University of New Mexico UNM Digital Repository

Pharmaceutical Sciences ETDs

Electronic Theses and Dissertations

8-30-2011

Pattern of multivitamin use : prevalence and predictors among pregnenat women in New Mexico

Sizhu Liu

Follow this and additional works at: https://digitalrepository.unm.edu/phrm etds

Recommended Citation

 $\label{linear_$

This Thesis is brought to you for free and open access by the Electronic Theses and Dissertations at UNM Digital Repository. It has been accepted for inclusion in Pharmaceutical Sciences ETDs by an authorized administrator of UNM Digital Repository. For more information, please contact disc@unm.edu.

Sizhu Liu	
Candidate	
College of Pharmacy	
Department	
This thesis is approved, and it is acceptable in quality a	nd form for publication
Approved by the Thesis Committee:	
Ludmila Bakhireya , Chairperson	
Ludmila Bakhireva , Chairperson	
Melanie A. Dodd	
Larry Georgopoulos	

PATTERN OF MULTIVITAMIN USE: PREVALENCE AND PREDICTORS AMONG PREGNANT WOMEN IN NEW MEXICO

 \mathbf{BY}

SIZHU LIU

BACHELOR OF SCIENCE

THESIS

Submitted in Partial Fulfillment of the Requirements for the Degree of

Master of Science

Pharmaceutical Science

The University of New Mexico Albuquerque, New Mexico

July, 2011

Pattern of Multivitamin Use: Prevalence and Predictors among Pregnant Women in New Mexico

By

Sizhu Liu

B.S., Economics, China Pharmaceutical University, 2009M.S., Pharmaceutical Science, University of New Mexico, 2011

ABSTRACT

Background: Prenatal vitamin use is recommended as a necessary supplement prior to conception and throughout pregnancy. Multivitamin use in early pregnancy can significantly reduce risks for birth defects: cardiovascular defects, ^{1, 2} limb defect, ³⁻⁵ urinary tract abnormalities, ^{4, 6} orofacial clefts, ^{7, 8} and neural tube defects (NTDs). ^{2, 9-16} Sixty-nine to seventy-eight percent of women take vitamins after pregnancy recognition, while only 23-35% of women in the U.S. start to use vitamins before pregnancy recognition. ¹⁷⁻¹⁹ Rates of NTDs by ethnicity demonstrat that Hispanics have the highest prevalence (4.2 per 10,000 births) than Non-Hispanics. (Non-Hispanic Black or African American: 3.2 per 10,000 births; Non-Hispanic Whites: 2.6 per 10,000 births). ²⁰ **Objectives:** To explore the prevalence rates of multivitamin use during pregnancy and to find out the predictors of vitamin use using an established cohort in New Mexico. **Methods:** This is a cross-sectional analysis which used data from the "Safety of Medication and Perception of Teratogenicity

(SMART)" study. Patients in the SMART study were recruited from 5 University of New Mexico affiliated prenatal clinics. Patients were considered vitamin users if they took vitamins at least 4 times a week. Based on the different time exposure to vitamins, vitamin use was further categorized into three groups: pre-conceptional vitamin users, vitamin users after pregnancy recognition, and vitamin non-users. Chi square and ANOVA were used to identify potential predictors including sociodemographic characteristics, lifestyle characteristics, medical and reproductive health, and medication use. For multivariate analysis, ordinal logistic regression and polychotomous logistic regression model were used. Results: Most pregnant women (71.9%) in the sample began to take vitamins regularly after pregnancy recognition. Earlier vitamin use in pregnancy was significantly associated with pregnancy planning independent of other maternal characteristics. Education level and health insurance status revealed a significant interaction with respect to vitamin use. Lastly, almost one third (32.1%) of women in this study had a history of adverse pregnancy outcomes and 45.0% of women had at least one medical condition. Vitamin non-users were more likely to have experience of adverse pregnancy outcomes in prior pregnancies compared to women who used vitamin after pregnancy recognition. Implication: This study highlights the importance of promoting vitamin use in New Mexico in women of childbearing years prior to conception and in early pregnancy, given the finding that less than one third of women used vitamins during the pre-conceptional period. Public health strategies should also include the promotion of pregnancy planning.

Key word: Vitamin, Pregnant women, Predictors, Prevalence, Ordinal Logistic Regression, Polychotomous Logistic Regression

TABLE OF CONTENT

CHAPTER I: INTRODUCTION	1
Background	1
Special Aims	
CHAPTER II: LITERATURE REVIEW	8
The Benefit of Multivitamin Use in Pregnancy	8
Cardiovascular Defects	8
Limb Defects	9
Urinary Tract Anomalies	10
Orofacial Clefts	11
NTD (Neural tube defects)	12
Other Defects and Adverse Perinatal Outcomes	13
The Prevalence of Multivitamin Use	
Prevalence of Multivitamin Use Anytime in Pregnancy	14
Prevalence of Multivitamin Use in Periconceptional Period	15
Multivitamins vs. Prenatal Vitamins	
Predictors of Multivitamin Use	17
Definition of Multivitamin Use in Periconceptional Period and Early	
Pregnancy	
Regular Use vs. Non-regular Use	18
Prescription vs. OTC Multivitamins	19
Multivitamin vs. Single Vitamin	20
The Timing of Measuring Multivitamin Exposure among Pregnant	
Women	
Economic evaluations on food fortification and folic acid use	21
CHAPTER III: RESEARCH DESIGN	
Study Population	23
Measurements	
Sociodemographic Characteristics	24
Lifestyle Characteristics	
Medical and Reproductive Health	
Vitamin and Dietary Supplement Use	27
Medication Use	28
Perinatal Outcomes	29
Data Modifications	29
Statistical Approaches	
Power Calculation	
CHAPTER IV: RESULTS	
Description of the Study Population	
Vitamin Use Patterns	38
Maternal Characteristics of Vitamin Use	
Ordinal Logistic Regression	
Polychotomous Logistic Regression	
CHAPTER IV: DISCUSSION	
Summary	
Prevalence of Vitamin Use as Compared to Other Studies	
Predictors of Vitamin Use as Compared to Other Studies	
Race/Ethnicity	
Maternal Education	53

	Health Insurance	54
	Marital Status	55
	Maternal Age	56
	Pregnancy Planning	
	History of Adverse Perinatal Outcomes	59
	Other Predictors	
	Limitations	61
	Strengths	64
	Future Directions & Recommendations	65
RE	FERENCES	
	T OF TABLES	
	Table 2.1. Prevalence of Multivitamin Use in Pregnancy	81
	Table 2.2. Predictors of Multivitamin Use in Pregnancy	.110
	Table 3.1. Sample size calculations for different effect sizes	.127
	Table 4.1 Characteristics of the Study Population (N=402)	.128
	Table 4.1 (Continued) Characteristics of the Study Population (N=402)129
	Table 4.2 Descriptive Statistics of Multivitamin Use Patterns (N=402)	130
	Table 4.3 Characteristics of the Study Participants by Vitamin Use	.131
	Table 4.3 (Continued) Characteristics of Study Participants by Vitamin	
	Use	.132
	Table 4.4 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis*	
	Table 4.5 Interaction between Predictors in Ordinal Logistic Regression	n
	and Polychotomous logistic Regression Model	.134
	Table 4.6 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis among Women without Health Insurance	
		.135
	Table 4.7 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis among Women with Private Insurance *	. 135
	Table 4.8 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis among Women with Public Insurance *	. 135
	Table 4.9 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis among Women without Insurance*	. 136
	Table 4.10 Predictors of Vitamin Use: Results of Multivariate Ordinal	
	Logistic Regression Analysis among Women with Insurance*	. 136
	Table 4.11 Predictors of Vitamin Use: Results of Univariate	
	Polychotomous Logistic Regression Analysis*	.137
	Table 4.12 Predictors of Vitamin Use: Results of Polychotomous	
	Multivariate Logistic Regression Analysis*	
AP	PENDICES	.139
	APPENDIX A Data Dictionary for SMART Study	.139
	APPENDIX R OLIESTIONNAIRE OF THE SMART STIIDY	164

CHAPTER I: INTRODUCTION

Background

Multivitamins are considered as a necessary supplement for pregnant women.²¹ They contain a wide range of supplements including water-soluble/fat soluble vitamins (e.g. vitamin A, vitamin B, folic acid) and minerals, most of which have been recommended for prenatal use in the U.S.^{22, 23} Women can benefit from using prenatal multivitamins during pregnancy. For instance, the risks of having preterm birth and preeclampsia can be reduced.^{24, 25} Prior evidence indicated that prenatal multivitamin supplementation can provide protective effects against a series of birth defects including cardiovascular defects, 1, 2 limb defects, 3-5 orofacial cleft, 7, 8 urinary tract anomalies, 4, 6 congenital hydrocephalus, 26 respiratory tract defects, 26 and omphalocele.²⁷ In addition, periconceptional use of prenatal multivitamins containing folic acid is associated with decreased risk for Neural Tube Defects (NTDs).2, 9-16

Among many birth defects, NTDs, a structural birth defect, are one of the most severe congenital malformations in human beings.²⁸ NTDs are a series of malformation of the developing brain and spinal cord, and it occurs when there is an incomplete closure of the neural tube within a month of fertilization.²⁹ Anencephaly and spina bifida are the most common NTDs, and they affected around 300,000 newborns worldwide.³⁰ In the U.S., the number of NTD affected pregnancies decreased after the introduction of food fortification in 1998. A CDC report analyzed data from 23-poplation-based surveillance systems and comparing the prevalence of NTD before and after folic acid

fortification (1995-1996 vs. 1999-2000), suggesting that the average annual number of NTD-affected pregnancies was 3,020, comparing to 4,130 before food fortification.³¹ However, the recent national data from 14 states showed that anencephaly and spina bifida still affected 859 and 1460 babies each year in the U.S., respectively.^{32, 33} The causes of NTDs can be either environmental or genetic, but the causative mechanism of NTDs is still unclear and poorly studied.^{28, 34} However, a majority of current clinical studies found that periconceptional multivitamin supplementation containing folic acid can have a protective effect against neural tube defects.^{2, 9-16} In addition, a meta-analysis summarized the findings and showed that the overall effect of periconceptional use of multivitamin plus folate was significantly effective in protecting against NTDs.²⁹

In the U.S., the prevalence of multivitamin use during pregnancy has been documented by previous studies and ranges from 68.8-78%.^{19, 35} Unfortunately, the prevalence of regular prenatal multivitamin use (at least 3 times per week) is reported 53.8%.³⁶ Although a study using data from the National Maternal and Infant Health Survey (NMIHS) showed that the prevalence rate of multivitamin regular use was 82.5% in pregnancy.³⁷ This prevalence combined mineral use with multivitamin use and therefore might amplify the estimation of prevalence of multivitamin use in pregnancy.

While the majority of women initiate multivitamin use after pregnancy recognition, only 24.0% of women reported having ever used multivitamins during periconceptional period³⁵- the most crucial period for organogenesis.

Multivitamin use during the first semester might be higher, but still in sufficient (29.3%).³⁸ With regard to multivitamin regular use during periconceptional period, the prevalence of periconceptional multivitamin regular users (at least 3 times per week) was reported around 21.0%.³⁶ Before recognition of pregnancy, studies showed that only 26% of women reported using multivitamins regularly (at least 3 times a week).³⁷ Two Centers for Disease Control and Prevention (CDC) reports using data from Pregnancy Risk Assessment Monitoring System (PRAMS) suggested that the prevalence of pre-conception multivitamin use was 23-45.2%.^{18, 39}

Unplanned pregnancies may account for the difference in prevalence rates of multivitamin use between periconceptional period and after the recognition of pregnancy.⁴⁰ Women with unintended pregnancies become aware of their pregnancies at a later gestational age than women who planned their pregnancies.⁴¹ As a result, the time of multivitamins exposure during periconceptional period may be delayed for women who did not plan a pregnancy.

Difference in prevalence of multivitamin use among pregnant women was also found among different racial groups. Only 25.6% and 67.0% African American use multivitamins in periconceptional period and later in pregnancy, respectively. On the other hand, 55.0% and 84% non-Hispanic white mothers use multivitamins in these two periods.^{37, 42} For Hispanics, a national investigation using data from the Behavioral Risk Factor Surveillance System (BRFSS) showed that 66.4% of Hispanic pregnant women reported having

ever used multivitamins since pregnancy.¹⁹ Data obtained from the National Birth Defects Prevention study showed that the prevalence of periconceptional intake of folic acid-containing supplements is only 30% among Hispanics, which is the lowest comparing to other racial groups (Non-Hispanic White: 66%; Non-Hispanic Black: 39%).⁴³

Some evidences show that Hispanic women might have a high rate of adverse pregnancy outcome comparing to non-Hispanic white women. Before the introduction of mandatory food fortification in 1998, it has been documented that the prevalence of NTD is almost 3 times higher among infants of Hispanic women than that for non-Hispanic white women. 44-46 Although the overall trend of NTD prevalence has declined since fortification, the prevalence of NTD is still greater in Hispanics (4.17 per 10,000), comparing to non-Hispanics (Non-Hispanic Black or African-American: 2.64 per 10,000; Non-Hispanic Whites: 3.22 per 10,000)²⁰ Also, a CDC report using data from the Pregnancy Risk Assessment Monitoring System (PRAMS) found that the prevalence of having a low birth weight infant was higher among Hispanics than Non-Hispanic white women. 47 Moreover, a cross-sectional study used data from the National Birth Defects Prevention Study and found that foreign-born Hispanics had a higher risk of delivering babies with spina bifida than US-born Hispanics, and the higher prevalence of NTDs was also found among recent immigrants from foreign countries.⁴⁸ In addition, a low or deficient level of folic acid are found among Hispanic women because they are more likely to become pregnant relatively soon after giving birth. 49 The national vital statistics

reports in 2005 showed that Hispanic women have a higher birth rate and often had children at an older age.⁵⁰

The lack of knowledge about benefits of vitamin use among Hispanic women might at least partially explain the higher prevalence of NTDs in Hispanics compared to Non-Hispanic Whites and African American women. However, there are also other factors that might account for a higher prevalence rate. Environmental exposures might be a contributing factor. For instance, pesticide exposure might be higher among Hispanic women, especially women involved in the farm labor, as demonstrated in some prior studies.⁵¹ A population-based case-control study among Mexican-born Hispanics demonstrated that cigarette smoke, including second-hand exposure, significantly increased risk of NTDs. 52 Cultural norms and fatalism could also be a potential explanation for high prevalence of NTDs at birth. Religious beliefs, one important component of Hispanic culture norms, make it possible that Hispanic women are less likely to abort their children regardless of possible birth defects. Moreover, in fatalistic attitudes among Hispanics, they might believe that life events are predetermined. Therefore, Hispanic mothers may be less likely to participate in prenatal screening. As a result, Hispanic mothers might be more likely to deliver babies with NTDs than Non-Hispanic mothers.

There is a great need to compare the prevalence of periconceptional and preconception multivitamin regular use among Hispanics vs. non-Hispanic Whites, which can help us estimate the difference in the vitamin use between

Hispanic women and other racial groups. In addition, understanding specific predictors can identify those Hispanic women who are at risk of not using prenatal multivitamin. This is important as culturally—sensitive interventions can be developed and promote early vitamin use in pregnancy among Hispanic women.

Special Aims

The main purpose of this research is to investigate the prevalence of regular multivitamin use among different racial/ethnic groups of pregnant women and to describe the maternal characteristics of regular vitamin users. Specifically, the following aims are proposed:

<u>Aim</u> 1) To determine the differences in the prevalence of multivitamin regular use (at least 4 times per week) in preconception period (at least one month before last menstrual period) and in pregnancy (after pregnancy recognition) between Hispanic and non-Hispanic pregnant women.

<u>Hypothesis 1:</u> we hypothesize that the prevalence of preconception multivitamin regular use will be lower in Hispanic than in Non-Hispanic women.

<u>Aim</u> 2) To examine the predictors (e.g. ethnicity, education level, insurance status, age, marital status, pregnancy intention, smoking status, alcohol use, recreational drug use, parity, gravidity, a family history of birth defects, the use of prescription medications) of not preconception multivitamin regular use from an established cohort of pregnant women in New Mexico.

Hypothesis 2: we hypothesize that pregnant women who did not use multivitamins in preconception period are more likely to be Hispanics, less educated, younger, uninsured, unmarried, had an unintended pregnancy intention, are foreign born, multiparous, multigravida, speaking Spanish, current smokers, periconceptional binge drinker, recreational drug user, without a family history of birth defects, and use less than 2 kinds of prescription medications.

CHAPTER II: LITERATURE REVIEW

The literature review for this study is organized into following sections: (1) the benefit of using multivitamin during pregnancy and periconceptional period; (2) the prevalence of multivitamin use among pregnant women; 3) predictors of multivitamin use; 4) definition of multivitamin use in periconceptional period and early pregnancy in different studies; 5) a review of economic evaluations on food fortification and folic acid use.

The Benefit of Multivitamin Use in Pregnancy

We systematically reviewed the benefits of multivitamin supplementation use in pregnancy and summarized contents as following categories: cardiovascular defects, limb defects, urinary defects, orofacial defects, neural tube defects, and other defects and adverse perinatal adverse outcomes.

<u>Cardiovascular Defects</u>

There are many types of cardiovascular defects, such as transposition of great arteries (TGA), hypoplastic left heart syndrome (HLHS), atrioventricular septal defect (ASD). In the U.S., it has been estimated that TGA, HLHS, and ASD occurred 1 cases in 3,333 births, 1 case in 2,122 births, and 1 case in 4,344 births. It also accounts for a major cause of infant deaths. Some studies showed a significant association between the use of multivitamin containing folic acid in pregnancy and occurrence of cardiovascular defects of infants. One randomized clinical trial from Hungary found that the risk of congenital cardiovascular malformations (e.g. ventricular septal defect, aortic stenosis, patent ductus arteriosus) was significantly lower among women daily used

periconceptional multivitamins with 0.8 mg folic acid at least one month before conception.² A population-based case-control study from the U.S. also confirmed that the periconceptional multivitamin use (any multivitamin use 3 month before to 3 months after conception) can reduce the risk of congenital heat defects (OR=0.60-0.97). In contrast, a case-control study collected data from Slone Epidemiology Unit Birth defects Study, tested a series of selected birth defects but did not find a significant association between preconceptional multivitamin supplementation (2 month before last menstrual period) and risks defects.4 of conotruncal This result is consistent with the Baltimore-Washington Infant Study (BWIS), a large case-control study, which showed no protective effect of multivitamin use during the first 5 weeks of gestation.⁵⁴ These two contradicting results may be due to differential maternal recall of multivitamin use in the Slone Epidemiology Study or the fact that the interview in BWIS study was conducted prospectively before the infant's birthday while other studies respectively interviewed mothers after the delivery of infants.

Limb Defects

There are two major types of limb defects: upper limb (e.g. arms) defects and lower limb (e.g. leg) defects. There are approximately 1,500 babied in the U.S. born with upper limb defects and 750 with lower limb defects for each year.³³ Periconceptional use of multivitamin can also provide a protective effect against limb defects. A few case-control studies suggested that women who used multivitamin periconceptionally (from 3 months before conception until 3 months after pregnancy) could significantly reduce (OR=0.3-0.64) the risk of

developing infants with limb defects.^{3, 5} One of the case-control studies specified that this protective effect was significant in transverse limb defects but not in other limb defects, such as preaxial and postaxial deficiencies.⁵ In addition, a meta-analysis conducted by Canada showed that use of multivitamin supplements before during the first trimester of pregnancy provided consistent protection against limb defects (OR=0.48, 95% Cl=0.30-0.76).⁵⁵ However, a case-control study conducted in Boston, Philadelphia, and Toronto during 1993-1996 did not find a significant reduction in limb defects among women who used multivitamins regularly before and in the first trimester use⁴ The reason might be due to non-population based subject ascertainment and selection bias.

Urinary Tract Anomalies

In the United States, congenital malformation of the genitourinary system caused 518 deaths per 100,000 live infant per year. Several studies have been done and demonstrated that periconceptional multivitamin use could significantly reduce the risk of urinary tract anomalies, such as rental agenesis, obstructive congenital abnormalities of the urinary system. Of note, the risk reduction become smaller when multivitamin use was limited to the second or third trimester, which was reported by a case-control study, using data from the Slone Epidemiology Unit Birth defects Study. A randomized double-blind controlled trial from Hungary was conducted and found that the daily periconceptional use (at least one month before conception and at least until the date of the second missed menstrual period) of multivitamin containing 0.8 mg of folic acid can significantly reduce the rate of urinary tract abnormalities.

However, the same author did not find this association in a follow up case-control study.² The protective effect of periconceptional multivitamin use might be attenuated due to potential selection bias.

Orofacial Clefts

cleft lip with or without cleft palate (CLP) and cleft palate alone (CP) are two kinds of main orofacial clefts, and it has been reported that CLP and CP affected approximately 1 in 941-1000 and 1 in 1574-2500 infants, respectively.32,58 Mixed findings have been found with regarding to the protective effect of multivitamin use and occurrence of orofacial clefts. Some studies reported that the risk reduction for CLP but not for not in CP among women who used multivitamins regularly during periconceptional period (3 months before through 3 months after conception), 10, 58 while other studies found a reduction for CP but not for CLP.⁵⁹ In addition, one case-control study reported that the greatest reduction in the risk of CLP occurred with periconceptional multivitamin use (28 days before through 28 days after conception).4 Also, there are other studies showing that the risks of both CP and CLP can be reduced by using multivitamin periconceptionally (one month before through two months after conception). 60 On the contrary, there is one case-control study which did not find any significant associations between the periconceptional multivitamin supplementation containing folic acid and reducing risk of CP or CLP.61 Recall bias might be a problem in this study when women were asked to report multivitamin use that occurred up to more than one year before the time of interview.

NTD (Neural tube defects)

NTD is a severe structural defect due to the incomplete closure of the neural tube within a month of fertilization.²⁹ The most common types of NTD are anencephaly and spina bifida. Infants born with anencephaly usually die within the first few hours/days after delivery, while a majority of infants born with spina bifida grow to adulthood exhibiting only physical disability but having normal neurobehavioral development.²² The medical costs for the first year of life for a child with spina bifida was \$52,415.62 It is still unclear of the causative mechanism, but there are extensive studies which suggested that the use of periconceptional multivitamin supplementation (from 3 months before and through 3 months after conception) can have a protective effect against NTDs.^{2, 9-16} A randomized double-blind prevention trial was conducted in 7 countries and find that periconceptional multivitamin containing folic acid (4 mg folic acid per day) can reduce recurrence risk of NTDs, but whether lower doses had same protective effect was unknown.9 Another randomized trial suggested that a daily dosage of 0.36 mg of folic acid could still be as protective as a daily dosage of 4 mg of folic acid 15. Moreover, one case-control study from California found that there was a significant reduction associated with periconceptional multivitamin use (from 3 months before and through 3 months after conception) for NTD. 10 Due to different study design and multivitamin supplementation measurement, some studies, however, did not find a protective effect of periconceptional multivitamin supplementation on preventing neural tube defects. 14, 63-65

Other Defects and Adverse Perinatal Outcomes

There are a number of studies which evaluated other birth defects and adverse pregnancy outcomes. A population-based case-control study found that periconceptional use (3 months before pregnancy through the first trimester of pregnancy) of multivitamin could reduce the risk for omphalocele, an abdominal wall defect. However, the precision of this study was limited by small sample size in case-infants, which only 72 cases were ascertained. A population-based Atlantic Birth Defects Case-Control study examined the associations of periconceptional multivitamin use (from 3 months before pregnancy through the first 3 months of pregnancy) and respiratory tract defects, pyloric stenosis, anal atresia, but none of them yielded a significant result. Each of the pregnancy is stenosis, and atresia, but none of them yielded a significant result.

The association of preterm birth and multivitamin use is also studied. Case-control studies suggested that its risk could be reduced approximately 1.6-2 folds with multivitamin supplement use during pregnancy.^{38, 67} Surprisingly, a case-control study reported that multivitamin use during the third trimester was increased risk of preterm birth.⁶⁸ The interpretation of this result, however, needs to be caution, and the mechanism is unclear.

The association between adverse pregnancy outcome (miscarriage, ectopic pregnancy, and stillbirth) and the use of multivitamin containing folic acid during periconceptional period was also studied, but none of them was reported significantly associated with periconceptional multivitamin supplementation use. ^{9, 69, 70}

Preeclampsia is considered to a pregnancy-related disorder, and it remains a significant health problem in obstetric population. A prospective cohort study was designed to detect factors of preeclampsia in the United States, and the results showed that the periconceptional multivitamin use was associated with 45% of reduction in preeclampsia risk comparing to nonusers.⁷¹ Another study found that regular multivitamin use in the periconceptional period was significantly associated with reduced risk of preeclampsia among normal-weight women.²⁴ However, more studies need to be done in this area in order to further confirm this result.

In addition to preeclampsia, periconceptional multivitamin supplementation (not specifically defined) can also significantly reduce the incidence of pregnancy-induced vertigo, nausea, and vomiting, which reported by a randomized placebo-controlled trial.⁷²

The Prevalence of Multivitamin Use

Prevalence of Multivitamin Use Anytime in Pregnancy

There is a variation in the prevalence of using multivitamins during pregnancy. In the United States, it was previously reported in 1998 that 53.8% of women from the National Maternal and Infant Health Survey reported using either multivitamin and minerals 3 times per week after pregnancy recognition.³⁶ After the introduction of mandatory folic acid fortification, the prevalence rate went up and ranges from 68.8-78% in 2009.^{19, 35}

In Europe, the prevalence of multivitamin use during pregnancy was reported as 76.2% in Portugal, which was highest comparing to other countries in Europe. The Countries in Europe. Unfortunately, the comparability of the data on the prevalence of multivitamin use was undermined by combining minerals use with multivitamins. Besides, there were another two studies estimating prevalence by combining multivitamins and minerals. One study suggested that 30.8% of participants using multivitamins minerals during pregnancy, while another study broken down the prevalence rate and specified it into three trimesters periods, revealing that 23.3%, 14.1%, and 18.6% of women used multivitamin-mineral during the first trimester, second semester, and third trimester, respectively. In addition, in Australia, the prevalence rates of multivitamin use during pregnancy were between 18-35% In Brazil, the prevalence of multivitamin use was 14% as reported.

