Evaluation of the Positive Prevention PLUS Teen Pregnancy Prevention Program

Final Impact Report for San Bernardino County Superintendent of Schools

Prepared by

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This publication was made possible by Grant Number TP2AH000007 from the Office of Adolescent Health, U.S. Department of Health & Human Services. The views expressed in this report are those of the author and do not necessarily represent the policies of HHS or the Office of Adolescent Health. Findings from an Innovative Teen Pregnancy Prevention Program

REVISED 2/22/16

Acknowledgements

This study was a collaborative effort between the San Bernardino County Superintendent of Schools (SBCSS) and California State University, San Bernardino (CSUSB). Dr. Kim R. Clark served as the Project Director. The author is grateful to Jessica Folmer, Research Associate, for her assistance in data collection, data entry, and assessing implementation fidelity, Carolyn Tillman at SBCSS for her assistance in project implementation, and to the students, teachers, and district staff who participated in the study. Finally, the author would like to thank Jaqueline Berman, PhD and Russell Cole, PhD at Mathematica Policy Research for their guidance, on-going support, technical assistance, and collegiality throughout the study.

Recommended Citation:

LaChausse, R. (2015). Evaluation of the Positive Prevention PLUS Teen Pregnancy Prevention Program. USDHHS, Office of Adolescent Health. TP2AH000007.

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I. Introduction

A. Introduction & Study Overview

The teen birth rate in the U.S. continues to decline and has dropped below 26.5 births for every 1,000 adolescent females ages 15-19.¹ While this reflects overall progress at achieving lower rates of teen pregnancy, progress in uneven. For example, Hispanic teens continue to have the highest birth rate at 42.7 per 1,000.² National data reveals that over 48 percent of all students in grades 9-12 have had sexual intercourse by the time they graduate and only 41 percent of teens had used a condom the last time they had sexual intercourse with Hispanic teens reporting slightly lower rates of condom use.³ Since a significant number of adolescents engage in sexual risk behaviors, the need for effective teen pregnancy programs cannot be overstated.

Although many teen pregnancy prevention programs can increase student knowledge about the consequences of becoming pregnant, only a few have demonstrated an impact on students' behaviors. In 2009, the U.S. Department of Health and Human Services (HHS) sponsored a systematic review examining the effectiveness of programs to reduce teen pregnancies and associated sexual risk behavior among teens. Of the approximately 2,000 potentially relevant studies published between 1989 and January 2011, 200 met screening criteria. Of these, 88 studies received a high or moderate rating and only 31 provided credible evidence demonstrating a statistically significant positive program impact on at least one sexual behavior or reproductive health outcome of interest (for example, sexual activity, contraceptive use, or pregnancy).⁴ The review concluded that there is a need for improved research quality and reporting to inform policy initiatives and programming decisions.

As a result, the Office of Adolescent Health (OAH) under HHS introduced a grant program to support the replication of evidence-based programs and to study the impact of new program models and strategies. With funding from OAH, the Positive Prevention PLUS, an 11-lesson schoolbased teen pregnancy prevention (TPP) program, was developed based on the existing literature surrounding school-based prevention programs that has supported programs that use experiential, interactive activities to emphasize abstinence and risk reduction techniques.⁵ Positive Prevention PLUS applied the findings from an earlier study of the Positive Prevention HIV/STD Curriculum for Students Grades 9-12 that employed clustered randomized controlled trial and met the criteria for a high rating under the HSS TPP evidence review. Findings from the study showed a positive, statistically significant impact on sexual initiation at the 6 month follow-up.⁶ However, the HHS TPP evidence review concluded that the impact estimates could have been biased because the analyses controlled for post-implementation variables that could have been affected by the program, so there was no evidence of effectiveness.⁷

Positive Prevention PLUS utilizes Social Cognitive Theory (SCT) as its theoretical basis. SCT posits that the likelihood of taking preventive action is determined by an understanding of what must be done to avoid pregnancy, a belief that one is able to use this method, and the belief that this method will successfully decrease the chance of getting pregnant.⁸ Past research has suggested that the use of social learning theory in pregnancy and STD prevention, particularly with adolescent youth, is far superior to other theoretical approaches.⁹

This report provides the results of an external evaluation of the Positive Prevention PLUS teen pregnancy prevention program for 9th grade students in Southern California. A clustered randomized controlled trial (CRCT) was employed in which participating high school sites were randomly assigned either to a treatment group that implemented the Positive Prevention PLUS program or a control group. Students completed a self-administered survey at baseline (prior to program implementation) and a 6 month follow-up survey (post program implementation). The

purpose of this study is to provide scientific evidence concerning the effectiveness of a schoolbased, theory-driven teen pregnancy prevention program.

B. Primary Research Questions

- 1. What is the impact of Positive Prevention PLUS relative to a control group on the initiation of sexual activity 6 months after the end of the intervention?
- 2. What is the impact of Positive Prevention PLUS relative to a control group on ever been pregnant 6 months after the end of the intervention?
- 3. What is the impact of Positive Prevention PLUS relative to a control group on having sexual intercourse <u>without</u> using birth control in the prior three months 6 months after the end of the intervention?

It was hypothesized that students receiving Positive Prevention PLUS would be less likely to initiate sexual intercourse six months after the end of the intervention than those in the control group. Further, it was expected that the program would decrease the likelihood that students would become pregnant (or get someone pregnant), and decrease the likelihood that adolescents would not use birth control.

II. Program and Comparison Programming

A. Description of Program as Intended

Positive Prevention PLUS was developed after a review of the literature of effective sexuality education programs that found that programs based on SCT can increase an adolescent's ability to use risk –reduction skills (e.g. assertive communication), and either abstain from sexual intercourse or use birth control when engaging in sexual activity. SCT posits that behavior change occurs through several constructs including observational learning, behavioral capability, and self-efficacy.^{6,9} SCT is used throughout the Positive Prevention

PLUS lessons. For example, lessons 1 and 2 explore reasons for teens to be sexually abstinent and make responsible decisions (observational learning), lesson 7 teaches students how to recognize and avoid risky situations and use condoms (behavioral capability), and lesson 8 uses interactive role-plays to increase confidence in using refusal skills in everyday life (selfefficacy). For a program description and logic model see Appendix A, Figure A.1.

