

# **Financial Incentives Increase Smoking Cessation and Improve Other Maternal-Infant Outcomes Among Pregnant & Newly Postpartum Women**

Stephen T. Higgins, PhD

Vermont Center on Behavior and Health

Departments of Psychiatry and Psychological Science

University of Vermont

<http://www.uvm.edu/medicine/behaviorandhealth/>

# Acknowledgements/Disclosures

- **Collaborators:** Tyler D. Nighbor, PhD, Allison A. Kurti, PhD, Sarah H. Heil, PhD, Eric P. Slade, PhD, Donald S. Shepard, PhD, Laura J. Solomon, PhD, Mary Ellen Lynch, RN, Harley K. Johnson, MA, Catherine Markesich, BS, Peter L. Rippberger BS, Joan M. Skelly, MS, Michael DeSarno, MS, Janice Bunn, PhD, Jessie B. Hammond, MPH, Maria L. Roemhildt, PhD, Rhonda K. Williams, MES, Deirdre M. O'Reilly, MD, Ira M. Bernstein, MD
- **Research Support:** NIGMS P20GM103644, NICHD R01R01HD075669
- **Disclosures:** Nothing to declare.

# Introduction

- Smoking during pregnancy is a leading preventable cause of poor pregnancy outcomes in the U.S. & other developed countries.
- Socioeconomically disadvantaged women are at high risk.
- Efficacious cessation interventions widely available to pregnant women but antepartum quit rates are unacceptably low (<15%).
- Usual care is typically a referral to a quitline; best practices includes follow-up and further referral for continuing smokers.
- There is broad consensus on need for improvements; I'll show evidence that financial incentives represent an innovation that can reliably increase quit rates antepartum and early postpartum.

# Methods

- **Current Trial:** RCT comparing best practices (BP) vs. BP + financial incentives (BP+FI) among 169 women still smoking at 1<sup>st</sup> AP visit and 80 SES-matched never smokers (NS)
- **Primary outcome:** antepartum abstinence
- **Secondary outcomes:** craving/withdrawal, birth outcomes, postpartum abstinence, breastfeeding, infant growth/development, cost-benefit analysis
- **Pooled data set:** to assess the reliability of positive trial outcomes we examined effects in a pooled data set wherein current trial (n=169) data were combined with four prior RCTs examining this FI model versus a non-contingent incentives control condition (n=245) for pooled total n=453 (FI=245; Controls=208)
- **Vermont-wide sample:** to assess external validity of trial results on relationship between maternal smoking status (never-smoked, smoked but quit, continued smoking) to small-for-gestational age (SGA) birth outcomes, and associated healthcare costs (i.e., all singleton deliveries in VT in 2019).

# Participants

- Recruited women still smoking at 1<sup>st</sup> prenatal visit from ObGyn clinics in Burlington VT and surrounding counties.
- Inclusion criteria: biochemically confirmed self-report smoking in past 7 days, gestational age  $\leq 25$  weeks, plans to remain in area for next 12 mos, English speaking.
- Exclusion criteria: incarceration, prior participation in incentives cessation study, residing with current trial participant, regular use of opioids, stimulant, antipsychotic meds
- 584 who reported smoking began screening; 126 failed to complete screening, 282 ineligible, 176 enrolled
- 759 never-smokers initiated screening; 21 failed to complete screening, 657 ineligible, 81 enrolled.
- Only reason for exclusion once enrolled was abortion/fetal demise (3 BP, 4 BP+FI, 1 NS)

# Trial Assessments

- At intake, participants completed questionnaires examining sociodemographic, smoking, and psychiatric conditions, provided breath and urine specimens
- Modified version of that battery completed one month after intake (early antepartum assessment), at  $\geq 28$ -weeks gestation (late-pregnancy assessment), 2-, 4-, 8-, 12-, 24-, 48-weeks postpartum. Also assessed breath CO and urine cotinine.
- Birth outcomes obtained from maternal medical record.