Prevalence of Multivitamin Use in Periconceptional Period

In the U.S., the prevalence of periconceptional multivitamin (begin before pregnancy and continue to use in pregnancy) was reported as 21-23.8%.^{35, 36} There were also studies which attempted to specify the time frame of multivitamin exposure during periconceptional period. A Pregnancy Exposure and Preeclampsia Prevention Study (PEPPS) found that 47% of women took multivitamins/prenatal vitamin at least once per week during periconceptional period (from 3 months before pregnancy through the first 3 months of pregnancy).⁷⁸ Recall bias might impact on the prevalence rate estimation in this study while women were asked to remember multivitamin use 3 months before pregnancy. A National Birth Defects Prevention (NBDP) Study further

examined periconceptional multivitamin use (one month before LMP and through the month after LMP) and pregnancy intention and found that 35.7% of women who did not plan for the pregnancy used periconceptional multivitamin, while the prevalence of periconceptional multivitamin use among women who had a pregnancy intention was reported as 59%. In Canada, a cross-sectional analysis conducted in Toronto Hospital and the Mount Sinai Hospital found that only 28% of women reported using multivitamin supplementation containing acid during periconceptional period (at least 4 weeks prior to conception until 8 weeks after conception). Additionally, preconception multivitamin use-multivitamin use during the month before pregnancy- was also studied. Data from PRAMS showed that the prevalence of preconception multivitamin regular users (≥ at least 4 times per week) was 23.0-43.6%. 18, 39, 47, 81

In Europe, a study from Demark using data from Danish National Birth Cohort and found that 65% of women had used multivitamins in periconceptional period (4 weeks before LMP through the 8 weeks after LMP).²⁴ The high prevalence is because of the fact that Caucasian women accounted for most of this population in Denmark and they were more prone to use periconceptional multivitamins. To the contrast, two other studies, one from Sweden using women who attended antenatal care units and another from Norway using Medical Birth Registry, found that 15.9-19% of women used vitamin during periconceptional period (not specifically defined: before pregnancy and during pregnancy).^{49, 82} Regarding to preconception multivitamin use, one study from Hungary reported that the prevalence of

preconception multivitamin regular use was 43.8%.⁸³ However, all the data on multivitamin use were based on self-reports from voluntary participants, which might have a potential selection bias.

In Australia, a cross-sectional analysis showed that 12.3% of women had used multivitamins during periconceptional period (3 months prior to pregnancy and during pregnancy).⁷⁶ Another cross-sectional survey from Australia found that 21.8% of women had used multivitamin during periconceptional period (not specifically defined: before and during pregnancy).⁷⁵

Multivitamins vs. Prenatal Vitamins

Most studies assumed that prenatal vitamins and multivitamins are similar in contents. Information of multivitamins and prenatal vitamins in other studies was mixed, and multivitamin users were identified if women reported having taken multivitamin or prenatal vitamins.^{35, 38, 47, 78, 79, 81} Therefore, we cannot separately specify the prevalence of prenatal vitamin use from these results. It is of note that there is one study from U.S. that investigated the prevalence of prenatal vitamin use among women at rural outreach clinics and reported that 92% of women had used prenatal vitamins during pregnancy.⁸⁴ For most of other studies, the use of prenatal vitamins was not separately specified to investigate the prevalence.

Predictors of Multivitamin Use

Studies identified examined the maternal characteristics of multivitamin users.

Prenatal multivitamin users (use multivitamin anytime during pregnancy) were

more likely to be White^{42, 47}, better educated,^{37, 42, 78, 85} older,^{37, 42, 77, 82} married or cohabiting with a partner,^{19, 37, 42, 78, 82} be primiparous,^{77, 82} have a higher income,^{19, 37, 42, 47} pregnancy intention,⁷⁹ and private health insurance.⁴⁷ Similar to multivitamin users during pregnancy, periconceptional multivitamin users are those who were non-smokers,^{42, 82, 86} married,^{42, 78, 82, 85, 86} older,^{42, 82, 86} have a high socioeconomics status,^{42, 86} better educated,^{42, 78, 85, 86} primiparous,^{75, 80, 82} normal BMI,⁷⁸ However, a population-based survey initiated by the Medical Birth Registry and the National Council on Nutrition and Physical Activity from Norway reported periconceptional multivitamin users were tended to be younger.⁸⁵ With regard to preconception, multivitamin use are more likely to be white,⁴⁷ older,^{39, 47} more educated,³⁹ intended pregnancy,^{47, 86} and have private health insurance.⁴⁷ To the contrary, the preconception non-multivitamin users are more likely to be Black,⁸¹ younger,⁸¹ less educated.³⁹ Medicaid recipients.^{39, 81}

Definition of Multivitamin Use in Periconceptional Period and Early Pregnancy

Regular Use vs. Non-regular Use

Multivitamin use is assessed in various ways. First, women are often identified as multivitamin users if they use multivitamins at the time of assessment or reported having ever used multivitamins in pregnancy. 19, 24, 35, 38, 49, 76, 77, 79, 82, 84 However, this assessment does not take into consideration the frequency and duration of multivitamin use, and this could lead to overestimation of multivitamin regular use. It could be possible that women who had just used multivitamin once were also regarded as multivitamin users. There were some studies estimating multivitamin use among regular users. 36, 37, 42, 74, 78, 85 The

change in the definition from multivitamin users to multivitamin regular users did not cause large variation in prevalence of using multivitamins during pregnancy. In the U.S., the prevalence of using multivitamins during pregnancy was reported between 78.0% and 82.5%. Regarding to the prevalence of periconceptional multivitamin use in the U.S., one study reported that 23.8% of women reported having used multivitamin during periconceptional period, while another study suggested that 21.0% of women reported regularly using multivitamins (3 days a week) during periconceptional period.

Prescription vs. OTC Multivitamins

The main difference between prescription and OTC multivitamins is the amount of folic acid: while OTC multivitamins contain 400 micrograms dose, prescription multivitamins have 1000 micrograms. A majority of studies asked women to report their multivitamin use but did not further asked women whether the multivitamins they took was prescription or OTC multivitamins. Although some studies collected information of brand of multivitamin supplementations, it is often not reported and still unclear about prescription multivitamin as well as OTC multivitamin use among pregnant women. A few of studies, however, investigated prescription and OTC multivitamins among multivitamin users. One study from U.S. investigated women at rural outreach clinics and reported that 92% of women had used prescription prenatal vitamins during pregnancy, while 10.9% of women had used OTC multivitamins.⁸⁴

Multivitamin vs. Single Vitamin

The use of single vitamin (e.g. vitamin A, vitamin B, vitamin C, vitamin D, vitamin E, folic acid) along with multivitamin use was studied. It seems that a small proportion of women who took multivitamin took additional single vitamins: a population-based study from UK assessed frequency and amount of all single vitamins during pregnancy including vitamin A, vitamin C, and vitamin E, all of which were below 5%. ⁶⁸ Contrary to other single vitamins, a large number of women who took multivitamin also took additional folic acid supplementations during pregnancy: it was reported that the prevalence of taking folic acid was between 71.4-78.7%. ⁶⁸ ⁷⁵

The Timing of Measuring Multivitamin Exposure among Pregnant Women

The time frames for multivitamin exposure during pregnancy and periconceptional period were defined differently. For pregnancy, multivitamin exposure was usually measured after women realized they were pregnant or used the last menstrual period as the beginning time point of pregnancy. There is a time gap between the last menstrual period and women's recognitions of pregnancies, and women may not realize they were pregnant even the fetus was developing. A population-based study combined these two time points by asking women if they used any multivitamins since their recognitions of pregnancies or last period menstrual.³⁵

Periconceptional period in studies is often defined 28 days prior to the last menstrual period to after 28 days after the last menstrual period.⁴² Some studies expanded the endpoint of periconceptional period to 8 weeks after last

menstrual period.^{24, 49} In addition, some studies did not define an exact time for periconceptional period but defined periconceptional multivitamin users as women who reported having used multivitamin before LMP and continued in pregnancy.^{35 36}

Economic evaluations on food fortification and folic acid use

Some studies evaluating the benefits of folic acid fortification in food in the U.S, showed that it was associated with an annual economic benefit of \$312-\$425 millions and that the cost savings (net reduction in direct costs) ranged from \$88 to \$145 millions per year. Another cost-effectiveness analysis from the U.S. analyzed folic acid fortification policy and found that it could achieve \$266,649 per QALY gains. However, this study did not focus on pregnant women and NTDs. The outcomes in this study included myocardial infarctions and color cancer for males.

In the Netherlands, a cost-effectiveness analysis conducted from a societal perspective found periconceptional use of folic acid supplementation was cost-effective (€1800-4500 per QALYS).⁸⁹ However, this study did not include productivity loss as a component in their cost estimation. There is another pharmacoeconomic study from the Netherlands which conducted a cost-effectiveness analysis to evaluate the folic acid food fortification and prevention of NTDs from a societal perspective. That study found that fortification of bulk food with 140µg folic acid per 100g results in cost savings if the cost of enrichment does not exceed €5.5 million.⁹⁰ However, the result of

this study might not be comparable, because the cost estimation was estimated under a different health care market and health care system.

CHAPTER III: RESEARCH DESIGN

Study Population

This is a cross-sectional study using data from the "Safety of Medications and Perception of Teratogenicity (SMART)" study. SMART is a prospective multi-clinics study conducted at the University of New Mexico (UNM). Eligible participants were women who aged 18 or older and attended UNM-affiliated prenatal care clinics (UNMH Fetal Monitoring Clinics, Triage, General Obstetric/Gynaecological, Satellite Clinic-West Mesa, Satellite Clinic-South Broadway, Satellite Clinic-North Valley) in Albuquerque, NM, during 2008-2009. A written informed consent in English or Spanish was obtained before each interview. If women agreed to participate, a semi-structured interview was administrated by a trained bi-lingual interviewer fluent in English and Spanish in a private setting at the clinics. If women did not want to participate, they would be asked to give reasons for non-participation. Institutional Review Board (IRB) approval for this study was obtained from the UNM Research Review Committee (HRRC).

On an initially screening, 493 pregnant women who were consecutively chosen from UNM main hospital and satellite clinics were offered participation in the study. Among those, 404 women enrolled in SMART study (response rate=81.5%). The major reasons for non-participation were the time constrains and lack of interest in the study. For the purpose of this study, women who did not report their multivitamin use were excluded (n=2). Therefore, 402 women were included for the present analysis.

Measurements

Sociodemographic Characteristics

Sociodemographic characteristics, including maternal age (continuous), marital status (single/never married, married/living with spouse, not married but living with partner, separated from spouse, divorced, widowed), race (white Non-Hispanic, white Hispanic, black or African American, American Indian or Alaskan native, Asian or Asian American or Pacific Islander, some other group, prefer not to report), ethnicity (Hispanic/Latino/Spanish descent, other ethnical groups), level of education (less than high school graduate, high school education or GED, some college or vocational school, college degree, master/doctorate or professional degree), health insurance status (no insurance, employer-based insurance, self-purchased insurance, Medicaid, other public insurances), place of born (U.S. born, foreign born), years of staying in U.S for immigrants, language predominantly spoken at home (English, Spanish, some other language) were ascertained. In addition, if women described themselves as American Indian or Alaskan Native, they were further asked to specify whether they can identify a specific tribe or pueblo. Women were also asked whether their health insurance covered prescription drug or not. For women who reported having other public health insurances, they would be asked to specify whether it is Indian Health Service, Veteran Affair (VA), First Choice, UNM/UNM Care, or other types.

<u>Lifestyle Characteristics</u>

Interviewer asked participants to report their lifestyle characteristics including smoking, alcohol use, and illicit drug use. We firstly asked women whether

they smoked cigarettes or used tobacco at the time of the interview. If they answered "yes", the number of cigarettes in a day was recorded. Otherwise, we asked them whether they have ever smoked more than 100 cigarettes and whether they stopped before or after recognition of pregnancy.

Regarding alcohol use, we asked women whether they have ever used alcohol. If women answered "yes", the interview would ask them to specify the number drinks typically she feels the first effect of alcohol and the number of drinks she can hold before passing out of falling asleep. We ascertained periconceptional binge drinker by asking women how many times participants drank four or more drinks during a month around the LMP. In addition, the standardized TWEAK (Tolerance, Worried, Eye-opener, Amnesia, Kut down) with "hold" and "high" versions of the questionnaire was administrated. 91-93 The difference between the two versions is the cutoff point of "Tolerance". In the "hold" version, tolerance was ascertained if a woman drank more than six drinks she could tolerate, while the cutoff point is three in the "high" version. For "Worried, we asked whether the woman's close friends or relatives worry or complain about her drinking; for "Eye-opener", we asked a woman whether she has taken a drink first thing in the morning to get herself going; for "Amnesia", we asked a woman whether her friend or family member tell her about things she said or did while she was drinking that she could not remember; for "Kut down", we asked a woman whether she felt a need to cut down on her drinking. To calculate TWEAK score, we gave each woman one point per question if they answered yes to a question regarding using alcohol as an eye opener in the morning, memory loss, or cutting down on alcohol. Two points would be given

to a woman if she responded yes to the question regarding "Worried" or she meets the cutoff point of "Tolerance".

Additionally, the interviewer asked women their recreational drug use including marijuana/hashish, heroin, cocaine/crack, inhalants, methamphetamines, other recreational drugs. For each type of recreational drug, recreational drug users were asked to specify the exposure time, whether before pregnancy or one month prior to LMP or during the current pregnancy. Women were also asked whether they had taken methadone treatment and whether they had completed treatment before pregnancy or still undergone treatment during pregnancy. Questions about recreational drug use were introduced in January 2009. Therefore, information on recreational use for women who were recruited in 2008 was missing.

Medical and Reproductive Health

Information on medical and reproductive health was queried and confirmed by electronic medical records review: pre-pregnant weigh (pounds), pre-pregnant height (feet), gestational age at the time of interview, presence of medical conditions (if yes, women would be asked to specify the following medical conditions: hypertension, depression, diabetes, anxiety, seizure disorder, migraine headaches, thyroid disorder, rheumatoid arthritis, asthma or allergies, heart disease, cancer, hepatitis).

Questions about reproductive health included pregnancy planning (i.e., "Did you plan to get pregnant with this child?" The answers consist of three options:

"Yes"; "No, not now"; "No, not at any time"). At the beginning of this study, the first 42 study participants were not asked about their pregnancy planning, and there were additional 8 women who did report their pregnancy planning during the interview.

Other questions regarding reproductive health included a history of miscarriage, stillbirth, pregnancy termination, and ectopic pregnancy in a prior pregnancy, gravidity (number of pregnancies including current pregnancy), parity (number of live-born children), the date of the last menstrual period, a family history of birth defects. For a family history of birth defects, women would be firstly asked whether she or her members of immediate family or the immediate family of her baby's father had any babies with birth defects. If such history was reported, participants were asked to specify the following birth defects: Down syndrome, cleft lip or palate, neural tube defects, cystic fibrosis, heart defect, or other defects. We also investigated the presence of complications for current pregnancy including bleeding, high blood pressure, diabetes, and other complication and asked women whether they had morning sickness during the current pregnancy.

Vitamin and Dietary Supplement Use

All enrolled women were asked whether they had taken any vitamin, including multivitamin or single vitamin, prescription or over-the-count (OTC) vitamin, at least 4 times a week since they became pregnant. Also, women who reported using single vitamin or multivitamin were asked to specify brand name of vitamin and the time they had started vitamin use. Periconceptional vitamin

use was ascertained by asking women if they had taken vitamin at least 4 times per week during the month around their LMP.

Regarding dietary supplements, we asked a woman whether she took dietary supplements (including iron supplements) or herbal products on a regular basis since her last menstrual period. If she response "Yes", we further asked her to specify the type of herbal products, the frequency of dietary supplement use, and reasons for taking it.

Medication Use

The use of medications during pregnancy was also ascertained. To begin with, we asked a woman whether she discussed safety of medications in pregnancy with any health care provider (physician, nurse-midwife, physician assistant, or pharmacist). Then, we asked women about the use of OTC and prescription medications since LMP. For OTC medication, following OTC medications were provided: acetaminophen, aspirin, ibuprofen, ketoprofen, chlorpheniramine, benadryl, pseudoephedrine, kaopectate/pepto bismiol. Questions about prescription medications included pain/fever medication, nasal decongestants/allergy/cough medications, antidiarrheal, medication, and heart burn/dyspersia/antiemetic/laxative medications. For the attitudes of each medication use, we provided a five-point Likert type scale was used to assess the perception of teratogenicity.

In additional to medication use, we also investigated vaccine exposure since LMP (e.g. flu vaccine, other vaccine) and asked women to specify the types of

cravings or non-food item/strange good they took. Women were also asked about the use of fertility medications with current pregnancy (e.g. Clomid, Metrodin, Fertinex, or Pergonal).

Perinatal Outcomes

Perinatal data were abstracted from electronic medical records including pregnancy outcomes, i.e. live-born infant, spontaneous abortion (<20 gestational weeks), stillbirth (≥20 gestational weeks), termination, lost to follow-up. Date of delivery and gestational age at the end of pregnancy were also recoded. Type of delivery (vaginal-vertex, vaginal-breech, vaginal-transverse, cesarian section-primary, cesarian section-repeat) was obtained from medical records. Maternal complications in pregnancy included preeclampsia, pregnancy induced hypertension, oligohydramnios, infection or fever at delivery, gestational diabetes, and others, all of which were recorded in this database. For the live-born children, we recorded gender of the infant, birth weight (grams), birth length (centimeters), and birth head circumference (centimeters). In addition, the database also captured the neonatal complications (respiratory distress, hypoglycemia, tachypnea, bradycardia, sepsis, and others), diagnosis of major structural anomaly, the number of days for infant stay in hospital, and diagnosis of any abnormalities/conditions in the neonatal period.

Data Modifications

A multivitamin regular user was identified if a woman reported having taken vitamins at least 4 times per week. Based on the timing of the vitamin

exposure, we further categorized women into three study groups: 1) pre-conceptional vitamin regular users (women who reported having taken vitamins ≥4 times per week at least one month before their LMP); 2) vitamin regular users after pregnancy recognition (women who reported beginning to take vitamins or single vitamin at least 4 times per week since they became pregnant); 3) vitamin non-users (women who reported not having taken any vitamins before their last menstrual periods or since they became pregnant).

Potential predictors were identified from previous studies. We categorized in the following manner: maternal age was categorized into predictors categories (≥21 yrs, 21-30 yrs, >30 yrs); level of education was categorized into three levels (high school education or less, some college or vocational school, college degree or higher); marital status was recategorized into three categories (single/never married. married/living with partner, separated/divorced/widowed); for health insurance status, we made three categories (uninsured, employer-based or self-purchased insurance, Medicaid or other public insurance); language speaking was dichotomized as "Spanish" and "English and other language" (there were 6 women who speak other languages other than English and Spanish); for body mass index, the reported prepregnancy height and weight were used to estimate the BMI (kg/m²) and then BMI was dichotomized as obese (BMI≥30) and nonobese (BMI<30).

For gravidity and parity, they were dichotomized into two categories (primigravida if gravidty=1 vs. multigravida if gravidity≥2) and (primiparous if parity=0 vs. multiparous if parity≥1), respectively. A woman was identified to

have a history of adverse pregnancy outcome if she reported having experienced miscarriage, stillborn, pregnancy termination, or ectopic pregnancy in previous pregnancies. For pregnancy planning, a woman was considered to have a planned pregnancy if she answered "yes" to the question "Did you plan to get pregnant with this child", while a woman was regarded not having a planned pregnancy if she answered "no, not now" or "no, not at any time".

For the family history of birth defects, a woman was identified to have a family history of birth defects if she reported at least one of the following birth defects in her family: Down syndrome, cleft lip or palate, NTD, cystic fibrosis, heart defect, or other defects. For BMI, we recoded it as a categorized variable. Women were classified as obese if BMI equal or larger than 30 or nonobese if BMI is smaller than 30.

For smoking, we categorized women into three groups: women who smoked less than 100 cigarettes in a lifetime and did not currently smoke were identified as non-smokers; women who smoked 100 or more cigarettes in a lifetime but stopped smoking before pregnancy recognition were identified as a former smoker who quit before pregnancy recognition; women who currently smoked or who smoked ≥100 cigarettes or more than 100 cigarettes in a lifetime but stopped cigarettes after pregnancy recognition were identified as smokers during pregnancy. For alcohol use, periconceptional binge drinkers were identified if women have drunk four or more on one occasion during a month around their LMP. For recreational drug use, women were identified as

recreational drug users if they reported using any of following recreational drugs one month prior to LMP or during pregnancy: marijuana, heroin/hashish, cocaine/crack, inhalant, methamphetamines, or other recreational drugs. In addition, we also investigated prescription medications use during pregnancy and categorized pill burdens as follow: no prescription medications at all, 1-2 prescription medications, more than 2 prescription medications.

Statistical Approaches

SAS 9.2 (Cary, NC) was used for all analysis. A descriptive statistics were performed to describe the maternal characteristics in the entire population. Also, the prevalence of vitamin use in each study group was estimated. For vitamin users, we further specified the type of vitamin whether it is OTC or prescription vitamins. The most common brands of vitamins were determined based on women's self reported information. The number of days of vitamin use per week was additionally calculated. For women who began to take vitamins after pregnancy recognition, the beginning gestational age of regular vitamin use was measured using the date of last menstrual period as a starting time point.

The distribution of maternal characteristics among study groups was compared by using chi square test for categorized variables and ANOVA for continuous variables. All the significant predictors (p<0.2) were then entered into a multivariate ordinal logistic regression model to determine the association between vitamin use and maternal characteristics after adjusting

for confounders. Cumulative odds ratio with 95% Confidence intervals were constructed.

In ordinal logistic regression, we treated the three study groups as categorical under the assumption that there is a natural ordering across different vitamin groups. Therefore, the analysis built a set of cumulative logits to interpret the outcome (vitamin use). In total, two logits were developed as follow: Group 1 versus (Group 2 and Group 3), (Group 1 and Group 2) versus Group 3. The final model was examined by performing a chi square score test in order to check the proportional odds assumption. If the assumption is violated, then an alternative model, polychotmous logistic regression, would be used.

A polychotomous logistic regression model is an extension of dichotomous logistic regression model when there are more than two categories in the dependent variable. Unlike an ordinal logistic regression model, the dependent variable is treated as a nominal and cannot be arranged in any meaningful natural ordering. In this case, the interest outcome measure in this analysis is the vitamin use in different time of pregnancy: pre-conceptional vitamin users, vitamin users after pregnancy recognition, and non-users. Therefore, a multinomial model was introduced to determine whether maternal characteristics differed across three study groups. We did not use a series of binary models because this approach will give less statistical power to the results and higher values of standard error.

In this model, vitamin regular users after pregnancy recognition were the reference group. The rational of choosing this group as a reference group is that it this group has the largest frequency of cases and it makes the greatest clinical sense. Based on the reference group, there were two multinomial logits for polychotomous logistic regression: pre-conceptional vitamin users relative to vitamin users after pregnancy recognition and non-users relative to vitamin users after pregnancy recognition. For each logit, odd ratios and 95% confidence intervals (CI) of each predictor were estimated from the regression parameters and their standard errors.

Additional analysis was conducted by using ordinal logistic regression and polychotomous logistic regression with vitamin use as the response variable in order to examine interaction terms between different predictors, including ethnicity and race, ethnicity and place of birth, ethnicity and language speaking, race and language speaking, race and place of birth, place of birth and language speaking, education and health insurance status, education and pregnancy plan. Significant interaction terms would then be involved for the multivariate analysis.

Power Calculation

The power calculation for this study was done in PASS 11 Software (Kaysville, Utah) for comparing two independent proportions. For this calculation, we set the alpha value (the probability of rejecting a true null hypothesis) as 0.05 and 0.2 for beta (the probability of accepting a false null hypothesis). Sample

allocation ratio was set as three, which is approximately the ratio between group 1 and group 2 in the present analysis.

Estimations of the required total sample size were based on previously reported prevalence of pre-conception multivitamin users and prevalence of multivitamin users during pregnancy. In reference to the former prevalence, data from PRAMS (Pregnancy Risk Assessment Monitoring System), the prevalence of preconception vitamin use (P1) is 23.0-43.6% in the U.S. 18, 39, 47 ⁸¹ For the prevalence of regular multivitamin users during pregnancy, the prevalence was previously estimated to be 68.8-78.0%. 19, 35 Therefore, we assume that the difference in the proportion of multivitamin use between the study groups would be between 20% and 60%. As presented in Table 3.1, the required total sample size might vary from 24 to 225 depending of the difference in prevalence between study groups. In sum, a total sample size of 24-225 subjects will achieve more than 80% power to defect a difference between the effect sizes of 20-60% using the two-sided Z test at an alpha level 0.05. In this present analysis, a sample of 402 subjects will be sufficient and can achieve a power that more than 80%. To detect 20% difference, a group sample sizes of 252 could achieve 80% power.

CHAPTER IV: RESULTS

Description of the Study Population

Maternal characteristics of the total population were summarized in Table 4.1. The study population included 80.4% of Hispanics, 6.7% American Indian or Alaskan Native, and 5.7% Black or African American women. In terms of other demographic characteristics, a majority of women were immigrants (59.6%), identified Spanish as their primary language (60.7%), were married or living with a partner (71.6%), reported education less than high school (35.1%), and did not have any health insurance (48.8%). Additional analysis assessed the association between insurance status and ethnicity. Results of this analysis indicate a statistically significant association (p<0.01). Among women without health insurance, 98.5% of women were Hispanics and 1.5% of women were Non-Hispanics. Among women with public health insurance, more than half of participants (66.8%) were Hispanic women. For women have private insurance, the difference in ethnic distribution was small (Hispanic women, 55.1%; Non-Hispanic women, 44.9%). Of note, even though there more than half of women in this sample did not have any health insurance, all participants received free prenatal care through the UNM clinics.

The average gestational age for women to initiate prenatal care was 8.5±4.4 weeks after the last menstrual period. Most of the women (89.4%) received prenatal care in their first trimester (1-11 weeks after LMP), and 10.3% of women initiated prenatal care during the second trimester (12-24 weeks after LMP). There is one woman who reported starting prenatal care 29 gestational weeks. There were no differences in the mean gestational age at which

prenatal care was initiated between Hispanic (8.6±4.3 weeks) and Non-Hispanic (8.2±4.6 weeks) women (p=0.45).

Almost half of women (45.0%) reported the presence of medical conditions with diabetes being the most prevalent (20.1%). One third of the study population (32.1%) had a history of adverse pregnancy outcomes, such as miscarriage (32.1%), stillbirth (27.6%), terminated birth (7.5%), and ectopic pregnancy (1.8%) Regarding tobacco use, most of women (87.0%) were not current smokers or quit before pregnancy recognition, while only 13.0% of participants reported smoking during some point of pregnancy.

Binge drinking (i.e., at least one episode of consuming ≥4 drinks per occasion) during a month around the last menstrual period was reported by 23.1% of women. Over half of women (53.4%) reported using prescription medications during pregnancy, and 29.1% of women who used medications did not discuss their safety in pregnancy with a health care provider.

Among 200 women who were asked questions about illicit drug use, 55 (27.5%) reported lifetime use of at least one type of drugs and 11 (5.5%) women reported use during the month prior to LMP or during the current pregnancy. Among 352 who answered a question about pregnancy planning, 181 women (51.4%) reported unplanned pregnancy.