Positive Prevention PLUS consists of eleven 45-minute lessons aimed at students in grade 9 delivered during the normal school day in science, health, or physical education courses. The program was implemented in public schools in Southern California. The lessons were taught by trained classroom teachers to students in group form during a regular class period. Teachers were given a 3 week period in the fall of 2013 to complete the 11 lessons in consecutive order. The curriculum lessons are as follows:

- *Getting Started*: Discussion on group agreements, the sexual health of teens, overview of the sexual health concerns of young men and women, and a review of ways to maintain reproductive health.
- *Lesson 1- Life Planning:* Review of the importance of personal goals and life plans.
- *Lesson 2- Healthy Relationships:* A discussion on intimacy and the characteristics of a quality relationship.
- *Lesson 3- Relationship Violence*: Signs of an abusive relationship, showing the relationship violence video, and the setting boundaries.
- Lesson 4- Family Planning and Contraceptives: Review of contraception, displaying and describing FDA-approved contraceptives, including their effectiveness, the California Safe Surrender law, and the benefits of abstinence.

- *Lesson 5- Myths and Stereotypes:* Video of persons infected by HIV and an activity on how HIV affects various people without regard to sexual orientation or family structure.
- *Lesson 6- HIV and AIDS:* A review of the HIV/AIDS epidemic, how a healthy immune system functions, and information on managing risk situations.
- *Lesson 7- Recognizing and Reducing Risk:* Steps in condom use, a condom demonstration, condom success and failure rates, and practicing negotiating condom use.
- *Lesson 8- Peer and Media Pressures*: Peer and media pressures, media analysis skills, and communication (assertiveness) and refusal skills practice.
- Lesson 9- HIV/STD Testing and Community Resources: An activity which focuses on demonstrating STD transmission and community resources.
- *Lesson 10- Steps to Success*: Review of personal goals and life plan and making a commitment to either abstaining from sexual intercourse or using birth control.

Teachers in participating public schools implemented the program. Each teacher must have at least a bachelor's degree and a single subject teaching credential in life science, physical science, physical education, or health education. Teachers were provided training about one month prior to program implementation from the program developer and his staff. Teachers participated in an initial two-day training consisting of an overview of teen pregnancy in the U.S. and local area, characteristics of effective teen pregnancy prevention programs (theory, pedagogy, fidelity, etc.), the purpose of the PP+, the program's logic model, and the need to implement the program with fidelity including the specified sequence. Program trainers demonstrated each of the eleven lessons. On day two, teachers were asked to teach back one of the eleven lessons (randomly selected) and received feedback from the trainers. After the initial training, teachers were asked to complete an online training program consisting of various modules to review the key components of lessons and to observe each lesson being taught by a veteran teacher. Approximately three weeks after the initial training (1week before implementation), teachers were brought back as a group for a one day refresher to emphasize key lessons, lesson activities, program quality and adherence, as well as to address any concerns teachers may have regarding program implementation. During program implementation, the project director periodically communicated with teachers to provide encouragement and technical assistance.

B. Description of Counterfactual Condition

Students in control group classrooms received the standard health, science or physical education curriculum. Control groups schools and teachers were asked to refrain from providing any sexuality- related classroom instruction or school wide teen pregnancy or STD prevention focused activities, however control group teachers were allowed to discuss human reproduction if relevant to their curriculum (for example, in a biology course).

III. Study Design

A. Sample Recruitment

The Project Director approached 6 school districts to assess each district's ability to participate in the study and to obtain a memorandum of understanding (MOU). School districts were approached based on previous data that indicated that there were a high number of teen births and/or STD rates and the lack of an existing evidence-based comprehensive sexuality education or teen pregnancy prevention program in the school district. For each school district, the feasibility of participating in a randomized controlled trial was assessed including the district's ability to implement a teen pregnancy prevention program like PP+, understand and maintain the treatment/control contrast, provide access to students for data collection, and ability to provide attendance data. Once MOUs were finalized, teacher training was provided and parent/guardian consent was requested. All six school districts agreed to participate in the study.

Twenty-two high schools in Southern California were approached to participate in this study. Eligibility to participate included interest in the program, a signed MOU, and having required science, health, or physical education courses for 9th grade students. Ultimately, twenty-two public high schools (with thirty-six teachers) across six school districts within Southern California met initial eligibility requirements for participation. All male and female 9th grade students in mandatory 9th grade health, science, or physical education (PE) classes at each school site (n= 22) were eligible for inclusion in the study. Prior to randomization, parental consent forms were distributed to teachers by the Principal Investigator (PI). Teachers distributed the consent forms to students. The consent form described the general purpose and nature of the study, issues of confidentiality of responses, and contact information for the PI. Additionally, the consent form included a statement that the survey questions are general in nature, including

questions on sexual behavior, but this does not imply that their child is sexually active. The consent form was provided in both English and Spanish. A small incentive (two movie tickets) was offered to the teacher with the highest percentage of parent consent forms returned at each school site regardless of whether parents agreed to have their child participate. The study was approved by the Institutional Review Board (IRB) at California State University, San Bernardino.

B. Study Design

A clustered randomized control trial (CRCT) was employed to determine program impacts. No stratification was employed. After parent/guardian consent was obtained, twenty one eligible school sites were randomly assigned into either the treatment or control condition using the select cases (RANDOM) function in SPSS 22.¹ School sites, students and their parent/guardian were not aware of their respective condition prior to consent. Treatment sites (n = 11) agreed to implement the Positive Prevention PLUS program in their 9th grade health (n = 7), PE (n = 1), or science (n = 4) classes. Ten control sites agreed to not provide any pregnancy prevention or STD prevention education within the study period.