# Trial Conditions

- All participants assigned to BP encouraged to choose a quit date in next two weeks; once a quit date was selected a signed referral faxed to Vermont quitline.
- Quitline offered perinatal-specific brief phone counseling (National Jewish Health) with quit coach (5 antepartum; 4 postpartum calls) based on stages of change, including motivational interviewing and cognitive-behavioral strategies.
- Quitline offered women \$65 in incentives for completing calls.
- Eligible for free nicotine replacement if their providers agreed.
- Women still smoking at scheduled assessments were referred again to the quitline.
- Women assigned to BP+FI encouraged to pick a Monday quit date in next two weeks; received everything above and started on financial incentives on their quit date.

# Incentives Model

- Vouchers exchangeable for retail items available antepartum through 12-weeks postpartum
- Voucher delivery contingent on biochemical test results: breath CO  $\leq 6$  ppm initial 5 days of the quit week; urine cotinine (onsite enzyme immunoassay  $\leq 80$  ng/ml) thereafter.
- Daily (M-F) of quit week, 2x weekly next 7 weeks, once weekly for 4 weeks, and then every other week till delivery; following delivery back to weekly through 12-weeks postpartum.
- Voucher value varied by baseline CPD;  $< 10$  CPD: began at \$6.25, escalated by \$1.25 each consecutive negative test to max \$45.00; positive test reset vouchers to initial low value; two negative tests restored vouchers to pre-reset value;  $\geq 10$  CPD: voucher same as above but \$ values doubled.
- Total mean earnings: \$510.02 $\pm$ 76.27 (\$467.70 $\pm$ 68.21 and \$560.34 $\pm$ 146.84 in  $< 10$  CPD and  $> 10$  CPD , respectively).

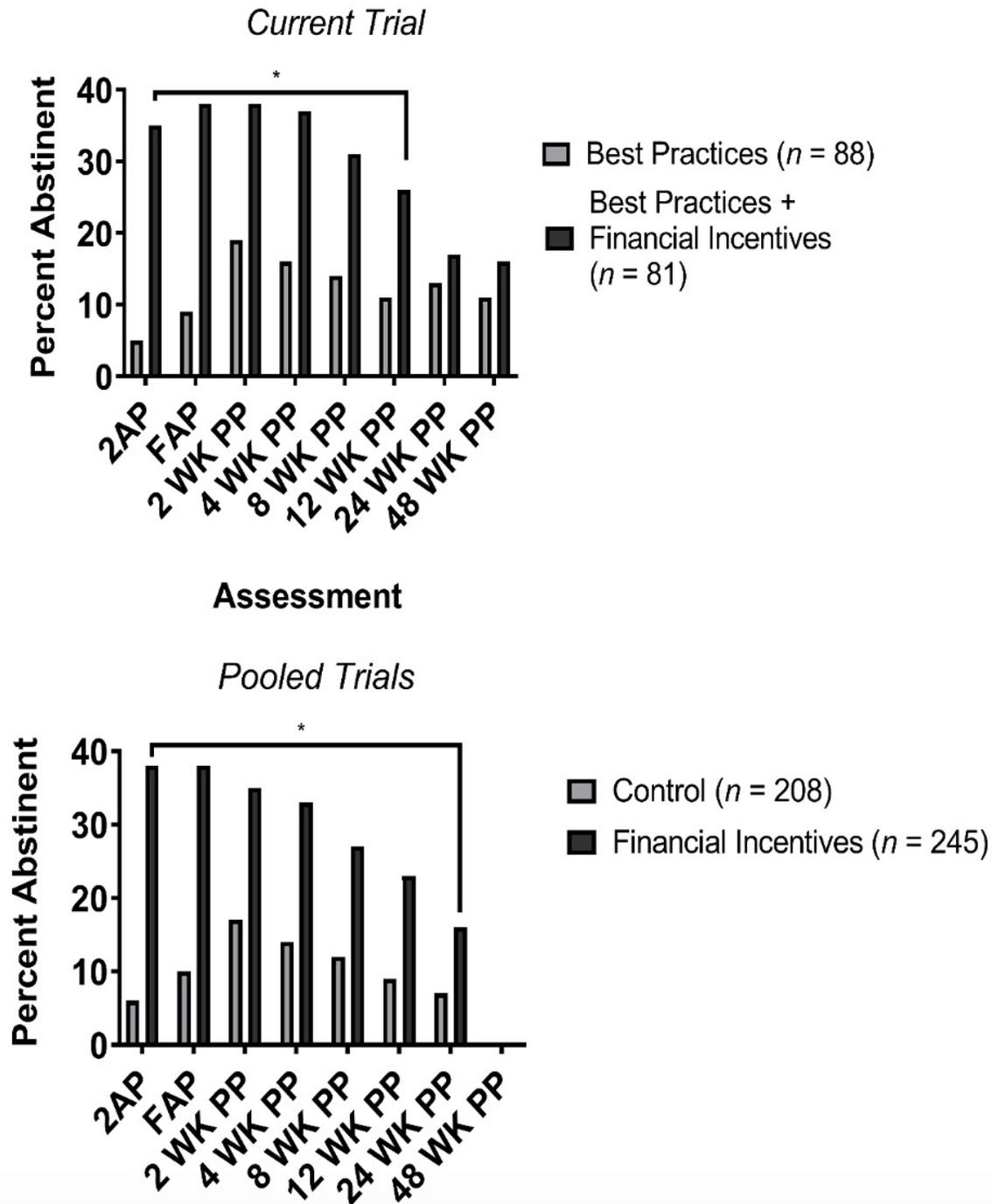
**Table 1.** Participant characteristics.