Vitamin Use Patterns

The vitamin use patterns were summarized in Table 4.2. In this population, 21.4% of women were identified as pre-conceptional vitamin users, 71.9% of women began to take vitamin after they realized they were pregnant, and 6.7% of women did not use any vitamins during pregnancy. Of note, there were three women who began to take vitamin regularly in pre-conceptional period but stopped taking after pregnancy recognition. All of the three women reported a family history of birth defects (two women with a family history of Down syndrome and the other woman with a family history of cleft lip/palate), but reasons for this behavior change were not captured by this study. Among vitamin users after pregnancy recognition, only one woman used a single vitamin (folic acid), while the rest reported multivitamin use. This woman was included into the study Group 2 for analysis. Among all vitamin users, 58.4% of women used OTC multivitamins and 30.9% of women reported using prescription multivitamins. The two most frequently identified brands of prenatal vitamin were Walmart® and Walgreen® prenatal vitamins. Both of these brands contain 800 microgram of folic acid. Other brands identified by participants included "One a day Prenatals", "Natalcare Plus", and "Flinestone vitamins". On average, vitamins were taken 6 days (s.d=1.9) per week during the week before the interview. Among women who took vitamin after pregnancy recognition, the average initiation time was 9 weeks after the LMP.

Maternal Characteristics of Vitamin Use

The distributions of maternal characteristics among the three study groups were summarized in Table 4.3. Only 17.3% of Hispanics took vitamin in

pre-conceptional period, while most of them began to take vitamins only after pregnancy recognition. There was a significant difference in the distribution of education level among the three study groups. A decreasing trend of the proportion of women with a college degree or higher education was identified: 50% of women with a college degree or higher education were identified among pre-conceptional vitamin users, followed by vitamin users after pregnancy recognition (47.5%) and vitamin non-users (2.5%). Interestingly, the largest proportion of women with some college or vocational school education was identified among vitamin users after pregnancy recognition (66.3%), followed by pre-conceptional vitamin users (23.6%). Association between the three study groups with respect to vitamin use and health insurance was significant. The proportion of women who had Medicaid or other public insurance among pre-conceptional vitamin users (44.9%) was closed to the proportion in vitamin users after pregnancy recognition (49.0%). In sum, there were significant associations among the three study groups with respect to ethnicity, education level, health insurance, language speaking, and pregnancy planning. Maternal age, race, marital status, primigravida, nulliparous, place of birth, history of adverse pregnancy outcomes, presence of medical conditions, a family history of birth defects or adverse pregnancy outcomes, use of prescription medications, smoking status, periconceptional drinking, periconceptional drug use, were not significantly associated with vitamin use in this sample.

Ordinal Logistic Regression

Results of ordinal multivariate logistic regression with cumulative odds ratios (ORs) and 95% confidence intervals (95% CI) for each predictor were summarized in Table 4.4. Cumulative ORs represent increasing odds of being early vitamin users. The results of multivariate analysis showed that maternal age, race, history of adverse pregnancy outcomes, country of birth, and use prescription medications were not significant associated with vitamin use. Moreover, ethnicity, education level, health insurance, and language speaking became not significant after adjusting for other variables in the model. The only significant predictor associated with vitamin use was pregnancy planning, suggesting that the odds of being vitamin users at an earlier stage in pregnancy were 1.76 times higher for women who had pregnancy planning than those women who did not have a pregnancy planning.

Interactions between predictors were examined using the ordinal logistic regression model and results were summarized in Table 4.5. Of note, there were no women who identified themselves as Non-Hispanics and spoke Spanish at the same time. Therefore, the interaction between ethnicity and language speaking could not be calculated. The only significant interaction was identified between education level and insurance status (p<0.05), meaning that the effect of education on vitamin use varied by the level of health insurance status. Thus, the association between education and vitamin use was examined after stratification by insurance status. Results are presented in Table 4.5 (women without health insurance), Table 4.6 (women

with employer-based or self-purchased insurance), and Table 4.7 (women with Medicaid or other public insurance).

Results of the stratified analysis demonstrate that there were no associations between maternal education and vitamin use among women without health insurance and women who had private or self-purchased health insurance. However, among women with public health insurance, having a college degree or higher was associated with greater odds of early vitamin use (OR=14.45; 95% CI: 1.78-117.66) compared to women with a high school education or less.

Due to the limited sample size in each category of health insurance status, the present analysis explored the interaction between education and health insurance by categorizing health insurance into a dichotomous variable (any type of insurance vs. none). Results were presented in Table 4.8 and Table 4.9, respectively. Among women without insurance, education level was not a significant predictor. However, among women with any type of insurance, the odds of earlier vitamin use were greater among women with a college degree or higher than women with a high school education or less (OR=5.90, 95% CI:1.89-18.44).

Polychotomous Logistic Regression

The test for the proportional odds assumption resulted in a p=0.15. The null hypothesis for the proportional odds assumption is that the assumption is met. Given that p=0.15 is greater that a usually used cut-off point of p=0.1, the H_0 is

not rejected and once can conclude that the proportional odds assumption is met. However, given that the p-value was close so close to the cut-off point, polychotomous logistic regression was also conducted in additional to the ordinal logistic regression.

All the significant covariates identified in univariate analysis at p<0.2 were considered as potential predictors of vitamin use and were included in a polychotomous logistic regression. As shown in Table 4.10, in the first logit of the unadjusted polychotomous logistic regression (pre-conceptional vitamin users vs. vitamin users after pregnancy recognition), significant predictors were ethnicity, education, health insurance, pregnancy plan, language speaking, and use of prescription medications during pregnancy. Specifically, women who were pre-conceptional vitamin users were less likely to be Hispanics (OR=0.33, 95% CI=0.19-0.57), speak Spanish (OR=0.43, 95% CI=0.27-0.71), more likely to have a college degree or higher education (OR=4.94, 95% CI=2.44-10.00), have employer-based or self-purchased insurance (OR=3.90, 95% CI=1.97-7.75), and use ≥2 prescription medications (OR=2.33, 95% CI=1.14-4.78) compared to women to initiated vitamin use later in pregnancy. Maternal age, race, and history of adverse outcomes were not significant in this analysis (p>0.05). In the second logit (non-users vs. vitamin users after pregnancy recognition), race was the only significant predictor. Specifically, non-users were more likely to be in the "other" racial groups (OR=3.35, 95% CI=1.13-9.92) compared to vitamin users after pregnancy recognition after adjusting for other risk factors.

Results of multivariate polychotomous logistic regression were present in Table 4.11. After adjusting for ethnicity, race, education level, language speaking, history of adverse pregnancy outcomes, place of birth, and use of prescription medications during pregnancy, pregnancy planning was the only significant predictor of using vitamins before pregnancy recognition as compared to vitamin user after pregnancy recognition (OR=2.29, 95% CI=1.29-4.09). Race was no longer significant in multivariate analysis for comparison of non-users vs. users after pregnancy recognition. However, a history of adverse pregnancy outcomes became a significant predictor: non-users are more likely to have a history of adverse pregnancy outcomes comparing to vitamin users after pregnancy recognition (OR=3.04, 95% CI=1.16-7.98).

For interaction terms, Table 4.5 summarized the results and showed that there were no significant interaction effects between ethnicity and race, ethnicity and place of birth, race and language speaking, ethnicity and place of birth, place of birth and language speaking, education and health insurance status, education and pregnancy plan.

CHAPTER IV: DISCUSSION

Summary

This study found that most pregnant women (71.9%) in the sample began to take vitamins regularly after pregnancy recognition. For women who started to use vitamins after pregnancy recognition, the mean gestational age of initiating vitamin use was 9 weeks. Women on average took vitamins 6 times per week. However, only 21.4% of study participants began to take vitamins before pregnancy, thus had vitamin exposure in early gestation even before pregnancy recognition. Most women reported using OTC vitamins (54.2%), with two most popular brands being Walgreen® and Walmart® prenatal vitamins, both containing 800mg of folic acid.

Hispanics were less likely to use vitamin early in pregnancy than non-Hispanics: however, ethnic difference in vitamin non-significant after adjusting for other maternal characteristics. Unplanned pregnancy is a big public health issue: half of participants, similar to national estimates. 94 in this study identified their pregnancies as unintended. Earlier vitamin use in pregnancy was significantly associated with pregnancy planning independent of other maternal characteristics. In addition, education level and health insurance status revealed a significant interaction with respect to vitamin use. Among women with public health insurance, the odds of early vitamin use were greater among women with a college degree or higher education than women with a high school education or less than high school. In addition, almost one third (32.1%) of women in this study had a history of adverse pregnancy outcomes and 45.0% of women had at least one medical condition. In multivariate analysis, vitamin non-users were more likely to have experience of adverse pregnancy outcomes in prior pregnancies compared to women who used vitamin after pregnancy recognition. This observation requires investigation in future studies.

Prevalence of Vitamin Use as Compared to Other Studies
In this study, 92.5% of women reported regularly using vitamin anytime during pregnancy. This prevalence rate is higher comparing to a cross-sectional study using data from the Behavior Risk Factor Surveillance System (BRFSS), in which 78% of women from 14 U.S. states and territories reported using multivitamins in pregnancy. For vitamin use among Hispanics, the present study identified that 76.2% of Hispanics were vitamin users anytime during pregnancy, while this prevalence in the BRFSS study was 66.4%. Although the estimation used a national sample, the BRFSS database did not contain information on the duration of multivitamin use, doses, contents, or frequency of use.

In additional to BRFSS, vitamin use during pregnancy was also investigated using data from the National Maternal and Infant Health Survey (NMIHS). This national survey defined vitamin use as vitamin/mineral supplement use at least 3 days a week. This definition was somewhat comparable to the present study (vitamin use at least 4 times per week). There were two cross-sectional studies which conducted analyses based on the NMIHS data. One of these studies was limited to mothers of live-born infants and reported that 82.5% of women used vitamin anytime during pregnancy, 37 while the prevalence was 74.2%

when the sample included women with miscarriages or women who terminated their pregnanciesl.³⁶ However, both of the prevalence estimates included mineral supplement use. Therefore, women who reported using mineral supplements only could be also included in these prevalence estimates. These studies have not reported the prevalence of vitamin use among Hispanics though, which comprised the majority of the sample in the present study.

Prescription claim databases were also used to estimate the prevalence of vitamin use during pregnancy. A population-based study analyzed data from the Kaiser Permanete Medical Care Program (KPMCP) in North California and reported that 69.0% of women used vitamin during pregnancy. As compared to the results of the present study, prevalence in the KPMCP was lower. The difference might be due to the different gestational age at the interview time. The KPMCP study recruited women at an earlier stage of pregnancy: the mean gestational age at study entry was less than 5 weeks, while the mean gestational age at current analysis was 30.7 weeks. The Pregnancy, Infection, and Nutrition (PIN) Study in North Carolina recruited women from four prenatal care clinics and reported that the prevalence of vitamin use anytime in pregnancy was 84%. However, neither KPMCP study nor PIN study reported vitamin use among Hispanics. Information on the duration of vitamin use and frequency was not reported in these two studies.

Regarding vitamin use after pregnancy recognition, 71.9% of women in this present analysis began to take vitamin after pregnancy recognition, which is higher comparing to two previous prospective cohort studies in the U.S

(44.8-48.9%).^{25, 35} One of these prospective cohort studies, however, interviewed women who were at 24-29 weeks of pregnancy, and it potentially excluded women at third trimester. Thus, the estimation of vitamin use might not include women at higher risk of adverse birth outcomes. For the other prospective cohort study, nearly half of women (49.4%) refused to participate in the study. Furthermore, among all the participants, there were 23.2% of women who did not complete an interview. Therefore, the result of vitamin use after pregnancy recognition might suffer from a selection bias.

With respect to vitamin use in early pregnancy/pre-conceptional period, only 21.4% of women reported using vitamin in this present study. Similar to this result, there were several Pregnancy Risk Assessment Monitoring System (PRAMS) reports which investigate the vitamin regular use (≥4 times per week) in pre-conceptional period. Centers for Disease Control and Prevention (CDC) introduced PRAMS in 1987, and PRAMS is an ongoing surveillance system in order to monitor maternal behaviors that occur before, during, and after pregnancy.³⁹ Several reports analyzed PRAMS data covered from 2000 to 2003 and reported that the prevalence of pre-conceptional vitamin regular user ranged from 25.0 to 45.2% based on 19 states in the U.S. 18, 39 Another PRAMS report limited the sample to women in Oklahoma and found the prevalence fell between this range (26.5%) from 2000 to 2003.81 Unfortunately, the prevalence remained relatively unchanged. The latest PRAMS report summarized data for 2003-2004 from 26 states and demonstrated that 35.1% of women on average reported using vitamin in pre-conceptional period.⁴⁷ Similar to the result of PRAMS, a state report using data from the Missouri

Pregnancy Related Assessment and Monitoring System (MoPRA) found that the prevalence of pre-conceptional daily vitamin use was 29.7%. ⁸⁶ These results are comparable to our findings.

However, the results from PRAMS and MoPRA analyses might be have limited generalizability since they only include women with live-born infants. Therefore, the prevalence of pre-conceptional vitamin users might be overestimated, given to a possibility that the sample did not include women who had a miscarriage or stillbirth and did not use prenatal vitamins during the pre-conceptional period. In addition, all the identified PRAMS reports and the MoPRA study did not investigate pre-conceptional vitamin use by race/ethnicity.

The prevalence of pre-conceptional vitamin use in this present analysis, to some extent, might reflect the prevalence of periconceptional vitamin use. The definition of periconceptional period varies across the studies and might refer to one month before and one month after the LMP or might include as many as three months before and after LMP. Women who reported using vitamin before pregnancy are more likely to continue using vitamin after they were pregnant. A cross-sectional study, which used data from the NMIHS, reported that the prevalence of periconceptional vitamin use (≥3 times per week during three months before and three months after pregnancy recognition) was 21%. Similarly, a Kaiser Permanete Medical Care Program (KPMCP) prospective cohort study reported 23.8% prevalence. Therefore, the prevalence of periconceptional vitamin use and pre-conceptional vitamin use were similar.

The difference in maternal characteristics might explain variability in the prevalence of periconceptional vitamin use. The PIN study reported that the prevalence of periconceptional vitamin use (before and during pregnancy) was 30.0%. The higher prevalence rate of pre-conceptional vitamin use in PIN study can be explained by the fact that this study included women who only spoke English and that most of women (64.5%) in this study were Non-Hispanic whites. Moreover, women who planned the pregnancy might be more likely to use vitamin in periconceptional period. The NBDPS study demonstrated that the prevalence of periconceptional vitamin use among women who planned their pregnancy was 64.3%. The PIN study reported that the prevalence of periconceptional vitamin use among women who planned their pregnancy was 64.3%.

There is also a difference in the prevalence of periconceptional vitamin use among different racial groups. For Hispanics, data obtained from the National Birth Defects Prevention Study (NBDPS) showed that the prevalence of periconceptional intake of folic acid-containing supplements is only 30% among Hispanics, which is the lowest compared to other racial groups (Non-Hispanic White: 66%; Non-Hispanic Black: 39%).⁴³ For other races, a secondary analysis using data from the Slone Epidemiology Center Birth Defects Study (SECBDS) showed that 55.0% of Non-Hispanic White women were periconceptional vitamin users (used vitamins ≥ 4 times per week during lunar months -1 to 1), while this prevalence was only 25.6% among African American women.⁴²

There are many potential reasons for lower prevalence of vitamin use among Hispanic women compared to Non-Hispanic Whites. One possible explanation might be the lower awareness about benefits of folic acid in prevention of birth defects and other adverse perinatal outcomes among Hispanic women. A household survey conducted in predominantly Hispanics neighborhoods of inner-city Hartford, Connecticut demonstrated that 78% of Hispanics (both women and men) are not aware of NTDs, including spina bifida. Another survey among Hispanic women of childbearing age conducted in Michigan and showed that nearly half of study participants (45%) did not know about the time window to prevent birth defects and 59% did not take daily multivitamins before pregnancy recognition.

With respect to the type of vitamins, more than half of women (54.2%) used OTC vitamins during pregnancy, and 28.7% of women reported using prescription vitamins. This result was quite different from a previous prospective cohort study, which recruited pregnant women from rural outreach clinics in the U.S. The result showed that 92.0% of pregnant women used prescription prenatal multivitamins and 10.9% of them used OTC multivitamins. This discrepancy might be due to the different sampling method. Glover's study recruited all the women from obstetric patients who were seen by private physicians, while a majority of study participants in our study came from the university-affiliated community clinics providing free prenatal care. The difference in the health insurance coverage could account for the discrepancy between the two studies. In Glover's study, all of the

participants had heath insurance (e.g. Medicaid, private insurance), while almost half of women (48.8%) in our study did not have any health insurance.

The average time of pregnancy recognition is 9.0 gestational weeks. The time interval between the LMP and the pregnancy recognition is a curial period for organogenesis. During the month of fertilization, cells along the dorsal surface of the embryo develops into a groove and then a hollow tube.²⁹ If this process is not completed, a neural tube defect develops. A number of prior studies suggested that periconceptional use of multivitamin supplementation containing folic acid (from 3 months before and through 3 months after conception) can significantly lower the risks of NTDs.^{2, 9-16} It has been strongly recommended that all women of childbearing age should consume 400 µg of folic acid daily given that 50% of pregnancies are unplanned.²² Regular multivitamin use (≥4 times per week) provides the recommended amount of folic acid.³⁹ Nevertheless, in the present study, the result showed that most women initiated vitamin regular use at 9 weeks after LMP, which is after the closure of neural tube (six weeks after the last menstrual period). In our study, there were only 21.4% of women who initiated vitamin use before pregnancy recognition.

Predictors of Vitamin Use as Compared to Other Studies

Race/Ethnicity

The sample in the present analysis included a large proportion of Hispanics women, but ethnic difference in vitamin use was non-significant after adjusting for other maternal characteristics. The review of published literature showed

that findings among different studies with respect to race/ethnicity are inconsistent. A cross-sectional analysis of the BRFSS data did not demonstrate that ethnicity was a significant predictor for vitamin use among pregnant women. 19 The analysis conducted in the MoPRA study also yielded non-significant results after adjusting for other predictors.86 However, other studies reported race/ethnicity to be a significant predictor. A study from the Slone Epidemiology Center Birth Defects Study suggested periconceptional vitamin users (-1 to +1 month around LMP) were more likely to be non-Hispanic Whites. 42 Another cross-sectional analysis, using data from the NMIHS, found that vitamin non-users were more likely to be African American and Asians. However, the NMIHS study mentioned above did not assess the effect of Hispanic ethnicity on vitamin use.³⁷

In terms of predominant language, Spanish was not a significant predictor of vitamin use in the present study. This result contradicted the findings from a cross-sectional study which used data from the National Birth Defects Prevention Program (NBDPP). In that study, women who did not use vitamins in periconceptional period (3 months before conception and one month after conception) tended to speak Spanish. However, the difference in the proportion of women who spoke Spanish may cause this difference. There were only less than 5% of Spanish-speaking women (n=80) in the NBDPP study, and the odds ratio was quite wide (OR=2.0-7.2), while our population included 60% Hispanics. In addition, the present study explored the relationship between place of birth and vitamin use. A population-based case-control study, using data from California birth cohort, found that women

who were Mexico descents had an increased risk of having a NTD-affected pregnancy.⁴⁶ However, this study did not specify the place of birth for the participants. In this present analysis, this demographic characteristic was not a significant predictor of vitamin use in pregnancy.

Maternal Education

Education was not a significant predictor of vitamin use during pregnancy or in pre-conceptional period after adjusting for other maternal characteristics. The result might due to the small sample of women with a college degree of higher in our samples. However, our results are consistent with several cross-sectional studies in the U.S. A cross-sectional analysis, using the 2004 data from the BRFSS, found that education was not a significant predictor of vitamin use during pregnancy. Another report using the MoPRA data also confirmed that education level was not significantly associated with pre-conceptional vitamin use (one month before pregnancy recognition) after adjusting other variables in the analysis. 86

However, other studies found an association between vitamin use and higher education level. For vitamin use anytime in pregnancy, a national survey initiated by the Medical Birth Registry in Brazil and a cross-sectional analysis using data from Norwegian Mothers and Child Cohort Study (MoBa) found that vitamin users during pregnancy tended to have a college/university degree. To periconceptional vitamin use, a cross-sectional analysis using data from the Slone Epidemiology Center Birth Defects Study and the 2001 New South Wales Child Health Survey in Australia found that periconceptional vitamin

user (-1 to +1 month around LMP vs. -2 to +3 month around pregnancy recognition) were those women with higher education level or more years of education. ^{42, 97} In addition, a PRAMS data and a cross-sectional analysis using data from NMIHS found that women with at least 12 years of education were significantly more likely to be pre-conceptional vitamin users (≥4 times per week during the month before pregnancy) than women with less than high school education (<12 years of education).³⁹

The present analysis observed a significant interaction between health insurance and education level in respect to vitamin use during early pregnancy. Among women with public insurance, patients with a college degree or higher education were more likely to be vitamin earlier users than women with only high school or less than high school education. Among women who had private insurance or did not have insurance, the present study did observe any significant association between education and vitamin earlier use in pregnancy. We did not find other studies which examined this interaction. Nevertheless, a PRAMS report used data from 19 states and found that pre-conceptional vitamin users (≥4 times per week during the month before pregnancy) were more likely to have a private health insurance and have more than 12 years of education, but it did not examine the interaction regarding pre-conceptional vitamin use.³⁹

Health Insurance

Even though there was a significant interaction between health insurance and maternal education, health insurance status by itself was not a significant

predictor of pre-conceptional vitamin use in the present analysis. Prior reports for this association are inconsistent. A cross-sectional analysis, using data from MoPRA, examined the insurance status of pregnant women, but this predictor was not significant. 86 This result did not concur with the results of two PRAMS reports, which used data during 2000-2003 in Oklahoma and 2003-2004 in 26 states, respectively. One of these reports found that the highest prevalence of pre-conceptional vitamin users were women with private health insurance. The other study further adjusted all other demographic characteristics and found that pre-conceptional vitamin users were more likely to be women with private health insurance than women who were recipients of Medicaid. 47, 81 It is worthy mentioning that the MoPRA study only asked women whether they had insurance or not, but it did not specify the type of insurance, which might cause the different result. The difference in findings between the PRAMS reports and current study might be due to various maternal characteristics of the study population between PRAM and the SMART study. There were no other studies from Europe or Australia that investigated the vitamin use in earlier pregnancy and health insurance status because of the different national health care system.

Marital Status

With respect to marital status, this present analysis did not observe any significant difference among the three study groups. A survey of postpartum women in Toronto Hospital also did not detect a significant association. However, a number of analyses presented a contrast to this result, suggesting that married women or women living with a partner were more likely to be

vitamin earlier users during pregnancy, ^{19, 42, 78, 82, 85, 86} even though the maternal characteristics varied by different studies. It is noteworthy that the present analysis included a small number of women who were separated or divorced. This might cause a non-significant result.

Maternal Age

Maternal age, measured as a continuous variable, was non-significant in the present study. A cross-sectional study, recruiting women in the antenatal clinics and birth center in Australia, also measured maternal age as a continuous variable, but did not find it to be significant. However, a prospective cohort study obtained data from the Pregnancy Exposure and Preeclampsia Prevention Study in the U.S. and found that maternal age as a continuous variable was a significant predictor of periconceptional vitamin use (-3 to +3 months around LMP) from a chi square test. However, this study did not calculate the odds ratio and did not adjust other maternal characteristics.

Even though the present analysis categorized this predictor into three age groups, the present analysis still did not detect a significant result. Another study in Canada surveyed postpartum women in Toronto Hospital and categorized maternal age as three groups (<16-25, 26-30, 31-35),⁸⁰ also suggesting a non-significant result and being consistent to an analysis using data from the BRFSS in 2004.¹⁹ On the contrary, there were also some studies pointing out that older age group was associated with vitamin earlier use in pregnancy.^{37, 39, 43, 47, 73, 81, 86} A potential reason for the different result between these studies and the present study is due to the various maternal

characteristics in different countries and areas. Also, the different way to categorize maternal age may account for the different result. A cross-sectional study in the U.S. conducted at MoPRA categorized women into two age groups (<20 years, ≥20 years) and detected a significant result with respect to pre-conceptional vitamin use.

Pregnancy Planning

Pregnancy planning identified by the present analysis was the only significant predictor that associated with earlier vitamin use. This finding confirmed the result from a cross-sectional study in the U.S. conducted at MoPRA. This study suggested that planning of pregnancy was significantly associated with pre-conceptional prenatal multivitamin intake (OR=2.04, 95% CI=1.45-2.94) when the study adjusted other maternal characteristics. ⁸⁶ In addition, a study obtained data from PRAMS during 2003-2004 reported that the highest prevalence of pre-conceptional vitamin users were among women with pregnancy intention. However, the PRAMS did not report the strength of association. ⁴⁷ For other studies in the U.S., analyses using data from BRFSS, NMIHS, and PRAMS did not involve pregnancy planning as a potential predictor of vitamin use. ^{19, 37, 39, 81}

Although there are differences in maternal characteristics between various populations in different countries, several studies from other countries also detected a significant association between earlier vitamin use and pregnancy planning. A cross-sectional study from Australia used data from a population-based survey (the 2001 New South Wales Child Health Survey)

and reported that women with an unplanned pregnancy (OR=0.15, 95% CI=0.08-0.26) were significantly less likely to take periconceptional folic acid supplementation (≥1 months before and 3 months after pregnancy recognition). A survey conducted at the Department of Obstetrics and Gynecology and in the Pregnancy Care Centre in Hungary also confirmed this result: planning of pregnancy (OR=4.22, 95% CI=2.61-6.84) was significantly associated with periconceptional vitamin intake (before and during pregnancy). Of note, in the survey, most participants (83.7%) had a pregnancy planning, which is higher than the present analysis. Also, the authors did not specifically defined periconceptional period. A cross-sectional analysis from Canada defined periconceptional period (≥ 4 weeks prior to conception until 8 weeks after conception) and surveyed a group of postpartum women in Toronto Hospital. The result also supported that unplanned pregnancy (OR=1.5, 95% CI=1.4-1.6) was significantly associated with a lack of folic acid use in periconceptional period.

Interestingly, pregnancy planning was not statistically associated with vitamin use after pregnancy recognition in the present study. This result may support the result of a cross-sectional study in Portugal. The study recruited mothers of live-born infants from two primary hospitals in the North of Portugal, but it did not observe a significant association between pregnancy planning and multivitamin use in pregnancy (OR= 0.96, 95% CI=0.89-1.03). Admittedly, there were other studies that also investigated the predictors of vitamin use in pre-conceptional or periconceptional period, 74, 75, 77, 82, 85 but none of these

studies measured pregnancy planning for women or did not involve this predictor for multivariate analysis.

