement error caused by day-to-day variability. To obtain the corrected distributions for total folate intake (food and supplement sources), we created a dataset from food folate observations from the distributions corrected for measurement error. We used a random process to draw from these food intake distributions based on the estimates of subgroup means and standard deviations, and summed each food folate observation with one chosen from the supplemental intake distribution. We assumed no correlation between supplemental and food folate intake on the basis of estimating Spearman rank correlation coefficients of 0.05 or less from the data.

Statistical Inference

To determine the significance of differences in folate intake before the pre- and postfortification time periods, we performed multiple linear regression analyses of the natural log of total daily folate (µg/day) on each time period. Estimates of the statistical significance of pre- to postfortification differences in supplement use were based on logistic regression. Both regression analyses were conducted using SUDAAN statistical software version 9.0 (Research Triangle Institute, Research Triangle Park, NC) and incorporated NHANES sampling weights, strata, and primary sampling units. Age (continuous), gender, and race/ethnicity were included in both models, and all possible 2-way interactions were considered. Manual forward selection was performed to find the best fitting models, and we also assessed the significance and validity of models by race/ethnicity, gender, and age-category subgroups. We used a 2-sample test for binomial proportions to determine the statistical significance of the pre- to postfortification differences in the proportions of the population for which total folate intake levels were more than 200 µg/day, 400 µg/day, and 1000 µg/day.

RESULTS

Measurement Error Correction

After correcting for measurement error, food folate intake distributions had smaller standard deviations both before and after implementation of the fortification policy. Larger percentages of both gender groups consumed higher levels of food folate after fortification than before, and the distribution modes shifted from approximately 200 µg/day to 300 µg/day.

Total Folate Intake

Total folate intake distributions before and after the fortification policy, corrected for measurement error, are shown in Figure 1.
FIGURE 1—Daily total folate intake distributions, pre- and postfortification, for non-Hispanic White men (a), non-Hispanic White women (b), non-Hispanic Black men (c), non-Hispanic Black women (d), Mexican American men (e), and Mexican American women (f), corrected for measurement error. Includes folate consumption from all food and supplement sources, with food data adjusted for measurement error. Points represent midpoints of categories, and plots are smoothed between points. Upper endpoint of 1600 µg/day represents all intake more than 1500 µg/day.

The distributions shifted to the right (higher intake levels) after fortification for all population subgroups. The mean increases were larger for Whites than for Blacks and Mexican Americans. Median total folate intakes for before and after fortification are shown in Figure 2 according to age, gender, and race/ethnicity.
Median folate intake for all Whites increased by more than 100 µg/day after fortification. All women of reproductive age increased their median intake by at least 100 µg/day. Among Blacks and Mexican Americans, the median intake among persons aged 65 years or older either increased by less than 100 µg/day (Black men and women), or decreased (Mexican American men and women) following fortification.

The effect of survey period (i.e., pre- vs postfortification) on the log of total daily folate consumption was significant ($P < 0.0001$) in a multiple linear regression model with control for age, age-squared, gender, race/ethnicity, and significant interactions.

**Threshold Estimates**

Table 1 shows the percentage of women of childbearing age who consumed more than 200 µg/day and 400 µg/day of total folate, and the percentage of older people who consumed more than 1000 µg/day. The within-person variation in food folate intake is shown, as well as the threshold estimates with and without the correction for this variation. Overall, Whites were reaching all thresholds in greater proportions than were Blacks and Mexican Americans; Blacks were least likely to have had intakes above these levels.

The FDA’s goal to have 50% of women of childbearing age consume at least 400 µg/day of folate has not been met for any racial/ethnic group, but the percentages have significantly increased since fortification. At 30% prefortification and 39% postfortification, more White women have reached the 400 µg/day threshold than have Black women (20% pre, 26% post) and Mexican-American women (17% pre, 28% post), both before and after fortification. There were also large increases in the percentage of women of all races/ethnicities who are consuming more than 200 µg/day of total folate: approximately 90% for both Whites and Mexican Americans, and 70% for Blacks. This represents an almost 100% increase for Mexican Americans.

Among people aged 65 years or older who may be at higher risk for B12 deficiency, the proportions of the population who consumed more than 1000 µg/day after fortification increased for White men and women and Black men, decreased for Mexican-American men and women, and remained constant for Black women. Corrected for measurement error, only 1%–4% of persons aged 65 and older were at high risk for masking after fortification.