C. Data Collection

1. Impact Evaluation

Baseline data was collected for both the treatment and control group in October 2013. Six month follow-up data collection occurred in May 2014 (See Table B.1). Data collection occurred for each school site within each district on the same day. Participants completed a self-administered, paper and pencil survey during their regular class period. All survey data

¹ Twenty two schools were part of the sample for random assignment, but one school was determined not to have any eligible 9th grade youth in health classes, and thus, was excluded from the study as ineligible.

collection was conducted by the Principal Investigator (PI). Project staff, teachers, site administrators, and other school staff were not allowed to provide any instructions, provide guidance, or answer student questions during data collection. Baseline data collection occurred approximately 8 weeks after parental consent forms were distributed and 1 week prior to the beginning of the intervention. For the 6 month follow-up, participating students were pulled into a central location (e.g. library) by their original class period at baseline. Data was collected on the same date for all study schools (both treatment and control) in each district. Each baseline survey was pre-printed with a random ID number to track individuals' survey responses across data collection periods. The same ID number was used for each follow-up survey to student responses over time. Data collectors were trained to respond to student questions with "Try your best to answer the question or leave it blank." Data collectors were not allowed to read aloud any survey items or response formats, or provide any other guidance. The data collection procedure was the same for both the treatment and control groups.

The student survey included a brief demographics section (gender, age, ethnicity) and outcome measures: 1) whether or not participants have ever had sexual intercourse, 2) whether or not participants have been pregnant (or gotten someone pregnant), and 3) whether or not participants had sex without using birth control in the prior 3 months (See Table C.1). The average time to complete the survey was 14 minutes and the reading level was 5.7 (Flesch-Kincaid grade level equivalent). Participants received one raffle entry for a chance to win two (2) movie tickets each time she/he completed a survey.

2. Implementation Evaluation

Adherence

Adherence to the program model was measured using the teacher curriculum log. This log was developed by the program evaluator based on other fidelity logs used in previous impact

evaluation studies.¹⁰ Program teachers completed this log after each class period and logs were collected by the PI after program implementation was completed. Intervention group teachers were asked to self-report the extent to which each topic, worksheet, or activity was covered completely, partially or not at all. Teachers were asked if any adaptations were made to any of the lesson activities.

Program reach was measured using school attendance data. This data was collected using each school district's student information system (SIS). The evaluator requested and received this data from each school district for the program implementation dates. The study considers a student "attended" if the teacher marked the student as "attended" or "tardy" for that class period per the district's SIS. If the student had an absence for that class period (for any reason), the student was marked as "not attended" for the study.

Quality of Implementation

The PI or a research assistant conducted a direct observation of each teacher using the classroom lesson observation form to measure lesson quality and student engagement. This observation assessed, the overall quality of the delivery of lesson activities including the teacher's explanation of lesson activities, teacher's ability to pace the lesson (i.e. keep on task), knowledge of the lesson, poise and confidence, level of enthusiasm, and student participation. One lesson observation was conducted during one class period for each teacher during the implementation period. The observation was conducted at random so the teacher would not know the day, period, or lesson the observation would be conducted. Each observer was trained in the curriculum during the teacher trainings and received additional training in conducting observational research.

Experiences of the counterfactual

Data regarding the difference between treatment and control experiences was collected on the student survey for both the treatment and control group participants at 6 month follow-up using the item "Which of the following health topics have you learned about this school year? pregnancy prevention, AIDS or HIV infection, human sexuality, taking care of a baby, using condoms, or abstinence." Respondents were asked to select each of the topics that they learned about (See Table E.1).

Context

Information regarding the context of program implementation was gained from surveys from teachers. Teachers in both the treatment and control group were asked "Which of the following health topics have you taught about in the last 6 months?- pregnancy prevention, AIDS or HIV infection, human sexuality, taking care of a baby, using condoms, or abstinence." on the teacher survey. Additionally, teachers were asked an open-ended question about whether there were any issues that occurred at the school site that may have affected program implementation.

D. Outcomes for Impact Analyses

The student survey assessed three outcome measures: 1) ever had sexual intercourse, 2) ever been pregnant (or gotten someone pregnant), and 3) ever had sexual intercourse <u>without</u> using birth control. Skip patterns are used so that if a participant reported that they had never had sex, she/he would skip the items that pertain to sexually active respondents and complete the remaining items in the survey. Sexual initiation was constructed from the survey question "Have you ever had sexual intercourse?" A dummy variable was created in which respondents who respond yes were coded as 1 and no were coded as 0. Missing data was coded as 999. The item "To the best of your knowledge, have you been pregnant or gotten somebody pregnant?" The

variable is constructed as a dummy variable with values where respondents who respond yes have been pregnant were coded as 1 and those who responded no are coded as 0. Missing data resulting from the skip pattern of the survey was logically imputed to 0 since one can infer that someone who had never had sex has never been pregnant. The variable "In the past 3 months, have you had sexual intercourse without you or your partner using any of these methods of birth control?"- Condoms, birth control pills, the patch, the ring (NuvaRing), IUD, Implant (Impanon). The variable was constructed as a dummy variable where respondents who respond yes have had sex in last 3 months without birth control are coded as 1 and those who responded no are coded as 0. Missing data due to the skip pattern was coded as 0 since one can infer that someone who had never had sex had also never had sex without birth control (See Table C.1).

E. Study Sample

Treatment sites (n = 11) agreed to implement the Positive Prevention PLUS program in their 9th grade health (n = 7), P.E. (n = 1), or science (n = 3) classes. Control sites (n = 10) agreed to not provide any teen pregnancy prevention education within the investigation period. A total of 7,042 students were eligible to participate in the study. Of the 4,969 students who returned the parent consent form, 4,267 had positive parental consent to participate. Of those students, 3,554 students participated in the baseline survey and 3,490 students participated in the 6th month follow-up survey. There were 2,113 participants in the intervention group and 1,377 in the control group that made up the final analytical sample (i.e. provided both baseline and follow-up data). See Table D.1.

F. Baseline Equivalence

The analytical sample is all students in the school sites that have parental consent and provide data at baseline and 6 month follow-up. Data is pooled across the school sites. Baseline

equivalence for the analytical sample is provided in Table G.1. This table displays the equivalence between the treatment and control groups on the following measures: age (in years), percent female, race, and percent Hispanic. Baseline equivalence was also examined on pretest data of the outcome measures including ever had sexual intercourse, ever been pregnant, and having had sex without using birth control in the prior 3 months. Equivalence was calculated using a linear regression model predicting the variable of interest from a grouping variable (dummy coded) for each measure adjusting for the clustering effect in the CRCT using Huber-White adjusted standard errors. Once adjusting for the clustered nature of the data, no statistically significant differences at baseline were observed.