History of Adverse Perinatal Outcomes

In this study, vitamin non-users were more likely to have a history of adverse pregnancy outcomes comparing to vitamin users after pregnancy recognition (OR=3.04, 95% CI=1.16-7.98). This result contradicted to a cross-sectional study using data from the National Birth Defects Prevention Study (NBDPS). In this analysis, women who did not take vitamins in periconceptional period (3 months before conception and 1 month after conception) tended to have no previous miscarriage (OR=0.7 95% CI=0.5-0.9).43 However, non-users in the present analysis were those who did not use vitamin anytime during pregnancy, while this NBDPS report limited vitamin non-users to women who did not use vitamin in periconceptional period. Moreover, the present analysis not only investigated miscarriage but also other previous adverse pregnancy outcomes, such as ectopic pregnancy, stillbirth, and elective pregnancy termination. Difference in maternal characteristics might also account for the discrepancy. All the women recruited from the NBDPS were mothers with live-born infants without major malformations. There were other studies which also investigated the association between vitamin use during pregnancy and previous adverse pregnancy outcomes. One report obtained data from the Birth Registry in Brazil examined the relationship between vitamin use during pregnancy and previous experience of miscarriages as well as stillbirths. 77 However, the result was insignificant. Also, a study surveyed women in the antenatal clinics and birth centre in Australia, but the association between periconceptional vitamin

use (before and during pregnancy) and a history of adverse pregnancy outcomes was not signflicant.⁷⁵

It is still unclear how previous adverse pregnancy outcomes associated with vitamin use in pregnancy. The most likely explanation is that those non-vitamin users did not use vitamin in the prior pregnancy either. However, the SMART study did not investigate the reasons of not taking vitamins. Therefore, neither of the two possibilities could be confirmed in this study.

Other Predictors

Other predictors associated with reproductive health included a family history of birth defects, presence of chronic conditions, parity, and gravidity. Unfortunately, none of them was significant in the present analysis. A prospective cohort study, analyzing data from the Pregnancy Exposure and Preeclampsia Prevention Study, also did not detect a significant relationship between a family history of preeclampsia or hypertension and periconceptional vitamin use (-3 to 3 months around LMP).²⁴ For the presence of chronic disease, although the SMART was inclusive of possible maternal medical conditions, the present analysis still did not detect a significant association. A prospective cohort study examined the presence of hypertension and periconceptional vitamin use, but the result was also not significant. A PRAM report examined data from 19 reporting areas during 2000 suggested that the association between the presence of gestational diabetes pre-conceptional vitamin use was not significant.³⁹ For parity, there were controversies about the relationship between parity and vitamin earlier use in

pregnancy. One study in the U.S. used data from the National Birth Defects Prevention Study found that women who did not take vitamin in periconceptional period (-3 months to 1 month around conception) tended to be nulliparous. Another study in Australia, analyzing data from the Victoria Survey of Recent Mothers 2000 and the 2001 NSW Child Health Survey, found a different result: periconceptional vitamin non-users (-2 to 3 months around pregnancy recognition) were multiparous. The difference in maternal characteristics between two countries might create this discrepancy.

Smoking during pregnancy was not a significant predictor. There were several studies in the U.S. found that women who were not current smokers were tended to be vitamin users in pregnancy, ^{42, 43, 74, 86} but none of these studies specified the time point when women stopped smoking. For alcohol and recreational drug exposure in periconceptional period, the present analysis did not observe any significant relationship with vitamin earlier use in pregnancy. A study, using data from the National Birth Defects Prevention Study, also examined yielded a similar result.⁴³ Of note, this study focused on any alcohol and recreational drug exposure from the three months before conception through the time of delivery, while the present study limited drink exposure to binge drink (≥4 drinks on one occasion) and recreational drug in periconceptional period (the month around the last menstrual period).

Limitations

The first limitation of this study was that we relied on the self-reported information from women. There is a possibility of outcome misclassification,

because women might have had difficulty remembering their behaviors regarding vitamin use before the interview or early in pregnancy. Recall bias might cause missing values for some maternal characteristics and information of vitamin use patterns. For instance, some women did not identify their family history of birth defects or adverse pregnancy outcomes. However, comparing to maternal characteristic, the present study had more missing values for vitamin use patterns. Some women failed to identify the type or brand of prenatal vitamins. There were also some women who could not specify the date when they began to take vitamins. Similarly, due to the sensitive nature of some questions (e.g. marital status, recreational drug use, periconceptional binge drink), there is a possibility that women felt reluctant to report or might not report their maternal behaviors accurately. In addition, potential interview bias might also occur and impact on the validity of self-report information.

A second limitation of this study was that it did not capture the maternal characteristics and vitamin use patterns for all the participants due to the nature of the SMART study. The late introduction of some questions also caused missing values for the present analysis. For instance, more than half of participants did not have information on recreational drug use. Also, the SMART study did not investigate pregnancy planning for the first 42 women. The incomplete information of maternal characteristics might cause loss of valid sample size for the analysis, thus reducing the power to detect significant predictors. Furthermore, the SMART study did not capture all the relevant maternal characteristics for vitamin use during pregnancy. For example, this study did not ask women about their house income, while this demographic

characteristic is a potential predictor of using vitamins during pregnancy. Given to the large proportion of women without any health insurance and recipients of Medicaid, it is likely that the study population included a large number of women at low socioeconomic status. For vitamin use patterns, the SMART study did not follow up vitamin use after the baseline interview. Therefore, this analysis could not estimate the duration of vitamin use during the entire process of pregnancy. In addition, the SMART study did not ask participants about the reason of vitamin use or non-use.

The inclusion criteria of the SMART study affected the estimation of vitamin use among all the pregnant women for the present analysis. The SMART study ascertained eligible women with known pregnancy status, but it did not include women with miscarriages that occur prior to this study. It is important to explore their vitamin use during pregnancy, for most of these women might be vitamin non-users.

In terms of generazability, the present study might limit the findings to the women in New Mexico. The ethnic distribution was quite different from that in the BRFSS and the PRAMS, for there were a large proportion of Hispanics women in the present analysis. Additionally, the SMART study might not be fully representative of all pregnant women in New Mexico, for it only captured women who did attend UNM-affiliated prenatal care clinics.

Strengths

The first strength of this study is that the validity of data from the SMART study is high. There is a high degree of intended cooperation, given to the fact that this study had a high participation rate (81.5%). For non-participation, the SMART study also asked them to specify the reasons. All the interviewers in the SMART study have previous experience of conducting interviews, and all of them received a standardized training before the study. Thus, we have reduced the interviews bias at the minimum level. Moreover, we realized that there were a large number of Spanish-speaking women in New Mexico. Hence, all the interviewers are bi-lingual and fluent in English in order to eliminate the language barrier of communication.

The second strength of this study is that it provides comprehensive information of vitamin use pattern. Firstly, we estimated the prevalence of vitamin use on a regular basis, which is meaningful. Prior CDC report demonstrates that multivitamin use at least four times per week can provide the recommended amount of folic acid.³⁹ Secondly, we categorized vitamin use in different time of pregnancy and created three mutually exclusive vitamin user groups. Based on this classification, we compared the different in maternal characteristics among three groups. In addition, the study showed the type and the most popular brands among vitamin users. Thirdly, for vitamin users after pregnancy recognition, we attempted to estimate the time gap between the last menstrual period and the initiate time of vitamin use. Knowing this time gap could help us determine whether women missed the appropriate time for vitamin intake. Lastly, although this study did not capture women who had a miscarriage

before enrollment, the present analysis included women who had pregnancy ended in a stillbirth (no live birth; gestational age<20 weeks).

Thirdly, this is the first study which adopted an ordinal logistic regression model and polychotomous logistic regression model for the analysis. With the ordinal logistic regression model, we assumed that there was a natural ordering across three vitamin user groups, and we were able to identify the significant predictor associated with earlier vitamin use. The use of polychotomous logistic regression enables this study to compare the difference in maternal characteristics between pre-conceptional vitamin users and vitamin users after pregnancy recognition. Additionally, the present analysis is the first one which attempted to examine interactions between two predictors.

The fourth strength of this study is that the population in SMART study has an ethnically and socially-diverse sample. The present study is the first one which investigated vitamin use in pre-conceptional period and after pregnancy recognition using a sample primarily with Hispanic women.

Future Directions & Recommendations

For vitamin use among pregnant women, future studies should clearly define "pre-conceptional period", "periconceptional period", and "anytime during pregnancy". The difference in these definitions can cause various results, including both prevalence and predictors. Also, future studies need to select the study population carefully. Only including mothers of live-born child might

overestimate the prevalence of vitamin use in pregnancy. It is highly possible that the prevalence could be lower if the sample also included women with a miscarriage. In addition, to ascertain vitamin use, it is meaningful to define a vitamin user as someone who takes multivitamins at least 4 times per week. This meets the recommended amount of folic acid to prevent NTDs, according to the CDC. Moreover, it is helpful to identify the type and brands of vitamins. Therefore, researchers can determine the components and estimate the amount of folic acid in each product. To ensure the compliance of vitamin intake during pregnancy, future studies need to follow up and estimate the duration of vitamin use. Last but not least, it is helpful to estimate the initiate time for vitamin use, especially for women who started vitamin use after pregnancy recognition. Women with late pregnancy recognition might miss the right time to initiate vitamin use for preventing NTDs.

With regard to the questionnaire design and interviews, future study may need to introduce Spanish or other foreign languages if the population includes minorities or foreign immigrants. Also, considering that some participants might fail to identify the type or brand of vitamins, researchers should provide a sample or bottles of vitamin products. Interviewers could also ask participants to show the vitamin products that they are using. To ascertain some sensitive maternal behaviors (e.g. alcohol use, recreational drug use), future studies can introduce biomarkers to certify the validity of self-report information. Lastly, the reasons of not taking vitamins need to be further studied. Previous experience of adverse pregnancy outcomes could mislead women into believing that vitamin use in pregnancy might account for adverse pregnancy outcomes.

Future studies need to assess the perception of vitamin use in pregnancy.

Public health campaign might incorporate appropriate education program in order to change attitudes towards an earlier vitamin use during pregnancy.

Although the association between some maternal characteristics (e.g. education, maternal age, BMI, health insurance, marital status) and vitamin use were not significantly in the present study, it does not mean that these factors do not account for the prevalence of vitamin use in early pregnancy. The findings of the present study should be interpreted with caution. Future studies need to explore these findings by recruiting a larger sample size.

There is a great need to develop prevention strategies to educate pregnant women about benefits of prenatal vitamin use. Our study indicated that ethnic minorities may particularly benefit from such education efforts. To begin with, improving access to prenatal care for all pregnant women will facilitate greater utilization of prenatal vitamins. For women who have financial or other barriers to obtain vitamins, publicly funded clinics with free access to prenatal care should be widely available. Health insurance companies can add multivitamin into their prescription plans and formulary. Also, healthcare providers should play a pivotal role in education of women at reproductive age. Specifically, they should provide counseling to pregnant women and women who plan a pregnancy in order to improve the patient's knowledge of preventing NTDs and to inform them about the benefits of taking vitamins. A March of Dimes survey suggested that 89% of women would be more likely to take folic acid if advised by health care providers. However, this survey also found that only 30% of

women who were aware of folic acid cite health care providers as the source of information about the benefits of folic acid use in pregnancy. 98 Therefore, there is a great need to incorporate healthcare providers as a part of the birth defects prevention programs. A physician-based intervention, conducted at Arkansas, suggested that a brief counseling from physicians can potentially reduce the risk of folate-preventable birth defects among their patients by as much as 11%.99 Besides physicians, pharmacists may also contribute to the promotion of folic acid use among women of reproductive age by encouraging them to take multivitamins. Moreover, the mass media, including printed media, audio and visual media, Internet, can also serve to increase the awareness of vitamin use at reproductive age. Lastly, additional research is needed to ascertain the reasons of using or not using vitamins among pregnant women or women who might become pregnant. Health behavior models, such as Health Belief Model, Theory of Planned Behavior, may help to better understand the patient's decision making process with respect to vitamin use during pregnancy and earlier initiation of prenatal care.

In sum, this study highlights the importance of promoting vitamin use in early pregnancy among pregnant women in New Mexico, given to the finding that only less than one third of women used vitamins during the pre-conceptional period. The findings of this study can also provide an implication that public health strategies to ensure early use of vitamin during pregnancy should include the promotion of pregnancy planning.

REFERENCES

- 1. Botto LD, Mulinare J, Erickson JD. Occurrence of congenital heart defects in relation to maternal mulitivitamin use. Am J Epidemiol 2000;151:878-84.
- 2. Czeizel AE, Dobo M, Vargha P. Hungarian cohort-controlled trial of periconceptional multivitamin supplementation shows a reduction in certain congenital abnormalities. Birth Defects Res A Clin Mol Teratol 2004;70:853-61.
- 3. Shaw GM, O'Malley CD, Wasserman CR, Tolarova MM, Lammer EJ.

 Maternal periconceptional use of multivitamins and reduced risk for
 conotruncal heart defects and limb deficiencies among offspring. Am J Med
 Genet 1995;59:536-45.
- 4. Werler MM, Hayes C, Louik C, Shapiro S, Mitchell AA. Multivitamin supplementation and risk of birth defects. Am J Epidemiol 1999;150:675-82.
- 5. Yang Q, Khoury MJ, Olney RS, Mulinare J. Does periconceptional multivitamin use reduce the risk for limb deficiency in offspring? Epidemiology 1997;8:157-61.
- 6. Li DK, Daling JR, Mueller BA, Hickok DE, Fantel AG, Weiss NS.

 Periconceptional multivitamin use in relation to the risk of congenital urinary tract anomalies. Epidemiology 1995;6:212-8.
- 7. Shaw GM, Zhu H, Lammer EJ, Yang W, Finnell RH. Genetic variation of infant reduced folate carrier (A80G) and risk of orofacial and conotruncal heart defects. Am J Epidemiol 2003;158:747-52.
- 8. van Rooij IA, Vermeij-Keers C, Kluijtmans LA, et al. Does the interaction between maternal folate intake and the methylenetetrahydrofolate reductase polymorphisms affect the risk of cleft lip with or without cleft palate? Am J Epidemiol 2003;157:583-91.

- 9. Prevention of neural tube defects: results of the Medical Research Council Vitamin Study. MRC Vitamin Study Research Group. Lancet 1991;338:131-7.
- 10. Khoury MJ, Shaw GM, Moore CA, Lammer EJ, Mulinare J. Does periconceptional multivitamin use reduce the risk of neural tube defects associated with other birth defects? data from two population-based case-control studies. Am J Med Genet 1996;61:30-6.
- 11. Milunsky A, Jick H, Jick SS, et al. Multivitamin/folic acid supplementation in early pregnancy reduces the prevalence of neural tube defects. JAMA 1989;262:2847-52.
- 12. Mulinare J, Cordero JF, Erickson JD, Berry RJ. Periconceptional use of multivitamins and the occurrence of neural tube defects. JAMA 1988;260:3141-5.
- 13. Seller MJ, Nevin NC. Periconceptional vitamin supplementation and the prevention of neural tube defects in south-east England and Northern Ireland. J Med Genet 1984;21:325-30.
- 14. Shaw GM, Schaffer D, Velie EM, Morland K, Harris JA. Periconceptional vitamin use, dietary folate, and the occurrence of neural tube defects. Epidemiology 1995;6:219-26.
- 15. Smithells RW, Sheppard S, Schorah CJ, et al. Possible prevention of neural-tube defects by periconceptional vitamin supplementation. Lancet 1980;1:339-40.
- 16. Werler MM, Shapiro S, Mitchell AA. Periconceptional folic acid exposure and risk of occurrent neural tube defects. JAMA 1993;269:1257-61.
- 17. Bank TW. WDI (World Development Indicators). In: Annual. 2011 ed: (Mimas) University of Manchester; 2011.

- 18. Suellentrop K, Morrow B, Williams L, D'Angelo D. Monitoring progress toward achieving Maternal and Infant Healthy People 2010 objectives--19 states, Pregnancy Risk Assessment Monitoring System (PRAMS), 2000-2003. MMWR Surveill Summ 2006;55:1-11.
- 19. Sullivan KM, Ford ES, Azrak MF, Mokdad AH. Multivitamin use in pregnant and nonpregnant women: results from the Behavioral Risk Factor Surveillance System. Public Health Rep 2009;124:384-90.
- 20. Boulet SL, Yang Q, Mai C, et al. Trends in the postfortification prevalence of spina bifida and anencephaly in the United States. Birth Defects Res A Clin Mol Teratol 2008;82:527-32.
- 21. Czeizel AE. Controlled studies of multivitamin supplementation on pregnancy outcomes. Ann N Y Acad Sci 1993;678:266-75.
- 22. Recommendations for the use of folic acid to reduce the number of cases of spina bifida and other neural tube defects. MMWR Recomm Rep 1992;41:1-7.
- 23. Institute of Medicine, Subcommittee for Clinical Applications Guide.
 Nutrition during pregnancy and lactaion: an implementation guide. Washington:
 National Academy Press 1992.
- 24. Catov JM, Nohr EA, Bodnar LM, Knudson VK, Olsen SF, Olsen J.

 Association of periconceptional multivitamin use with reduced risk of preeclampsia among normal-weight women in the Danish National Birth Cohort. Am J Epidemiol 2009;169:1304-11.
- 25. Vahratian A, Siega-Riz AM, Savitz DA, Thorp JM, Jr. Multivitamin use and the risk of preterm birth. Am J Epidemiol 2004;160:886-92.

- 26. Correa A, Botto L, Liu Y, Mulinare J, Erickson JD. Do multivitamin supplements attenuate the risk for diabetes-associated birth defects? Pediatrics 2003;111:1146-51.
- 27. Botto LD, Erickson JD, Mulinare J, Lynberg MC, Liu Y. Maternal fever, multivitamin use, and selected birth defects: evidence of interaction? Epidemiology 2002;13:485-8.
- 28. Elwood M LJ, Elwood JH. Epidemiology and control of neural tube defects: Oxford: Oxford University Press; 1992.
- 29. Lumley J, Watson L, Watson M, Bower C. Periconceptional supplementation with folate and/or multivitamins for preventing neural tube defects. Cochrane Database Syst Rev 2001:CD001056.
- 30. Botto LD, Moore CA, Khoury MJ, Erickson JD. Neural-tube defects. N Engl J Med 1999;341:1509-19.
- 31. Spina bifida and anencephaly before and after folic acid mandate--United States, 1995-1996 and 1999-2000. MMWR Morb Mortal Wkly Rep 2004;53:362-5.
- 32. Parker SE, Mai CT, Canfield MA, et al. Updated National Birth Prevalence estimates for selected birth defects in the United States, 2004-2006. Birth Defects Res A Clin Mol Teratol 2010;88:1008-16.
- 33. Canfield MA, Honein MA, Yuskiv N, et al. National estimates and race/ethnic-specific variation of selected birth defects in the United States, 1999-2001. Birth Defects Res A Clin Mol Teratol 2006;76:747-56.
- 34. Kondo A, Kamihira O, Ozawa H. Neural tube defects: prevalence, etiology and prevention. Int J Urol 2009;16:49-57.

- 35. Ammon Avalos L, Kaskutas LA, Block G, Li DK. Do multivitamin supplements modify the relationship between prenatal alcohol intake and miscarriage? Am J Obstet Gynecol 2009;201:563 e1-9.
- 36. Wu T, Buck G, Mendola P. Maternal cigarette smoking, regular use of multivitamin/mineral supplements, and risk of fetal death: the 1988 National Maternal and Infant Health Survey. Am J Epidemiol 1998;148:215-21.
- 37. Yu SM, Keppel KG, Singh GK, Kessel W. Preconceptional and prenatal multivitamin-mineral supplement use in the 1988 National Maternal and Infant Health Survey. Am J Public Health 1996;86:240-2.
- 38. Scholl TO, Hediger ML, Bendich A, Schall JI, Smith WK, Krueger PM. Use of multivitamin/mineral prenatal supplements: influence on the outcome of pregnancy. Am J Epidemiol 1997;146:134-41.
- 39. Williams LM, Morrow B, Lansky A, et al. Surveillance for selected maternal behaviors and experiences before, during, and after pregnancy. Pregnancy Risk Assessment Monitoring System (PRAMS), 2000. MMWR Surveill Summ 2003;52:1-14.
- 40. Than LC, Honein MA, Watkins ML, Yoon PW, Daniel KL, Correa A. Intent to become pregnant as a predictor of exposures during pregnancy: is there a relation? J Reprod Med 2005;50:389-96.
- 41. Santelli J, Rochat R, Hatfield-Timajchy K, et al. The measurement and meaning of unintended pregnancy. Perspect Sex Reprod Health 2003;35:94-101.
- 42. Burris HH, Mitchell AA, Werler MM. Periconceptional multivitamin use and infant birth weight disparities. Ann Epidemiol 2010;20:233-40.

- 43. Carmichael SL, Shaw GM, Yang W, et al. Correlates of intake of folic acid-containing supplements among pregnant women. Am J Obstet Gynecol 2006;194:203-10.
- 44. Canfield MA, Annegers JF, Brender JD, Cooper SP, Greenberg F.

 Hispanic origin and neural tube defects in Houston/Harris County, Texas. I.

 Descriptive epidemiology. Am J Epidemiol 1996;143:1-11.
- 45. Feuchtbaum LB, Currier RJ, Riggle S, Roberson M, Lorey FW, Cunningham GC. Neural tube defect prevalence in California (1990-1994): eliciting patterns by type of defect and maternal race/ethnicity. Genet Test 1999;3:265-72.
- 46. Shaw GM, Velie EM, Wasserman CR. Risk for neural tube defect-affected pregnancies among women of Mexican descent and white women in California.

 Am J Public Health 1997;87:1467-71.
- 47. D'Angelo D, Williams L, Morrow B, et al. Preconception and interconception health status of women who recently gave birth to a live-born infant--Pregnancy Risk Assessment Monitoring System (PRAMS), United States, 26 reporting areas, 2004. MMWR Surveill Summ 2007;56:1-35.
- 48. Ramadhani T, Short V, Canfield MA, et al. Are birth defects among Hispanics related to maternal nativity or number of years lived in the United States? Birth Defects Res A Clin Mol Teratol 2009;85:755-63.
- 49. Kallen B, Lundberg G, Aberg A. Relationship between vitamin use, smoking, and nausea and vomiting of pregnancy. Acta Obstet Gynecol Scand 2003;82:916-20.
- 50. Martin JA, Hamilton BE, Sutton PD, et al. Births: final data for 2005. Natl Vital Stat Rep 2007;56:1-103.

- 51. Brender JD, Felkner M, Suarez L, Canfield MA, Henry JP. Maternal pesticide exposure and neural tube defects in Mexican Americans. Ann Epidemiol;20:16-22.
- 52. Suarez L, Felkner M, Brender JD, Canfield M, Hendricks K. Maternal exposures to cigarette smoke, alcohol, and street drugs and neural tube defect occurrence in offspring. Matern Child Health J 2008;12:394-401.
- 53. Gillum RF. Epidemiology of congenital heart disease in the United States.

 Am Heart J 1994;127:919-27.
- 54. Scanlon KS, Ferencz C, Loffredo CA, et al. Preconceptional folate intake and malformations of the cardiac outflow tract. Baltimore-Washington Infant Study Group. Epidemiology 1998;9:95-8.
- 55. Goh YI, Bollano E, Einarson TR, Koren G. Prenatal multivitamin supplementation and rates of congenital anomalies: a meta-analysis. J Obstet Gynaecol Can 2006;28:680-9.
- 56. Heron M, Hoyert D, Murphy S, Xu J, Kochanek K, Tejada-Vera B. Death: Final data for 2006. Natl Vital Stat Rep 2009;57.
- 57. Czeizel AE. Reduction of urinary tract and cardiovascular defects by periconceptional multivitamin supplementation. Am J Med Genet 1996;62:179-83.
- 58. Itikala PR, Watkins ML, Mulinare J, Moore CA, Liu Y. Maternal multivitamin use and orofacial clefts in offspring. Teratology 2001;63:79-86.
- 59. Czeizel AE. Prevention of congenital abnormalities by periconceptional multivitamin supplementation. BMJ 1993;306:1645-8.

- 60. Shaw GM, Lammer EJ, Wasserman CR, O'Malley CD, Tolarova MM. Risks of orofacial clefts in children born to women using multivitamins containing folic acid periconceptionally. Lancet 1995;346:393-6.
- 61. Hayes C, Werler MM, Willett WC, Mitchell AA. Case-control study of periconceptional folic acid supplementation and oral clefts. Am J Epidemiol 1996;143:1229-34.
- 62. Ouyang L, Grosse SD, Armour BS, Waitzman NJ. Health care expenditures of children and adults with spina bifida in a privately insured U.S. population. Birth Defects Res A Clin Mol Teratol 2007;79:552-8.
- 63. Bower C, Stanley FJ. Periconceptional vitamin supplementation and neural tube defects; evidence from a case-control study in Western Australia and a review of recent publications. J Epidemiol Community Health 1992;46:157-61.
- 64. Mills JL, Rhoads GG, Simpson JL, et al. The absence of a relation between the periconceptional use of vitamins and neural-tube defects.

 National Institute of Child Health and Human Development Neural Tube Defects Study Group. N Engl J Med 1989;321:430-5.
- 65. Thompson SJ, Torres ME, Stevenson RE, Dean JH, Best RG.

 Periconceptional multivitamin folic acid use, dietary folate, total folate and risk of neural tube defects in South Carolina. Ann Epidemiol 2003;13:412-8.
- 66. Botto LD, Mulinare J, Erickson JD. Occurrence of omphalocele in relation to maternal multivitamin use: a population-based study. Pediatrics 2002;109:904-8.

- 67. Czeizel AE, Puho EH, Langmar Z, Acs N, Banhidy F. Possible association of folic acid supplementation during pregnancy with reduction of preterm birth: a population-based study. Eur J Obstet Gynecol Reprod Biol 2010;148:135-40. 68. Alwan NA, Greenwood DC, Simpson NA, McArdle HJ, Cade JE. The relationship between dietary supplement use in late pregnancy and birth outcomes: a cohort study in British women. BJOG 2010;117:821-9. 69. Czeizel AE, Dudas I, Metneki J. Pregnancy outcomes in a randomised controlled trial of periconceptional multivitamin supplementation. Final report. Arch Gynecol Obstet 1994;255:131-9.
- 70. Kirke PN, Daly LE, Elwood JH. A randomised trial of low dose folic acid to prevent neural tube defects. The Irish Vitamin Study Group. Arch Dis Child 1992;67:1442-6.
- 71. Bodnar LM, Tang G, Ness RB, Harger G, Roberts JM. Periconceptional multivitamin use reduces the risk of preeclampsia. Am J Epidemiol 2006;164:470-7.
- 72. Czeizel AE, Dudas I, Fritz G, Tecsoi A, Hanck A, Kunovits G. The effect of periconceptional multivitamin-mineral supplementation on vertigo, nausea and vomiting in the first trimester of pregnancy. Arch Gynecol Obstet 1992;251:181-5.
- 73. Lunet N, Rodrigues T, Correia S, Barros H. Adequacy of prenatal care as a major determinant of folic acid, iron, and vitamin intake during pregnancy. Cad Saude Publica 2008;24:1151-7.
- 74. Haugen M, Brantsaeter AL, Alexander J, Meltzer HM. Dietary supplements contribute substantially to the total nutrient intake in pregnant Norwegian women. Ann Nutr Metab 2008;52:272-80.