**Supplemental Folate Intake**

Figure 3 shows the percentage of men and women who take supplements that contain folic acid before and after fortification. More Whites took supplements containing folic acid than did Blacks and Mexican Americans, more people older than 44 years took supplements than did younger people, and more than half of the subgroups showed postfortification
decreases in the proportion of people taking supplements. For all women of childbearing age, fewer people were taking supplements after fortification than before; the same result was found for Mexican American men of all ages. The largest absolute change was among Mexican American men older than 64 years, for whom the percentage taking supplements decreased by more than 50% after fortification. In a logistic regression model, the effect of survey period (i.e., pre- vs postfortification) on consumption of folic acid supplements was found to be significant, and the effect was modified by age (P<.0001) and race/ethnicity (P<.05). The model controls for age, age-squared, gender, race/ethnicity, and significant interactions.

DISCUSSION

We used population-based survey data adjusted for measurement error to examine the effect of the FDA folic acid fortification policy. Although our analysis showed a substantial increase of total folate intake levels after fortification among US adults, we found considerable variations in intake by age, gender, and race/ethnicity. Although all women of childbearing age—the target population for the policy—increased their median total folate intake by at least 100 µg/day following fortification, Black and Mexican American women still lagged behind White women in total consumption and may be target populations for additional fortification or supplementation policies. Among older people who may be at risk for B₁₂ deficiencies, there are higher percentages of Whites and Blacks consuming more than 1000 µg/day of folate (the “tolerable upper intake level”) since fortification. At the same time, higher folate intake levels among these populations may help to reduce risks of heart disease and colon cancer.

The total intake distributions found in our analysis describe the typical American diet: most people consume approximately 200–300 µg/day (pre- to postfortification) of folate from food sources (legumes, leafy greens, fortified cereals, etc.), which is depicted in Figure 1 by the first and larger mode in the distributions. The second, smaller peaks at approximately 600 µg/day represent those people who consume 400 µg/day from supplemental sources in addition to an average diet. Peaks at 1200–1300 µg/day in women aged 15–44 years may represent women who take a supplement (400 µg) in addition to a prenatal vitamin (800 µg); that this peak exists primarily among Whites but not Blacks and Mexican Americans of this age group may indicate that the latter groups are not receiving comparable prenatal care.

Several studies have estimated the changes in folate intake as a result of fortification. Our results corroborate the findings from those studies and show that folate intake levels have increased for the average person by approximately 100 µg/day. However, our analysis also corrects for dietary intake data measurement error when estimating folate intake levels. Because NHANES uses a single 24-hour dietary recall per individual when collecting food and nutrient intake data, estimated distributions of population intakes will have artificially high standard deviations. Although these incorrect values may not affect the estimated means, the error can have important effects on estimates of the proportion of a population above or below certain thresholds.

Our results show that failing to take measurement error into account would have overestimated the percentage of women of childbearing age who reached the 400 µg/day goal set by the US Public Health Service in folate acid fortification. For example, we would have incorrectly concluded that the FDA target had already been met (50% of White women; 52% consuming more than 400 µg/day). At the same time, we would have overestimated the percentage of older people consuming more than 1000 µg/day, and incorrectly concluded that more people may be at risk for masking B₁₂ deficiency than truly is the case.

After correcting for measurement error, we found that the percentage of women of childbearing age now consuming more than 400 µg/day of total folate has increased to between 26% and 38%, but has not yet reached 50% for any racial/ethnic group. Such results are consistent with data that indicate NTD rates have decreased 20%–32% after the policy. Compared with the potential to prevent 50%–70% of NTDs by additional folate intake, By correcting for measurement error among age, racial, and gender subgroups, our analysis is able to more accurately determine the percentages of target and at-risk populations who have reached the thresholds of interest for drawing policy-level conclusions.

One unexpected finding from our analysis is that within all racial/ethnic groups, fewer women of childbearing age are now taking supplements that contain folic acid than were taking them before fortification. This finding indicates a potential need for increased awareness among these populations and their physicians regarding multivitamins. Ray and
colleagues found evidence of suboptimal use of periconceptional folic acid supplements globally among women of childbearing age.\textsuperscript{51}

The substantial racial/ethnic variations in folate consumption found in our analysis are also seen in serum and red blood cell folate levels. Population-level analyses of NHANES surveys indicate that serum and red blood cell folate concentrations for women of childbearing age (both before and after fortification) are lowest among Blacks and highest among Whites; Mexican Americans fall in between those groups.\textsuperscript{39,52} Our results suggest that differences in supplement use are driving the disparities, because food folate intake means and medians are very similar among all three racial/ethnic groups both before and after fortification. Further understanding of the basis for these disparities will provide policymakers more insight into ways to achieve their target: 50% of all women of childbearing age consuming at least 400 µg/day of folate.