G. Methods

1. Impact Evaluation

Analyses were conducted on student-level data and used an intent-to-treat framework. To answer the primary research questions and estimate possible program impacts, hierarchical linear regression models using a random effects approach were used. A two-level model was expressed in a multilevel framework using a random effects ANOVA to determine what portion of the variance in the outcome variable of interest is due to site level differences (i.e. school sites) as compared to individual differences. This approach accounted for clustering using maximum likelihood to estimate parameters that specified the structure of the covariance between individuals in clusters.¹¹ The model was written as:

$$\beta 1_{1ij} = \beta 0_j + e_{ij}$$
$$\beta 0_j = y_{00} + \delta_{0j}$$

where $\beta 1_1$ is the outcome of interest (e.g. ever had sex at follow-up) for an individual in a site, $\beta 0_i$ was the mean score for a site and y_{00} was the grand mean (i.e. the mean across all individuals and sites). The level 1 error term (e_{ij}) indicates how an individual's score deviates from the mean in the site in which the individual resides. The level 2 error term $(\delta 0_j)$ indicated how the mean score in a particular site deviates from the grand mean. This was calculated using the XTMIXED function in Stata 8.0 by first estimating the percent variance due to differences across sites by examining the interclass correlation (ICC) and the amount of variance attributable to individual differences. Using the XTREG function in Stata 8.0, a random effects model was calculated using the Group variable to determine the amount of variance that is explained by the Group variable (treatment vs. control). Finally, the XTREG command was used to predict the outcome variable of interest from Group with the baseline score, gender, age, and ethnicity (Hispanic vs. non-Hispanic) used as covariates. Findings are considered statistically significant if the p-value is less than 0.05, using a two-sided test. Because of the number of a priori hypotheses (3), the Benjamini-Hochberg adjustment was made to control for the false discovery rate.¹²

2. Implementation Evaluation

There were four sources of data used to assess adherence, quality of implementation, counterfactual experiences, and context: teacher curriculum logs, classroom lesson observation forms, teacher surveys, and participant attendance data. This data is limited as it relies on the self-report of teachers.

Adherence

Adherence to the program components was calculated by summing the number of completed lesson activities divided by the number of activities assigned. Lesson completion is defined as completing all lesson components (e.g. activities). The number of scheduled activities was calculated by multiplying the number of activities within each lesson by the number of class sections. The number of completed activities were identified from the self-reported teacher

curriculum logs and summed for each lesson. An adaptation is defined as any changes to the lesson activity or pedagogical strategy. The percent of activities completed was calculated by dividing the number of completed activities by the number of scheduled activities. The number of lessons taught with adaptations were identified from the self-reported teacher curriculum logs and summed for each lesson. The percent of activities completed with an adaptations was calculated by dividing the number of lessons taught with at least one adaptation to the total number of class sections.

Attendance data was also collected using each school district's student information system (SIS). The study considers a student "attended" if the teacher marked the student as "attended" or "tardy" for that class period per the district's SIS. If the student had an absence for that class period (for any reason), the student was marked as "not attended" for the purposes of this study.

Quality

Lesson quality was calculated by dividing the number of high quality lesson observations by the number of total observations. An average score of four or greater on a scale ranging from 1= poor to 5=excellent across individual session elements was considered a high quality observation. The quality of student participation was captured through classroom observations (e.g., "How exactly did the group members participate in the discussions and activities?" 1= Little participation, 3= Some participation, 5= Active participation). High student engagement was measured by summing the participation observation scores of 4 or better and dividing it by the total number of observations.

Experiences of the Counterfactual

To measure the counterfactual, students were asked which TPP health related topics they had learned about in the past 6 months including pregnancy prevention, AIDS or HIV infection,

taking care of a baby, using condoms, human sexuality, and abstinence. This was measured on the 6 month follow-up survey. Likewise, a teacher survey was conducted at the 6 month followup assessing the extent to which teachers had taught various health related topics outside of the PP+ program in the past 6 months and the presence of any school-wide teen pregnancy, STD, or HIV/AIDS prevention activities. The percent of TPP health topics received by students in each group and from the teacher's self-report was calculated by summing the responses from each item (1 = yes, 0 = no) and dividing them by the total number of possible responses. Table E.1 outlines the methods used to collect information about implementation fidelity.

Context

Teachers were asked if their school site had conducted any school-wide activities like assemblies, club events, or guest speakers regarding pregnancy prevention, sexuality, HIV/AIDS, or reproductive health in the past 6 months and asked to explain the type and duration of that activity. The percent of TPP health topics taught by teachers in each group (not in the PP+ program) was calculated by summing the responses from each teacher (1 = yes, 0 = no) and dividing them by the total number of possible responses. This information was captured through the curriculum fidelity log and informal conversations with treatment group teachers.

IV. Study Findings

A. Implementation Study Findings

Adherence

Eleven school sites within six school districts were assigned to the treatment group. Thirty-four teachers with a total of 123 class sections (periods) implemented the program. Teachers had 18 days between the dates of October 28th, 2013 to November 14th, 2013 to implement the 11 lessons in consecutive order. Ninety-five percent of the scheduled 1,353 lessons were completed. A total of 6,124 lesson activities were completed of the 6,396 lesson activities scheduled to be delivered (95%). Some lessons were completed with adaptions. For example, 36% of the completed lesson 4 (Family Planning and Contraceptives) had adaptions. Table F.1 shows the findings from the implementation fidelity data.

The number of students who participated in the program was 2,139. Overall, average program attendance across all sessions was 91 percent. The average number of lessons attended by each student was 10.01 of 11 lessons (SD = 1.64).