- 75. Forster DA, Wills G, Denning A, Bolger M. The use of folic acid and other vitamins before and during pregnancy in a group of women in Melbourne, Australia. Midwifery 2009;25:134-46.
- 76. Maats FH, Crowther CA. Patterns of vitamin, mineral and herbal supplement use prior to and during pregnancy. Aust N Z J Obstet Gynaecol 2002;42:494-6.
- 77. Wehby GL, Castilla EE, Lopez-Camelo JS, Murray JC. Predictors of multivitamin use during pregnancy in Brazil. Int J Public Health 2009;54:78-87.
- 78. Catov JM, Bodnar LM, Ness RB, Markovic N, Roberts JM. Association of periconceptional multivitamin use and risk of preterm or small-for-gestational-age births. Am J Epidemiol 2007;166:296-303.
- 79. Dott M, Rasmussen SA, Hogue CJ, Reefhuis J. Association between pregnancy intention and reproductive-health related behaviors before and after pregnancy recognition, National Birth Defects Prevention Study, 1997-2002.

 Matern Child Health J 2010;14:373-81.
- 80. Tam LE, McDonald SD, Wen SW, Smith GN, Windrim RC, Walker MC. A survey of preconceptional folic acid use in a group of Canadian women. J Obstet Gynaecol Can 2005;27:232-6.
- 81. Surveillance of preconception health indicators among women delivering live-born infants--Oklahoma, 2000-2003. MMWR Morb Mortal Wkly Rep 2007;56:631-4.
- 82. Nilsen RM, Vollset SE, Rasmussen SA, Ueland PM, Daltveit AK. Folic acid and multivitamin supplement use and risk of placental abruption: a population-based registry study. Am J Epidemiol 2008;167:867-74.

- 83. Paulik E, Csaszar J, Kozinszky Z, Nagymajtenyi L. Preconceptional and prenatal predictors of folic acid intake in Hungarian pregnant women. Eur J Obstet Gynecol Reprod Biol 2009;145:49-52.
- 84. Glover DD, Amonkar M, Rybeck BF, Tracy TS. Prescription, over-the-counter, and herbal medicine use in a rural, obstetric population. Am J Obstet Gynecol 2003;188:1039-45.
- 85. Vollset SE, Lande B. Knowledge and attitudes of folate, and use of dietary supplements among women of reproductive age in Norway 1998. Acta Obstet Gynecol Scand 2000;79:513-9.
- 86. Garikapaty VP, Feyerharm R, Zhu B. Folic acid consumption among Missouri women in the periconceptional period. Mo Med 2008;105:504-9.
- 87. Grosse SD, Waitzman NJ, Romano PS, Mulinare J. Reevaluating the benefits of folic acid fortification in the United States: economic analysis, regulation, and public health. Am J Public Health 2005;95:1917-22.
- 88. Bentley TG, Weinstein MC, Willett WC, Kuntz KM. A cost-effectiveness analysis of folic acid fortification policy in the United States. Public Health Nutr 2009;12:455-67.
- 89. Postma MJ, Londeman J, Veenstra M, de Walle HE, de Jong-van den Berg LT. Cost-effectiveness of periconceptional supplementation of folic acid. Pharm World Sci 2002;24:8-11.
- 90. Jentink J, van de Vrie-Hoekstra NW, de Jong-van den Berg LT, Postma MJ. Economic evaluation of folic acid food fortification in The Netherlands. Eur J Public Health 2008;18:270-4.
- 91. Chang G, Carroll KM, Behr HM, Kosten TR. Improving treatment outcome in pregnant opiate-dependent women. J Subst Abuse Treat 1992;9:327-30.

- 92. Dawson DA, Das A, Faden VB, Bhaskar B, Krulewitch CJ, Wesley B. Screening for high- and moderate-risk drinking during pregnancy: a comparison of several TWEAK-based screeners. Alcohol Clin Exp Res 2001;25:1342-9.
- 93. Russell M, Martier SS, Sokol RJ, et al. Screening for pregnancy risk-drinking. Alcohol Clin Exp Res 1994;18:1156-61.
- 94. Finer LB, Henshaw SK. Disparities in rates of unintended pregnancy in the United States, 1994 and 2001. Perspect Sex Reprod Health 2006;38:90-6.
 95. Perez-Escamilla R, Himmelgreen D, Segura-Millan S, Gonzalez A, Mendez I, Haldeman L. Knowledge of folic acid and neural tube defects among inner-city residents: have they heard about it? J Am Diet Assoc 1999;99:80-3.
 96. Kannan S, Menotti E, Scherer HK, Dickinson J, Larson K. Folic acid and the prevention of neural tube defects: A survey of awareness among Latina women of childbearing age residing in southeast Michigan. Health Promot Pract 2007;8:60-8.
- 97. Watson LF, Brown SJ, Davey MA. Use of periconceptional folic acid supplements in Victoria and New South Wales, Australia. Aust N Z J Public Health 2006;30:42-9.
- 98. March of Dimes and Gallup Organization. Folic acid and the prevention of birth defects: A national survey of pre-pregnancy awareness and behavior among women of childbearing age 1995-2003: Princeton (NJ): Gallup Organization; 2003.
- 99. Robbins JM, Cleves MA, Collins HB, Andrews N, Smith LN, Hobbs CA.
 Randomized trial of a physician-based intervention to increase the use of folic acid supplements among women. Am J Obstet Gynecol 2005;192:1126-32.

LIST OF TABLES

Table 2.1. Prevalence of Multivitamin Use in Pregnancy

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date		•	Use		
Dott M	Case-control	N=4,094 mothers of	Periconceptional period:	Periconceptional non-	Potential selection bias
2010	study	live born infants	-1 month to +1 month	multivitamin users:	exists when this study
U.S.		born during	around conception	41% (992/2437) of	did not capture mothers
		1997-2002		women without	whose current pregnancy
			During pregnancy: not	pregnancy intent;	ended in a miscarriage,
	*Data from National Birth Defects		studied	64.3% (1059/1648) of	stillbirth, or an infant with
	Prevention study		Multivitamin: multivitamin	women with pregnancy	a major birth defect,
	1997-2002		containing folic	intent	potentially resulting in an
			acid/prenatal		underestimate of the
			multivitamin		impact of pregnancy
					intention on maternal
			Frequency: not studied		behaviors. Also, this
					study may have recall
			OTC & Rx: not studied		bias.
			Single vitamin: not		
			studied		

Author & Date	Study Design	Рорг	ulation	Assessment of Vitamin Use	Prevalence	Comments
Burris HH	Cross-sectional	Control:	random	Periconceptional period:	Periconceptional	
2009	study	sample	of	-1 to +1 around LMP	multivitamin users:	
U.S.		Massach	usetts		55.0% of white women	
		births		During pregnancy: not	vs. 25.6% of black	
				studied	women	
		Case: N	Nothers of			
		malforme	ed infants	Multivitamin: at least 2		
	*Data source: the			water-soluble vitamins+2		
	Slone Epidemiology Center Birth Defects			fat-soluble vitamins		
	Study					
				Frequency: ≥4		
				times/week		
				OTC & Rx: not mention		
				Single vitamin: not		
				mention		
				Dua matal vita vain vast		
				Prenatal vitamin: not		
				mention		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Sullivan K 2009 U.S	*Data from Behavioral Risk Factor Surveillance System(BRFSS)	N= 788 pregnant women who were ascertained from 20, 263 women who aged from 18 to 44 years by 14 states and U.S territories	Periconceptional period: Not studied During pregnancy: studied but not defined Multivitamin: not defined Frequency: not studied OTC & Rx: not studied Single vitamin: not studied Prenatal multivitamin: not studied	Multivitamin users in pregnancy: 78% of pregnant women reported multivitamin use.	Women were asked whether they currently used multivitamin or not. Information about the duration of multivitamin use, dose, contents, and frequency was not collected

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Avalos LA 2009 U.S.	*Data: women members in the Kaiser Permanente Medical Care Program (KPMCP)	N=1,061 women who was at their first pregnancy, speaking English, have pregnancy intentness and whose gestational age at the pregnancy test was less than or equal to 10 complete weeks	Periconceptional period: began prior pregnancy and in pregnancy (not specifically defined) During pregnancy: since become pregnant or LMP Multivitamin: multivitamin/prenatal vitamin Frequency: any use (not specifically defined) OTC & Rx: not studied but not presented	Periconceptional multivitamin users: 23.8% (252/1061) During pregnancy: 68.8% (730/1061) Prenatal multivitamin users who begin to use multivitamin during pregnancy: 44.8% (475/1061) Multivitamin non-users: 31.2% (331/1061)	Multivitamin status is an important modifier in the relationship between pregnancy drinking and miscarriage: the risk of miscarriage was greatest for women who drank alcohol and reported no multivitamin supplementation

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Venkata P.S 2008 U.S.	Cross-sectional study	N=1,789 mothers in Missouri who delivered the previous 2-6 months	Pre-conceptional period: one month before became pregnant During pregnancy: not studied	Pre-conceptional multivitamin users: 29.7% (every day a week), 5.0% (4-6 times per week), 10% (1-3 times per week)	
	*Data: Missouri Pregnancy Related Assessment and Monitoring System (MoPRA)		Multivitamin: multivitamin/prenatal vitamin		
			Frequency: 1-3/4-6/7 times per week		
			OTC & Rx: not studied		
			Single vitamin: studied but not presented		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Catov JM 2007 U.S.	Catov JM Prospective 2007 Cohort study	N=1,823 women who enrolled at less than 16 weeks and followed through the postpartum visit	Periconceptional period: 6 months which included prior to conception, at the time of conception immediately after conception (not specifically defined)	Periconceptional multivitamin users: 47%	The study relied on self-reported multivitamin use. Also, This study lack of information on the brand or dose of supplement
	Preeclampsia Prevention Study		During pregnancy: not studied Multivitamin: multivitamin/prenatal vitamin		periconceptional vitamin supplementation is associated with preterm birth (<34 weeks) has been reported by others
			Frequency: at least once/week		
			OTC & Rx: not studied		
			Single vitamin: not studied		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
D'Angelo D	Cross-sectional	Women from 26	Preconceptional period:	Pre-conceptional	
et al	study	PRAMS reporting	before pregnancy (1	multivitamin users:	
2007		areas	month before, 3 months,	35.1% (26.7% in	
U.S.			12 months before	Arkansas – 43.6%	
	*Data from Pregnancy Risk Assessment		pregnancy)	Rhode Island)	
	Monitoring		Periconceptional period		
	System(PRAMS)		&		
	During 2003-2004		During pregnancy: not		
			studied		
			Multivitamin: multivitamin/prenatal vitamin		
			Frequency: at least 4 times/week		
			OTC & Rx: not studied		
			Single vitamin/prenatal		
			vitamin: not studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
CDC (not listed) 2007 U.S.	Cross-sectional study	Women from PRAMS in Oklahoma	Preconceptional period: during the month before pregnancy	Pre-conceptional multivitamin users: 26.5%	
0.3.	*Data from Pregnancy Risk Assessment Monitoring System(PRAMS) During 2000-2003		Periconceptional period & During pregnancy: not studied Multivitamin: multivitamin/prenatal vitamin		
			Frequency: at least 4 times/week		
			OTC & Rx: not studied		
			Single vitamin/prenatal vitamin: not studied		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
Suellentrop et al 2006 U.S.	*Data from Pregnancy Risk Assessment Monitoring System(PRAMS) During 2000-2003	Women from 19 PRAMS reporting areas	Pre-conceptional period: during the month before pregnancy Periconceptional period & During pregnancy: not studied Multivitamin: not defined Frequency: at least 4 times/week OTC & Rx: not studied Single vitamin/prenatal vitamin: not studied	Pre-conceptional multivitamin users: 23.0% in Arkansas to 45.2% in Maine	Sectional bias may exist because only women with live infants were included

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Carmichael	Cross-sectional	N=2518 women	Pre-conceptional period:	Periconceptional	Participants were
2006	analysis	with estimated	not studied	vitamin non-users:	mothers of live-born
U.S.	*Data from the	delivery dates from		53%	infants without major
	*Data from the National Birth	1997-2000	Periconceptional period:		malformation
	Defects Prevention		3 month before	Earlier pregnancy:	
	Study		conception/one month	35%	
			after conception	Later pregnancy: 8%	
			Earlier pregnancy: the	Later pregnancy. 070	
			second and third month		
			after conception		
			Later pregnancy: during		
			the fourth month or later		
			during pregnancy		
			Multivitamin:		
			multivitamin/prenatal		
			vitamin: folic acid		
			supplementation		
			Frequency: not defined		
			OTC & Rx/prenatal		
			vitamin: not studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Glover et al 2003 U.S	Prospective cohort study	N=578 participants from four rural outreach clinics between August 1999 and November 2001	Periconceptional period: not studied During pregnancy: since LMP Multivitamin: not defined Frequency: studied but not presented OTC & Rx: separately reported (Rx multivitamin was defined as a prenatal vitamin) Single vitamin: vitamin C was reported Prenatal multivitamin: not studied	Multivitamin users in pregnancy: 92% (prescribed prenatal vitamin) vs. 10.9% (OTC multivitamins) Vitamin C users in pregnancy: 2.4% (14)	Information about the duration of multivitamin use, dose, contents, and frequency was not mentioned

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
Vahratian	Cross-sectional	Women who were	Periconceptional period:	Periconceptional	Frequency of
2004	analysis	at 24-29 weeks of	Before and during	multivitamin users:	multivitamin use, type of
U.S.		pregnancy from 4	pregnancy (not	30%	multivitamin was not
		prenatal care clinics	specifically defined)	Prenatal multivitamin	studied in this analysis
	*Data Dasanasa	in central North		users: 54%	
	*Data: Pregnancy, Infection, and	Carolina	During pregnancy: any		
	Nutrition (PIN) study		time during pregnancy		
			up until the time of interview or recruitment		
			(24-29 gestational		
			weeks)		
			weeks)		
			Multivitamin: not defined		
			Frequency: studied but		
			not presented		
			OTC & Dy/propotal		
			OTC & Rx/prenatal vitamin: not studied		
			vitariiri. Hot studied		
			Single vitamin: folate,		
			vitamin A, vitamin C was		
			studied but not reported		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date	-	-	Use		
Williams LM	Cross-sectional	N=32,479 women	Pre-conceptional period:	Pre-conceptional	
2000	study	who were identified	during the month before	multivitamin users:	
U.S.		from 19 states and	pregnancy	25.0%40.7%	
	*Data from	U.S territories			
	Pregnancy Risk Assessment		Periconceptional period		
	Monitoring System		&		Because data are
	(PRAMS) 2000		During pregnancy: not		self-reported 28
			studied		months after delivery,
					responses might be
			Multivitamin: not defined		subject to recall bias,
					particularly for behaviors
			Frequency: at least 4		and experiences that
			times/week		occurred before the
			0.70 0.0 / / /		pregnancy
			OTC & Rx/prenatal		
			vitamin: not studied		
			Cinalo vitamini nat		
			Single vitamin: not		
			studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Wu T	Cross-sectional	N=12,465 women	Periconceptional period:	Periconceptional	Multivitamin and mineral
1998	study	were identified from	-3 to +3 months around	multivitamin/mineral	supplements were
U.S		livebirth, fetal	pregnancy recognition	users: 21%	combined and could not
		death, infant death			be separated for further
		certifications from	During pregnancy: 3	Multivitamin/mineral	analysis
		the National	months after pregnancy	users during	
	*Data from the National Maternal	Maternal and Infant	recognition	pregnancy: 53.8%	
	and Infant Health	Health Survey			
	Survey		Multivitamin: not defined		
			Fraguerow et laget 2		
			Frequency: at least 3 times/week		
			times/week		
			OTC & Rx/prenatal		
			vitamin: not studied		
			vitariiii. Hot stadica		
			Single vitamin: vitamin		
			A, vitamin C, folic acid		
			were studied but not		
			reported		

Author &	Study Design	Population	Assessment of	Prevalence	Comments
Date	_	-	Vitamin Use		
Scholl TO 1997 U.S.	Prospective cohort study	N=1,430 low income, urban women who entered prenatal care during the first and second trimesters and who had singleton pregnancies.	Periconceptional period: Not studied During pregnancy: after pregnancy recognition Multivitamin: prenatal multivitamin/mineral supplements Frequency: not studied OTC & Rx: not studied	Multivitamin/supplements users during pregnancy: 80.4% 29.3% (418/1430) at the first trimester vs. 730 51.0% (730/1430) during the second trimester	Multivitamin and mineral supplements were combined, and there was insufficient information on the use of multivitamin such as type of multivitamin, frequency, or length of ingestion
			Single vitamin: not studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Milunsky	Cross-sectional	N=22,776 pregnant	Periconceptional	Periconceptional	
1989	analysis	women who had a	period: -3 to +3 months	multivitamin users:	
U.S.		either maternal serum	around LMP	31.9% (n=7261)	
		a-fetoprotein(MSAF	During pregnancy: 3	Multivitamin non-users:	
		P) screen or an amniocentesis	months after LMP	12.9% (n=2927)	
		around 16 weeks of	Multivitamin: not		
		gestation	defined		
			Frequency: studied but not reported		
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: vitamin		
			A, vitamin C, vitamin E,		
			folic acid were studied		
			but not reported		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Yu SM 1996 U.S.	Cross-sectional study	N=9,953 women who delivered live infants in the 1988 National Maternal and Infant Health Survey	Periconceptional period: not studied During pregnancy: after pregnancy recognition Multivitamin: not defined	Multivitamin-mineral supplement users: 82.5% (67% of Black mothers vs. 84% of White mothers)	Multivitamin and mineral supplements were combined,
	*Data from the National Maternal and Infant Health Survey		Frequency: ≥3 times/week OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: vitamin A, vitamin C, vitamin E, folic acid were studied but not reported		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
Tam LE	Cross-sectional	N=383 postpartum	Periconceptional period:	Periconceptional	
2004	analysis	women in Toronto	at least 4 weeks prior to	multivitamin users:	
Canada		Hospital and the	conception until 8 weeks	28%	
		Mount Sinai	after conception		
		Hospital	and conseption		
		rioopitai	During pregnancy: not		
			studied		
			Studied		
			Multivitamin: not defined		
			Multivitariiii. Hot deiined		
			Fraguesay: not atudied		
			Frequency: not studied		
			OTO 9 Durkanan atal		
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: not		
			studied		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date Alwan NA 2010 United Kingdom	prospective cohort study	N=1,274 women who aged at 18-45 years with low-risk pregnancies were prospectively recruited at 8-12 weeks of gestation from the Leeds Teaching Hospital	Periconceptional period: not studied During pregnancy: from the first trimester to the third trimester Multivitamin: not defined	Multivitamin-mineral supplement users during pregnancy: 23.3 % (293/1274) in the first trimester, 14.1% (177/1274) in the second trimester, and 18.6% (79/425)*.	Data on using multivitamin-mineral preparation was mixed and could not be separated.
		maternity unit between 2003 and 2006	Frequency: not studied OTC & Rx/prenatal vitamin: not studied Single vitamin: vitamin A, B6, B12, C, D, E were studied but not prevalence rate was reported	* Only 425 women had information on supplement intake in the third trimester.	

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
KÄLLÉN	Prospective	N=3675 women	Periconceptional period:	Periconceptional	The study explored the
2003	Cohort study	who were at their	before pregnancy and	multivitamin users:	relationship between
Sweden		first visit to the antenatal care unit	the first two months after pregnancy (not	19.3% (2525/3675)	vitamin use, smoking, and nausea and vomiting
		and return questionnaire	specifically defined)	Multivitamin user during early	of pregnancy, but it does not focus on the
		around gestational	During early pregnancy:	pregnancy: 7.8%	predictors of the use of
		week 28.	first two months	(286/3675) of women used vitamin only in	multivitamin/vitamin
			Multivitamin: not defined	early pregnancy.	
			Frequency: not studied		
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: not		
			studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Catov JM	Prospective	N=26,601	Periconceptional period:	Periconceptional	Regular periconceptional
2009	Cohort study	women who were	-4 wks to +8 wks around	multivitamin users:	multivitamin use is
Denmark	_	recruited in Danish	LMP	65% (18,551/26601)	associated with a
	*Data from Danish	National Birth			reduced risk of
	National Birth Cohort	Cohort and	During pregnancy: not		preeclampsia among
		completed revised	studied		normal-weight women
		recruitment form			
			Multivitamin: not defined		
			Frequency: at least once/week		
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: vitamin A, B1, B12, B6, B12, C, D,		
			E, and folic acid were		
			studied but prevalence		
			rates were not reported		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
	One an estimat	N. 240 salsatan		December	
Paulik E	Cross-sectional	N=349 voluntary	Periconceptional period:	Preconception	
2009	study	pregnant women	not studied	multivitamin regular	
Hungary		who were at the		users: 43.8%	
	*Data from a	Department of	During pregnancy: not	(153/349)	
	self-administrated questionnaire	Obstetrics and	defined		
	(response	Gynecology and in		Preconception folic	
	rate=69.8%)	the Pregnancy	Preconception period:	acid regular users:	
		Care Center	Not defined	31.5% (110/349)	
			Multivitamin: not defined	Multivitamin regular	
			Water reasons 1100 dointed	users in pregnancy:	
			Frequency: not defined	48.4% (141/349)	
			i requency. Not defined	40.470 (1417349)	
			OTC & Rx: not studied	Folic acid regular users	
				in pregnancy: 40.4%	
			Single vitamin: folic acid	(141/349)	

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date		-	Use		
Haugen M	Cross-sectional	N=40,817 Pregnant	Periconceptional period:	Multivitamin users in	
2008	study	women in Norway	not studied	the first 4-5 months of	
Norway		who were 17-18		pregnancy: 16.3%	
	*Data from Norwegian Mothers	weeks of gestation	During pregnancy: first		
	and Child Cohort Study(MoBa)	and participates in study between	4-5 months		
		February 2002 and February 2005	Multivitamin: not defined		
		,	Frequency: never/<1/1-7 times per week		
			OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: vitamin A, B1, B12, B6, B12, C,		
			D, E, and folic acid were studied but prevalence		
			rates were not reported		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Vollest SE	Prospective	N=1,140 Norwegian	Periconceptional period:	Periconceptional	
2000	cohort study	women who aged	before or during the first	multivitamin users	
Norway		18-45 years	2-3 months of their last recent pregnancies	(daily or almost daily): 19.8%	
			During pregnancy: not defined	Periconceptional Vitamin B users: 3.6%	
			Multivitamin: not defined	Periconceptional folic acid users: 2.4%	
			Frequency: daily or almost daily/less		
			frequent than daily/ almost daily/never		
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: vitamin B,		
			folic acid		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
Lunet N	Cross-sectional	N=836 mothers	Periconceptional period:	Multivitamin uses in	Frequency of
2008	analysis from a	who were	not studied,	pregnancy: 76.2%	multivitamin use and the
Portugal	case-control	interviewed 24-72			content of multivitamin
	study	hours after delivery	During pregnancy:	Folic acid users in	was not mentioned in
		at two major public hospitals in the North of Portugal	without exactly refer to exact trimester	pregnancy: 55.4%	this study;
		rtorur or r ortagar	Multivitamin: not defined		
			Frequency: not studied		
			OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: folic acid		

Author &	Study Design	Population	Assessment of Vitamin	Prevalence	Comments
Date			Use		
Nilsen RM 2008 Norway	Cross-sectional Analysis *Data from Medical birth registry of Norway	N=226,724 women who have been recorded by Medical Birth Registry of Norway from 1998 through 2004	Periconceptional period: before and during pregnancy (not specifically defined) During pregnancy: not defined Multivitamin: multivitamin alone/folic acid alone/both folic acid and multivitamin Frequency: not studied OTC & Rx: not studied Single vitamin/prenatal vitamin: not studied	Periconceptional multivitamin and/or folic acid users: 15.9% Multivitamin and/or folic acid users in pregnancy: 27.3%	The prevalence of using each supplement type can not be separated from the data

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Forster D 2009 Australia	Cross-section al survey	N=588 women were approached in the antenatal clinic and the birth centre at around 36-38 weeks gestation	Periconceptional period: before and during pregnancy (not specifically defined) During pregnancy: not defined Multivitamin: folic acid alone/folic acid with multivitamin/folic acid with other vitamin Frequency: does/day and length of time were recorded OTC & Rx/prenatal vitamin: not studied	Periconceptional multivitamin users: 7.8% (46/587) Multivitamin users not before pregnancy but during pregnancy: 26.9% (158/587)	A combination of folic acid and multivitamin intake in different time of pregnancy was provided.
			Single vitamin: folic acid, vitamin B6		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Maats FH 2002 Australia	Cross-section al study(Hospital -based sample)	N=211 women who were 26 weeks gestation or more	Periconceptional period: during the three months prior to pregnancy During pregnancy: from the 1 st trimester to the 3 rd trimester.	Periconceptional multivitamin users: 26 (12%) Multivitamin users in pregnancy: 33(16%) in the first trimester,	
			Multivitamin/: not defined Frequency: studied but	37(18%) in the second trimester, 39(18%) in the third trimester, respectively.	
			not reported OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: folic acid alone or folic acid combined with multivitamins		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Prevalence	Comments
Wehby G 2009 Brazil	Cross-sectional analyses(Birth registry data)	N=1,774 mothers of liveborn infants without congenital malformations between 1995 and 2001 in Brazil	Periconceptional period: not studied During pregnancy: not defined Multivitamin: multivitamin/prenatal vitamin Frequency: any use during pregnancy (not specifically defined) OTC & Rx: not studied Single vitamin: not studied	Multivitamin users in pregnancy: 14%	This study evaluate the demand for multivitamins both pooled and stratified by African ancestry based on maternal report of the ancestries of the infant