Characteristics	BP <sup>1</sup> (n=88)	BP+FI <sup>2</sup> (n=81)	NS <sup>3</sup> (n=80)	BP vs BP+FI p-value <sup>a</sup>	All groups p-value <sup>b</sup>
<b>Demographics:</b>					
Age (years)	26.61 ±5.47	25.40 ± 4.96	25.6 ± 5.0	.13	.25
Education				.08	.001
% <12 years	16	28	10 <sup>c</sup>		
% 12 years	64	48	80 <sup>d</sup>		
% >12 years	20	23	10 <sup>c</sup>		
% Non-Latino White	93	91	90	.66	.76
% Married	20	16	42 <sup>d</sup>	.46	<.001
% Private insurance	28	27	40	.86	.15
% Work outside of home	50	59	66	.23	.10
% 1st pregnancy	42	53	49	.15	.35
Weeks pregnant at intake	11.14± 4.07	12.37 ± 4.22	15.2 ± 6.4 <sup>d</sup>	.05	<.001
Pre-pregnancy BMI	29.60 ±8.45	28.20 ± 7.54	28.4 ± 8.3	.26	.47

**Table 1.** Participant characteristics.

<b>Characteristics</b>	<b>BP<sup>1</sup> (n=88)</b>	<b>BP+FI<sup>2</sup> (n=81)</b>	<b>NS<sup>3</sup> (n=80)</b>	<b>BP vs BP+FI p-value<sup>a</sup></b>	<b>All groups p-value<sup>b</sup></b>
<b>Smoking Characteristics:</b>					
Cigs/day pre-pregnancy	18.27± 9.42	19.25 ± 9.87	NA	.51	
Cigs/day at 1 <sup>st</sup> AP	9.92 ± 6.18	8.99 ± 5.21	NA	.29	
Age started smoking (yrs)	15.47± 2.96	15.10 ± 2.92	NA	.41	
% Living with other smoker(s)	77	79	22 <sup>d</sup>	.75	<.001
% With no smoking allowed in home	70	68	91 <sup>d</sup>	.72	<.001
% With none or few friends/family who smoke	26	25	78 <sup>d</sup>	.83	<.001
% Attempted to quit pre-pregnancy	73	70	NA	.73	
Number of quit attempts during pregnancy	0.73 ± 2.35	0.57 ± 0.97	NA	.56	
Nicotine withdrawal questionnaire total scores	1.60 ± 0.75	1.37 ± 0.76	NA	.05	
Fagerström total scores	4.23 ± 2.30	3.98 ± 2.07	NA	.46	