Lesson Quality

Twenty-six classroom lesson observations were conducted. An average score of four or greater was considered a high quality observation. Seventy-three percent (73%) of the observed lessons received a high quality observation. Most teachers received high marks for her/his explanation of lesson activities while the lowest scores were associated with teacher's ability to pace the lesson (i.e. keep on task) or low levels of student participation in the lesson. Only 65% of the observed lessons were rated with high student participation. Observers noted that in most cases, teachers' inability to correctly pace the lesson led to adaptations in a way that made the lesson activity less interactive. For example, a teacher would model refusal skills in front of the class rather than give students an opportunity to practice refusal skills.

Counterfactual experiences

The treatment group students reported learning 60 percent of the TPP health topics, whereas the control group reported learning approximately 17 percent of the TPP health topics. Additionally, teachers in the treatment group and control group were surveyed at the 6 month follow-up and teachers in the treatment group reported teaching 58 percent of the TPP health topics, whereas control teachers taught 10 percent of the TPP health topics. None of the teachers in either group reported any additional school-wide activities related to pregnancy prevention, sexuality, HIV/AIDS or reproductive health.

Context

Six treatment teachers within one school were not allowed to perform condom demonstrations per their district's school board policy. Teachers could explain but not demonstrate the steps in condom use. No other issues were documented.

B. Impact Study Findings

The results of the impact of the Positive Prevention PLUS program on participants' likelihood to engage in sexual intercourse, become pregnant, or have sex without birth control are presented in Table G.2. There was a significant effect of the Positive Prevention PLUS program on delaying sexual activity. Relative to the control group, participants in the treatment group were approximately 4 percentage points less likely to have had sex at 6 month follow-up (b = -.04, t = -2.38, p = .01). There was no impact of the Positive Prevention PLUS program on getting pregnant at 6 month follow-up (b = -.01, t = -1.87, p = .07). There was a significant effect of the program on ever having sex without birth control in the prior 3 months measured at the 6 month follow-up (b = -.02, t = -2.61, p = .01). Relative to the control group, participants in the treatment group were approximately 2 percentage points less likely to have had sex without birth control at 6 month follow-up. Table G.3 shows the results of the analyses using the Benjamini-Hochberg adjustment.

To explore the sensitivity of results from the benchmark sample, two different sensitivity analyses were conducted. First, because data on one or more of the outcome variables of interest were missing for some participants, this could lead to biased impact estimates including causality bias. To address missing data from partially completed surveys, multiple imputation (MI) was used. Five multiple imputed datasets were created and the regression models used in the impact

analysis were computed on the each of the 5 imputed data sets and on the pooled estimates. Variables used in the imputation included the same study outcomes of interest and covariates (gender, age, and Hispanic (binary)) used in the benchmark analysis. Imputations were conducted separately for treatment and control groups. The impact regression model used in the benchmark analysis was tested in the MI sample providing pooled estimates of the coefficients from the 5 imputed data sets.

The second sensitivity analysis used logical imputation. That is, imputing a missing score based on how each respondent should have answered. For example, if a participant had reported having sex in lifetime at pretest but reported had not having sex at posttest, their score on posttest was changed to having had sex in lifetime (i.e. carried through to follow-up). The impact regression model used in the benchmark analysis was tested in the logical imputation sample. Further explanation of the logical imputation approach is presented in Table H.1.

Table H.2 shows the results of the sensitivity analyses compared to the benchmark analysis. When impact estimation used the logical imputation sample, estimated impacts were similar to those in the benchmark analysis with the impact of the program on delaying sexual activity remaining robust. When compared to the impact estimation using the multiple imputation (MI) sample, impact of the program on delaying sexual activity only approaches statistical significance. This may be due to the way the impact estimates are calculated on the MI sample. That is, the impact regression model tested in the MI sample uses pooled estimates of the coefficients from the 5 imputed data sets. The standard error tends to be much larger than those in each individual imputed dataset because there are two sources of variation in the calculation: between iterations and within each iteration. It should be noted that the impact regression models

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computed on each of the 5 imputed datasets (not presented) in the MI sample individually yielded similar results to one another as well as to the benchmark analysis (See Table G.2).

V. Conclusion

This report focused on the impact of the Positive Prevention PLUS (PP+) teen pregnancy prevention program on delaying sexual intercourse, pregnancy, and using birth control among those students who were offered the program. This study indicates that the program has a small but statistically significant impact on delaying sexual intercourse and birth control use at the sixmonth follow-up. However, the program did not affect whether or not participants would become pregnant (or get someone pregnant). These findings are consistent with a previous study on an earlier version of Positive Prevention.⁶

The use of a clustered randomized control trial with an analytic sample equivalent at baseline provides rigorous evidence concerning the effectiveness of the program. While the results of this are promising, several methodological limitations should be noted. First, data were collected using self-reported surveys. Although it is impossible to be completely confident of the validity of self-report responses, there is some evidence that supports the general validity of adolescents' self-report of health behaviors.¹⁰ Second, the survey items asking student about which TPP-related topics they had learned about in the past 6 months is only reflective of student experiences after the intervention occurred. As a result, this may not reflect the experience of the intervention even though the intervention is relatively short. Third, variation in program implementation may have affected the strength of the treatment received. Prevention programs are seldom implemented perfectly and several studies have revealed the extent to which program fidelity occurs and how it affects program outcomes.¹²

Since a significant number of a youth engage in sexual risk behaviors, the need for effective school-based teen pregnancy curriculums cannot be overstated. Although many sexuality education programs can increase student knowledge, only a few have demonstrated an impact on students' behaviors. Findings from this study suggest that the Positive Prevention PLUS program is effective in the short term on reducing sexual initiation and unprotected sex. These results have implications for both health educators and researchers. Future studies should investigate the mechanisms by which teen pregnancy prevention programs affect behavior. This could include an examination of the possible relationships between program activities, determinates of behaviors (e.g. self-efficacy, attitudes, behavioral capability) and behavioral outcomes (e.g. birth control use). Additionally, the literature would be enhanced by exploring the long term impacts of the Positive Prevention PLUS program on adolescent sexual risk-taking behaviors and teen pregnancy.