Table 2.2. Predictors of Multivitamin Use in Pregnancy

Author &	Study Design	Popul		Assessment of Vitamin	Predictors	Comments
Date		•		& Folic acid Use		
Burris HH 2009 U.S.	*Data source: the Slone Epidemiology Center	Control: sample Massachu births Case: Mo malforme	others of	Periconceptional period: -1 to +1 month around LMP During pregnancy: not studied	Periconceptional multivitamin users: white women, non-smokers, married, older, wealthier (annual household≥\$45,000),	**Periconceptional: 28 days prior to the last menstrual period to after 28 days after the last menstrual period
	Birth Defects Study			Multivitamin: at least 2 water-soluble vitamins+2 fat-soluble vitamins Frequency: ≥4 times/week	and better educated (above high school education)	
				OTC & Rx: not mention Single vitamin: not mention Prenatal multivitamin: not		
				Prenatal multivitamin: not mention		

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Sullivan K	Cross-sectional	N=20, 263 women	Periconceptional period:	Multivitamin users	Based on 14 U.S
2009	study	who aged from 18	Not studied	during pregnancy:	states, the
U.S.		to 44 years by 14		Higher Income (annual	generalizability of the
		states and U.S	During pregnancy: studied	household≥\$50,000)	results is limited.
	*Data from Behavioral	territories	but not defined	Marital status (married	Information about the
	Risk Factor Surveillance System(BRFSS)			or previously married)	duration of
	Gyotom(Erti GG)		Multivitamin: not defined		multivitamin use,
					dose, contents, and
			Frequency: not studied		frequency was not
					collected
			OTC & Rx/prenatal		
			vitamin: not studied		
			Single vitamin: not studied		
			Prenatal multivitamin: not		
			studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Predictors	Comments
Venkata P.S 2008	Cross-sectional study	N=1,789 mothers in Missouri who delivered the	Pre-conceptional period: one month before became pregnant	Periconceptional multivitamin regular users (≥4 times/week):	
U.S.		previous 2-6 months	During pregnancy: not	Intended pregnancy, high house income	
		1110111115	studied	(≥35,000), married, non-smoker, more	
	*Data: Missouri Pregnancy Related Assessment and Monitoring System (MoPRA)		Multivitamin: multivitamin/prenatal vitamin	educated (≥12 years), older (≥20 years)	
			Frequency: 1-3/4-6/7 times per week		
			OTC & Rx: not studied		
			Single vitamin: studied but not presented		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Predictors	Comments
D'Angelo D et al 2007 U.S.	*Data from Pregnancy Risk Assessment Monitoring System(PRAMS) During 2003-2004	Women from 26 PRAMS reporting areas	Preconceptional period: before pregnancy (1 month before, 3 months, 12 months before pregnancy) Periconceptional period & During pregnancy: not studied Multivitamin: not defined Frequency: at least 4 times/week OTC & Rx: not studied Single vitamin/prenatal vitamin: not studied	Preconceptional multivitamin users: older (≥35 years), white women, intended pregnancy, had private health insurance	

Author & Date	Study Design	Population	Assessment of Vitamin Use	Predictors	Comments
CDC (not listed) 2007 U.S.	*Data from Pregnancy Risk Assessment Monitoring System(PRAMS) During 2000-2003	Women from PRAMS in Oklahoma	Preconceptional period: during the month before pregnancy Periconceptional period & During pregnancy: not studied Multivitamin: multivitamin/prenatal vitamin Frequency: at least 4 times/week OTC & Rx: not studied Single vitamin/prenatal vitamin: not studied	Preconception Multivitamin non-users: Younger (<20 & 20-24) or, unmarried, < 12 years of education, no health insurance, enrolled in Medicaid	

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Carmichael	Cross-sectional	N=2518 women	Preconceptional period:	Periconceptional	Participants were
2006	analysis	with estimated	not studied	vitamin non-users:	mothers of live-born
U.S.		delivery dates from		race (nonwhite),	infants without major
	*Data from the	1997-2000	Periconceptional period:	language speaking	malformation
	National Birth Defects Prevention		3 months before	(Spanish), education	
	Study		conception/one month	(low education),	
			after conception	maternal age	
			·	(younger<25), parity	
			Earlier pregnancy: the	(nulliparous), smoke	
			second and third month	(current smoker), a	
			after conception	history of miscarriage	
			·	(no history of	
			Later pregnancy: during	miscarriage)	
			the fourth month or later	<u> </u>	
			during pregnancy		
			Multivitamin:		
			multivitamin/prenatal		
			vitamin: folic acid		
			supplementation		
			Frequency: not defined		
			OTC & Rx/prenatal		
I			vitamin: not studied		

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date		-	& Folic acid Use		
Williams LM 2000 U.S.	*Data from Pregnancy Risk Assessment Monitoring System(PRAMS)	N=32,479 women who were identified from 19 states and U.S territories	Preconceptional period: during the month before pregnancy Periconceptional period & During pregnancy: not	Preconceptional users: Education (>12 years), age (>35 years), non-Medicaid recipients	Because data are self-reported 28 months after delivery, responses might be subject to recall bias, particularly for
			studied Multivitamin: multivitamin/prenatal vitamin		behaviors and experiences that occurred before the pregnancy
			Frequency: at least 4 times/week		
			OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: not studied		

Author & Date	Study Design	Population	Assessment of Vitamin & Folic acid Use	Predictors	Comments
Catov JM 2007 U.S.	*Data from the Pregnancy Exposure and Preeclampsia Prevention Study	N=1,823 women who enrolled at less than 16 weeks and followed through the postpartum visit	Periconceptional period: 6 months which included prior to conception, at the time of conception immediately after conception (not specifically defined) During pregnancy: not studied Multivitamin: multivitamin/prenatal vitamin Frequency: at least once/week OTC & Rx: not studied Single vitamin: not studied	Periconceptional multivitamin users: higher level of education, married, of normal body mass index (18.5-24.9), and to have participated in moderate or vigorous physical activity in the year before becoming pregnancy	The study relied on self-reported multivitamin use. Also, This study lack of information on the brand or dose of supplement

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Yu S	Cross-sectional	N=9953 women who	Periconceptional period:	Multivitamin non-users	The self-report
1996 U.S.	study	delivered live infants in the 1988 National	not studied	before pregnancy: black, unmarred,	nature of the mothers'
		Maternal and Infant	During pregnancy: after	age<30 or	questionnaire data
		Health Survey	pregnancy recognition	20 <age<34, less="" td="" than<=""><td>poses certain</td></age<34,>	poses certain
				or only have a high	limitations to the
	*Data: National		Multivitamin: not defined	school education,	study
	Maternal and Infant Health Survey			Multivitamin non-users	
			Frequency: ≥3 times/week	during pregnancy:	
				black/Asia, younger,	
			OTC & Rx/prenatal	low education,	
			vitamin: not studied	unmarried, smoking	
				status.	
			Single vitamin: vitamin A,	age, race (white),	
			vitamin C, vitamin E, folic	socioeconomics	
			acid were studied but not	married status	
			reported	(married), education	

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Tam LE 2004 Canada	Cross-sectional analysis	N=383 postpartum women in Toronto Hospital and the Mount Sinai Hospital	Periconceptional period: at least 4 weeks prior to conception until 8 weeks after conception During pregnancy: not studied Multivitamin: not defined	folic acid non users in pregnancy: unplanned pregnancy; lack of knowledge about folic acid Periconceptional folic acid/multivitamin users are more likely to be Jewish decent,	marital status, education, family income were not significant
			Frequency: not studied OTC & Rx/prenatal vitamin: not studied	parity (≥1)	
			Single vitamin: not studied		

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			Use		
Paulik E 2009 Hungary	*Data from a self-administrated questionnaire (response rate=69.8%)	N=349 voluntary pregnant women who were at the Department of Obstetrics and Gynecology and in the Pregnancy Care Center	Periconceptional period: not studied During pregnancy: not defined Pre-conceptional period: Not defined Multivitamin: not defined Frequency: not defined	Pre-conceptional folic acid users: age (older), pregnancy planning Folic acid users in pregnancy: age (older), gestational age (younger)	Multivitamin use in pregnancy or preconception was correlated with folic acid use
			OTC & Rx: not studied		
			Single vitamin: folic acid		

sign Population	Assessment of Vitamin	Predictors	Comments
	& Folic acid Use		
y women who have	Periconceptional period: before and during	Periconceptional multivitamin users: order (≥25), married or cohabiting, primiparous, non-smoker multivitamin users during pregnancy have the same predictors	The prevalence of using each supplement type can not be separated from the data
k	birth N=226,724 try women who have been recorded by Medical Birth Registry of Norway from 1998 through	birth try N=226,724 women who have been recorded by Medical Birth Registry of Norway from 1998 through 2004 Multivitamin: multivitamin alone/folic acid alone/both folic acid and multivitamin Frequency: not studied OTC & Rx: not studied	birth try N=226,724 women who have been recorded by Medical Birth Registry of Norway from 1998 through 2004 Multivitamin: multivitamin alone/folic acid and multivitamin Frequency: not studied Dirth try N=226,724 women who have before and during pregnancy (not specifically defined) During pregnancy: not defined Multivitamin: multivitamin alone/folic acid alone/both folic acid and multivitamin Frequency: not studied OTC & Rx: not studied Single vitamin/prenatal

Author & Date	Study Design	Population	Assessment of Vitamin & Folic acid Use	Predictors	Comments
Vollest SE	Cross-sectional	N=1,140	Periconceptional period:	Periconceptional	Household was not
2000	study	Norwegian	before or during the first	multivitamin users:	significant(p=0.25) in
Norway.		women who aged	2-3 months of their last	young (18-24), high	multiple logistic
	*Data: a national survey initiated by the Medical Birth Registry and the	18-45 years	recent pregnancies	education (university/college	regression result
	National Council on Nutrition and Physical Activity		During pregnancy: not defined	degree), married	
			Multivitamin: not defined		
			Frequency: daily or almost daily/less frequent than daily/ almost daily/never		
			OTC & Rx/prenatal vitamin: not studied		
			Single vitamin: vitamin B, folic acid		

Author & Date	Study Design	Population	Assessment of Vitamin & Folic acid Use	Predictors	Comments
Forster D 2009 Australia	Cross-sectional survey	N=588 women were approached in the antenatal clinic and the birth centre at around 36-38 weeks gestation	Periconceptional period: before and during pregnancy (not specifically defined) During pregnancy: not defined Multivitamin: folic acid alone/folic acid with multivitamin/folic acid with other vitamin Frequency: does/day and length of time were recorded OTC & Rx/prenatal vitamin: not studied Single vitamin: folic acid, vitamin B6	Multivitamin non-users during pregnancy: unmarried/living with a partner, low income (annual house income<\$30,000), Low education, parity (≥1)	The author did not undertook regression analysis to explore the predictor of the use of multivitamin during periconceptional period

Author & Date	Study Design	Population	Assessment of Vitamin & Folic acid Use	Predictors	Comments
Waston	Cross-sectional	N=1240 women	Periconceptional period:	Folic acid non-users in	
2006	survey	who gave birth in	1-2 months before	periconceptional	
Australia		Victoria	pregnancy recognition+ 3	period: age (younger),	
	45 (())(()		months after pregnancy	education (lower	
	*Data: the Victoria Survey of Recent		recogntiion	education), income	
	Mothers 2000 and the			(less income),	
	2001 NSW Child Health		During pregnancy: after	language speaking	
	Survey		pregnancy recognition	(Non-English	
			Multivitamin, falia acid	speaking), marital	
			Multivitamin: folic acid	status (unmarried),	
			supplementation	parity (multiparous), pregnancy planning	
			Frequency: daily (7 times	(unplanned), living	
			per week)	area (rural area)	
			OTC & Rx/prenatal		
			vitamin: not studied		

Author &	Study Design	Population	Assessment of Vitamin	Predictors	Comments
Date			& Folic acid Use		
Haugen M	Cross-sectional	N=40,817	Periconceptional period:	Multivitamin users in	Multivitamin use was
2008	study	Pregnant women	not studied	pregnancy: age (≥25),	combined with
Norway		in Norway who		BMI(≤24.9), smoking	mineral use and other
		were 17-18 weeks	During pregnancy: first 4-5	status during	dietary supplements.
	*Data from Norwegian	of gestation and	months	pregnancy	It could not be
	Mothers and Child Cohort Study(MoBa)	participates in		(non-smokers), parity	separated from data
	, , , , , , , , , , , , , , , , , , , ,	study between	Multivitamin: not defined	(primiparous),	
		February 2002		education (≥10 years	
		and February	Frequency: never/<1/1-7	of education)	
		2005	times per week		
			OTC & Rx: not studied		
			Single vitamin: vitamin A,		
			B1, B12, B6, B12, C, D, E,		
			and folic acid were studied		
			but prevalence rates were		
			not reported		
			prenatal vitamin: not		
			studied		

Author & Date	Study Design	Population	Assessment of Vitamin Use	Predictors	Comments
Lunet N 2008 Portugal	Cross-sectional analysis from a case-control study	N=836 mothers who were interviewed 24-72 hours after delivery at two major public hospitals in the North of Portugal	Periconceptional period: not studied, During pregnancy: without exactly refer to exact trimester Multivitamin: not defined Frequency: not studied OTC & Rx/prenatal vitamin: not studied Single vitamin: folic acid	Folic acid supplementation non-users during pregnancy: marital status (single), pregnancy planning (unplanned); Multivitamin users: education (less educated)	Frequency of multivitamin use and the content of multivitamin was not mentioned in this study;

Table 3.1. Sample size calculations for different effect sizes

Effect size	N1	N2	Total N	Power
20%	63	189	225	0.80
30%	28	84	112	0.82
40%	13	39	52	0.81
50%	9	27	36	0.83
60%	6	18	24	0.82

Two-sided Z tests were used with 1:3 ratio at alpha level 0.05
N1: required sample size for pre-conception multivitamin regular users
N2: required sample size for multivitamin regular users during pregnancy

Total N: the total required sample size in both groups
Effect size: the difference in the proportion of multivitamin user between the study groups

Table 4 1	Characteristics	of the	Study	Population	(N=402)
I UDIC T. I	Orial actoristics		Oluav	i obalation	111 TOE1

Table 4.1 Characteristics of the Study Population (N=402)				
Maternal	N (%) ^a			
Characteristics	222 (00.4)			
Hispanics	323 (80.4)			
Race*	40 (42 2)			
White Non-Hispanic	49 (12.2)			
White Hispanic	298 (74.3)			
Black or African American	23 (5.7)			
American Indian or Alaskan Native	27 (6.7)			
Asian or Asian American or Pacific Islander	3 (0.8)			
Some other group	1 (0.3)			
Educational Level	111 (07.1)			
Less than high school graduate	141 (35.1)			
High school graduate or GED	132 (32.8)			
Some college or vocational school	89 (22.1)			
College degree	28 (7.0)			
Master, doctorate or professional degree	12 (3.0)			
Marital Status	2- (2- 2)			
Single, never married	95 (23.6)			
Married, living with spouse	168 (41.8)			
Not married, but living with partner	120 (29.9)			
Separated	15 (3.7)			
Divorced	4 (1.0)			
Widowed	0			
Health Insurance Status	100 (100 0)			
No insurance	196 (48.8)			
Employer-based insurance	45 (11.2)			
Self-purchased insurance	4 (1.0)			
Medicaid	110 (27.4)			
Other public insurance ^b	47 (11.7)			
Language Speaking				
Spanish speaking	244 (60.7)			
English speaking	152 (37.8)			
Other language speaking	6 (1.5)			
Country of Birth: U.S.	163 (40.6)			
Primigravida	89 (22.2)			
Nulliparous	130 (32.5)			
History of Adverse Pregnancy Outcomes	129 (32.1)			
Miscarriage (<20 wk of gestation)	110 (27.6)			
Stillborn child (≥ 20 wk of gestation)	17 (4.3)			
Terminated birth	30 (7.5)			
Ectopic pregnancy	7 (1.8)			
Presence of Chronic Condition(s)	181 (45.0)			
Diabetes	81 (20.1)			
Depression	26 (6.5)			
Asthma or Allergies	35 (8.8)			

Table 4.1 (Continued) Characteristics of the Study Population (N=402)

Table 4.1 (Continued) Characteristics of the	
Maternal	N (%) ^a
Characteristics	
A Family History of Birth Defect(s) or Adverse	76 (19.3)
Pregnancy Outcome	
Down syndrome	12 (3.0)
Cleft lip or palate	4 (1.0)
Neural tube defect	5 (1.2)
Cystic fibrosis	0
Heart defect	6 (1.4)
Other ^c	63 (15.7)
Obesity (BMI≥30)	104 (25.9)
Pregnancy Planning	
Yes	171 (48.6)
No, not now	110 (31.3)
No, not at any time	71 (20.2)
Smoking Status	
Non-smoker	294 (73.3)
Past smoker, quit before pregnancy	55 (13.7)
recognition	
Past smoker, quit after pregnancy recognition	41 (10.2)
Current smoker	11 (2.7)
Periconceptional Binge Drinker	93 (23.1)
Periconceptional Recreational Drug User	11 (5.5)
Use of Prescription Medications during	
Pregnancy	
None	187 (46.6)
1-2 medications	165 (41.2)
More than 2 medications	49 (12.2)
Mean±s.d	
Maternal Age (yrs)	27.6±6.1
Gestational Age at Enrollment (wks)	30.7±8.0

^a Sample size might vary due to missing values

b Other public insurance: UNM/UNM Care (7.0%, 28/402), Indian Health Service (1.0%, 4/402), First Choice (1.0%, 3/402)

^c Other birth defect or adverse perinatal outcome: miscarriage (9.7%, 39/402), stillbirth (1.7, 7/402)

Table 4.2 Descriptive Statistics of Multivitamin Use Patterns (N=402)

Pattern of Use	N (%)
Three Study Groups	
Group 1-Pre-conceptional vitamin users	86 (21.4)
Group 2-Vitamin users after pregnancy	289 (71.9)
recognition*	
Group 3-Vitamin non-users	27 (6.7)
Prescription vs. OTC Multivitamin	
Prescription vitamins	116 (28.7)
OTC vitamins	219 (54.2)
Non-specified	40 (6.7)
The Most Common Brands of Vitamin**	
Wal-Mart prenatal vitamin	87 (21.6)
Walgreen prenatal vitamin	50 (12.4)
Mean±s.d	
Initiation of regular vitamin use (gestational	9.0±7.2 wks
weeks)***	
Number of days of vitamin use per week	6.0±1.9 days

^{*} There is only one woman who used single vitamin (folic acid) after pregnancy recognition, and she was still included in the Group 2.

** The sample size is limited to group 1 and group 2

^{***} The sample size is limited to group 2

Table 4.3 Characteristics of the Study Participants by Vitamin Use

Table 4.3 Characteristics of the				se
Maternal	Group 1	Group 2	Group 3	P
Characteristics	(n=86) ^a	(n=289) ^a	(n=27) ^a	value
	%	%	%	
Hispanics	17.3	76.2	6.5	<0.01
Maternal age (yrs)				0.16
≤21 yrs	13.8	75.0	11.3	
21-30 yrs	24.7	69.6	5.7	
>30 yrs	21.1	73.4	5.5	
Race*				0.20
Whites	21.0	72.9	6.1	
Black or African American	17.4	78.3	4.4	
Some other groups	21.0	72.9	6.1	
Educational Level				<0.01
High school education or less	16.5	77.3	6.2	
Some college or vocational school	23.6	66.3	10.1	
College degree or higher	50.0	47.5	2.5	
Marital Status				0.44
Single, never married	20.0	72.6	7.4	
Married or with a partner	22.9	70.8	6.3	
Separated/divorced/widowed	5.3	84.2	10.5	
Health Insurance Status				<0.01
No insurance	17.7	76.0	6.1	
Medicaid or other public insurance	44.9	49.0	6.1	
Employer-based or self-purchased	18.5	73.9	7.6	
Spanish-Speaking	16.0	77.9	6.2	<0.01
Primigravida	23.6	70.8	5.6	0.79
Nulliparous	25.4	70.0	4.6	0.23
History of Adverse Pregnancy	25.4	65.9	8.7	0.15
Outcomes				
Presence of Chronic Condition(s)	22.7	69.1	8.3	0.40
A family History of Birth Defect(s)	26.3	67.1	6.6	0.53
Country of Birth: U.S	27.0	66.3	6.8	0.07
Obesity (BMI≥30)	25.4	67.7	6.9	0.32
Planned Pregnancy	26.9	66.7	6.4	0.04
Smoking Status				0.40
Non-smoker	20.8	72.1	7.1	
Past, quit before pregnancy	30.9	61.8	7.3	
recognition				
Smoker in pregnancy	15.4	80.8	3.9	
Periconceptional Binge Drinker	21.5	69.9	8.6	0.66
Periconceptional Recreational Drug	18.2	81.8	0	0.52
User				
	•		•	•

Table 4.3 (Continued) Characteristics of Study Participants by Vitamin Use

Maternal Characteristics	Group 1 (n=86) ^a	Group 2 (n=289) ^a	Group 3 (n=27) ^a	P value
	%	%	%	
Use of Prescription Medications				0.16
during Pregnancy*				
None	17.1	74.9	8.0	
1-2 types of medications	23.0	71.5	5.5	
More than 2 types of medications	32.7	61.2	6.1	
<u>N</u>	<u>1ean±s.d</u>			
Maternal Age (yrs)	28.5±5.5	27.4±6.2	26.3±6.7	0.18
Gestational Age at Enrollment	31.2±7.8	30.5±8.1	31.6±7.3	0.63
(wks)				
^a Sample size might vary due to missing va	lues			

Table 4.4 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis*

Maternal Characteristics	ORs	95% CL	P value
Hispanics	0.68	0.29-1.61	0.38
Maternal age (yrs)	0.00	0.23-1.01	0.58
			0.02
≤21 yrs	1.00	0.05.2.71	0.07
21-30 yrs	1.88	0.95-3.71	0.07
>30 yrs	1.27	0.61-2.65	0.53
Race			
Whites			
Black or African American	0.77	0.28-2.13	0.61
Some other groups	0.33	0.11-1.04	0.06
Educational Level			
High school education or less			
Some college or vocational school	1.06	0.56-2.04	0.86
College degree or higher	2.47	0.91-6.72	0.08
Health Insurance Status			
No insurance			
Employer-based or self-purchased	1.42	0.46-4.46	0.54
Medicaid or other public insurance	0.82	0.38-1.78	0.61
Spanish-speaking	0.69	0.22-2.13	0.52
History of Adverse Pregnancy	0.87	0.52-1.47	0.61
Outcomes			
Country of Birth: U.S	0.76	0.24-2.44	0.64
Planned Pregnancy	1.77	1.08-2.89	0.02
Use of Prescription Medications			
during Pregnancy			
None			
1-2 types of medications	1.52	0.89-2.59	0.13
More than 2 types of medications	2.17	0.94-4.99	0.07

^{*} All Odds ratios are adjusted for all variables in the table

Table 4.5 Interaction between Predictors in Ordinal Logistic Regression and Polychotomous logistic Regression Model

Interaction	Ordinal Logistic Regression	Polychotomous Logistic Regression	
	P-value	P-value	
Ethnicity*Race	0.20	0.91	
Ethnicity*Place of Birth	0.13	0.48	
Ethnicity*Language Speaking	N.A	N.A	
Race*Language Speaking	0.29	0.89	
Race*Place of Birth	0.44	0.68	
Place of Birth*Language Speaking	0.17	0.69	
Education*Health Insurance Status	0.01	0.96	
Education*Pregnancy Plan	0.69	0.85	

Table 4.6 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis among Women without Health Insurance*

Maternal Characteristics	ORs	95% CL	P value
Educational Level			
High school education or less			
Some college or vocational school	2.11	0.70-6.34	0.18
College degree or higher	0.24	0.03-2.17	0.20

^{*} Odds ratio was adjusted for ethnicity, maternal age, race, language speaking, history of adverse pregnancy outcomes, country of birth, pregnancy planning, and use of prescription medications were adjusted.

Table 4.7 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis among Women with Private Insurance *

- J J <u></u>			
Maternal Characteristics	ORs	95% CL	P value
Educational Level			
High school education or less			
Some college or vocational school	0.09	0.01-1.32	0.08
College degree or higher	0.33	0.02-6.73	0.47

^{*} Odds ratio was adjusted for ethnicity, maternal age, race, language speaking, history of adverse pregnancy outcomes, country of birth, pregnancy planning, and use of prescription medications were adjusted.

Table 4.8 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis <u>among Women with Public Insurance</u>*

Maternal Characteristics	ORs	95% CL	P value
Educational Level			
High school education or less			
Some college or vocational school	1.04	0.39-2.80	0.94
College degree or higher	14.45	1.78-117.66	0.01

^{*} Odds ratio was adjusted for ethnicity, maternal age, race, language speaking, history of adverse pregnancy outcomes, country of birth, pregnancy planning, and use of prescription medications were adjusted.

Table 4.9 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis among Women without Insurance*

Maternal Characteristics	ORs	95% CL	P value
Educational Level			
High school education or less			
Some college or vocational school	2.11	0.70-6.34	0.18
College degree or higher	0.24	0.03-2.17	0.20

^{*} Odds ratio was adjusted for ethnicity, maternal age, race, language speaking, history of adverse pregnancy outcomes, country of birth, pregnancy planning, and use of prescription medications were adjusted.

Table 4.10 Predictors of Vitamin Use: Results of Multivariate Ordinal Logistic Regression Analysis among Women with Insurance*

<u> </u>			
Maternal Characteristics	ORs	95% CL	P value
Educational Level			
High school education or less			
Some college or vocational school	0.92	0.39-2.17	0.85
College degree or higher	5.90	1.89-18.44	<0.01

^{*} Odds ratio was adjusted for ethnicity, maternal age, race, language speaking, history of adverse pregnancy outcomes, country of birth, pregnancy planning, and use of prescription medications were adjusted.