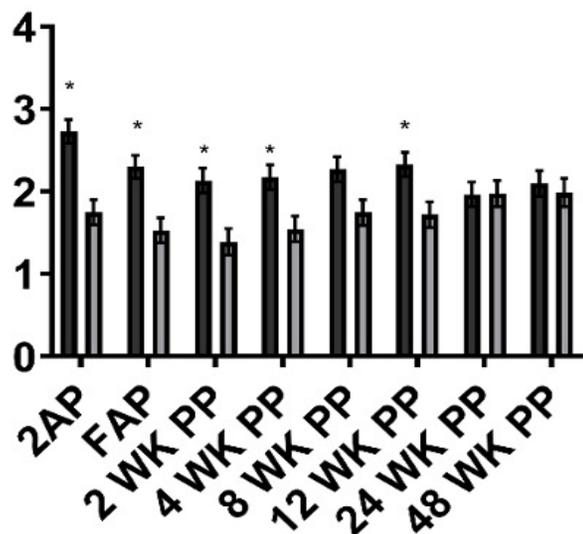
# 7-Day Point Prevalence Abstinence



# Least Squares Means

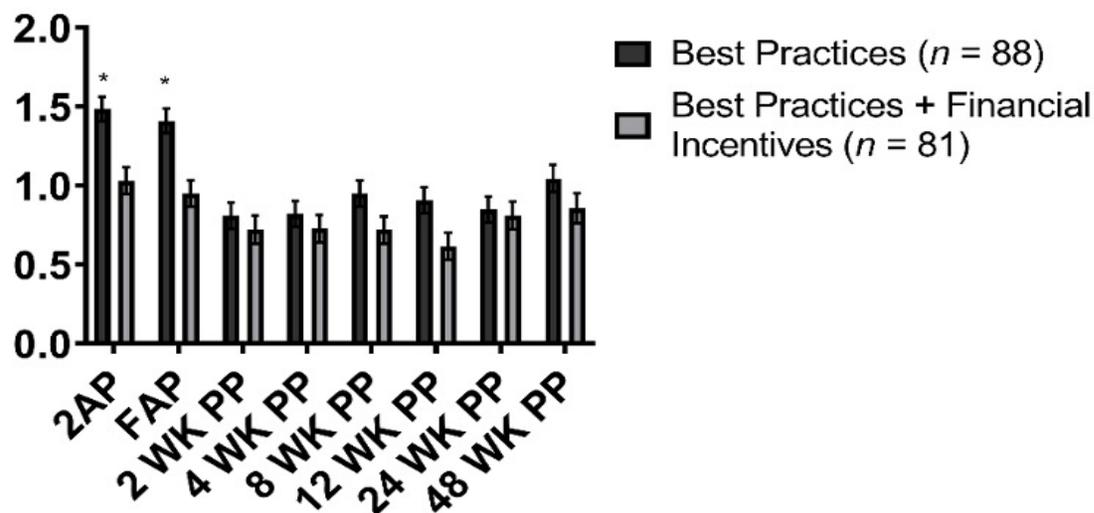
## Craving

Current Trial

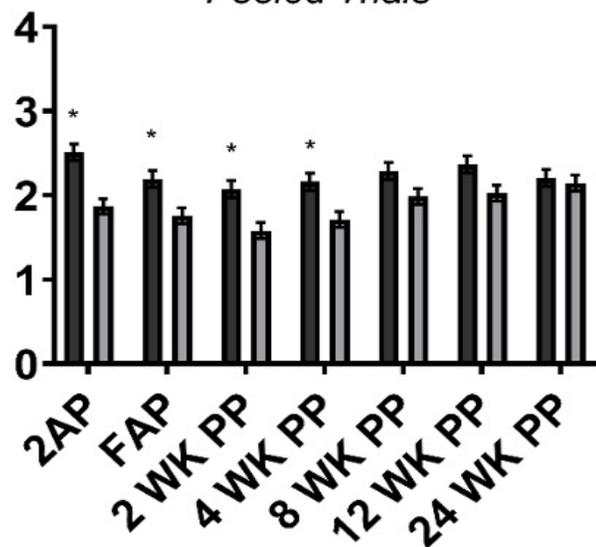


## Total Scores

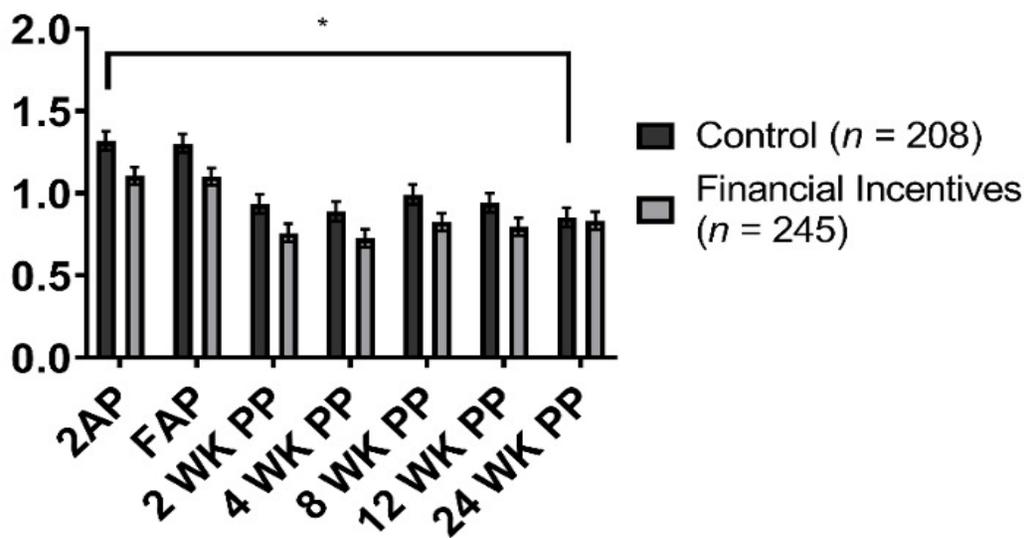
Current Trial



Pooled Trials



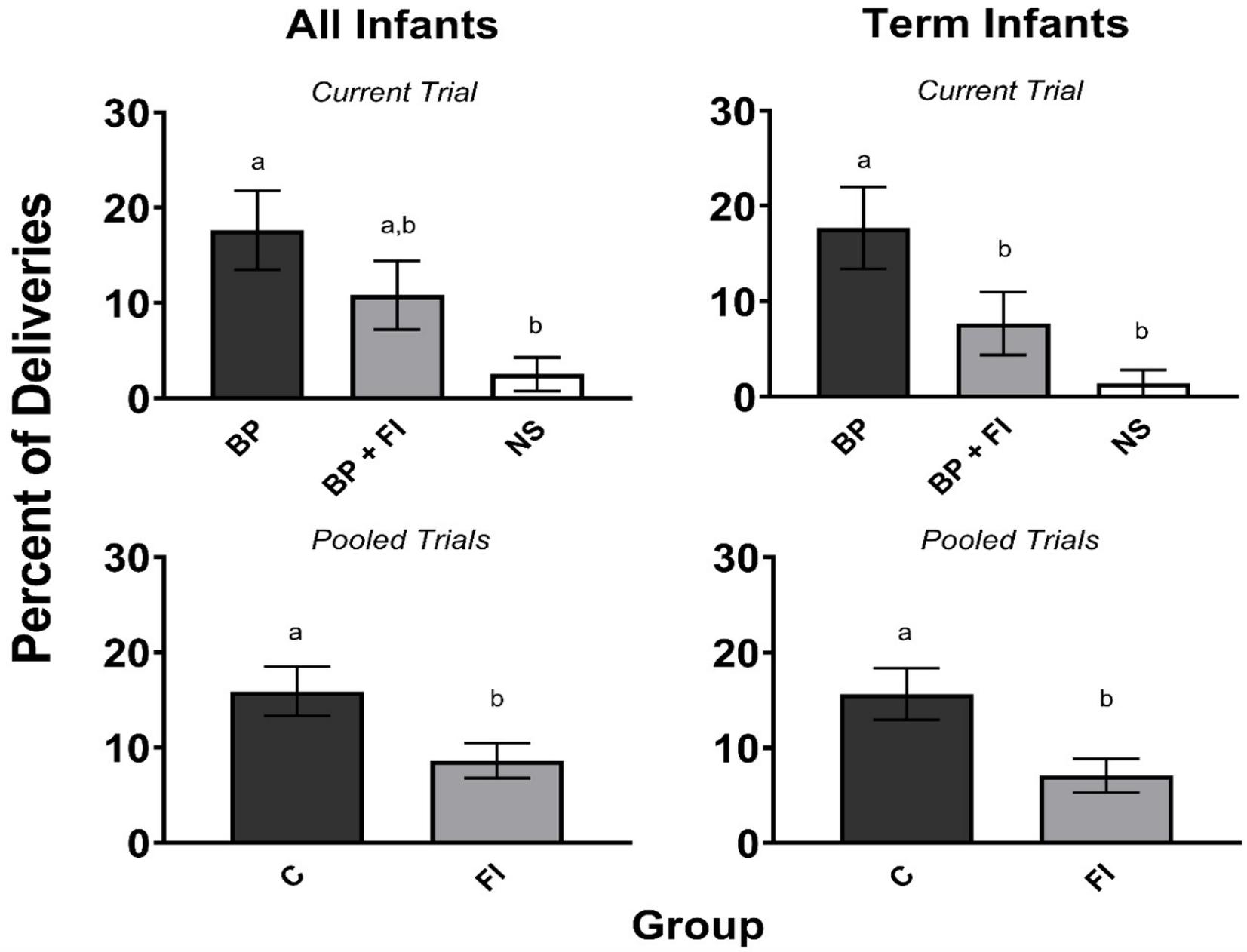
Pooled Trials



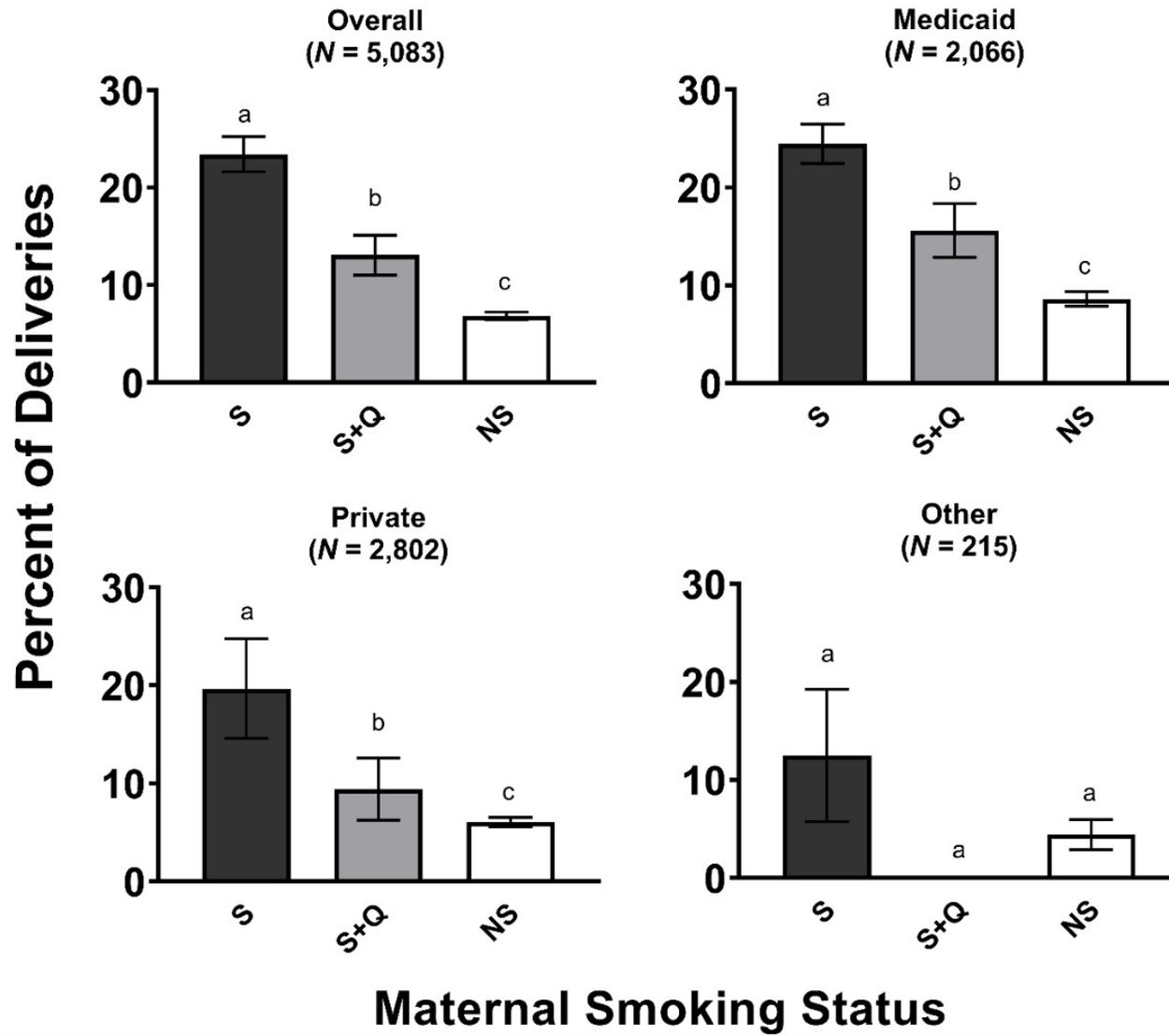
Assessment

Assessment

# Small for Gestational Age Deliveries



# Vermont Small for Gestational Age Deliveries



# Breastfeeding

<b>Assessments</b>	<b>% Breastfeeding*</b>		
	<b>BP<sup>1</sup> (n=88)</b>	<b>BP + FI<sup>2</sup> (n=81)</b>	<b>NS<sup>3</sup> (n=80)</b>