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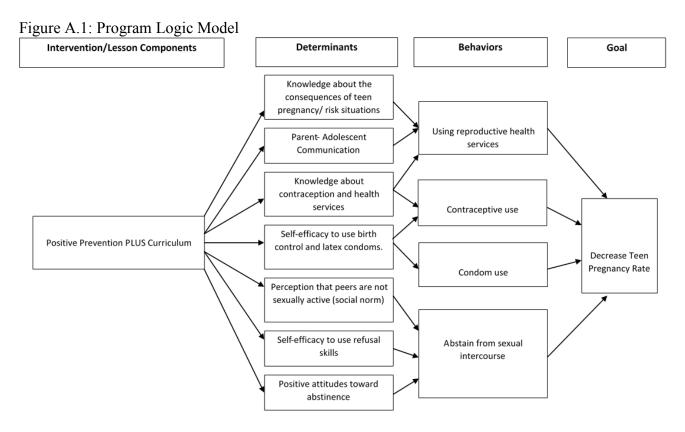
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Appendix A. Program Description and Logic Model

Table A.1: Application of Social Cognitive Theory (SCT) Constructs in the Positive Prevention	
PLUS Program	

SCT Construct	Definition	Applications in <i>Positive Prevention PLUS</i> (including lesson #)
Environment		Seeing data on teen sexual abstinence (Getting Started) Exploring myths and stereotypes re: HIV-infected
		persons (4)
	Factors physically external to	Identifying teen-friendly reproductive health services in the community (2,6,9)
	the person (e.g. social norm).	Analyzing media pressures (8)
Situation	Perception of the environment; correct misperceptions and promote healthful forms	Decision-making re: an unplanned pregnancy (3) Recognizing warning signs in relationships (1) Recognizing forms of intimacy (1) Risk-recognition and risk-reduction strategies (7)
Expectations		Understanding family planning, using contraception, being abstinent (2)
	Anticipatory outcomes of a behavior	Safely surrendering a newborn baby (3) Using assertiveness and refusal skills (8)
Behavioral Capability	Knowledge to perform a given behavior (e.g. refusal skills, contraceptive use)	Practicing assertiveness and refusal skills (8) Recognizing and avoiding risk situations, incl. universal precautions, condom use (7)
Self-Efficacy	The person's confidence in performing a particular	Practicing assertiveness and refusal skills (8) Recognizing and avoiding risk situations (5,7), using condoms (7)
Desirenceal	behavior	Knowing how & where to access STD testing (9)
Reciprocal Determinism	The dynamic interaction of the person, the behavior, and the environment	Utilizing teen-friendly reproductive health services (2, 6, 9) Recognizing the existence of Safe Surrender Sites in the community (3) Youth discuss sexual attitudes and behaviors with parents (1-10)
Reinforcement	Responses to a person's behavior that increase or decrease the likelihood of reoccurrence.	Youth receive praise for proper practice of the refusal model (8), for identifying ways to help HIV-infected persons (4), for identifying reasons to be sexually abstinent (2), for making responsible decisions (3), for recognizing pressure situations (8).
Observational Learning	Behavior change occurs by watching the actions and outcomes of others' behavior	Youth observe others (receive praise for) proper practice of the refusal model (8), for identifying reasons to be sexually abstinent (2), for making responsible decisions (3), recognizing pressure situations (8).



Appendix B: Data Collection Periods

Table B.1. Data Collection Efforts Used in the Impact Analysis of Positive Prevention PLUS and Timing.

Data collection effort	Timing
Parental Consent	8/23/13-8/30/13
Random Assignment	9/3/13
Initial Teacher Training (2 Days)	9/10-9/11/13
Follow-up Training (1 day)	10/15/13
Baseline Survey	10/21-10/25/13
Program Implementation	10/28-11/15/13
6 Month Follow-Up	5/08-5/20/14

Notes: Data collection occurred during one time period and was pooled across sites.

Appendix C: Primary Research Questions

Outcome name	Description of outcome	Timing of measure relative to program
Ever had sexual intercourse	The variable is a yes/no measure of whether a person has ever had sexual intercourse. The measure is taken directly from the following item on the survey:	6 months after program ends
	• "Have you ever had sexual intercourse?"	
	The variable is constructed as a dummy variable where respondents who respond yes they have had sex are coded as 1 and no are coded as 0. Missing data was coded as 999.	
Ever been pregnant	The variable is a yes/no measure of whether a person has ever been pregnant or gotten someone pregnant. The measure is taken directly from the following item on the survey:	6 months after program ends
	• "To the best of your knowledge, have you been pregnant or gotten somebody pregnant?"	
	The variable is constructed as a dummy variable where respondents who respond yes have been pregnant were coded as 1 and those who responded no are coded as 0. Missing data resulting from the skip pattern of the survey was logically imputed to 0 since one can infer that someone who had never had sex has never been pregnant.	
Had sex in last 3 months WITHOUT	The variable is a yes/no measure of whether a person has sex in the past 3 months without using birth control. The measure is taken directly from the following item on the survey:	6 months after program ends
birth control	• "In the past 3 months, have you had sexual intercourse without you or your partner using any of these methods of birth control?"	
	-Condoms	
	-Birth control pills	
	-The shot (Depo Provera)	
	-The patch	
	-The ring (NuvaRing)	
	-IUD (Mirena or Paragard)-Implant (Impanon)	
	The variable was constructed as a dummy variable where respondents who respond yes have had sex in last 3 months without birth control are coded as 1 and those who responded no are coded as 0. Missing data due to the skip pattern was coded as 0 since one can infer that someone who had never had sex had also never had sex without birth control.	