Table 4.11 Predictors of Vitamin Use: Results of Univariate Polychotomous Logistic Regression Analysis*

Maternal Characteristics	Group 1 vs. Group 2		Group 3 vs. Group 2	
	ORs	95% CL	ORs	95% CL
Hispanics	0.33	0.19-0.57	0.61	0.23-1.60
Maternal age (yrs)				
≤21 yrs	1.00		1.00	
21-30 yrs	1.94	0.94-3.99	0.54	0.21-1.38
>30 yrs	1.57	0.72-3.39	0.50	0.18-1.40
Race				
Whites	1.00		1.00	
Black or African American	0.77	0.25-2.35	0.67	0.09-5.26
Some other groups	1.54	0.64-3.69	3.35	1.13-9.92
Educational Level				
High school education or less	1.00		1.00	
Some college or vocational	1.67	0.92-3.02	1.89	0.80-4.47
school				
College degree or higher	4.94	2.44-10.00	0.65	0.08-5.18
Health Insurance Status				
No insurance	1.00		1.00	
Employer-based or	3.90	1.97-7.75	1.28	0.56-2.96
self-purchased				
Medicaid or other public	1.06	0.62-1.84	1.55	0.41-5.91
insurance				
Spanish-speaking	0.43	0.27-0.71	0.65	0.29-1.45
History of Adverse Perinatal	0.67	0.41-1.10	1.17	0.49-2.79
Outcomes				
Country of Birth: U.S	1.76	1.08-2.82	1.15	0.52-2.57
Planned Pregnancy	1.95	1.15-3.30	1.13	0.48-2.65
Use of Prescription medications				
during pregnancy				
None	1.00		1.00	
1-2 medications	1.41	0.83-2.39	0.71	0.30-1.69
More than 2 medications	2.33	1.14-4.78	0.93	0.25-3.43

^{*} All Odds ratios are adjusted for all variables in the table

Table 4.12 Predictors of Vitamin Use: Results of Polychotomous Multivariate Logistic Regression Analysis*

Maternal Characteristics	eristics Group 1 vs. Group 2		Group 3	3 vs. Group 2
	ORs	95% CL	ORs	95% CL
Hispanics	0.62	0.24-1.59	0.87	0.13-5.70
Maternal age (yrs)				
≤21 yrs	1.00		1.00	
21-30 yrs	1.57	0.68-3.63	0.41	0.13-1.29
>30 yrs	0.95	0.38-2.40	0.46	0.13-1.56
Race				
Whites	1.00		1.00	
Black or African American	0.50	0.14-1.81	0.45	0.05-4.09
Some other groups	0.64	0.16-2.56	3.90	0.71-21.37
Educational Level				
High school education or less	1.00		1.00	
Some college or vocational	1.34	0.63-2.86	1.36	0.46-4.05
school				
College degree or higher	2.73	0.92-8.07	0.70	0.06-7.81
Health Insurance Status				
No insurance	1.00		1.00	
Employer-based or	1.88	0.52-6.80	2.38	0.31-18.47
self-purchased				
Medicaid or other public	0.83	0.33-2.09	1.03	0.26-4.04
insurance				
Spanish-speaking	0.37	0.09-1.58	0.34	0.03-3.46
History of Adverse Prenatal	1.32	0.73-2.40	3.04	1.16-7.98
Outcomes				
Country of Birth: U.S	0.32	0.07-1.47	0.20	0.02-1.91
Planned Pregnancy	2.29	1.29-4.09	1.31	0.52-3.29
Use of Prescription Medications				
during Pregnancy				
None	1.00		1.00	
1-2 types of medications	1.52	0.81-2.84	0.75	0.28-2.02
More than 2 types of	2.00	0.80-5.01	0.46	0.05-4.19
medications				

^{*} All Odds ratios are adjusted for all variables in the table

APPENDICES

APPENDIX A Data Dictionary for SMART Study

Question	Variable name	Categories	Value
	GENERAL INF		
Subject ID	subjID	Text	
Date of interview	dateint	Date (mm/dd/yy)	
Location of interview	locat	UNMH Fetal	1
		Monitoring Clinic	
		(FMC)	
			2
		Triage	3
		General	4
		Satellite Clinic-West	5
		Mesa	
		Satellite Clinic- South	6
		Broadway	7
		Satellite Clinic C	8
		Satellite Clinic D	
Algorithm for	First two digits	Other First two digits for	
assigning Subject IDs	First two digits -Locat1009036	location:	
assigning Subject ibs	5ion	UNMH Fetal	10
	Next two	Monitoring Clinic	10
	digits- Year of	(FMC)	
	interview	(1.11.5)	20
	Last four	Triage	30
	digits- Serial	General	41
	no.	Satellite Clinic-West	42
		Mesa	
		Satellite Clinic- South	43
		Broadway	
		Satellite Clinic –North	44
		Valley	50
		Satellite Clinic D	
		Other	
		Novt two digits for	08
		Next two digits for year of interview:	00
		2008	0001-01
		2000	24
		Last four digits for	
		serial no.:	
If location is "Other",	locatot	Text	
please specify			
Prenatal care provider's	obgyn	Text	
last name			
Examiner's last name	examln	Text	

Question	Variable name	Categories	Value		
Patient's phone number	patpn	Text			
	<u>DEMOGRAPHICS</u>				
How old are you?	momage	Continuous			
What is your marital	mommarit	Single, never married	1		
status now?		Married, living with	2		
		spouse	3		
		Not married, but living			
		with partner	4		
		Separated from	5		
		spouse	6		
		Divorced			
		Widowed			
Are you Hispanic,	mometh	Yes	1		
Latino or of Spanish		No	0		
descent?		NA/Intia Nia a Lita a anta	0		
How do you describe	race	White Non-Hispanic	0		
yourself?		White Hispanic	1		
		Black or African American	2		
		American Indian or	3		
		Alaskan Native	4		
		Asian or Asian	4		
		American or Pacific	5		
		Islander	6		
		Some other group			
		Prefer not to report			
If race is American	race3	Tribe	1		
Indian or Alaskan		Pueblo	2		
Native, then specify					
If race is others, then specify	raceoth	Text			
What is the highest	momedlev	Less than high school	1		
level in school you have		graduate			
completed?		High school graduate or GED	2		
		Some college or	3		
		vocational school	4		
		College degree Masters, doctorate or	4 5		
		professional degree]		
What is your health	momins	No insurance	1		
insurance status?		Employer-based	2		
		insurance	3		
		Self-purchased	4		
		insurance	5		
		Medicaid			
		Other public			

Question	Variable name	Categories	Value
		insurances	
If you have other public	momins5	Indian Health Service	1
health insurance, then		VA	2
specify		First Choice	3
-		UNM/UNM Care	4
Does your insurance	prescov	Yes	1
cover prescription		No	0
drugs?			
Were you born in the	usborn	Yes	1
Unites States?		No	0
How long have you	momyr	Continuous	
lived in the United			
States?			
What language do you	momlang	English	1
mostly use at home?		Spanish	2
		Some other language	3
If some other language	momlang3	Text	
is used mostly at home,			
then specify			
Do you currently smoke	smkmom	Yes	1
cigarettes or use		No	0
tobacco?			
If yes, how many	smkpreg	Continuous	
cigarettes do you			
usually smoke in a day?			
If no, have you smoked	smkmp1	Yes	1
>100 cigarettes in your		No	0
life?			
If you have smoked	smkstop	Before I became	1
>100 cigarettes in your		pregnant	2
life, when did you stop		After I realized that I	
smoking?		was pregnant	
Have you ever drank	havdrnk	Yes	1
alcohol?		No	0
How many drinks	HIGH	Continuous	
typically can you hold			
before you feel high?			
How many drinks	HOLD	Continuous	
typically can you hold			
before passing out or			
falling asleep?		<u> </u>	
What was the first day	Imenst	Date (mm/dd/yy)	
of your last menstrual			
period?			
During a month or so	drink4m	Continuous	
around your last			
menstrual period before			

Question	Variable name	Categories	Value
you got pregnant, how			
many times did you			
drink 4 or more drinks			
on one occasion?			
During the year before	worry	Yes	1
you got pregnant, did		No	0
close friends or relatives			
worry or complain about			
your drinking habits?			
During the year before	eyem	Yes	1
you got pregnant, did		No	0
you ever take a drink			
first thing in the morning			
to get yourself going?			
During the year before	amnesm	Yes	1
you got pregnant, did a		No	0
friend or family member			
tell you about things you			
said or did while you			
were drinking that you			
could not remember?			
During the year before	cutm	Yes	1
you got pregnant, did		No	0
you feel you need to cut			
down on your drinking?			
TWEAK SCORE	tweak1 (High)	Continuous	
	tweak2 (Hold)		
Algorithm for calculating	TWEAK HIGH		
TWEAK score			
	TOLERANCE:		
	2, when no. of		
	drinks >=3; 0 if		
	no. of drinks <3		
	WORRY:		
	Yes= 2 points;		
	No= 0 pts.		
	EYEM:		
	Yes= 1 pt., No=		
	0 pt		
	AMNESM:		
	Yes= 1 pt., No=		
	0 pt		
	CUT:		
	Yes= 1 pt., No=		
	0 pt		
	TWEAK HOLD		
	TWEAK HOLD		
			[

Question	Variable name	Categories	Value
	TOLERANCE:		
	2, when no. of		
	drinks >=6; 0 if		
	no. of drinks <6		
	WORRY:		
	Yes= 2 points;		
	No= 0 pts.		
	EYEM:		
	Yes= 1 pt., No=		
	0 pt		
	AMNESM:		
	Yes= 1 pt., No=		
	0 pt		
	CUT:		
	Yes= 1 pt., No=		
	0 pt		
MEDICAL	AND REPROD	UCTIVE HEALTH	
What was your	wtmpre	Continuous	
pre-pregnancy weight?			
What was your	htmpre	Continuous	
pre-pregnancy height?			
Researcher Calculated	BMI	Continuous	
BMI			
Do you have a medical	medcon	Yes	1
condition or problem		No	0
that requires ongoing,			
periodic, or occasional			
treatment?			
If yes, specify			
III a a da a air a duli da DD)		. Vara	
Hypertension (High BP)	medcon1	Yes	1
		No	0
Depression		Van	4
Depression	medcon2	Yes	1
		No	0
Diabotos	modoon?	Voc	
Diabetes	medcon3	Yes	1
		No	0
Anxiety	medcon4	Yes	1
Allalety	111606011 4	No	0
		INU	
Seizure Disorder	medcon5	Yes	1
OCIZUIE DISOIUEI	IIIGUCUIIJ	No	0
		INO	
	1	1	

Question	Variable name	Categories	Value
Migraine headaches	medcon6	Yes No	1
Thyroid disorder	medcon7	Yes No	1 0
Rheumatoid Arthritis	medcon8	Yes No	1 0
Asthma or Allergies	medcon9	Yes No	1 0
Heart Disease	medcon10	Yes No	1 0
Cancer	medcon11	Yes No	1 0
Hepatitis	medcon12	Yes No	1 0
Other (s)	medcon13	Yes No	1 0
Specify the type of diabetes	diabetes	Gestational Type I Type II	1 2 3
If you have other medical condition, not listed, please specify	othcon	Text	
If diabetic, how likely do you think uncontrolled high blood sugar could harm your developing baby by causing birth defects or other serious health problems?	diabet1	Likert Scale	1-5
How likely do you think uncontrolled asthma could harm your developing baby by causing birth defects or other serious health problems?	asthma	Likert Scale	1-5

Question	Variable name	Categories	Value
Have you ever had gestational diabetes?	gestdia	Yes, in a previous pregnancy only Yes, in the current pregnancy only	2
		Yes, in a previous pregnancy and in the current pregnancy No, never had	3
		gestational diabetes No, never been pregnant before	4
			5
Did you plan to get	planchld	Yes	1
pregnant with this child?		No, not at any time	2
			3
Were you or your partner doing anything to try to prevent becoming pregnant with this child?	bcontr	Yes No	0
If Yes, which method were you using?	bcontr1	Condoms Diaphragm Birth control pills Withdrawal IUD Rhythm Depo Provera, Implanon or Norplant Other	1 2 3 4 5 6 7
If other method used, please specify	othmtd	Text	
Did you take any fertility drugs to help you get pregnant with this child, like Clomid, Metrodin, Fertinex, or Pergonal?	fertdrg	Yes No	1 0
If fertility drug taken, please specify	fertdrg1	Text	
Have you or members of your immediate family (mother or	bdefh	Yes No	1 0

Question	Variable name	Categories	Value
sisters) or the			
immediate family of			
your baby's father had			
any babies with birth			
defects (including			
babies that might not			
have survived)?			
If 'Yes', please specify			
Down syndrome	bdefhs1	Yes	1
		No	0
Cleft lip or palate	bdefhs2	Yes	1
		No	0
Neural tube defect	bdefhs3	Yes	1
		No	0
0 (5)			4
Cystic fibrosis	bdefhs4	Yes	1
		No	0
Llagut dafaat	h d a fla a F	Vac	
Heart defect	bdefhs5	Yes	1
		No	0
Other	bdefhs6	Yes	1
Other	Dueiliso	No	0
If other, please specify	bdoth	text	U
What is the date your	bdate	Date (mm/dd/yy)	
baby is due to be born?	Saaro		
How was your due date	dateest1, 2, and	Last menstrual period	1
estimated	3	Ultrasound	2
		Physical exam	3
What is the gestational	gestage	Continuous	
age of your baby?	3 3 -		
How many times	gravid	Continuous	
(including this			
pregnancy) have you			
been pregnant?			
How many live-born	parity	Continuous	
children have you had?	-		
Have you ever had a	miscrg	If no, then '0'	
miscarriage (<20 wk of		If yes, then Continuous	
gestation). If yes, how			
many?			
Have you ever had a	stillbrn	If no, then '0'	
stillborn child (≥ 20 wk		If yes, then Continuous	
of gestation). If yes,			
how many?			
Have you ever had a	termin	If no, then '0'	
pregnancy terminated?		If yes, then Continuous	

Question	Variable name	Categories	Value
If yes, how many?			
Have you ever had an	ectop	If no, then '0'	
ectopic pregnancy. If		If yes, then Continuous	
yes, how many?			
For this pregnancy, how	realize	Continuous	
many weeks after your			
last menstrual period			
did you first think you			
were pregnant?			
For this pregnancy, how	prenwk	Continuous	
many weeks after your			
last menstrual period			
did you first go to see			
a doctor or other health			
care provider or go to			
the clinic for prenatal			
care?			
Have you had any complications in this			
pregnancy so far?			
Bleeding	bleed	Yes	1
Dieeding	Dieeu	No	0
		140	o l
High blood pressure	highbp	Yes	1
l light blood procedure	gp	No	0
Diabetes	diabet	Yes	1
		No	0
Other	othcom	Yes	1
		No	0
If other, please specify	othcoms	Text	
Have you experienced	mornsick	Yes	1
morning sickness		No	0
during this pregnancy?			
USE OF MEDIC		UPPLEMENTS DURI	<u>NG</u>
	<u>PREGNAN</u>	<u>CY</u>	
Did you take a	multivit	Yes	1
multivitamin regularly (4		No	0
times a week or more)			
during the month before			
your last menstrual			
period?			
Have you taken any	vitreg	Yes, multivitamins	1
VITAMINS regularly (4		Yes, a single vitamin	2
times/week or more)		No	3

Question	Variable name	Categories	Value
since you became			
pregnant?			
If "Yes", then specify	vitRx	Prescription OTC	1 2
If Brand name vitamin	vitregname	Text	
taken, then specify			
If "yes" when did you	vitdate	Date (mm/dd/yy) or 0	
start taking vitamins?	vitwk	Continuous or 0	
How many days during	vitdays	Continuous	
the last week did you			
take vitamins?	la a ula a	No.	4
Have you taken any DIETARY SUPPLEMENTS (including iron	herbsup	Yes No	0
supplements) or HERBAL PRODUCTS			
on a regular basis since your last menstrual period?			
If 'Voo' to borbal	horboun1	Horbo	1
If 'Yes' to herbal	herbsup1	Herbs	1 2
products, please specify		Tablets or capsules Teas	3
		Other	4
If other, please specify	herboth	Text	
How often do you take	herboft	Regularly	1
them?	11010010	When I feel sick	2
If taken regularly,	herbreg	Text	
specify time	· ·		
Please specify any		Text	
other dietary			
supplements or			
products and reason for			
taking it	aa.14a.a.4		
Broduct1 Boscon/Condi	suppl1;reas1		
Product1,Reason/Condition	suppl2;reas2 suppl3;reas3		
Produc2,Reason/Condit	Suppio,ieaso		
ion			
Product3,Reason/Condi			
tion			
Have you had any	crave	Yes	1
cravings for non-food		No	0
items or really "strange"			
foods?			
If 'yes' what did you	item 1	If no, then 0	
crave, do you eat it, and		If yes , then text	

Question	Variable name	Categories	Value
how often do you eat it?			
	howoft1	Text	
	eat1	Yes	1
	- 3	No	0
	item2	If no , then 0 If yes , then text	
	howoft2	Text	
	eat2	Yes	1
		No	0
	item3	If no, then 0	
		If yes, then text	
	h #2	Tand	
	howoft3	Text	
	eat3	Yes	1
		No	0
Have you ever taken	recdrug	Yes	1
recreational drugs?	recurug	No	0
If Yes, specify the recreational drug that			
you took Marijuana/Hashish	hashish	Before pregnancy	1
Wangaana/naomon	ridoriiori	Boloro programoy	
		1 month prior to LMP	2
		or during this	
Heroin	heroin	pregnancy	1
		Before pregnancy	
		4	2
		1 month prior to LMP or during this	
		pregnancy	
Have you gone through	methtrt	F - 3	1
methadone treatment?		Neves	
		Never	2
		Completed treatment	
		before pregnancy	3
		Undergoing treatment	
Cocaine/Crack	cocaine	Undergoing treatment during current	1
		pregnancy	
			2

Inhalants (Glue, solvent) Inhalants (Glue, solvent) Inhalant I month prior to LMP or during this pregnancy 2 Before pregnancy 2 Before pregnancy 1 month prior to LMP or during this pregnancy 2 Tecdrug1 Before pregnancy 1 month prior to LMP or during this pregnancy 2 Other (name) Inhalant I month prior to LMP or during this pregnancy 2 Continuous Tecdrug2 Before pregnancy 1 month prior to LMP or during this pregnancy 2 Continuous 1 month prior to LMP or during this pregnancy 2 Continuous Before pregnancy 1 month prior to LMP or during this pregnancy 2 Continuous Before pregnancy 1 month prior to LMP 1	Question	Variable name	Categories	Value
solvent) or during this pregnancy Before pregnancy 1 month prior to LMP or during this pregnancy 2 Other (name) recdrug1 Before pregnancy 1 month prior to LMP or during this pregnancy 1 month prior to LMP or during this pregnancy Continuous recdrug2 when2 Before pregnancy 1 month prior to LMP or during this pregnancy 1 month prior to LMP or during this pregnancy Continuous Before pregnancy 1 month prior to LMP or during this pregnancy Continuous Before pregnancy Continuous Before pregnancy			Before pregnancy	
methamp 1 month prior to LMP or during this pregnancy 2 Tecdrug1 Before pregnancy 2 Tecdrug1 Before pregnancy 1 month prior to LMP or during this pregnancy 1 month prior to LMP or during this pregnancy 2 Continuous Pefore pregnancy 1 month prior to LMP or during this pregnancy 2 Tecdrug2 Before pregnancy 1 month prior to LMP or during this pregnancy 1 month prior to LMP or during this pregnancy 2 Continuous Before pregnancy Continuous Before pregnancy		inhalant	•	1
Methamphetamines methamp 1 month prior to LMP or during this pregnancy Performed and the proof of the proo	,		_	2
Methamphetamines or during this pregnancy 2 Other (name) recdrug1 Before pregnancy 1 month prior to LMP or during this pregnancy Continuous recdrug2 When2 Before pregnancy 1 month prior to LMP or during this pregnancy 2 Continuous 1 month prior to LMP or during this pregnancy Continuous Before pregnancy Continuous Before pregnancy Continuous Before pregnancy			Before pregnancy	
Other (name) recdrug1 Before pregnancy 1 month prior to LMP or during this pregnancy Continuous recdrug2 when2 Before pregnancy 1 month prior to LMP or during this pregnancy 1 month prior to LMP or during this pregnancy Continuous Before pregnancy Continuous Before pregnancy	Methamphetamines	methamp	or during this	
Other (name) when1 I month prior to LMP or during this pregnancy Continuous recdrug2 when2 Before pregnancy 1 month prior to LMP or during this pregnancy Continuous 1 continuous Continuous Before pregnancy Continuous Before pregnancy			pregnancy	2
when1	Other (name)	recdrug1	Before pregnancy	
Other (name) recdrug2 when2 Before pregnancy 1 1 month prior to LMP or during this pregnancy Continuous Before pregnancy 2	Other (hame)	when1	•	1
Other (name) recdrug2 when2 Before pregnancy 1 1 month prior to LMP or during this pregnancy Continuous Before pregnancy			pregnancy	2
Other (name) when2 Before pregnancy 1 1 month prior to LMP or during this pregnancy Continuous Before pregnancy			Continuous	
when2 1 month prior to LMP or during this pregnancy Continuous Before pregnancy		recdrug2		
or during this pregnancy Continuous Before pregnancy	Other (name)	when2	Before pregnancy	1
Before pregnancy			or during this	2
			Continuous	
1 month prior to I MP			Before pregnancy	
or during this pregnancy			_	
If respondent used >=1 recdrugpreg Yes No 0 LMP or during this	drug(s) 1 month prior to LMP or during this	recdrugpreg		
pregnancy Did you discuss the safdis Yes 1	· · ·	eafdie	Ves	1
safety of medications in No 0		Garais		l -
pregnancy with any	pregnancy with any			
health care provider (physician,	I			
nurse-midwife,	nurse-midwife,			
physician assistant, or				
pharmacist)? Have you had any vaccine Yes 1		vaccine	Yes	1

Question	Variable name	Categories	Value
vaccinations since your		No	0
last menstrual period			
If yes to vaccinations,			
then please specify			
Flu	vaccineF	Yes	1
Olla e e		No	0
Other	vaasina O	Vee	1
	vaccineO	Yes No	1 0
If vaccination other than	Othvacc	Text	0
flu, then specify	Ollivacc	Text	ļ
Have you taken any	presmed	Yes	1
medications	presined	No	0
PRESCRIBED by your			
doctor or any other			
health care provider			
since your last			
menstrual period, even			
if you stopped taking			
them once you knew			
you were pregnant?			
Desire a Maio sono ano ano	-4	Vaa	1
During this pregnancy, did you take any	otcmed	Yes No	1 0
OVER-THE-COUNTER		NO	0
MEDICATIONS (sold			
without prescription)?			
If yes, please specify			
Pain/fever medications	pain	Yes	1
		No	0
Need Decements	alless :	Vaa	
Nasal Decongestants,	allrgy	Yes	1
Allergy, Cough Medications		No	0
IVICUICALIUIIS	antdiar	Yes	1
Antidiarrheal	aritalar	No	Ö
Medications			
	digesmed	Yes	1
		No	0
Heartburn, Dyspepsia,			
Antiemetic, Laxative			
Medications	antfungl	Yes	1
		No	0
Antifungal Medications			
(taken for vaginal yeast	nicorcth	Vac	
infection or thrash)	nicoreth	Yes	1
		No	0

Question	Variable name	Categories	Value
Nicotine Replacement			
Therapy (for smoking cessation)			
Cessalion)			
OTC Medications and			
their perceptions			
Acetaminanhan	tylenol	Yes	1
Acetaminophen (Tylenol)	tylerioi	No	0
(13101101)			
	tylenolP	Likert Scale	1-5
	aspirin	Yes	1
Aspirin	азрин	No	0
/ topii iii			
	aspirinP	Likert Scale	1-5
	ibuprof	Yes	1
Ibuprofen (Advil,		No	0
Motrin)	'l (D	L'I a d O a da	4.5
	ibuprofP	Likert Scale	1-5
	ketopro	Yes	1
Katanyafan (Oyudia)		No	0
Ketoprofen (Orudis)	ketoproP	Likert Scale	1-5
	naprox	Yes	1
Naproxen (Aleve)		No	0
Tapionon (Moto)	naproxP	Likert Scale	1-5
	chlorph	Yes No	0
Chlorpheniramine		140	
(Chlor-Trimeton)	chlorphP	Likert Scale	1-5

Question	Variable name	Categories	Value
	benadrl	Yes	1
		No	0
Benadryl	benadrlP	Likert Scale	1-5
Dogudoonhodrino	sudafed	Yes No	1 0
Pseudoephedrine (Sudafed)	sudafedP	Likert Scale	1-5
	clarzyr	Yes No	1 0
	clarzyrP	Likert Scale	1-5
	pepbsml	Yes No	1 0
Kaopectate, Pepto Bismol	pepbsmIP	Likert Scale	1-5
If you took prescription medications regularly before you got pregnant, did you change the use of these medications when you realized you are pregnant?			
Did not take prescription medications regularly before pregnancy	usechngA	Yes No	0
Discontinued the use upon recognition of	usechngB	Yes No	1 0
pregnancy	usechngC	Yes No	1 0
Decreased the use (dose or frequency)	usechngD	Yes No	1 0
Increased the use	usechngE	Yes No	1 0
Stayed the same, continued without any			

Question	Variable name	Categories	Value
change			
If discontinued the use upon recognition of pregnancy, please specify medication name	usechng2	Text	
If decreased the use (dose or frequency), please specify medication name	usechng3	Text	
If Increased the use, please specify medication name	usechng4	Text	
If medication stayed the same, specify medication name	usechng5	Text	
If you changed the use of a medication upon recognition of pregnancy, why?	medchng	Provider recommendation Family or friend suggestion Self-initiated Financial constraints Other	1 2 3 4 5
If other, then specify	chngoth	Text	

Question	Variable name	Categories	Value
If a women plans a pregnancy or finds out that she is currently pregnant, she should	pregplan	Stop taking all medications immediately to protect the baby	1
		Continue taking only those medications that are absolutely necessary and check with her doctor to see if the medications are safe for the baby Continue taking	2
		necessary medications but reduce the dose or the number of days you take them to limit the amount that gets to the baby Continue with all medications as needed since	3
		medications are safe for the baby	
When a woman uses medications regularly during pregnancy, how often can medications cause birth defects?	medbdef	Never Sometimes Often Very Often Always	1 2 3 4 5
Which statement best describes your view about women drinking alcohol during pregnancy?	pregalc	Pregnant women should abstain from drinking any alcohol (even small amounts) during pregnancy	1
		It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than once a week	2
		It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than one	3

Question	Variable name	Categories	Value
		drink per day	4
		It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than two drinks per day	5
		It is OK for a woman to drink during pregnancy as long as she does not drink hard liquor (i.e., vodka, whiskey, brandy) but only drinks wine or beer	
During your current pregnancy, have you ever asked with anyone about the safety of medications you are taking for your baby?	safemed	Yes No	1 0
If yes, please specify: Your primary care doctor or provider	safemed1	Yes No	1 0
Your OB/GYN doctor or midwife	safemed2	Yes No	1
A pharmacist	safemed3	Yes No	1
A member of your family, spouse	safemed4	Yes No	1 0
A friend, partner	safemed5	Yes No	1
Other	safemed6	Yes No	1
Any other Hx provider	safemed7	Yes No	1 0
If other, please specify Please check any sources below in which you have looked for	safeoth	Text	