The results of our analysis must be considered in light of its limitations. Because of restrictions in data availability, we used within-person variance estimates from NHANES III data to correct for measurement error in NHANES 1999–2000. This may have caused the analysis to underestimate the degree of measurement error correction needed postfortification, because the consumption of fortified grains will increase the within-person variability. Our results may have thereby overestimated the percentages of our target populations that reached the 400 µg/day and 1000 µg/day thresholds.

There is also some evidence that when estimating the true between-person variance from repeat measurements, more accurate results may be derived by including all subjects (those with and without repeat measurements) in the repeated-measures analysis of variance.\textsuperscript{53} We tested the affect of this assumption on the percentages of women of childbearing age who reached the 400 µg/day threshold. We found that the new distributions did have different standard deviations, but there was no pattern among subgroups as to whether the change in methods increased or decreased the distribution spread. The changes in percentage of women of childbearing age who reached the 400 µg/day threshold before and after fortification decreased for Whites and Mexican Americans but increased for Blacks. However, neither the significance of the changes nor the conclusions drawn regarding racial/ethnic variations were affected.

Other factors that may cause imprecise estimates include the use of interview data and uncertainty about the actual amounts of folate added to foods. Possible underreporting biases and underestimation of food folate content\textsuperscript{27,27} could cause our results to underestimate true intake levels, which could be accentuated after fortification as a result of the added effect of fortified foods. Estimating dietary folate intake alone to evaluate the fortification program is also limited in that total folate consumption is not the only factor that influences folate status and its affect on disease, both of which can also be influenced by smoking, body mass index, history of NTD, and variations in individual responses to folate intake.\textsuperscript{38,54,55} Future analyses of the folate biochemical indicator data from

![FIGURE 3—Percentage of population taking supplements that contain folic acid, pre- and postfortification, among men (a) and women (b). The change after fortification was significant and dependent on age ($P < .001$) and race ($P < .05$), after control for age, age-squared, gender, race/ethnicity, and the interaction between gender and race/ethnicity.](image-url)
NHNES III and NHANES 1999–2000, in conjunction with folic acid intake data, may ultimately provide better estimates of the effects of folic acid fortification.52

Given the uncertainty of the minimum effective dose of folate for preventing NTDs, the safety concerns for the elderly, and the magnitude of a national fortification mandate, there is great interest in evaluating folate intake and health outcomes after folic acid fortification.24,30,31,56–58 Although most of the FDA policy’s focus has been on NTDs and masking anemia caused by vitamin B12 deficiency, increased folate consumption is also believed to reduce the risk of coronary heart disease and colon cancer. It would help to better guide policy evaluation if future research could incorporate the existing epidemiological evidence on the relation of folate intake with myocardial infarctions and colon cancer, into an analysis that weighs the risks versus benefits of folic acid fortification on these multiple health outcomes. Such knowledge could contribute to the continuing debate about fortification levels,23,38,46,57–59 as would a better understanding of the dose–response group relationships between folate intake and the progression of these health outcomes.

Since implementation of folic acid fortification in the United States, folate intake among the general population has increased, but there are substantial variations by age, gender, and race/ethnicity. Although the goal of the fortification policy (50% of women of childbearing age) has not been met, the policy has successfully increased the percentage of women aged 14–44 years who consume at least 400 µg/day of folate) has not been met, the policy has successfully increased the percentage of women aged 14–44 years who consume at least 400 µg/day of folate). Targeted interventions to promote use of supplements among women of childbearing age may be needed to further increase these proportions and to further reduce the risk of NTDs in the United States.


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Race and Research Perspectives on Minority Participation in Health Studies

Edited by Bettina Beech, DrPH, MPH, and Maurine Goodman, MA, MPH

Race and Research: Perspectives on Minority Participation in Health Studies is a teaching text and resource guide for students, health professionals, public health researchers, and the general public that extends the discussion of environmental factors that influence ethnic minority participation in health studies. This book examines the lack of minority participation in health studies from social, historical, and scientific perspectives. This book is divided into three main sections: 1) The Meaning of Race, Culture and Ethnicity in Research; 2) Health Studies and Ethnic Minority Populations and 3) The Impact of Revolutionary Changes in Medicine and Health Care on Minority Participation in Health Studies.

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