Table C.1: Behavioral Outcomes Used for Primary Impact Analyses Research Questions

Appendix D: Intervention Status

Table D.1: Sample Size by Intervention Status

Number of:	Time period	Total sample size	Intervention sample size	Comparison sample size	Total response rate	Intervention response rate	Comparison response rate
Clusters: At beginning of study		21 ^a	11	10			
Clusters: Contributed at least one youth at baseline	<i>October 21st-</i> 25th, 2013	21	11	10	=1.00	=1.00	=1.00
Clusters: Contributed at least one youth at follow-up	May 8th- 20th, 2014	21	11	10	=1.00	=1.00	=1.00
Youth: In non-attriting clusters/sites at time of assignment		4,267	2,483	1,784			
Youth: Contributed a baseline survey		3,554	2,149	1,405	=.83	=.87	=.79
Youth: Contributed a follow-up survey	May 8th- 20th, 2014	3,490	2,113	1,377	=.82	=.85	=.77

Appendix E: Implementation Fidelity Methods

Table E.1: Methods Used to Address Implementation Research Questions

Implementation element	Methods used to address each implementation element
Adherence: How often were sessions offered? How many were offered?	The teacher curriculum log measured the date each lesson was taught, adherence to each curriculum component or lesson activity, and adaptations made to lessons. Teachers in the treatment group were asked to self-report the extent to which each topic, worksheet, or activity was covered completely or if any changes were made and why after teaching each lesson.
Adherence: What and how much was received?	This was measured using attendance data from each school district's student information system (SIS).
Adherence: What content was delivered to youth?	The teacher curriculum log measures the date each lesson was taught, adherence to each curriculum component or lesson activity, and if any adaptations were made to a lesson activity. Teachers in the treatment group were asked to self-report the extent to which each topic, worksheet, or activity was covered completely or if any changes were made.
Adherence: Who delivered material to youth?	Demographic information on program implementers (teachers) was captured on the teacher survey.
Quality: Quality of staff- participant interactions	Classroom observations were completed on one lesson per teacher at random using the classroom observation form. Scores from each of the observations ranged from one to five $(1 = Poor Understanding, 3 = Some understanding, 5 = Good understanding)$. To calculate each teacher's average quality score, the sum of each teacher's observation scores were divided by the number of observation questions (n=10). An average high score of 4 or better indicates a high-quality rating.
Quality: Quality of youth engagement with program	The quality of youth engagement was captured through classroom observations (e.g., "How exactly did the group members participate in the discussions and activities?" $1 = Little participation$, $3 = Some participation$, $5 = Active participation$). Student engagement was measured by dividing the student participation observation scores of 4 or better and dividing it by the total number of observations.

Implementation element	Methods used to address each implementation element
Counterfactual: Experiences of counterfactual condition	Teachers in both the treatment and control groups were asked questions (teacher survey) about their background, comfort teaching health related topics and if they had taught any health related topics in the past six months (i.e., Which of the following health topics have you taught about in the last six months? Drugs/tobacco/alcohol, abstinence, nutrition, decision making etc. Students were given a similar surveys asking "Which of the following health topics have you learned about this school year? -pregnancy prevention, AIDS or HIV infection, human sexuality, taking care of a baby, using condoms, or abstinence." The percent of TPP health topics received by students in each group was calculated by summing the responses from each student ($1 = yes$, $0 = no$) and dividing them by the total number of possible responses from each teacher ($1 = yes$, $0 = no$) and dividing them by the total number of possible responses.
Context: Other TPP programming available or offered to study participants (both intervention and counterfactual)	Teachers were asked if their school site had conducted any school-wide activities like assemblies, club events, or guest speakers regarding pregnancy prevention, sexuality, HIV/AIDS, or reproductive health in the past 6 months and asked to explain the type and duration of that activity. The percent of TPP health topics taught by teachers in each group (not in the PP+ program) was calculated by summing the responses from each teacher ($1 = yes$, $0 = no$) and dividing them by the total number of possible responses.
Context: External events affecting implementation	This information was captured through the curriculum fidelity log and informal conversations with treatment group teachers.
Context: Substantial unplanned adaptation(s)	This information was captured through the curriculum fidelity log and informal conversations with treatment group teachers.

TPP = Teen Pregnancy Prevention

Appendix F: Implementation Fidelity Data

Table F.1: Implementation adherence summary

Table F.1: Implementation adherence summary				
Implementation element	Summary of Findings			
Adherence: How often were sessions offered? How many were offered?	There were 1,353 lessons offered in the program (123 sections of the 11 lesson program). Each lesson was approximately 40 minutes in length. The average weekly frequency was 455 lessons within the 18 school day implementation period.			
Adherence: What and how much was received?	The average number of lessons attended by each student was 10.01 (SD = 1.63). Ninety-one percent of the entire program was attended (21,418 attended lessons of the 23,529 possible lessons (2,139 students with 11 lessons each)).			

Table F.1a: Details of program attendance (Adherence)

Student attendance by lesson	<i>n</i> = 2,139	%
PP+ Getting Started	1,712	80.03
Lesson 1: Life Planning	1,912	89.39
Lesson 2: Healthy Relationships	2,117	98.97
Lesson 3: Relationship Violence	2,128	99.49
Lesson 4: Family Planning and Contraceptives	2,130	99.58
Lesson 5: Myths and Stereotypes	2,129	99.53
Lesson 6: HIV Disease and AIDS	2,129	99.53
Lesson 7: Recognizing and Reducing Risk	2,129	99.53
Lesson 8: Peer and Media Pressures	2,129	99.53
Lesson 9: HIV/STD Testing and Community Resources	2,128	99.49
Lesson 10: Steps to Success	2,114	98.83

Implementation element	Summary of Findings
Adherence: Who delivered material to youth?	Thirty-six teachers were trained in the program by project staff. Ninety-four percent (thirty- four teachers) of the trained teachers implemented the program. Thirty-three teachers participated in a teacher survey before their first training session.
Quality: Quality of staff-participant interactions	Seventy-three percent of the 26 observed sessions received a high quality observation.
Quality: Quality of youth engagement with program	Sixty-five percent of the observed sessions were rated with high student engagement (17 of the 26 observed sessions).

Table F.2: Characteristics of program staff and interactions

Table F.2a: Demographic characteristics of teachers in treatment condition

Treatment Teacher Demographics	<i>n</i> = 33	%
Gender		
Female	27	81.80
Male	6	18.20
Ethnicity		
White	19	63.30
African American	2	6.70
Asian	1	3.30
Hispanic	5	16.70
Other race	3	10.00
Teaching Credential		
Health Education	17	51.50
Other credential	16	48.50
Degree		
Health Education	8	27.60
Physical Education	3	10.30
Science	13	44.80
Other degree	5	17.20
Years Teaching (M/SD)	10.58	7.33

Implementation element	Summary of Findings
Counterfactual: Experiences of comparison condition	The percent of TPP health topics received by students in each group was calculated by summing the responses from each student ($1 = yes$, $0 = no$) and dividing them by the total number of possible responses.
	The percent of TPP health topics taught by teachers in each group was calculated by summing the responses from each teacher $(1 = yes, 0 = no)$ and dividing them by the total number of possible responses.