Question	Variable name	Categories	Value
information about the			
safety of medications			
for your baby?			
	babyinfA	Yes	1
I have never looked at		No	0
any of these sources about the safety of			
medications for my			
baby.	babyinfB	Yes	1
A minternet web site(s)		No	0
An internet web site(s).	babyinfC	Yes	1
	babyiiiio	No	0
A book.			
	babyinfD	Yes	1
A magazine		No	0
Amagazine	babyinfE	Yes	1
		No	0
Pregnancy information			
telephone service/hotline (i.e.,			
OTIS, Nurse Advisory	babyinfF	Yes	1
Line)		No	0
Others	h ah sinfO	Vaa	
Other	babyinfG	Yes No	0
		140	O
I have not had any			
questions about the			
safety of medications for my baby and have	babyinfH	Yes	1
not looked at any of	babyiiiii	No	0
these sources.			
Clinic pamphlet or brochure			
If checked an internet	babyinf2	Text	
website, please specify	,		
the name of the website	habita fo	Taxet	
If used any other source, please specify	babyinf6	Text	
Is the patient currently	insulin	Yes	1
on Insulin		No	0
Is the patient currently	oralhypo	Yes	1
on any oral hypoglycemic?		No	0
hypogryceniic :			

Question	Variable name	Categories	Value
If yes, specify the type	sulfnylu	Yes	1
		No	0
	biguanid	Yes	1
		No	0
	TZD	Yes	1
	120	No	0
Perception regarding	insulinP	Likert scale	1-5
Insulin use		Emore oddio	'
Perception regarding	sulfnylP	Likert scale	1-5
Sulfonylurea (Oral			
Hypoglycemic) use			
Perception regarding	bguanidP	Likert scale	1-5
Biguanid (Oral			
Hypoglycemic) use	T700	12 - 4 1-	4.5
Perception regarding	TZDP	Likert scale	1-5
Thiazolidinedione (Oral Hypoglycemic) use			
Is the patient currently	ICS	Yes	1
on ICS	100	No	0
Is the patient currently	BetaA	Yes	1
on BetaA		No	0
Is the patient currently	Steriod	Yes	1
on steriod		No	0
Perception regarding	ICSP	Likert scale	1-5
ICS use	DataAD	Liberteede	4.5
Perception regarding BetaA use	BetaAP	Likert scale	1-5
Perception regarding	SteriodP	Likert scale	1-5
steroid use	Oterioui	LINCIT SCAIC	1-5
Is the patient currently	Antipsychotics	Yes	1
on Antipsychotics	.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	No	0
Is the patient currently	Nortiptyline	Yes	1
on Nortiptyline		No	0
Is _the patient currently	Bupropion	Yes	1
on Bupropion	0	No	0
Is the patient currently	Citalopram	Yes	1
on Citalopram	Facitalogram	No Voc	0
Is the patient currently on Escitalopram	Escitalopram	Yes No	0
Is the patient currently	Fluoxetine	Yes	1
on Fluoxetine	. 100,101110	No	0
Is the patient currently	Paroxetine	Yes	1
on Paroxetine		No	0
Is the patient currently	Sertraline	Yes	1
on Sertraline		No	0
Is the patient currently	Venlafaxine	Yes	1

Question	Variable name	Categories	Value
on Venlafaxine		No	0
Perception regarding Nortiptyline use	AntipsychoticsP	Likert scale	1-5
Perception regarding Antipsychotics use	NortiptylineP	Likert scale	1-5
Perception regarding Bupropion use	BupropionP	Likert scale	1-5
Perception regarding Citalopram use	CitalopramP	Likert scale	1-5
Perception regarding Escitalopram use	EscitalopramP	Likert scale	1-5
Perception regarding Fluoxetine use	FluoxetineP	Likert scale	1-5
Perception regarding Paroxetine use	ParoxetineP	Likert scale	1-5
Perception regarding Sertraline use	SertralineP	Likert scale	1-5
Perception regarding Venlafaxine use	VenlafaxineP	Likert scale	1-5
\Any comments?	comment	Text	
	NATAL DATA A		
Pregnancy outcome	pregoutc	Live-born infant Spontaneous abortion (no live birth; gestational age less than 20 weeks)	2
		Stillbirth (no live birth; gestational age 20 weeks or greater)Ectopic pregnancy	3
		Termination Lost to follow-up	4
		·	5
			6
Was this a multiple birth?	multbir	Yes, twins Yes, triplets	1
		No	2
			3
Date of delivery or end of pregnancy?	pregend	Date (mm/dd/yy)	

Question	Variable name	Categories	Value
Cocaine present in	CocianeMR	Yes	1
Medical Record		No	0

Question	Variable	Categories	Value
Gestational age at end of pregnancy	gestageE	Continuous	
Type of delivery?	deltype	Vaginal – vertex Vaginal – breech Vaginal – transverse Cesarian section- primary Cesarian section – repeat	1 2 3 4 5
If cesarian section, reason for surgical delivery?	csectn	Emergency Failure to progress through labor Elective (pre-planned) Not applicable Breech presentation	1 2 3 4 5
Maternal		·	
complications: Preeclampsia or toxemia	pretox	Yes No	1
High blood pressure	НВР	Yes No	1
Oligohydramnios	oligohyd	Yes No	1 0
Infection or fever at delivery	infect	Yes No	1
Gestational diabetes	gediabts	Yes No	1
Other	otcomp	Yes No	1 0

If "Other", specify	otcomps	Text	
What was mother's	endwt	Continuous	
weight at the end of			
pregnancy?			
What was mother's	wtgain	Continuous	
weight gain during	331		
pregnancy?			
Is mother breastfeeding	brstfeed	Yes	1
the infant?		No	0
		N/A	2
Infant ID	infntID	Text	
Question	Variable	Categories	Value
	name		
Sex of a child	sex	Boy	1
		Girl	2
Birth weight	bweight	Continuous	
Birth length	blength	Continuous	
Birth head circumference	bheadcf	Continuous	
Apgar Score 1 minute	Apgar1	Continuous	
Apgar Score 5 minute	Apgar5	Continuous	
Neonatal			
complications:			
	respdist	Yes	1
Respiratory distress		No	0
	hypglycm	Yes	1
Hypoglycemia		No	0
	tachypn	Yes	1
Tachypnea		No	0
	la manalura al	Vaa	
Dradvoordia	bradycd	Yes	1
Bradycardia		No	0
	concic	Yes	1
Sepsis	sepsis	No	Ö
Sepsis		INO	
	otcompn	Yes	1
Other	Otoompii	No	Ö
If "Other", specify	otcompns	Text	
Major structural anomaly	stranom	Yes	1
diagnosed during the		No	Ö
hospital stay?			
How many days was	hospdays	Continuous	
infant in the hospital?			
Was the infant in	inflCU	Yes	1
Intensive Care Unit?		No	0

Did infant go home with	infhome	Yes	1
the mother?		No	0
If 'No', what is the reason?	infhomeN	Still in nursery	1
		Neonatal death	2
		Orphanage	3
		Other	4
If 'Other', specify reason	othomes	Text	
Any	abncon	Yes	1
abnormalities/conditions		No	0
diagnosed in neonatal			
period			
If Yes, specify	abncons	Text	
Notes	Notes	Text	

MAJOR DRUG CATEGORIES

Question	Variable	Categories	Value
	name		
Nutrients and Nutritional	nutrients	Yes	1
Agents		No	0
Hematological Agents	hematolog	Yes	1
		No	0
Endocrine and Metabolic	endocmetab	Yes	1
Agents		No	0
Cardiovasculars	cardiovasc	Yes	1
		No	0
Renal and Genitourinary	renalgenitour	Yes	1
Agents		No	0
Respiratory Agents	respiratory	Yes	1
		No	0
Central Nervous System	cns	Yes	1
Agents		No	0
Gastrointestinal Agents	gastro	Yes	1
		No	0
Anti-Infectives, Systemic	antiinfect	Yes	1
		No	0
Biological and	bioimmun	Yes	1
Immunological Agents		No	0
Dermatological Agents	dermatol	Yes	1
		No	0
Ophthalmic and Otic	ophthal	Yes	1
Agents		No	0
Antineoplastic Agents	antineoplast	Yes	1
		No	0
Herbalife(Herbal Life)	Herbalife	Yes	1
		No	0

Fish Oil(Omega-3l DHA;	Fish_Oil	Yes	1
Expecta; DHA Expecta;	_	No	0
Omega fats)			
Chamomile	Chamomile	Yes	1
(Manzanilla, German		No	0
chamomile, Roman			
Chamomile)			
Valerian	Valerian	Yes	1
(Valeriana)		No	0
Primrose Oil	Primrose_Oil	Yes	1
(Evening primrose oil;		No	0
Primrose pill; primrose oil			
capsule)			
Cranberry	Cranberry	Yes	1
(Cranberry extract,		No	0
cranberry tea)			
Raspberry	Raspberry	Yes	1
(Red Raspberry;		No	0
Raspberry Leaf)			
Flax Seed	Flax_Seed	Yes	1
		No	0
Other Herb drug	Other_Herb	Yes	1
		No	0
If Yes, specify the name of herb	Herb_specify	Text	
0			

APPENDIX B QUESTIONNAIRE OF THE SMART STUDY

GENERAL INFORMATION

1.	Date of interview:/_/_ (month/day/year)	
2.	Location of interview:	
3.	Prenatal care provider's last name:	
4.	Examiner's last name:	
5.	Patient's phone number:	
	DEMOGRAPHIC / LIFESTYLE INFORMATION	
6.	How old are you? (years)	
7.	What is your marital status now?	
	[] Single, never married	
	[] Married, living with spouse	
	[] Not married, but living with partner	
	[] Separated from spouse	
	[] Divorced	
	[] Widowed	
8.	Are you Hispanic, Latino or of Spanish descent? [] Yes	[] No
9.	How do you describe yourself: (check all that apply)	
	[] White, non-Hispanic or [] White, Hispanic	
	[] Black or African American	
	[] American Indian or Alaskan Native - Please specify [] [] pueblo	ribe or
	[] Asian or Asian American or Pacific Islander	

	Some other group(s) – please specify:	
	[] Prefer not to report	
10.	What is the highest level in school you have completed?	
	[] Less than high school graduate	
	[] High school graduate or GED	
	[] Some college or vocational school	
	[] College degree	
	[] Masters, doctorate or professional degree	
11.	What is your health insurance status?	
	[] No insurance	
	[] Employer-based insurance	
	[] Self-purchased insurance	
	[] Medicaid	
])	[] Other public insurances] Indian Health Service, [] VA, [] First Choice, [] UNN	M/UNMCARE)
	11a. Does your insurance cover prescription drugs? [] Yes	[] No
12.	Were you born in the Unites States? [] Yes	[] No
	If 'Yes', go to question 13. If 'No', please answer questions 12d	a and 12b.
	12a. Did you move to the United States: [] With your parents when you were a child [] When you were an adult (≥18 years old)	
	12b. How long have you lived in the United States:	years
13.	What language do you mostly use at home?	
	[] English	
	[] Spanish	
	[] Some other language – specify:	

[] Yes	[] No
14a. How many cigarettes do you usually smoke in one day?	
	[] Yes [] No
14c. When did y * Go to the next	ou stop smoking? question
[] Before I becam	ne pregnant
[] After I realized	that I was pregnant
15. Have you ever drank alcohol in your life (drinks)?	e.g., beer, wine, hard liquor, mixed] Yes [] No
If 'yes,' continue to questions 15a and 150	b. If 'no,' continue to 21a.
15a. How many drinks does it take before you	begin to feel the first effect of alcohol?
15b. How many drinks typically can you hold	before passing out or falling asleep?
a. What was the first day of your last n (mm/dd/yy)? I would like you to think back to that n	nenstrual period// eriod and tell me about your drinking at
that time.	erioa ana ieu me aboui your arinking ai
16. During a month or so around your last men how many times did you drink 4 or more drin	
Now I want you to think of <u>12 months before y</u> LMP)	ou got pregnant (a year prior to your
17. During the year before you got pregnant, decomplain about your drinking habits? [] Yes [] No	lid close friends or relatives worry or

14. Do you currently smoke cigarettes or use tobacco?

18. During the year before you got pregnant, did you ever take a drink first thing in the morning to get yourself going? [] Yes [] No
19. During the year before you got pregnant, did a friend or family member tell you about things you said or did while you were drinking that you could not remember? [] Yes [] No
20. During the year before you got pregnant, did you feel you need to cut down on your drinking?
[] Yes [] No
[TWEAK High:; TWEAK Hold:]
MEDICAL AND REPRODUCTIVE HEALTH
21a. What was your pre-pregnancy weight? pounds
21b. What was your pre-pregnancy height? feet/inches
[Researcher Calculated BMI:]
22. Do you have a medical condition or problem that requires ongoing, periodic, or occasional treatment? [] Yes [] No
22a. If yes, check all that apply:
[] Hypertension (high blood pressure) [] Depression [] Diabetes: [] Gestational [] Type I [] Type II [] Anxiety [] Seizure disorder (i.e., epilepsy) [] Migraine headaches [] Thyroid disorder [] Rheumatoid arthritis [] Asthma or allergies [] Heart disease [] Cancer [] Hepatitis [] Other(s) problem - specify:
If 'Yes' to diabetes, please answer questions 23 and 24. If 'No', skip to question 26. If 'Yes' to asthma, please answer question 25. If 'No,' skip to question 26.
23. Have you ever had gestational diabetes?
[] Yes, in a previous pregnancy only
[] Yes, in the current pregnancy only
[] Yes, in a previous pregnancy and in the current pregnancy

[] No	o, never had gestatio	nal diabetes		
[] No	o, never been pregna	ant before		
	y do you think uncoaby by causing birth			
1 Not at all	2 Unlikely likely to harm		2	5 Very likely to harm to cause harm
	y do you think asthi clinic visits could ha			
l Not at all	2 Unlikely likely to harm		4 Likely likely to harm	2 2
26. Did you p	lan to get pregnant	with this child?		
[] Yes	[] No, not not	w[] No, not at a	any time	
27. Were you this child?	or your partner doin	ng anything to try	to prevent become	ning pregnant with
[] Yes	[] No)		
27a. I	f Yes, which method	were you using?		
	[] Condoms	[] Diaphr	agm []	Birth control pills
	[] Withdrawal	[] IUD	[] R	hythm
	[] Depo Provera	, Implanon or No	rplant [] C	other:
•	ake any <u>fertility drug</u> odin, Fertinex, or Po [] No		pregnant with th	is child, like
28a It	f <i>Yes</i> , which drugs d	id vou use?		

29. Have you or members of your immediate family of your baby's fababies that might not have survived	ather had any babies	
	[] Yes	[] No
* If 'No', go to question 30	. If 'Yes', please spe	ecify:
Down syndrome	[] Yes	[] No
Cleft lip or palate	[] Yes	[] No
Neural tube defect	[] Yes	[] No
Cystic fibrosis	[] Yes	[] No
Heart defect	[] Yes	[] No
Other	[] Yes	[] No
If "Yes," ple	ease specify:	
30. What was the first day of your dd/yy)	last menstrual perio	od?//(mm/
31. What is the date your baby is $\frac{d}{dd} / yy$)	ue to be born:	//(mm/
31a. What is the gestationa	l age of your baby?	weeks
[] Ultı	t menstrual period	
32. How many times (including the	is pregnancy) have y	you been <u>pregnant</u> ?
If this is the 1^{st} pregnancy put "1" question 37.	for q. 32 and "0" fo	or questions 33-37 & skip to
33. How many live-born children is <i>If no live-born children or i</i>		nancy, then put "0"
34. Have you ever had a miscarriagmany?	ge (<20 wk of gestar	tion). If yes, how
If never had a miscarriage,	put "0"	

35. Have you ever had a stillborn child (\geq 20 wk of gestation). If yes, how many?
If never had a stillborn child, put "0"
36. Have you ever had a pregnancy terminated? If yes, how many? If never had a termination, put "0"
37. Have you ever had an ectopic pregnancy. If yes, how many? If never had an ectopic pregnancy, put "0"
38. For this pregnancy, how many weeks after your last menstrual period did you first think you were pregnant?
39. For this pregnancy, how many weeks after your last menstrual period did you first go to see a doctor or other health care provider or go to the clinic for prenatal care?
40a. Have you had any complications in this pregnancy so far? (* Please check yes or no for each complication)
- Bleeding [] Yes [] No - High blood pressure [] Yes [] No - Diabetes [] Yes [] No - Other [] Yes [] No
40b. If "other", please specify:
41. Have you experienced morning sickness during this pregnancy? [] Yes[] No
USE OF MEDICATIONS AND SUPPLEMENTS DURING PREGNANCY
42. Did you take a multivitamin regularly (4 times a week or more) during the month before your last menstrual period?
[] Yes [] No
43. Have you taken any VITAMINS regularly (4 times/week or more) since you became pregnant? 43a. [] Yes, multivitamins [] Yes, a single vitamin [] No
If 'Yes,' answer questions 43b-43e.
43b. [] Prescription [] OTC
43c. [] Brand name:
43d. When did you start taking vitamins? (mm/dd/yy) (gestational weeks)

43e. How many days during the last week did you take vitamins? (days/week)
44a. Have you taken any DIETARY SUPPLEMENTS (including iron supplements) or HERBAL PRODUCTS on a regular basis since your last menstrual period?
[] Yes [] No
44b. If 'Yes' to herbal products, please specify: [] Herbs
[] Tablets or capsules
[] Teas
[] Other:
44c. How often do you take them? [] Regularly: times per or [] When I feel sick
Please specify any other dietary supplements or products and reason for taking it:
Product 1: Reason/Condition:
Product 2:Reason/Condition:
Product 3:Reason/Condition:
44d. Have you had any cravings for non-food items or really "strange" foods? [] Yes [] No
If 'yes' what did you crave, do you eat it, and how often do you eat it?
Item 1: Eat it? [] Yes [] No; How often?
Item 2:Eat it? [] Yes [] No; How often?
Item 3:Eat it? [] Yes [] No; How often?
45. Have you ever taken any recreational drugs? [] Yes [] No
If 'Yes' please specify the recreational drug name(s) and when it was used:
Check if taken:
[] Marijuana/Hashish: [] Before pregnancy

		[] 1 month prior to LMP or during this pregna	anc	y
[]	Heroin:	[] Before pregnancy[] 1 month prior to LMP or during this pregnance	anc	y
		nrough methadone treatment?		
	[] Never [] Completed	reatment before pregnancy		
		treatment during current pregnancy		
[] Cocaine/Crack:	[] Before pregnancy[] 1 month prior to LMP or during this pregnance	anc	y
[]	Inhalants (glue, solve	ent): [] Before pregnancy [] 1 month prior to LMP or during this	pre	gnancy
[]	Methamphetamines:	[] Before pregnancy[] 1 month prior to LMP or during this	pre	egnancy
[]	Other:	[] Before pregnancy[] 1 month prior to LMP or during this	pre	egnancy
[]	Other:	[] Before pregnancy[] 1 month prior to LMP or during this	pre	egnancy
		afety of medications in pregnancy with any healt e-midwife, physician assistant, or pharmacist)?	h ca	are
		[] Yes	[] No
47a.	. Have you had any va	accinations since your last menstrual period?		
		[] Yes	[] No
	47b. If <i>Yes</i> to vaccina	ations, please specify: [] Flu		
		[] Other:		
heal	-	nedications PRESCRIBED by your doctor or a your last menstrual period, even if you stopped pregnant?	-	
		[] Yes	[] No

If 'Yes' please specify the medication name, reason for taking it, and <u>your perception</u> of how likely it is that this medication might be harmful for your baby if taken during pregnancy:

a. Medicatio	on 1:	Indication:				
	likely do you think by causing birth de ber)					
1	2	3	4	5		
Not at all		Somewhat	Likely	Very likely		
b. Medicati	on 2:		Indication:			
	likely is it that this ing birth defects or o					
1	2	3	4	5		
	Unlikely		Likely	Very likely		
c. Medicatio	on 3:		Indication:			
	likely is it that this ing birth defects or o		-			
1	2	3	4	5		
	Unlikely		Likely	Very likely		
d. Medicati	on 4:	Indication:				
	likely is it that this ing birth defects or o					
1	2	3	4	5		
Not at all	Unlikely	Somewhat	Likely	Very likely		
	likely to harm	to cause harm	likely to harm	to harm to cause		

e.	Medicatio	on 5:			Indication	n:	
		likely is it that this n			•		
1 2 Not at all Unlikely likely to harm			3 Somewhat to cause harm			5 Very likely to harm to cause harm	
		nis pregnancy, did yo IONS (sold withou [] Yes	t prescrip	tion)?	R-THE-CO	OUNT.	ER
ev m	ven if you st edications	nedications that you topped taking them of you took since pregnetion is to cause birth	once you kn ancy, pleas	ew you se spec	i were preg ify <u>your pe</u>	nant. '. rceptio	Then for on of how likely
	ain/Fever l ate all med	Medications:					
N	1 ot at all	2 Unlikely likely to harm	Somewh to cause		4 Likely likely to	•	5 Very likely to harm to cause harm
[] Acetam	inophen (Tylenol)	1	2	3	4	5
[] Aspirin		1	2	3	4	5
[] Ibuprofe	en (Advil, Motrin)	1	2	3	4	5
[] Ketopro	ofen (Orudis)	1	2	3	4	5
[] Naproxe	en (Aleve)	1	2	3	4	5
[-	nedication — fy:	1	2	3	4	5
	asal Decor	ngestants, Allergy, (Cough Med	licatio	ns:		
	1	2	3		4		5
Not at all Unlikely likely to harm		Somewhat to cause harm		Likely to	•	Very likely to harm to cause harm	

[] Chlorpheniramin	ne (Chlor-Tri	meton)	1	2	3	4	5
[] Benadryl			1	2	3	4	5
[] Pseudoephedrin	e (Sudafed)		1	2	3	4	5
[] Claritin, Zyrtec			1	2	3	4	5
Other medication specify:			1	2	3	4	5
Not at all Un	: 2		what se harm		cely	to harm	
Check if taken:							
[] Kaopectate, Pep	to Bismol	1	2	3	4	5	
[] Loperamide (Im	odium)	1	2	3	4	5	
Other medication specify:		1	2	3	4	5	
Heartburn, Dyspepsia, Antiemetic, Laxative Medications: Rate all medications:							

Ra	Rate all medications:							
	1	2	3		4		5	
No	ot at all	Unlikely	Somewha	at	Likely	7	Very likely	
		likely to harm	to cause l	narm	likely to l	narm	to harm to cause	į
							harm	
<u>Cł</u>	neck if take				_		_	
L] Maalox,	Mylanta Gas	1	2	3	4	5	
F	1.00		1	2	2	4	~	
L] Tums		1	2	3	4	5	
Г	1 Tagamat	t, Zantac, Axid, Pepci	d 1	2	3	4	5	
L] Tagaine	., Zamac, Axiu, Pepci	u i	2	3	4	3	
Г] Colace		1	2	3	4	5	
L] coluce		•	_	J	•		
Γ] Correcto	l, Dulcolax, Ex-Lax	1	2	3	4	5	
-	-							
[] Senna, f	iber products	1	2	3	4	5	
[] Unisom		1	2	3	4	5	

specif	.y:					
Antifungal M	Iedications (taken f	or vaginal	yeast in	nfection or	thrash	n):
Rate all medi	cations:					
1 Not at all	2 Unlikely likely to harm				-	-
Check if take	<u>n</u> :					iiwiiii
	cream or suppositori stat, Vagistat, Fems		2	3	4	5
	edication – y:			3	4	5
Nicotine Rep	lacement Therapy	(for smok	ing cess	sation):		
Rate all medi	cations:	2		4		_
-	Unlikely likely to harm	Somewh to cause	at harm	4 Likely likely to	y harm	5 Very likely to harm to cause harm
Check if take	<u>n</u> :					
[] Nicotin	ne gum, spray or inh	aler 1	2	3	4	5
[] Nicotir	ne patch	1	2	3	4	5
[] Other 1	medication –	1	2	3	4	5
specif	ÿ:					
Other over-t	he-counter medicat	tions you l	nave ta	ken while	pregn	ant:
Rate all medi		2				_
l Not at all	2 Unlikely	3 Somewh	at	4 Likel	y	5 Very likely
	likely to harm				-	to harm to cause harm

[] Other medication –

Check if taken:						
Other medication – specify:		2	3	4	5	
Other medication – specify:		2	3	4	5	
Other medication – specify:		2	3	4	5	
50. If you took prescription change the use of these me						did you
[] Did not take p	rescriptio	on medic	ations reg	gularly be	efore pregnan	cy
[] Discontinued to Medication		pon reco	gnition of	f pregnar	ncy.	
[] Decreased the	use (dos	e or frequ	uency). M	1edicatio	n:	
[] Increased the t	use. Med	ication: _				
[] Stayed the san						
50a. If you changed th			-	_		
Provider recomme			vien uper		61 p1 -8	~11 0 j ,
[] Family or friend s	suggestio	on				
[] Self-initiated						
[] Financial constra	ints					
[] Other:						
Now I'm going to ask you pregnancy in general. Pleaseach question.	•	0			•	-
51. If a woman plans a pre should:	gnancy o	or finds o	ut that sh	e is curre	ently pregnan	t, she
[] Stop taking all	medicat	ions imn	nediately	to protec	et the baby	
[] Continue takir	ng only th	nose med	ications t	hat are a	bsolutely nec	essary and

	check with her doctor to see if the medications are safe for the baby						
[Continue taking necessary medications but reduce the dose or the number of days you take them to limit the amount that gets to the baby						
[] Continue with all medications as needed since medications are safe for the baby						
	a woman uses medications regularly during pregnancy, how often can ons cause birth defects?						
[] Never	[] [] [] Sometimes Often Very Often Always						
53. Which	h statement best describes your view about women drinking alcohol during y?						
[] Pregnant women should abstain from drinking any alcohol (even small amounts) during pregnancy.						
[] It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than once a week.						
[] It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than one drink per day.						
[] It is OK for a woman to have an occasional drink during pregnancy as long as it is not more often than two drinks per day.						
[[] It is OK for a woman to drink during pregnancy as long as she does not drink hard liquor (i.e., vodka, whiskey, brandy) but only drinks wine or beer.						
medicatio	ng your current pregnancy, have you ever asked anyone about the safety of ons you are taking for your baby?] Yes [] No						
	If yes, check any individuals who you have asked a question about the safety y medications for your baby: (Check all that apply to you)						
[] Your primary care doctor or provider						
[] Your OB/GYN doctor or midwife						
[] A pharmacist						
[] A member of your family, spouse						
[] A friend, partner						

[] Other - specify:
[] Any other heath care provider
	e check any sources below in which you have looked for information about of medications for your baby? (<i>Check all that apply to you</i>)
_] I have never looked at any of these sources about the safety of edications for my baby
[Specify:_] An internet web site(s).
[] A book.
[] A magazine
[] Pregnancy information telephone service/hotline (i.e., OTIS, Nurse Advisory Line)
[] Other - specify:
[] I have not had any questions about the safety of medications for my baby and have not looked at any of these sources.
[] Clinic pamphlet or brochure
56. NOTI	ES/COMMENTS: