

RESEARCH ARTICLE

Open Access



Associations between gestational weight gain and rate of infancy weight gain in Hawai'i and Puerto Rico WIC participants

Cheryl L. K. Gibby¹, Cristina Palacios², Maribel Campos³, Eunjung Lim⁴ and Jinan Banna^{1*} 

Abstract

Background: Excessive gestational weight gain and rapid infancy weight gain (RIWG) are associated with increased susceptibility to childhood obesity. Since low-income and minority children are particularly at risk, investigation of the associations between gestational weight gain and rate of infancy weight gain may inform childhood obesity prevention. This study investigated the associations between gestational weight gain and rate of infancy weight gain during the first four to six months postpartum in participants from the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC) in Hawai'i and Puerto Rico.

Methods: This was a cross-sectional secondary data analysis from a text message-based intervention in WIC participants in Hawai'i and Puerto Rico. The analysis included 80 mother/infant pairs from the control group who completed the follow-up visit when infants were four to six months old. Maternal weight, height, and gestational weight gain were self-reported. Infant weight was measured at baseline and follow-up. A proportional odds model was used to investigate the association between gestational weight gain and infancy weight gain rate (rapid or extremely rapid, on-track, or slow), adjusting for maternal age, pregravid body mass index (BMI) status, parity, and being up-to-date with infant vaccinations.

Results: In comparison to recommended gestational weight gain, excessive and inadequate (under the recommended amount) gestational weight gain was associated with 77% decreased (adjusted odds ratio [AOR] = 0.23; 95% confidence interval [CI] = 0.08, 0.70; $p = 0.01$) and 71% decreased (AOR = 0.29; 95% CI = 0.09, 0.94; $p = 0.04$) odds of RIWG versus on-track or slow infant weight gain, respectively. In comparison to women with one child, women with two children (AOR = 0.31; 95% CI = 0.11, 0.87; $p = 0.03$) or three or four children (AOR = 0.24; 95% CI = 0.07, 0.88; $p = 0.03$) had significantly lower odds of RIWG versus on-track or slow infancy weight gain.

Conclusions: Women with excessive or inadequate gestational weight gain had lower proportional odds of RIWG and were more likely to have slower infant weight gain than women who gained the recommended amount of weight.

Trial registration: [ClinicalTrials.gov](https://clinicaltrials.gov/ct2/show/study/NCT02903186) Identifier; [NCT02903186](https://clinicaltrials.gov/ct2/show/study/NCT02903186); September 16, 2016.

Keywords: Gestational weight gain, Rapid infancy weight gain, WIC

* Correspondence: jcbanna@hawaii.edu

¹Department of Human Nutrition, Food and Animal Sciences, College of Tropical Agriculture and Human Resources, University of Hawai'i at Mānoa, Agricultural Sciences 216, 1955 East-West Rd, Honolulu, HI 96822, USA
Full list of author information is available at the end of the article



Background

Childhood overweight and obesity is a global public health concern and has increased from 32 to 41 million children from 1990 to 2016 [1]. In the US, according to the 2011–2012 National Health and Nutrition Examination Survey (NHANES), 7.1% of infants and toddlers under the age of two years were at or above the 97.7th percentile, signifying high weight for length [2]. The 2011–2014 NHANES further indicated that 8.9% of US children aged two to five years and 17% of youth aged two through 19 years were obese [3]. Low-income and minority children are particularly at risk for obesity [4]. In 2014, 40% of US one-year-olds and 30% of two to five-year-olds participating in the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC), a federally funded program for low-income women and children with gross household income at or below 185% of the US Poverty Income Guidelines [5], were overweight or obese [6, 7]. Among minority groups, Hispanic and Black children have increased odds of rapid infancy weight gain (RIWG) [8], and Native Hawaiian or Other Pacific Islander (NHOPi) children have higher weight status between two and four months of age compared to other races [9].

Although infants display the greatest variation in rates of weight gain during the first and second years after birth [10], the first six months are a critical time of rapid growth during which metabolic programming can occur, leading to increased susceptibility to obesity later in life [11–13]. Rapid weight gain during the first two years is associated with higher body mass index (BMI), greater percentage of body fat, and more total fat mass [10], as well as other problems later in life, such as overweight, obesity, high blood pressure, and diabetes [14–17]. Children who displayed catch-up growth, a compensatory mechanism for intrauterine growth restraint [10, 18] or low birth weight [19], during the first two years were heavier, taller, and had more central fat than other children [10]. Similarly, children displaying RIWG (defined as $> +0.67$ change in weight-for-age z-scores) in the first two years had about 7 kg more total body fat and 3 kg greater abdominal adipose tissue at age 30–61 years (mean age 46.5 years) [20]. In Hawai'i, it was shown that a greater change in weight between two and 24 months was associated with higher BMI at age five [9] and that rapid growth from 12 to 23 months was associated with obesity at four to five years in low-income Filipino and NHOPi children [21].

Similarly, maternal weight gain during pregnancy may have significant effects on weight of offspring [22]. Genetic, dietary, or other behavioral factors that increase gestational weight gain may program offspring weight gain by altering the fetal intrauterine environment [22]. Excessive gestational weight gain is associated with increased

neonatal fat mass and body fat percentage, possibly indicating that neonatal adiposity is influenced by maternal fat accumulation and the intrauterine environment during early and mid-pregnancy [23]. One study reported mothers with greater weight gain during pregnancy had children with greater adiposity at age three as measured by BMI and skinfold thickness [22]. Other studies have concluded that excessive gestational weight gain is associated with abnormal weight in offspring, with higher risk of high BMI at ages two and eight [24], obesity at age eight [25], and increased risk of obesity in adolescence and early adulthood [26].

Therefore, excessive maternal gestational weight gain and RIWG are two important factors associated with childhood overweight and obesity. Preventing excessive gestational weight gain, a modifiable maternal risk factor, may allow for prevention of RIWG. Examination of this association may be informative for obesity prevention strategies. A positive association or trend between excessive gestational weight gain and RIWG has been reported in other studies [27–31]; however, no studies have been conducted in Hawai'i or Puerto Rico. To our knowledge, this is the first study to assess the association in low-income women in these locations. We hypothesized that excessive gestational weight gain was positively associated with RIWG in the first four to six months postpartum in this population.

Methods

Participants

We conducted a cross-sectional secondary data analysis using data collected from a four-month text message-based intervention aimed at preventing excessive weight gain in low-income infants at Hawai'i and Puerto Rico WIC clinics [32]. Eligibility criteria for mothers/caregivers included the following: at least 18 years old, able to read, owned a mobile phone with unrestricted texting capabilities, were responsible for caring for the infant, were willing to be randomized and to complete the entire study, and were willing to sign, for themselves and their infant, the written informed consent form which included a description of the study and information such as procedures, voluntary withdrawal, risks, benefits, compensation, and confidentiality. Participants were presented with the informed consent form prior to data collection. Additionally, inclusion criteria required the infant to have been born after 37 weeks of gestation, to be on a normal diet, and to be free from disabilities that hinder movement. Since the main study investigated infant feeding practices, at baseline, infants were required to have been no more than two months old to ensure that they were no more than six months old at the end of the intervention when complementary foods are recommended to be introduced in the diet. Also, infants

must have had birthweight at or between the 10th and 90th percentiles as indicated by the World Health Organization (WHO) growth charts [33], thereby excluding infants who were small for gestational age who could experience compensatory growth. A convenience sample of infants ($n = 202$) and their mother/caregiver were recruited from four WIC clinics in Hawai'i and two WIC clinics in Puerto Rico from January to April, 2016. Participants were assigned to control or intervention groups by block randomization. Detailed methods used in the trial have been published elsewhere [32].