	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>
2 weeks	68.06 (49/72) <sup>a</sup>	78.13 (50/64) <sup>a</sup>	78.67 (59/75) <sup>a</sup>
4 weeks	52.63 (40/76) <sup>a</sup>	60.61 (40/66) <sup>a</sup>	72.73 (56/77) <sup>a</sup>
8 weeks	44.74 (34/76) <sup>a</sup>	49.25 (33/67) <sup>a</sup>	63.16 (48/76) <sup>a</sup>
12 weeks	36.84 (28/76) <sup>a</sup>	47.06 (32/68) <sup>a</sup>	56.58 (43/76) <sup>a</sup>
24 weeks	28.00 (21/54) <sup>a</sup>	35.48 (22/62) <sup>a</sup>	50.00 (36/72) <sup>a</sup>
48 weeks	12.16 (9/74) <sup>a</sup>	15.00 (9/60) <sup>a</sup>	32.86 (23/70) <sup>a</sup>

## **% Breastfeeding and abstinent<sup>#</sup>**

<b>Assessments</b>	<b>BP (n=88)</b>	<b>BP + FI (n=81)</b>	<b>NS (n=80)</b>
	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>
2 weeks	17.05 (15/88) <sup>a</sup>	35.80 (29/81) <sup>b</sup>	78.67 (59/75) <sup>c</sup>
4 weeks	12.50 (11/88) <sup>a</sup>	29.63 (24/81) <sup>b</sup>	72.73 (56/77) <sup>c</sup>
8 weeks	10.23 (9/88) <sup>a</sup>	23.46 (19/81) <sup>b</sup>	63.16 (48/76) <sup>c</sup>
12 weeks	7.95 (7/88) <sup>a</sup>	18.52 (15/81) <sup>b</sup>	56.58 (43/76) <sup>c</sup>
24 weeks	7.95 (7/88) <sup>a</sup>	11.11 (9/81) <sup>b</sup>	50.00 (36/72) <sup>c</sup>
48 weeks	3.41 (3/88) <sup>a</sup>	6.12 (5/81) <sup>b</sup>	32.86 (23/70) <sup>c</sup>

# Breastfeeding

## % Breastfeeding<sup>#</sup>

<u>Assessments</u>	<u>Controls<sup>4</sup> (n=208)</u>	<u>FI<sup>5</sup> (n=245)</u>
	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>
2 <u>wks</u>	61.18 (104/ <u>170</u> ) <sup>a</sup>	67.31 (140/ <u>208</u> ) <sup>a</sup>
4 <u>wks</u>	47.73 (84/ <u>176</u> ) <sup>a</sup>	52.58 (112/ <u>213</u> ) <sup>a</sup>
8 <u>wks</u>	38.64 (68/ <u>176</u> ) <sup>a</sup>	43.98 (95/ <u>216</u> ) <sup>a</sup>
12 <u>wks</u>	29.05 (52/ <u>179</u> ) <sup>a</sup>	37.79 (82/ <u>217</u> ) <sup>a</sup>
24 <u>wks</u>	22.78 (41/ <u>180</u> ) <sup>a</sup>	24.76 (52/ <u>210</u> ) <sup>a</sup>
48 wks		