Table F.3: Experiences of the treatment and comparison conditions

Table F.3a: Health topic coverage, as described by students

"Which of the following health topics have you learned about this school year?"	Treatment group youth N	%	Control group youth N	%
Pregnancy Prevention	1,620	86.30	354	28.30
AIDS or HIV Infection	1,878	92.90	475	38.00
Human Sexuality	1,257	66.90	300	24.00
Taking Care of a Baby	277	14.70	92	7.40
Using Condoms	1,643	87.50	97	7.80
Abstinence	1,234	65.70	128	10.20

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"Which of the following health topics have you taught about in the last 6 months?"	Treatment group teachers N	%	Control group teachers N	%
Pregnancy Prevention	21	67.70	1	4.80
AIDS or HIV Infection	23	74.20	6	28.60
Human Sexuality	17	54.80	2	9.50
Taking Care of a Baby	6	19.40	0	0.00
Using Condoms	20	64.50	1	4.80
Abstinence	20	64.50	3	14.30

Table F.3b: Health topic coverage, as described by teachers

Table F.4: Implementation context

Implementation element	Summary of Findings
Context: Other TPP programming available or offered to study participants (both intervention and comparison)	None of the teachers in either group reported any additional school-wide activities related to pregnancy prevention, sexuality, HIV/AIDs or reproductive health.
Context: External events affecting implementation	None
Context: Substantial unplanned adaptation(s)	Six teachers at one school site were not allowed to conduct the condom demonstration per their school district policy.

Baseline measure	Intervention mean or % (standard deviation)	Comparison mean or % (standard deviation)	Intervention versus comparison mean difference	Intervention versus comparison <i>p</i> -value of difference
Age or grade level	14.63 (.50)	14.63 (.48)	0	.99
Gender (female)	.52	.56		.78
Race: Asian/Pacific Islander	.04	.05	01	.41
Race:Black	.07	.07	0	1.00
Race: White	.14	.16	02	.29
Race: Native American	.03	.03	0	1.00
Race: Multiple Races	.60	.59	.01	.55
Race: Unkown	.12	.10	.02	.29
Ethnicity: Hispanic	.74	.73		.48
Ever Had Sex	.12	.12	0	.76
Ever Been Pregnant	.06	.06	0	.73
Had Sex WITHOUT BC in Prior 3 Months	.02	.03	.01	.39
Sample size ^b	1,902-1,886	1,238-1,197		

Notes: ^a Estimate with Huber-White Robust Standard Errors; ^b Due to item non-response, N's vary slightly by variable.

Outcome measure	Intervention mean or % (N)	Comparison mean or % (N)	Intervention compared to comparison mean difference (<i>p</i> -value of difference)
Ever had sexual intercourse	.14 (1,900)	.18 (1,238)	04 (.01)
Ever been pregnant (or gotten someone pregnant)	.02 (1,902)	.03 (1,243)	01 (.07)
Ever had sexual intercourse without using birth control	.04 (1,886)	.06 (1,197)	02 (.01)

Table G.2 Post-intervention Estimated Effects using Data from PP+ Student Survey to Address the Primary Research Questions

Source: Follow-up surveys administered 6 months after the program.

Outcome measure	Reported p value from Benchmark Analysis of Analytical Sample	p value rank	$\infty_i^{'} = \frac{i^* \infty}{M}$	$p_i \langle X_i$	Statistical Significance After BH Correction
Ever had sexual intercourse	.01	1.5	0.025	Yes	Significant
Ever been pregnant (or gotten someone pregnant)	.07		0.050	No	Not Significant
Ever had sexual intercourse without using birth control	.01	1.5	0.025	Yes	Significant

Table G.3 Benjamini- Hochberg Correction for Multiple Comparisons

Source: Follow-up surveys administered 6 months after the program.

Appendix H: Sensitivity Analyses

This appendix presents impacts of the Positive Prevention PLUS program estimated using the benchmark analytic sample and two alternative samples concerning the treatment of missing data: (1) impact models involving data where logical imputation was made and (2) impact models involving data where multiple imputation was made. The primary impacts are predominantly robust to alternative specifications. When impact estimation used the logical imputation sample, estimated impacts were similar to the primary impact estimates in magnitude and identical in statistical significance. The data decision rules for logical imputation are shown in Table G.1.

	IF	THEN	
The following data decision rules were applied consistently across both groups and for baseline and follow-up data:	Had sex in past 3 months= Yes and Ever had sex= No.	Impute both to missing value.	
	Ever been pregnant=Yes and Ever Had Sex= No	Impute both to missing value.	
	Having had sex without birth control in prior 3 months = Yes and Ever had sex= No	Impute both to missing value.	
	Between survey inconsistencies (e.g. if Ever had Sex at baseline=Yes and Ever had Sex at 6 month follow-up= No or missing.)	Impute Ever had Sex at 6 month follow-up=Yes.	

Table H.1: Logical Imputation Rules

Intervention compared with comparison	Benchmark approach difference ^a	Benchmark approach <i>p-</i> value	Data Applying Logical Imputation Using Data Decision Rules difference	Data Applying Logical Imputation Using Data Decision Rules <i>p</i> - value	Data Applying Missing Data Imputation (Using Multiple Imputation) difference ^b	Data Applying Missing Data Imputation (Using Multiple Imputation) <i>p</i> -value
Ever had sex	04	.01	03	.01	04	.10
Ever been pregnant	01	.07	03	.19	05	.53
Having had sex without birth control in prior 3 months	02	.01	06	.22	11	.28

Table H.2: Sensitivity of Impact Analyses using Data from PP+ Student Survey to Address the Primary Research Questions

Source: Follow-up surveys administered 6 months after the program.

Notes: ^a Using estimated adjusted means; ^b Using pooled estimates of coefficients from MI regression; imputations made separately for the treatment and control groups. PP+= Positive Prevention PLUS.