Measures

As described previously [34], the baseline assessment included questionnaires and infant anthropometric measurements. Data collected from the questionnaires included the following: socio-demographic information such as race, mother's age, and infant's gender; pregnancy- and health-related information such as maternal pregravid weight and height, weeks of gestation at delivery, weight gained during pregnancy, and infant vaccinations; food frequency information; and infant feeding practices information such as breastfeeding initiation and duration, reasons for discontinuing breastfeeding, and complementary foods used. Questionnaires and anthropometric measurements were repeated at the follow-up visit four months after the completion of the baseline assessment.

Analyses for the current study included the control group only ($n = 100$). At baseline, gestational weight gain (exposure) was assessed by self-report on the general demographics questionnaire using item, "How much weight did you gain during pregnancy?" Pregravid BMI was calculated using self-reported weight and height. Pregravid BMI was defined based on the National Institutes of Health classifications for BMI (kg/m^2): underweight (<18.5), normal ($18.5\text{--}24.9$), overweight ($25\text{--}29.9$), obese (≥ 30) [35]. Based on these BMI ranges, excessive gestational weight gain was defined as weight gain above the 2009 Institute of Medicine (IOM) gestational weight gain guidelines: underweight women are recommended to gain 12.7–18.2 kg; normal weight women are recommended to gain 11.4–15.9 kg; overweight women are recommended to gain 6.8–11.3 kg; obese women are recommended to gain 5–9.1 kg [36]. Inadequate gestational weight gain was defined as weight gain under the recommended amount.

Infant weight and length were measured in duplicate by trained researchers and WIC employees at baseline and follow-up. The Hawai'i team used Doran infant pan scales (model DS4100) and Easy Glide Bearing Infantometers (Perspective Enterprises, PE-RILB-BRG2), which were provided by the WIC clinics. The Puerto Rico team used Detecto mechanical pediatric scales

(models 450, 451, 459, 459HC) and Seca Infantometer 417. Weight was measured with light or no clothes, no shoes, and a clean diaper. Infant age at follow-up ranged from four to six months. As the preferred system for analysis of anthropometric data [37], z-scores were used for assessment. Z-scores are independent of sex and age, and different age and sex groups may be combined for growth evaluation [38]. Infancy weight gain (main outcome of interest) z-scores are defined as the change in weight-for-age z-scores from baseline to follow-up, calculated using the WHO AnthroPlus online calculator [39], which is based upon WHO reference values [40, 41]. Rapid infancy weight gain is defined as a change in weight gain z-score of $+0.67$ to $+1.28$, representing upwards crossing of at least one percentile line on WHO standard growth charts; extremely rapid infancy weight gain is $> +1.28$, representing upwards crossing of two or more percentile lines on WHO standard growth charts [10, 42–45]. Similarly, change in weight-for-age z-score between -0.67 and 0.67 is defined as on-track infancy weight gain in which there is no crossing of percentile lines, and slow infancy weight gain occurs when weight gain z-score < -0.67 , representing downward crossing of one or more percentile lines [42].

Figure 1 describes inclusion in the analysis group. Eighty-two participants from the control group who completed the follow-up visit and had complete data were included in the final analysis group. Eighteen participants were lost to follow up: Hawai'i ($n = 13$) and Puerto Rico ($n = 5$).

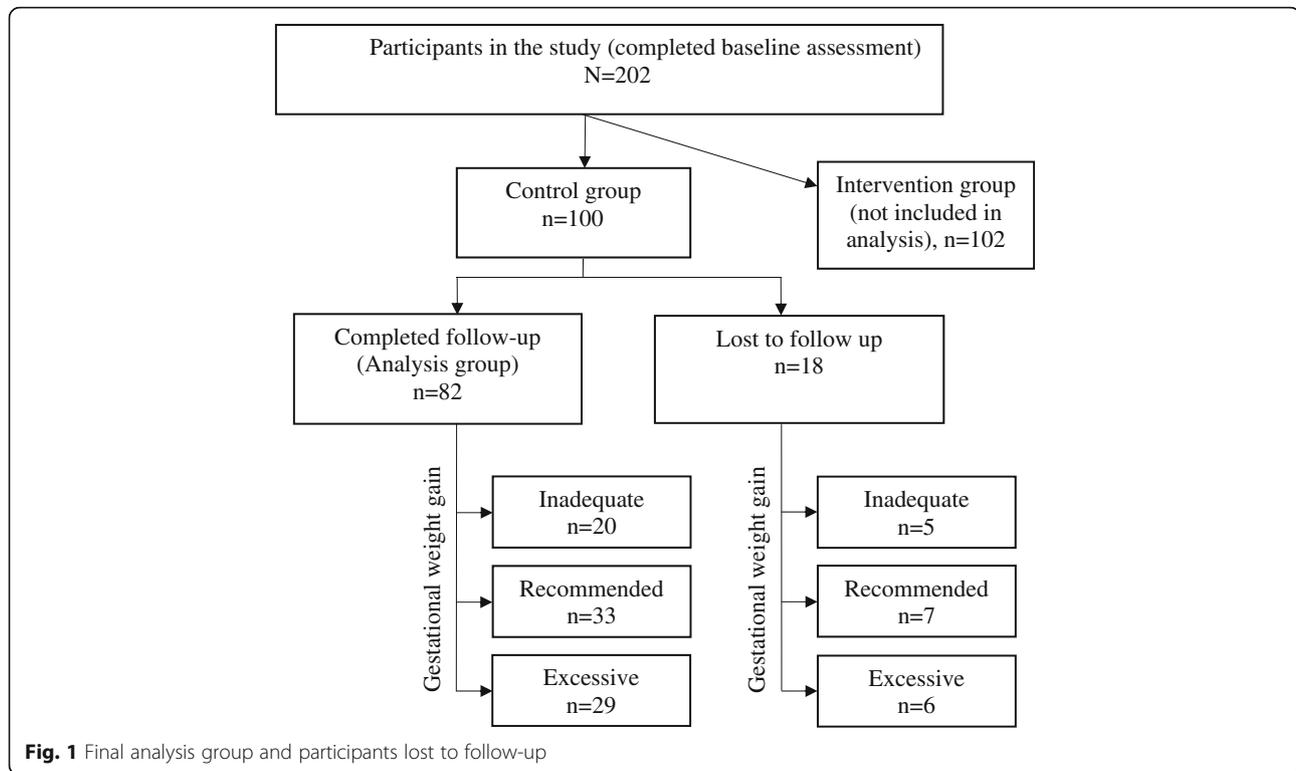
Variables

Dependent variable

Rate of infancy weight gain, determined by the change in weight-for-age z-score, during the first four to six months postpartum was defined as a ternary outcome in mothers from the control group ($n = 100$) who completed the study ($n = 82$). Rapid ($n = 12$) and extremely rapid ($n = 7$) infancy weight gain were combined into one category, simply referred to as rapid infancy weight gain [43, 44], due to the small sample size. In addition, the distributions between rapid and extremely rapid infancy weight gain on the following covariates were comparable to each other.

Independent variable

Maternal gestational weight gain was categorized as "recommended," "inadequate," or "excessive," based on the IOM gestational weight gain guidelines [36]. Excessive gestational weight gain was defined as weight gain above the IOM gestational weight gain guidelines, and inadequate gestational weight gain was defined as weight gain under the recommended amount.



Possible covariates

Variables tested as possible covariates included site (Hawai'i or Puerto Rico), mother's race/ethnicity, mother's age (by years: 18–24, 25–31, 32–39), mother's education level (less than college, some college, college degree or higher), parity (1, 2, 3 or 4), use of prenatal vitamins during pregnancy, pregnancy complications (such as diabetes, hypertension, or anemia), use of vitamins while breastfeeding, weeks of gestation at delivery, pregravid BMI status (underweight, normal, overweight, obese), infant gender, and being up-to-date with infant vaccinations. Based upon the US Office of Management and Budget standards [46], race/ethnicity was categorized into the following groups: Hispanic, NHOPI, American Indian or Alaska Native, Asian, Black or African American, and White. Most of these covariates were selected based upon their inclusion in previous studies that investigated similar associations [31, 47]. However, in regards to the vaccination covariate, it was reported that, in rats, the pertussis vaccine increased glucose-induced insulin secretion [48]. Therefore, since infants routinely receive the pertussis vaccine during the first six months after birth [49], being up-to-date with infant vaccinations was included as a possible covariate.

Statistical analyses

For baseline characteristics, descriptive statistics were presented using frequencies and percentages, or using

means and standard deviations. Chi-squares tests, Fisher's exact tests, and analysis of variance (ANOVA) were conducted to investigate the bivariate associations or comparison between baseline characteristics and rate of infancy weight gain. Fisher's exact test was used to examine the bivariate association between rate of infancy weight gain (rapid, on-track, or slow) and gestational weight gain (recommended, inadequate, or excessive).

To determine the association between gestational weight gain and RIWG, an adjusted proportional odds regression was conducted.

From the bivariate analyses, demographic variables with a p value of < 0.15 were included in the adjusted proportional odds regression [50, 51]. The covariates in the proportional odds model were pregravid BMI status (combined as underweight/normal (UN) or overweight/obese (OWOB)), being up-to-date with infant vaccinations, parity, and maternal age. For all other analyses, including a chi-square score test for the proportional odds assumption to determine goodness of fit, a p -value of < 0.05 was considered statistically significant. Analyses were performed using SAS 9.4 (Cary, NC).

Results

Thirty-seven participants (45.1%) from Hawai'i and 45 participants (54.9%) from Puerto Rico were included in the final analysis ($n = 82$). Characteristics of the control group, final analysis group, and group that was lost to

follow-up are shown in Table 1. Comparing participants in the final analysis group ($n = 82$) with those who did not complete the study ($n = 18$), no statistically significant differences were found in subject characteristics except site, education, and use of prenatal vitamins. Women who were in Hawai'i ($p = 0.04$), who did not take prenatal vitamins during pregnancy ($p = 0.03$), or

who had less than a college education ($p = 0.02$) were more likely to have been lost to follow-up.

Forty-four women (53.7%) were underweight or normal weight prior to pregnancy, and 38 women (46.3%) had overweight or obese pregravid BMI. The majority of women were Hispanic (62.2%) and among all races, White was the most prevalent (48.8%). Fifteen percent of

Table 1 Distribution of select maternal and infant characteristics, n (%)

	Total control ($n = 100$)	Final analysis ($n = 82$)	Lost to follow-up ($n = 18$)	p -value
Site				0.037
Hawai'i	50 (50.0)	37 (45.1)	13 (72.2)	
Puerto Rico	50 (50.0)	45 (54.9)	5 (27.8)	
<i>Maternal factors</i>				
Pregravid BMI (mean [SD])	26.1 [6.9]	26.3 [7.0]	24.8 [6.6]	0.340
Underweight or normal	56 (56.0)	44 (53.7)	12 (66.7)	
Overweight or obese	44 (44.0)	38 (46.3)	6 (33.3)	
Age (mean [SD])	27.0 [5.0]	26.9 [4.7]	27.4 [6.5]	0.958
18–24 years	33 (33.0)	27 (32.9)	6 (33.3)	
25–31 years	47 (47.0)	39 (47.6)	8 (44.4)	
32–39 years	20 (20.0)	16 (19.5)	4 (22.2)	
Race ^a /ethnicity				
Hispanic	60 (60.0)	51 (62.2)	9 (56.3)	0.655
Native Hawaiian or Other Pacific Islander	17 (17.0)	12 (14.6)	5 (27.8)	0.179
Asian	20 (20.0)	14 (17.1)	6 (33.3)	0.118
American Indian or Alaska Native	4 (4.0)	3 (3.7)	1 (5.6)	0.555
Black or African American	12 (12.0)	11 (13.4)	1 (5.6)	0.689
White	45 (45.0)	40 (48.8)	5 (27.8)	0.105
Education				0.019
Less than college	42 (42.0)	31 (37.8)	11 (61.1)	
Some college	22 (22.0)	22 (26.8)	0 (0)	
College degree or higher	36 (36.0)	29 (35.4)	7 (38.9)	
Parity				0.112
1	38 (38.0)	32 (39.0)	6 (33.3)	
2	43 (43.0)	34 (41.5)	9 (50.0)	
3 or 4	19 (19.0)	16 (19.5)	3 (16.7)	
Use of prenatal vitamins	98 (98.0)	82 (100.0)	16 (88.9)	0.031 ^b
Pregnancy complications	41 (41.0)	31 (37.8)	10 (55.6)	0.166
Took vitamins while breastfeeding	53 (53.0)	42 (51.2)	11 (68.8)	0.319
Gestational age (weeks; mean [SD])	38.9 [1.0]	38.9 [1.0]	38.9 [1.1]	0.844
GWG (kg; mean [SD])	12.7 [5.5]	12.6 [5.5]	13.0 [5.9]	0.759
<i>Infant factors</i>				
Male	52 (52.0)	40 (48.8)	12 (66.7)	0.169
Female	48 (48.0)	42 (51.2)	6 (33.3)	
Up-to-date with vaccinations	81 (81.0)	65 (79.3)	16 (88.9)	0.512

Note: Column percentages; p -value was obtained from comparison analysis between final analysis vs lost to follow-up. For categorical variables, chi-square tests or Fisher exact tests were conducted and two-sample t -tests were conducted for continuous variables

^aIncludes all races for mixed participants

mothers were NHOPI. All of the women used prenatal vitamins and most (79.3%) were up-to-date with infant vaccinations.

Mean gestational weight gain was 12.6 kg (SD = 5.5). Thirty-three women (40.2%) gained the recommended amount of weight during pregnancy, 20 women (24.4%) gained inadequate amounts, and 29 (35.4%) women had excessive gestational weight gain. At baseline, all except one infant's weight-for-age z-score fell within -2 to $+2$ on the WHO growth chart, which is the range considered "normal" based upon the statistical definition of the central 95% on a normal curve. [38] During the first four to six months postpartum, 19 infants (23.2%) experienced RIWG, 42 infants (51.2%) experienced on-track weight gain, and 21 infants (25.6%) had slow weight gain.

As shown in Table 2, almost half (48.5%) of women who gained the recommended amount of weight during pregnancy had infants who displayed on-track weight gain that did not cross over percentile lines on the WHO growth charts (change in weight-for-age z-score between -0.67 and 0.67). More than half (58.6%) of women with excessive gestational weight gain had infants who displayed on-track weight gain. Of mothers who gained excessive weight during pregnancy, more infants displayed slow infancy weight gain (31.0%) than RIWG (10.3%). The majority of infants who displayed RIWG (63.2%) came from mothers who gained the recommended amount of weight during pregnancy.

Table 2 also presents the results for the bivariate analysis, which showed the association between rate of infancy weight gain and the following variables: site, race/ethnicity, pregnancy complications, took vitamins while breastfeeding, education, gestational age at birth, and infant gender (none of these associations were statistically significant). Variables with $p < 0.15$ were included in the final adjusted proportional odds cumulative logit model: pregravid BMI ($p = 0.067$), parity ($p = 0.142$), maternal age ($p = 0.047$), and up-to-date with infant vaccinations ($p = 0.061$).

Table 3 presents the results from the final proportional odds model. In the bivariate analysis, Fisher's exact test showed the association between gestational weight gain and the rate of infancy weight gain was not statistically significant ($p = 0.108$). However, in the adjusted model, the association between gestational weight gain and RIWG was statistically significant. In comparison to recommended gestational weight gain, excessive gestational weight gain was associated with 77% decreased odds of RIWG versus on-track or slow infant weight gain (adjusted odds ratio (AOR) = 0.23; 95% CI = 0.08, 0.70; $p = 0.01$). That is, women with excessive gestational weight gain were more likely (4.3 times the odds) to have slower infant weight gain than women who gained the recommended amount of gestational weight.

In comparison to recommended gestational weight gain, inadequate gestational weight gain was associated with 71% decreased odds of RIWG versus on-track or slow infant weight gain (AOR = 0.29; 95% CI = 0.09, 0.94; $p = 0.04$). In other words, women with inadequate gestational weight gain were more likely (3.5 times the odds) to have slower infant weight gain than women who gained the recommended amount of gestational weight.

The adjusted model (Table 3) also indicated that, in comparison to women with one child, women with two children (AOR = 0.31; 95% CI = 0.11, 0.87; $p = 0.026$) or three or four children (AOR = 0.24; 95% CI = 0.07, 0.88; $p = 0.031$) had significantly lower odds of RIWG versus on-track or slow infancy weight gain. Although not statistically significant, women who were up-to-date with infant vaccinations had 68% decreased odds of RIWG versus on-track or slow infancy weight gain (AOR = 0.32; 95% CI = 0.10, 1.03; $p = 0.056$).

The goodness-of-fit test for the proportional odds assumption was used to validate our model; the p -value was 0.64, indicating good fit.

Discussion

In contrast to our hypothesis, excessive gestational weight gain was associated with decreased, not increased, odds of RIWG. The majority of infants with RIWG (63.2%) were from mothers who achieved recommended gestational weight gain. Therefore, preventing excessive gestational weight gain may not be associated with the odds of RIWG. The latter would need to be addressed separately in prevention programs. This finding was in contrast to the results from a number of studies in diverse populations that revealed a positive association between the two [27–31]. For example, a study in China reported excessive gestational weight gain was associated with RIWG across all BMI categories [27]. The difference in results between the studies could have resulted from differences related to geography and/or ethnicity/race, such as diet, environment, or physical stature. For example, Asian adults have higher percentage of body fat than Whites of the same BMI, age, and sex [52]. In the current study, almost half of the participants were White and only 17% were Asian, whereas 100% of the population were Chinese in the aforementioned study. Other studies have reported differences in the relationship between percent body fat and BMI not only among different ethnic groups, but also among ethnicities/races in different geographic regions, such as between American Caucasians and European Caucasians and between Chinese in New York, Beijing, and Hong Kong [53, 54]. Therefore, the results of the current study may be reflective only of the study population, and more research on the factors and mechanisms behind cultural, racial, and geographical differences in the association between gestational weight gain and rate of infancy

Table 2 Results of bivariate analysis by rate of infancy weight gain

Variable	Rate of infancy weight gain, n(%)			p-value
	Rapid 19 (23.2)	On-track 42 (51.2)	Slow 21 (25.6)	
Site				0.615
Hawai'i	7 (18.9)	21 (56.8)	9 (24.3)	
Puerto Rico	12 (26.7)	21 (46.7)	12 (26.7)	
Gestational weight gain				0.108
Recommended	12 (36.4)	16 (48.5)	5 (15.2)	
Inadequate	4 (20.0)	9 (45.0)	7 (35.0)	
Excessive	3 (10.3)	17 (58.6)	9 (31.0)	
Pregravid BMI				0.067
Underweight	2 (28.6)	2 (28.6)	3 (42.9)	
Normal	8 (21.6)	21 (56.8)	8 (21.6)	
Overweight	6 (31.6)	12 (63.2)	1 (5.3)	
Obese	3 (15.8)	7 (36.8)	9 (47.4)	
Race ^a /ethnicity				
Hispanic	12 (23.5)	26 (51.0)	13 (25.5)	0.995
Native Hawaiian	2 (28.6)	2 (28.6)	3 (42.9)	0.441
Other Pacific Islander	3 (50.0)	3 (50.0)	0 (0)	0.161
Native Hawaiian or Other Pacific Islander	4 (33.3)	5 (41.7)	3 (25.0)	0.654
Asian	2 (14.3)	8 (57.1)	4 (28.6)	0.741
American Indian or Alaska Native	1 (33.3)	1 (33.3)	1 (33.3)	0.796
Black or African American	1 (9.1)	7 (63.6)	3 (27.3)	0.580
White	11 (27.5)	20 (50.0)	9 (22.5)	0.622
Parity				0.142
1	12 (37.5)	15 (46.9)	5 (15.6)	
2	5 (14.7)	19 (55.9)	10 (29.4)	
3 or 4	2 (12.5)	8 (50.0)	6 (37.5)	
Use of prenatal vitamins				
Yes	19 (23.2)	42 (51.2)	21 (25.6)	
Pregnancy complications				0.693
No	11 (21.6)	28 (54.9)	12 (23.5)	
Yes	8 (25.8)	14 (45.2)	9 (29.0)	
Took vitamins while breastfeeding				0.921
No	7 (20.6)	17 (50.0)	10 (29.4)	
Yes	11 (26.2)	21 (50.0)	10 (23.8)	
(No response given)	1 (16.7)	4 (66.7)	1 (16.7)	
Education				0.274
Less than college	9 (29.0)	11 (35.5)	11 (35.5)	
Some college	4 (18.2)	13 (59.1)	5 (22.7)	
College degree or higher	6 (20.7)	18 (62.1)	5 (17.2)	
Maternal age (years)				0.047
18–24	11 (40.7)	11 (40.7)	5 (18.5)	
25–31	5 (12.8)	20 (51.3)	14 (35.9)	
32–39	3 (18.8)	11 (68.8)	2 (12.5)	

Table 2 Results of bivariate analysis by rate of infancy weight gain (*Continued*)

Variable	Rate of infancy weight gain, n(%)			p-value
	Rapid 19 (23.2)	On-track 42 (51.2)	Slow 21 (25.6)	
Infant gender				0.257
Male	10 (25.0)	23 (57.5)	7 (17.5)	
Female	9 (21.4)	19 (45.2)	14 (33.3)	
Up-to-date with vaccinations				0.061
No	6 (35.3)	10 (58.8)	1 (5.9)	
Yes	13 (20.0)	32 (49.2)	20 (30.8)	

Note: Row percentages; p-value was obtained from chi-square test or Fisher's exact test for categorical variable and analysis of variance for continuous variable

^aIncludes all races for mixed participants. Each race/ethnicity variable was defined as Yes if a participant marked the race/ethnicity on the questionnaire

weight gain may lead to public health strategies that are more effective and better tailored to meet the needs of different populations.

In comparison to recommended gestational weight gain, both excessive gestational weight gain and inadequate gestational weight gain, separately, were associated with higher odds of slow infancy weight gain versus RIWG or on-track infancy weight gain. Another study of normal birthweight infants reported that slow

infancy weight gain between ages eight weeks and nine months was followed by RIWG; however, children did not return to mean reference weights until age 13 years and they remained lighter and shorter than their peers throughout childhood [55]. Since the current study followed infant weight gain only up until six months of age, future studies should observe rates of weight gain beyond this period to see if similar patterns hold true in this population.

Table 3 Proportional odds model using rate of infancy weight gain as the dependent variable

Independent variable	Estimate (SE)	p-value	AOR (95% CI)	
			Rapid vs on-track or slow	Slow vs rapid or on-track
Intercept				
Rapid vs on-track or slow	1.16 (0.69)	0.093		
Rapid or on-track vs slow	4.00 (0.83)	<0.001		
Gestational weight gain				
Recommended	Reference			
Excessive	-1.46 (0.57)	0.010	0.23 (0.08, 0.70)	4.32 (1.42, 13.14)
Inadequate	-1.24 (0.60)	0.039	0.29 (0.09, 0.94)	3.45 (1.07, 11.17)
Maternal age (years)				
18-24	Reference			
25-31	-0.60 (0.57)	0.289	0.55 (0.18, 1.67)	1.83 (0.60, 5.54)
32-39	0.02 (0.67)	0.975	1.02 (0.28, 3.77)	0.98 (0.27, 3.61)
Pregravid BMI				
Underweight/normal	Reference			
Overweight/obese	0.30 (0.50)	0.555	1.35 (0.50, 3.61)	0.73 (0.28, 1.99)
Parity				
1 child	Reference			
2 children	-1.16 (0.52)	0.026	0.31 (0.11, 0.87)	3.20 (1.15, 8.93)
3 or 4 children	-1.43 (0.66)	0.031	0.24 (0.07, 0.88)	4.17 (1.14, 15.28)
Up-to-date with infant vaccinations				
No	Reference			
Yes	-1.15 (0.60)	0.056	0.32 (0.10, 1.03)	3.15 (0.97, 10.21)

Estimate: Ordered log-odds regression coefficient, SE Standard error of the estimate, AOR adjusted odds ratio, CI confidence interval

The current study found that 25.6% of infants experienced slow infancy weight gain. Nine infants with slow infancy weight gain (including four infants from mothers with excessive gestational weight gain and three infants from mothers with inadequate gestational weight gain) fell into the category of “failure to thrive” (FTT), defined as a child under two years whose weight crosses two major percentiles downward on a standardized growth chart [56]. Like RIWG, slow infancy weight gain may be associated with adverse health conditions and must be addressed to prevent unfavorable outcomes later in life. Slow infancy weight gain, especially when severe enough to be considered FTT, may be a sign of feeding problems, malnutrition, or of other physiological conditions that lead to inadequate calorie intake, inadequate calorie absorption, or increased calorie requirements [56]. FTT is associated with decreased developmental skills, lower height and weight, lower verbal intelligence, lower social maturity, and increased behavioral disturbances [56, 57]. Therefore, excessive or inadequate gestational weight gain may be an important risk factor for developing adverse health conditions related to FTT, and are important to address in pregnant women.

In comparison to first-time mothers with only one child, multiparous women with two to four children had approximately 3–4 times the odds of slow infancy weight gain versus RIWG or on-track infancy weight gain. First-time mothers had higher odds of rapid infancy weight gain. Likewise, a study conducted in the United Kingdom reported infants of first-time mothers displayed RIWG during the first year and, thereafter, were heavier and taller than infants from multiparous mothers [58]. A similar finding was reported in primiparous mothers in Denmark whose infants gained more weight during the first year than infants of multiparous women [59]. Different parenting behaviors between first-time mothers and multiparous mothers may influence infant nutrition [58]. A study in Australia reported that first-time mothers failed to demonstrate an understanding of the rationale supporting the WHO guidelines to exclusively breastfeed for the first six months, nor did they comprehend well the signs of readiness for solid foods [60]. Formula feeding and feeding-to-schedule may lead to overfeeding and are associated with RIWG [61]. Likewise, early introduction of solid foods along with shorter duration of breastfeeding may partially explain greater infant weight gain [59]. Therefore, women may benefit from nutrition and infant care education during their first pregnancies to prevent RIWG and during subsequent pregnancies to prevent slow infancy weight gain and FTT.

Many studies have shown that excessive gestational weight gain is associated with increased risk for obesity [24–26, 62]. This association has public health implications, as excessive gestational weight gain is not

uncommon in the US. In 2015, 48% of uniparous women who gave birth to a full-term infant in the US had excessive gestational weight gain [63]. The current study found that more than one-third (35.4%) of the women in the final analysis gained excessive gestational weight, making it an important risk factor to address in this population.

Along with epidemiological evidence, the physiological mechanisms driving excessive gestational weight gain to metabolically program offspring weight gain towards overweight or obesity [62] are important to consider in obesity prevention strategies. Potential mechanisms involved in metabolic imprinting through excessive gestational weight gain include the following: 1) higher levels of maternal circulating insulin during the third trimester [64] leads to increased levels of adipocyte development and fat storage in the fetus, 2) excessive maternal adipose tissue accumulation increases low-grade chronic inflammation and circulating adipokines, possibly creating an inflammatory environment for the fetus and promoting excess body fat accumulation [65], 3) fetal metabolic programming is influenced by epigenetic changes brought on by maternal diet [66, 67]. Based upon these mechanisms, one prevention strategy may involve providing at-risk women with vigorous nutrition/diet counseling and accountability along with frequent blood glucose monitoring, if applicable. Although mainly a measure of energy balance, weight-for-age has been used most often as the metric in studies on infancy growth and later obesity [68]. However, a study that examined the differences in association between adult overweight/obesity and infant weight-for-age versus weight-for-length measurements suggested that weight-for-length during the first two years is a more comprehensive metric for infant growth and provides a stronger, more consistent association with overweight/obesity in young adulthood [69]. However, using weight-for-length versus weight-for-age measurements, the study reported similar odds ratios (OR) for being overweight at age 20–29 years based on rapid growth ($\geq +0.67$ z-score) compared to non-rapid growth from birth to six months (OR 1.67 vs. 1.69) and from birth to three months (OR 1.13 vs. 1.07) [69]. Therefore, since the current study only examined infants up to six months of age, and considering the preponderance of previous studies using weight-for-age, weight-for-age was chosen as the metric. However, future studies that look at growth beyond six months may benefit from using weight-for-length measurements.