## % Breastfeeding and abstinent\*

<u>Assessments</u>	<u>Controls (n=208)</u>	<u>FI (n=245)</u>
	<u>Percent (n/d)</u>	<u>Percent (n/d)</u>
2 <u>wks</u>	12.98 (27/ <u>208</u> ) <sup>a</sup>	27.76 (68/ <u>245</u> ) <sup>b</sup>
4 <u>wks</u>	9.62 (20/ <u>208</u> ) <sup>a</sup>	22.86 (56/ <u>245</u> ) <sup>b</sup>
8 <u>wks</u>	7.21 (15/ <u>208</u> ) <sup>a</sup>	18.37 (45/ <u>245</u> ) <sup>b</sup>
12 <u>wks</u>	4.81 (10/ <u>208</u> ) <sup>a</sup>	14.69 (36/ <u>245</u> ) <sup>b</sup>
24 <u>wks</u>	3.37 (7/ <u>208</u> ) <sup>a</sup>	7.35 (18/ <u>245</u> ) <sup>b</sup>
48 <u>wks</u>		



**Table 4.** Estimates of the Economic Costs and Benefits of BP+FI compared to BP (\$ in 2020 dollars)

Costs & Benefits	BP+FI			BP Only			Average Cost Differences: BP+FI versus BP Only		
	Mean Cost Per Visit (\$)	Mean Cost Per Participant (\$)	Std. Err.	Mean Cost Per Visit (\$)	Cost Per Participant (\$)	Std. Err.	Mean Cost Difference Per Participant (\$)	95% CI LL	UL
C. Total Intervention Cost (A+B)	\$98.28	\$1,486.26	\$99.35	\$43.48	\$124.93	\$1.97	\$1,361.33	\$1,163.65	\$1,559.01
D. Medicaid Cost (Delivery and Newborn Care Months 0-12)	\$1,426.04	\$21,566.68	\$1,281.00	\$7,742.84	\$22,260.67	\$1,594.23	-\$693.99	-\$3,242.77	\$1,854.79
E. Medicaid Cost Plus Smoking Intervention Cost (C+D)	\$1,524.32	\$23,052.94	\$1,284.83	\$7,786.30	\$22,385.60	\$1,594.23	\$667.34	-\$2,111.35	\$3,380.87
F. Value of Reduced SUID Mortality from Quitting or Reducing Smoking During Pregnancy	\$269.02	\$4,068.53	\$1,117.13	\$894.22	\$2,570.89	\$121.08	\$1,497.64	\$1,094.99	\$1,952.25
G. Societal Net Benefit of BP+FI vs. BP							\$830.30	-\$2,285.88	\$4,063.60
Societal Return on Investment in BP+FI vs. BP (100% * H / C)							61.0%	-196.4%	260.7%



# Summary and Conclusions

- Overwhelming evidence that financial incentives increase antepartum abstinence (largest effect sizes in RCTS).
- Effects on abstinence remain robust through 12-weeks postpartum; effects after discontinuation of incentives remained above controls across current and pooled trials; only significant in the latter where power was greater.
- Effect of incentives on SGA is consistent across the current and pooled trials with VT-wide study supporting external validity. SGA increases infant and childhood morbidity and mortality risk and later-in-life risk for metabolic-disorder.
- Effects of incentives on continuing to breastfeed while abstinent illustrates the multifaceted ways in which increasing abstinence can foster health improvements.
- Economic analysis supports the cost-benefit of BP+FI over BP alone, but with wide CIs--uncertainty. Future economic impact studies examining beyond 1<sup>st</sup> year of infant life and with larger samples are needed.
- We have developed and pilot-tested the efficacy of a smart-phone translation to increase reach (Kurti et al., 2020, *Prev Med*) and have a recently completed RCT on the same that will be submitted in the near future.