Weight gain during infancy plays an important role in overall growth and disease risk throughout life [10, 55]. Considering the results presented, weight gain in excess of or below the recommended amounts set forth by the 2009 IOM guidelines may lead to adverse health outcomes related to slow infancy weight gain and subsequent growth patterns. Preventing excessive or inadequate gestational weight gain may work simultaneously to prevent slow

infancy weight gain in this population of low-income WIC participants. These findings may be helpful for health promotion and obesity prevention strategies and programs for women in WIC in Hawai'i and Puerto Rico.

Limitations of the study

Our study has several limitations. First, although the majority of participants in the final analysis group was Hispanic, making 77% of the final analysis group Hispanic or partly NHOPI, the sample size was small and may have caused bias in the results, such as under or overestimation of the associations. Therefore, larger studies, especially with a greater percentage of NHOPI participants, should be conducted to further investigate the associations described in this study. Second, our results are not generalizable to women beyond those participating in the study due to the inherent bias of convenience sampling. However, this study provides new information that may inform future studies. Third, although 95% of American adults in 2018 own cell phones [70] and 73% of cell phone owners use text messaging [71], according to the Pew Research Center, it is possible that the participants in the current study were at a slightly higher income level than the average WIC population since one of the eligibility criteria was that participants owned a mobile phone. However, household income information was not collected during the main study and our analysis could not include stratification within the WIC low-income bracket. Fourth, maternal height, pregravid weight, and gestational weight gain were self-reported. Fifth, although researchers were trained in proper techniques and documentation of measurements, having multiple researchers taking anthropometric measurements may have introduced human error. Additionally, 11 participants from Hawai'i (29.7%) had infant anthropometric measurements taken in-part or in-full by WIC staff, whose training differed from the research team. All measurements in Puerto Rico were taken by the research team.

Lastly, it is possible that other factors that were not considered in the current analysis mediated the association between excessive gestational weight gain and RIWG. It was recently reported that, as mediators, breastfeeding duration and birth weight accounted for up to 76% of the total effect of gestational weight gain on weight-for-age z-scores [47]. The current study investigated the change in weight-for-age z-scores as indicative of infant weight gain, rather than single point measurements, but it is notable that these variables could have mediated the current observations. It is also possible that the associations observed in the current study may have changed had it been conducted over a longer period. In the aforementioned study, it was reported that the direct effects of gestational weight gain on weight-for-age z-scores and BMI z-scores were not significant until children were six years old [47].

Therefore, since birth weight and breastfeeding duration may be important mediators between gestational weight gain and anthropometric measures during infancy and early childhood [47], it would be prudent to include these variables in future analyses and studies.

Conclusions

In low-income WIC participants in Hawai'i and Puerto Rico, excessive gestational weight gain did not increase susceptibility to RIWG; however, both variables are risk factors for childhood obesity and collectively affected more than half of the women/infants in this sample. Preventing excessive and inadequate gestational weight gain may benefit infant health by preventing slow infancy weight gain and FTT. Achieving the recommended amount of gestational weight gain and keeping an infant's weight gain on track are important to prevent malnutrition and undesirable metabolic programming. The results of this study should be interpreted within the context of the limitations noted but may provide direction for the development of strategies for obesity prevention and gestational weight gain management.

Abbreviations

ANOVA: Analysis of variance; AOR: Adjusted odds ratio; BMI: Body mass index; CI: Confidence interval; FTT: Failure to thrive; IOM: Institute of Medicine; NHANES: National Health and Nutrition Examination Survey; NHOPI: Native Hawaiian or Other Pacific Islander; OWOB: Overweight/obese; RIWG: Rapid infancy weight gain; UN: Underweight/normal; WHO: World Health Organization; WIC: Special Supplemental Nutrition Program for Women, Infants, and Children

Acknowledgments

The authors acknowledge Linda Chock (Branch Chief), Iris Takahashi (Clinic Operations Section Chief), Laura Morihara (Wahiawa WIC Coordinator), Wendy Baker (Wahiawa WIC Office Assistant), Jean Kanda (Leeward WIC Coordinator), Dawn Fujimoto-Redoble (Leeward WIC Nutritionist), and Christina Mariano (Pearl City WIC Coordinator) from the Hawai'i Department of Health WIC Services Branch. They also acknowledge the Puerto Rico WIC Program, including Dana Miró (Executive Director), Blanca Sastre (Interim supervisor of Nutrition and Lactation Division), Alexandra Reyes (Nutrition Education Coordinator), Iris Roldan (Breastfeeding Coordinator), Ivelisse Bruno (Breastfeeding Peer Support Coordinator), Sherley M. Panell (Nutrition Supervisor in Plaza las Americas WIC Clinic) and her team, and Marta Meaux (Nutrition Supervisor in Trujillo Alto WIC Clinic) and her team. They also thank the WIC participants and undergraduate and graduate research assistants.

Funding

The current study was supported by the National Institute of Minority Health and Health Disparities (NIMHD) of the National Institutes of Health under award number U54MD008149 and in part by grant U54MD007584 (RMATRIX) and U54MD007601 (Ola Hawaii). Infrastructure support was also provided in part by the NIMHD RCMI Grant: 8G12MD007600.

Availability of data and materials

Data used for the current study are available from the corresponding author.

Authors' contributions

CG and JB developed the research objectives and plan. JB and CP oversaw the research teams in Hawai'i and Puerto Rico, respectively. EL and CG developed the statistical analysis plan and analyzed the data. CP, MC, and JB provided guidance for data analysis and interpretation. CG and JB wrote the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Study procedures were approved by the institutional review boards at the University of Hawai'i at Mānoa and the University of Puerto Rico, Medical Sciences Campus, and written informed consent was obtained prior to data collection.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interest.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Department of Human Nutrition, Food and Animal Sciences, College of Tropical Agriculture and Human Resources, University of Hawai'i at Mānoa, Agricultural Sciences 216, 1955 East-West Rd, Honolulu, HI 96822, USA.

²Department of Dietetics and Nutrition, Robert Stempel College of Public Health & Social Work, Florida International University, 11200 SW 8th Street, AHC 5-313, Miami, FL 33199, USA. ³Dental and Craniofacial Genomics Unit, Endowed Health Services Research Center, Medical Sciences Campus, University of Puerto Rico, PO Box 365067, San Juan, PR 00936-5067, USA.

⁴Department of Complementary and Integrative Medicine, Biostatistics Core Facility, John A. Burns School of Medicine, University of Hawai'i at Mānoa, Medical Education Building 411, 651 Ilalo Street, Honolulu, HI 96813, USA.

Received: 7 June 2018 Accepted: 19 November 2018

Published online: 03 December 2018

References

- WHO Facts and figures on childhood obesity. WHO. 2014. <http://www.who.int/end-childhood-obesity/facts/en/>. Accessed 25 Oct 2016.
- Ogden C, Carroll M, Kit B, Flegal K. Prevalence of childhood and adult obesity in the United States, 2011–2012. *JAMA*. 2014;311:806–14.
- Ogden C, Carroll M, Fryer C, Flegal K. Prevalence of obesity among adults and youth: United States, 2011–2014. 2015. <https://www.cdc.gov/nchs/data/databriefs/db219.pdf>. Accessed 10 Aug 2017.
- Pan L, Blanck HM, Sherry B, Dalenius K, Grummer-Strawn LM. Trends in the prevalence of extreme obesity among US preschool-aged children living in low-income families, 1998–2010. *JAMA*. 2012;308:2563–5.
- United States Department of Agriculture. WIC Income Eligibility Guidelines Food and Nutrition Service 2017. <https://www.fns.usda.gov/wic/frequently-asked-questions-about-wic#1>. Accessed 24 Oct 2017.
- Thorn B, Tadler C, Huret N, Ayo E, Trippe C. WIC participant and program characteristics final report. 2015. <http://www.fns.usda.gov/sites/default/files/ops/WICPC2014.pdf>. Accessed 23 Oct 2016.
- Segal L, Rayburn J, Martin A. Childhood obesity trends: the state of obesity. The State of Obesity <http://stateofobesity.org/childhood-obesity-trends/>. Accessed 23 Oct 2016.
- Taveras E, Gillman M, Kleinman K, Rich-Edwards J, Rifas-Shiman S. Racial/ethnic differences in early-life risk factors for childhood obesity. *Pediatrics*. 2010;125:686–95.
- Oshiro C, Novotny R, Grove J, Hurwitz E. Race/ethnic differences in birth size, infant growth, and body mass index at age five years in children in Hawaii. *Child Obes*. 2015;11:683–90.
- Ong KKL, Ahmed ML, Emmett PM, Preece MA, Dunger DB. Association between postnatal catch-up growth and obesity in childhood: prospective cohort study. *BMJ*. 2000;320:967–71.
- Young B, Johnson S, Krebs N. Biological determinants linking infant weight gain and child obesity: current knowledge and future directions. *Adv Nutr*. 2012;3:675–86.
- Singhal A, Lanigan J. Breastfeeding, early growth and later obesity. *Obes Rev*. 2007;8:51–4.
- Singhal A, Lucas A. Early origins of cardiovascular disease: is there a unifying hypothesis? *Lancet*. 2004;363:1642–5.
- Monteiro PO, Victora CG. Rapid growth in infancy and childhood and obesity in later life—a systematic review. *Obes Rev*. 2005;6:143–54.
- Baird J, Fisher D, Lucas P, Kleijnen J, Roberts H, Law C. Being big or growing fast: systematic review of size and growth in infancy and later obesity. *BMJ*. 2005;331:929.
- Gardner DSL, Hosking J, Metcalf BS, Jeffery AN, Voss LD, Wilkin TJ. Contribution of early weight gain to childhood overweight and metabolic health: a longitudinal study (EarlyBird 36). *Pediatrics*. 2009;123:e67–73.
- Gillman MW. Early infancy - a critical period for development of obesity. *J Dev Orig Health Dis*. 2010;1:292–9.
- Ekelund U, Ong K, Linné Y, Neovius M, Brage S, Dunger DB, et al. Upward weight percentile crossing in infancy and early childhood independently predicts fat mass in young adults: the Stockholm weight development study (SWEDES). *Am J Clin Nutr*. 2006;83:324–30.
- Dewey KG. Cross-cultural patterns of growth and nutritional status of breast-fed infants. *Am J Clin Nutr*. 1998;67:10–7.
- Demerath EW, Reed D, Choh AC, Soloway L, Lee M, Czerwinski SA, et al. Rapid postnatal weight gain and visceral adiposity in adulthood: the Fels longitudinal study. *Obesity*. 2009;17:2060–6.
- Okhiro M, Davis J, White L, Derauf C. Rapid growth from 12 to 23 months of life predicts obesity in a population of Pacific Island children. *Ethn Dis*. 2012;22:439–44.
- Oken E, Taveras EM, Kleinman KP, Rich-Edwards JW, Gillman MW. Gestational weight gain and child adiposity at age 3 years. *Am J Obstet Gynecol*. 2007;196:322.e1–8.
- Starling AP, Brinton JT, Glueck DH, Shapiro AL, Harrod CS, Lynch AM, et al. Associations of maternal BMI and gestational weight gain with neonatal adiposity in the healthy start study. *Am J Clin Nutr*. 2015;101:302–9.
- Mourtakos SP, Tambalis KD, Panagiortakos DB, et al. Association between gestational weight gain and risk of obesity in preadolescence: a longitudinal study (1997–2007) of 5,125 children in Greece. *J Hum Nutr Diet*. 2017;30:51–8. <https://doi.org/10.1111/jhn.12398>
- Mourtakos SP, Tambalis KD, Panagiortakos DB, Antonogeorgos G, Arnaoutis G, Karaterliotis K, et al. Maternal lifestyle characteristics during pregnancy, and the risk of obesity in the offspring: a study of 5,125 children. *BMC Pregnancy Childbirth*. 2015;15:66.
- Rooney BL, Mathiason MA, Schaubberger CW. Predictors of obesity in childhood, adolescence, and adulthood in a birth cohort. *Matern Child Health J*. 2011;15:1166–75.
- Li N, Liu E, Guo J, Pan L, Li B, Wang P, et al. Maternal prepregnancy body mass index and gestational weight gain on offspring overweight in early infancy. *PLoS One*. 2013;8. <https://doi.org/10.1371/journal.pone.0077809>.
- Subhan FB, Colman I, McCargar L, Bell RC. Team TAPs. Higher pre-pregnancy BMI and excessive gestational weight gain are risk factors for rapid weight gain in infants. *Matern Child Health J*. 2017;21:1396–407.
- Heerman WJ, Bian A, Shintani A, Barkin SL. The interaction between maternal pre-pregnancy BMI and gestational weight gain shapes infant growth. *Acad Pediatr*. 2014;14:463–70.
- Deierlein AL, Siega-Riz AM, Adair LS, Herring AH. Effects of prepregnancy body mass index and gestational weight gain on infant anthropometric outcomes. *J Pediatr*. 2011;158:221.
- Deierlein AL, Siega-Riz AM, Herring AH, Adair LS, Daniels JL. Gestational weight gain and predicted changes in offspring anthropometrics between early infancy and 3 years. *Pediatr Obes*. 2012;7:134–42.
- Banna J, Campos M, Gibby C, Graulau RE, Meléndez M, Reyes A, et al. Multi-site trial using short mobile messages (SMS) to improve infant weight in low-income minorities: Development, implementation, lessons learned and future applications. *Contemp Clin Trials*. 2017;62:56–60.
- Centers for Disease Control and Prevention. Growth Charts WHO Child Growth Standards. National Center for Health Statistics. http://www.cdc.gov/growthcharts/who_charts.htm#The%20WHO%20Growth%20Charts. Accessed 9 Nov 2016.
- Gibby CLK, Palacios C, Campos M, Lim E, Banna J. Breastfeeding discontinuation not associated with maternal pregravid BMI but associated with Native Hawaiian or Other Pacific Islander race in Hawaii and Puerto Rico WIC participants. *Matern Child Health J*. 2018. <https://doi.org/10.1007/s10995-018-2587-x>
- Losing Weight, Body Mass Index. http://www.nhlbi.nih.gov/health/educational/lose_wt/BMI/bmi_dis.htm. Accessed 26 Oct 2016.
- Institute of Medicine, National Research Council Committee to Reexamine IOM Pregnancy Weight Guidelines. *Weight Gain During Pregnancy: Reexamining the Guidelines*. Washington: National Academies Press; 2009. <https://www.ncbi.nlm.nih.gov/books/NBK32815/>
- World Health Organization. *Physical status: the use and interpretation of anthropometry*. Report of a WHO Expert Committee. Geneva: WHO

- Technical Report Series; 1995. http://www.who.int/childgrowth/publications/physical_status/en/. Accessed 27 Oct 2016.
38. World Health Organization. The Z-score or standard deviation classification system. Global Database on Child Growth and Malnutrition. <http://www.who.int/nutgrowthdb/about/introduction/en/index4.html>. Accessed 27 Oct 2016.
 39. World Health Organization. WHO Application tools. WHO Application Tools. <http://www.who.int/growthref/tools/en/>. Accessed 20 Dec 2016.
 40. World Health Organization. WHO Z-scores for girls. http://www.who.int/childgrowth/standards/chts_girls_z.pdf?ua=1. Accessed 27 Oct 2016.
 41. World Health Organization. WHO Z-scores for boys. http://www.who.int/childgrowth/standards/chts_boys_z.pdf?ua=1. Accessed 27 Oct 2016.
 42. Wang G, Johnson S, Gong Y, Polk S, Divall S, Radovick S, et al. Weight gain in infancy and overweight or obesity in childhood across the gestational spectrum: a prospective birth cohort study. *Sci Rep*. 2016;6:29867.
 43. Ong KK, Loos RJF. Rapid infancy weight gain and subsequent obesity: systematic reviews and hopeful suggestions. *Acta Paediatr*. 2006;95:904–8.
 44. Zhou J, Dang S, Zeng L, Gao W, Wang D, Li Q, et al. Rapid infancy weight gain and 7- to 9-year childhood obesity risk: A prospective cohort study in rural Western China. *Medicine (Baltimore)*. 2016;95:e3425.
 45. Wang Y, Hsin Jen C. Chapter 2, Use of Percentiles and Z-Scores in Anthropometry. In: Preedy V, ed. *Handbook of Anthropometry: Physical Measures of Human Form in Health and Disease*. New York: Springer; 2012. p. 29–48.
 46. US Census Bureau. Race. <https://www.census.gov/topics/population/race/about.html>. Accessed 1 Aug 2017.
 47. Liu JX, Xu X, Liu JH, et al. Association of maternal gestational weight gain with their offspring's anthropometric outcomes at late infancy and 6 years old: mediating roles of birth weight and breastfeeding duration. *Int J Obes* (2005). 2018;42:8–14. <https://doi.org/10.1038/ijo.2017.183>
 48. Smith M, Furman BL. Augmentation of glucose induced insulin secretion by pertussis vaccine, phentolamine and benextramine: involvement of mechanisms additional to prevention of the inhibitory actions of catecholamines in rats. *Acta Endocrinol (Copenh)*. 1988;118:89–95.
 49. Centers for Disease Control and Prevention. Birth-18 Years Immunization Schedule CDC. 2018. <https://www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html>. Accessed 8 Aug 2018.
 50. Yen TE, Allen JC, Rivelli SK, Patterson SC, Metcalf MR, Flink BJ, et al. Association between serum IGF-I levels and postoperative delirium in elderly subjects undergoing elective knee arthroplasty. *Sci Rep*. 2016;6:20736.
 51. Afifi A, May S, Clark VA. *Practical Multivariate Analysis*. 5th ed. New York: Chapman and Hall; 2011.
 52. WHO expert consultation. Appropriate body-mass index for Asian populations and its implications for policy and intervention strategies. *The Lancet*. 2004;363:157–63.
 53. Deurenberg P, Deurenberg-Yap M, Guricci S. Asians are different from Caucasians and from each other in their body mass index/body fat percent relationship. *Obes Rev*. 2002;3:141–6.
 54. Deurenberg P, Yap M, van Staveren WA. Body mass index and percent body fat: a meta analysis among different ethnic groups. *Int J Obes Relat Metab Disord*. 1998;22:1164–71.
 55. ud DZ, Emmett P, Steer C, Emond A. Growth outcomes of weight faltering in infancy in ALSPAC. *Pediatrics*. 2013;131:e843–9.
 56. Zenel JA. Failure to thrive: a general pediatrician's perspective. *Pediatr Rev*. 1997;18:371–8.
 57. Oates RK, Peacock A, Forrest D. Long-term effects of nonorganic failure to thrive. *Pediatrics*. 1985;75:36–40.
 58. Ong KKL, Preece MA, Emmett PM, Ahmed ML, Dunger DB. ALSPAC study team. Size at birth and early childhood growth in relation to maternal smoking, parity and infant breast-feeding: longitudinal birth cohort study and analysis. *Pediatr Res*. 2002;52:863–7.
 59. Baker JL, Michaelsen KF, Rasmussen KM, Sørensen TI. Maternal prepregnant body mass index, duration of breastfeeding, and timing of complementary food introduction are associated with infant weight gain. *Am J Clin Nutr*. 2004;80:1579–88.
 60. Walsh A, Kearney L, Dennis N. Factors influencing first-time mothers' introduction of complementary foods: a qualitative exploration. *BMC Public Health*. 2015;15. <https://doi.org/10.1186/s12889-015-2250-z>.
 61. Mhrshahi S, Battistutta D, Magarey A, Daniels LA. Determinants of rapid weight gain during infancy: baseline results from the NOURISH randomised controlled trial. *BMC Pediatr*. 2011;11:99.
 62. Hillier TA, Pedula KL, Vesco KK, Oshiro CES, Ogasawara KK. Impact of maternal glucose and gestational weight gain on child obesity over the first decade of life in normal birth weight infants. *Matern Child Health J*. 2016;20:1559–68.
 63. QuickStats. Gestational weight gain among women with full-term, singleton births, compared with recommendations - 48 states and the District of Columbia, 2015. *Morb Mortal Wkly Rep*. 2016;65:1121.
 64. Perichart-Perera O, Muñoz-Manrique C, Reyes-López A, Tolentino-Dolores M, Sosa SEY, Ramírez-González MC. Metabolic markers during pregnancy and their association with maternal and newborn weight status. *PLoS One*. 2017;12. <https://doi.org/10.1371/journal.pone.0180874>.
 65. Segovia SA, Vickers MH, Gray C, Reynolds CM. Maternal obesity, inflammation, and developmental programming. *Biomed Res Int*. 2014;2014. <https://doi.org/10.1155/2014/418975>.
 66. Chango A, Pogribny IP. Considering maternal dietary modulators for epigenetic regulation and programming of the fetal epigenome. *Nutrients*. 2015;7:2748.
 67. Reynolds CM, Gray C, Li M, Segovia SA, Vickers MH. Early life nutrition and energy balance disorders in offspring in later life. *Nutrients*. 2015;7:8090.
 68. Stettler N, Iotova V. Early growth patterns and long-term obesity risk. *Curr Opin Clin Nutr Metab Care*. 2010;13:294–9.
 69. Odegaard AO, Choh AC, Nahhas RW, Towne B, Czerwinski SA, Demerath EW. Systematic examination of infant size and growth metrics as risk factors for overweight in young adulthood. *PLoS One*. 2013;8. <https://doi.org/10.1371/journal.pone.0066994>.
 70. Pew Research Center. Mobile Fact Sheet. Pew Research Center: Internet, Science & Tech. 2018. <http://www.pewinternet.org/fact-sheet/mobile/>. Accessed 11 Aug 2018.
 71. Smith A. Americans and text messaging. Pew Research Center: Internet, Science & Tech. 2011. <http://www.pewinternet.org/2011/09/19/americans-and-text-messaging/>. Accessed 21 Sept 2016.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

