

Leveraging loss aversion and present bias to improve incentives for smoking cessation

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- I have never received any support from any industry of any kind, including the tobacco industry

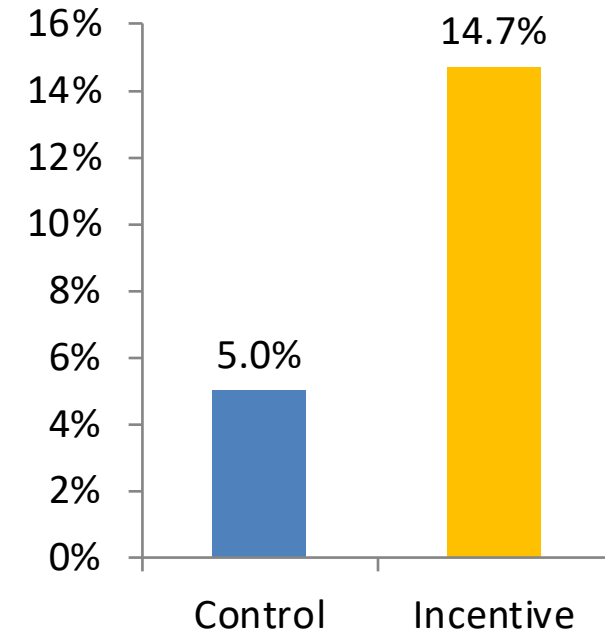
SPECIAL ARTICLE

A Randomized, Controlled Trial of Financial Incentives for Smoking Cessation

Kevin G. Volpp, M.D., Ph.D., Andrea B. Troxel, Sc.D., Mark V. Pauly, Ph.D., Henry A. Glick, Ph.D., Andrea Puig, B.A., David A. Asch, M.D., M.B.A., Robert Galvin, M.D., M.B.A., Jingsan Zhu, M.B.A., Fei Wan, M.S., Jill DeGuzman, B.S., Elizabeth Corbett, M.L.S., Janet Weiner, M.P.H., and Janet Audrain-McGovern, Ph.D.

878 General Electric employees, assigned to usual care (access to cessation counseling) or usual care + incentives worth \$750

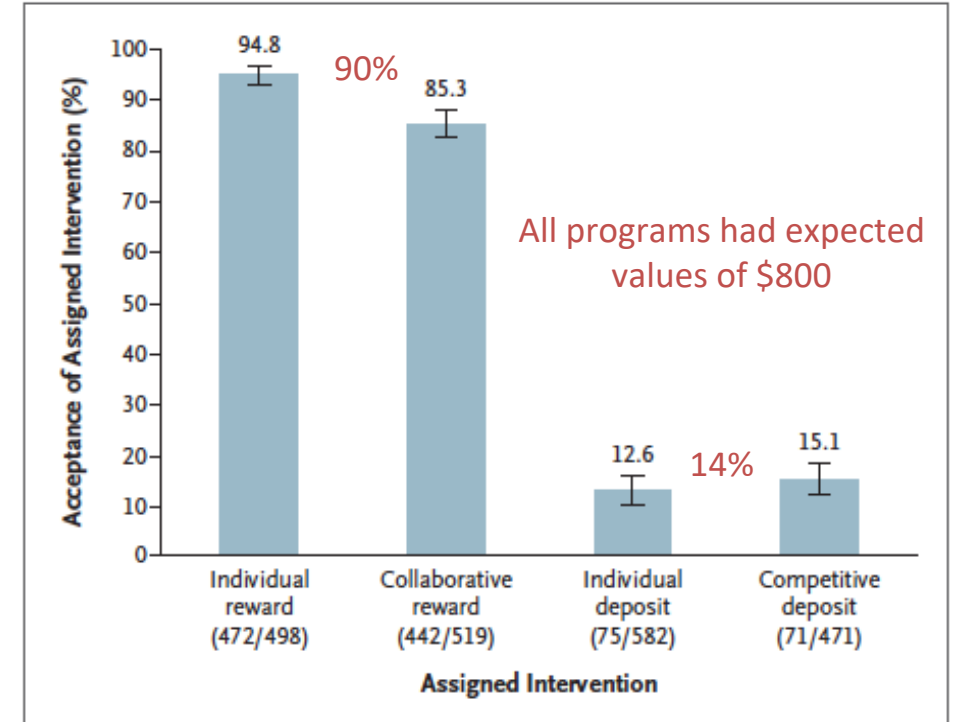
Sustained abstinence through 6 months

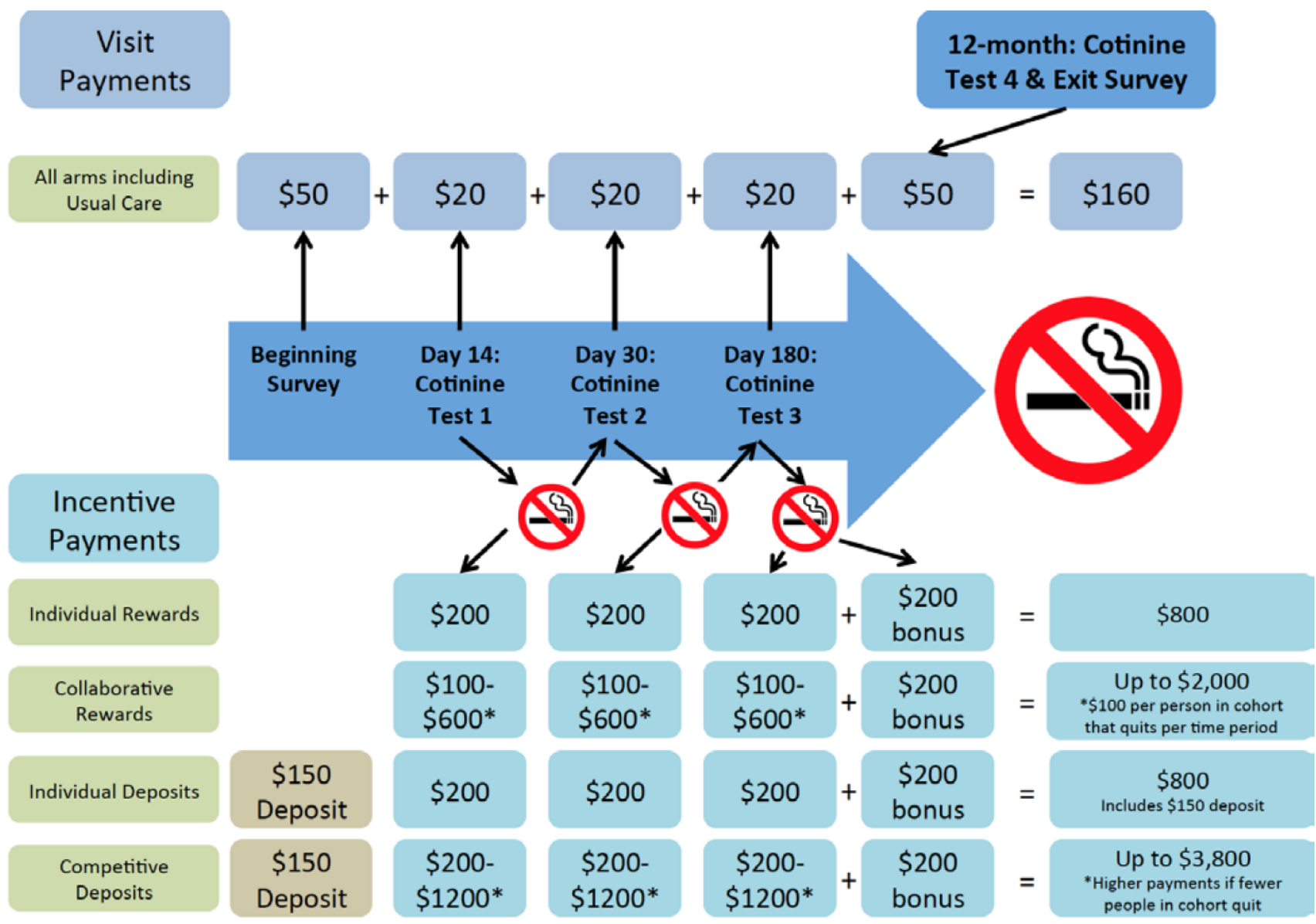


Randomized Trial of Four Financial-Incentive Programs for Smoking Cessation

Scott D. Halpern, M.D., Ph.D., Benjamin French, Ph.D., Dylan S. Small, Ph.D., Kathryn Saulsgiver, Ph.D., Michael O. Harhay, M.P.H., Janet Audrain-McGovern, Ph.D., George Loewenstein, Ph.D., Troyen A. Brennan, M.D., J.D., David A. Asch, M.D., M.B.A., and Kevin G. Volpp, M.D., Ph.D.

- Sample: 2,538 CVS Health employees or friends or family
- Usual care: access to information about benefits of cessation & to nicotine replacement therapy (NRT) & behavioral counseling





No differences across arms in actual payments

Table S1: Observed incentive payouts to participants in the four incentive arms

| | N | Mean | Median | Interquartile Range | Full Range |
|---|----|----------|--------|---------------------|------------|
| <i>All participants achieving abstinence for 6 months</i> | | | | | |
| Individual Reward | 76 | \$800 | \$800 | \$800-800 | \$0-800 |
| Collaborative Reward | 83 | \$890.36 | \$800 | \$700-1100 | \$0-1,700 |
| Individual Deposit | 56 | \$557.14 | \$800 | \$0-800 | \$0-800 |
| Competitive Deposit | 52 | \$839.62 | \$900 | \$630-1100 | \$0-1,940 |

Sustained abstinence rates (ITT)

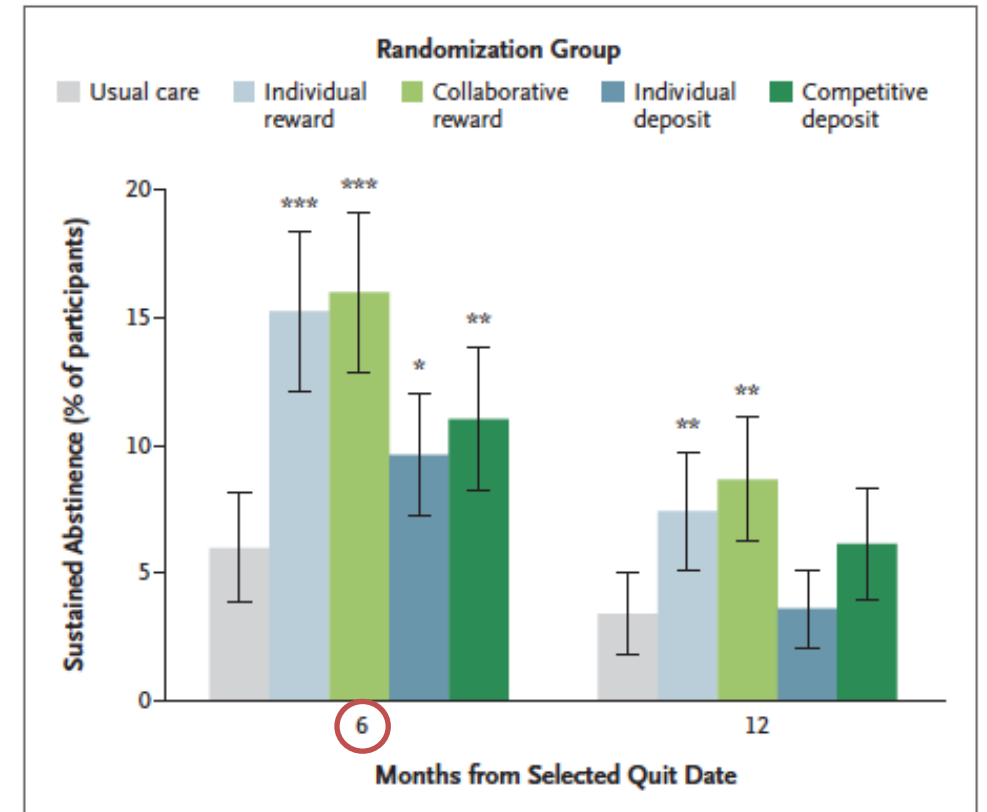
Rewards (15.7%) vs. Deposits (10.2%)

$p < 0.001$

Group (13.7%) vs. Individual (12.1%)

$p = 0.29$

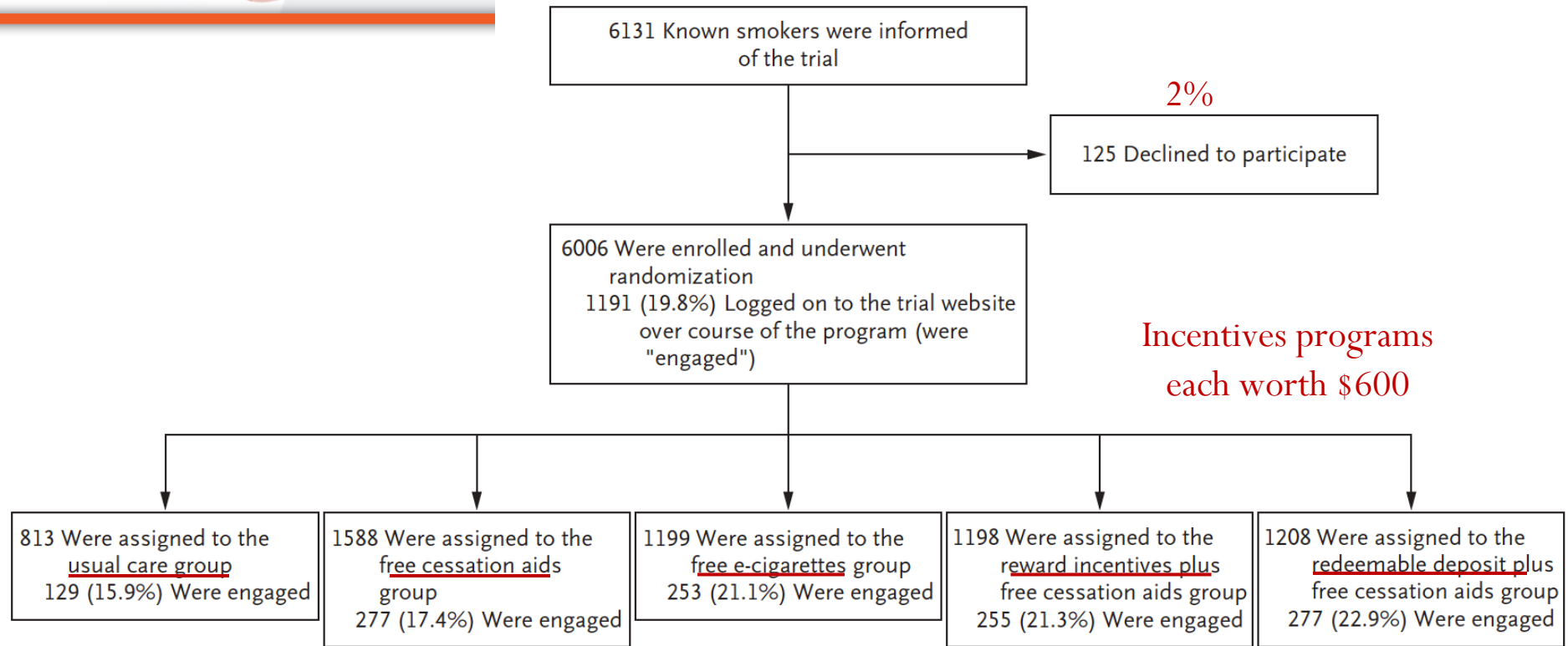
Complier average treatment effect analysis shows that among people who would have accepted deposits, **deposits were more efficacious than rewards** (29% vs. 16%)



SPECIAL ARTICLE

A Pragmatic Trial of E-Cigarettes, Incentives, and Drugs for Smoking Cessation

Scott D. Halpern, M.D., Ph.D., Michael O. Harhay, Ph.D.,
Kathryn Saulsgiver, Ph.D., Christine Brophy, Andrea B. Troxel, Sc.D.,
and Kevin G. Volpp, M.D., Ph.D.



All smokers at 54 U.S. companies who identified as smokers on health-risk assessment in prior year



2. E-cigarette Arm
Basic Vitality program AND free e-cigarettes

3. Choice Cessation Aids Arm
Basic Vitality program AND choice of free NRT, varenicline, bupropion; e-cig option if fail

4. Reward Incentive Arm
Basic Vitality program, choice of any free aid, AND \$600 incentives for confirmed tobacco cessation

Opted out = 125

Never engaged = 4,815

Engaged = 1,191



8 contrasts specified *a priori*,
with significance thresholds
adjusted using Holm method

Statistically significant

**Not statistically
significant**

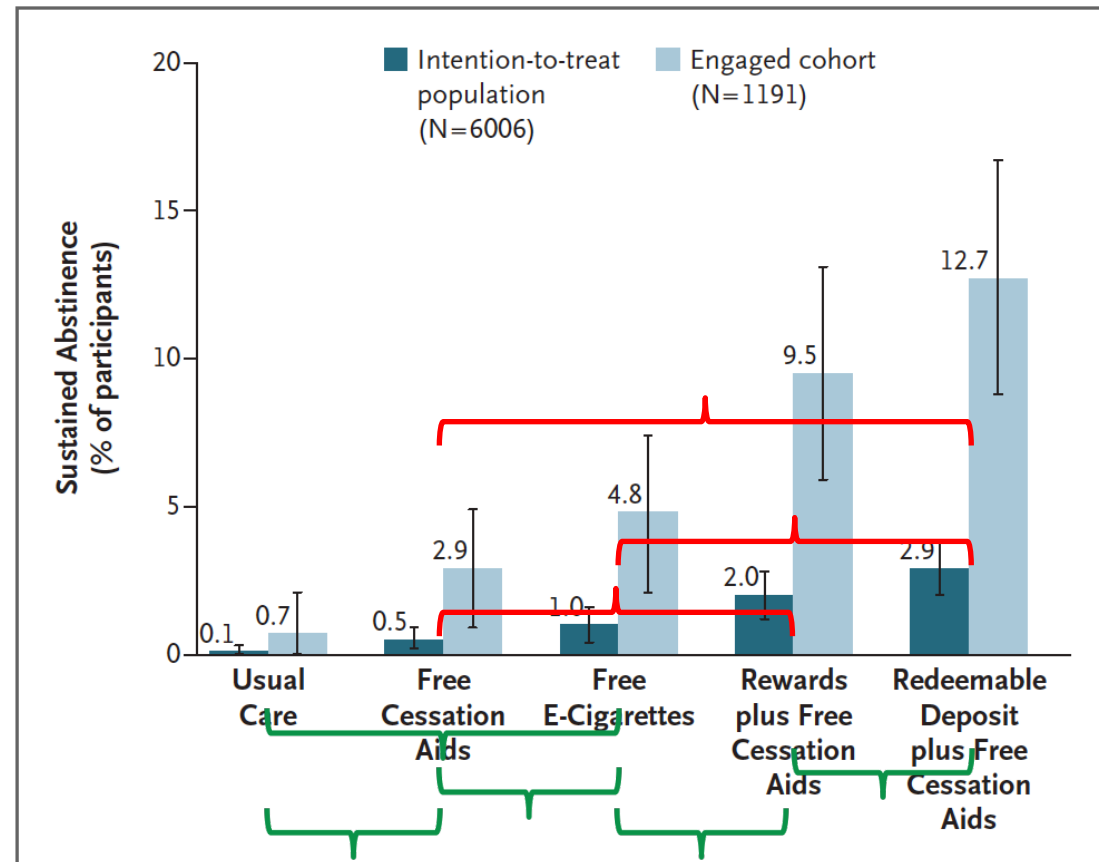


Figure 2. Sustained Smoking Abstinence at 6 Months after the Target Quit Date.

Answers to our questions

1. How successful are workplace smoking-cessation programs among all people to whom they are offered?

Not very, but cost-effective

2. How effective are incentives when added to free nicotine-replacement therapy and pharmacotherapy (bupropion or varenicline)?

Still triple quit rates

3. How effective are free e-cigarettes or free cessation aids when added to smoking cessation information without assistance on how to use?

Not effective

4. Do deposit contracts that are funded in advance without participant contributions, but from which money is removed if abstinence milestones are not met, achieve higher quit rates than reward incentives?

No

Discussion

The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

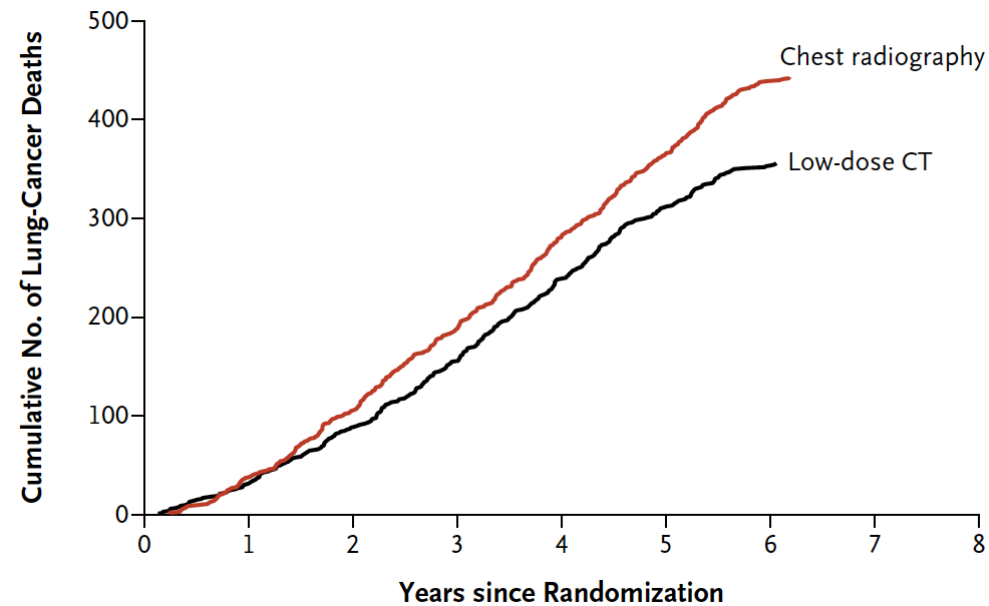
AUGUST 4, 2011

VOL. 365 NO. 5

Reduced Lung-Cancer Mortality with Low-Dose Computed Tomographic Screening

The National Lung Screening Trial Research Team*

B Death from Lung Cancer



Smoking cessation & lung cancer screening

1. USPSTF defines high risk as: 55-80 years old; 30+ pack-year history; active smoker or quit within past 15 years (2020 changes: 50-80 years, and 20+ pack-years)
2. Several studies estimate that 50% of patients meeting these criteria are actively smoking, and that there are ~ 5 million eligible active smokers in U.S.*
3. CMS began reimbursement for LDCT February 5, 2015
4. Requirements: (a) shared decision-making visit; (b) smoking cessation counseling
5. Lung cancer screening sites report: lack of patient interest; lack of staff training or time; complexities of reimbursement for smoking cessation services; lack of knowledge of what works best, let alone what is most cost-effective in this setting

Smoking cessation and SCALE collaboration

1. Patients who quit during LCS estimated to derive a 4-year increase in life expectancy
2. SCALE: NCI and VA fund 8 RCTs of smoking cessation interventions within LCS
3. Characteristics of 8 trials:
 - a. Six have 7 sites or fewer (max 26)
 - b. Sample sizes range from 500-1,650
 - c. All use traditional informed consent
 - d. Anticipated enrollment of 19% or fewer Black patients in 7 trials (37% in 1)
 - e. All test ask-advise-refer, behavioral counseling, and/or pharmacologic interventions
4. In light of evidence for above interventions, diverse stakeholder panel recommends testing “mobile health applications” and “financial incentives”

Eligible Population



Current smokers



Order for LDCT scan

- Ages ≥ 18 years



Underserved

- Black, or
- Hispanic, or
- Rural residence, or
- Low SES (no greater than HS education or 2x fed poverty)



In 4 Health Systems



Geisinger

4-arm Pragmatic Trial



1. Ask-advise-refer



2. *and* free medications



3. *and* money to quit



4. *and* episodic future thinking (EFT) tool

Discussion

ORIGINAL ARTICLE

Explanatory and Pragmatic Attitudes in Therapeutical Trials

Daniel Schwartz, Joseph Lellouch

Unité de Recherches Statistiques, Institut National de la Santé et de la Recherche Medicale, 94 Villejuif, France

The “comparison between two treatments” is a problem which is inadequately specified even in its over-all characteristics. It may imply one of at least two types of problem which are basically different.

The first type corresponds to an explanatory approach, aimed at *understanding*. It seeks to discover whether a difference exists between two treatments which are specified by strict and usually simple definitions. Their effects are assessed by bio-

The second type corresponds to a pragmatic approach, aimed at *decision*. It seeks to answer the question—which of the two treatments should we prefer? The definition of the treatments is flexible and usually complex; it takes account of auxiliary

Combining the ‘forward thinking’ of lung cancer screening programs and episodic future thinking to supercharge incentives in underserved populations

Figure 1: Conceptual model of barriers to smoking cessation that will be addressed by interventions in this RCT

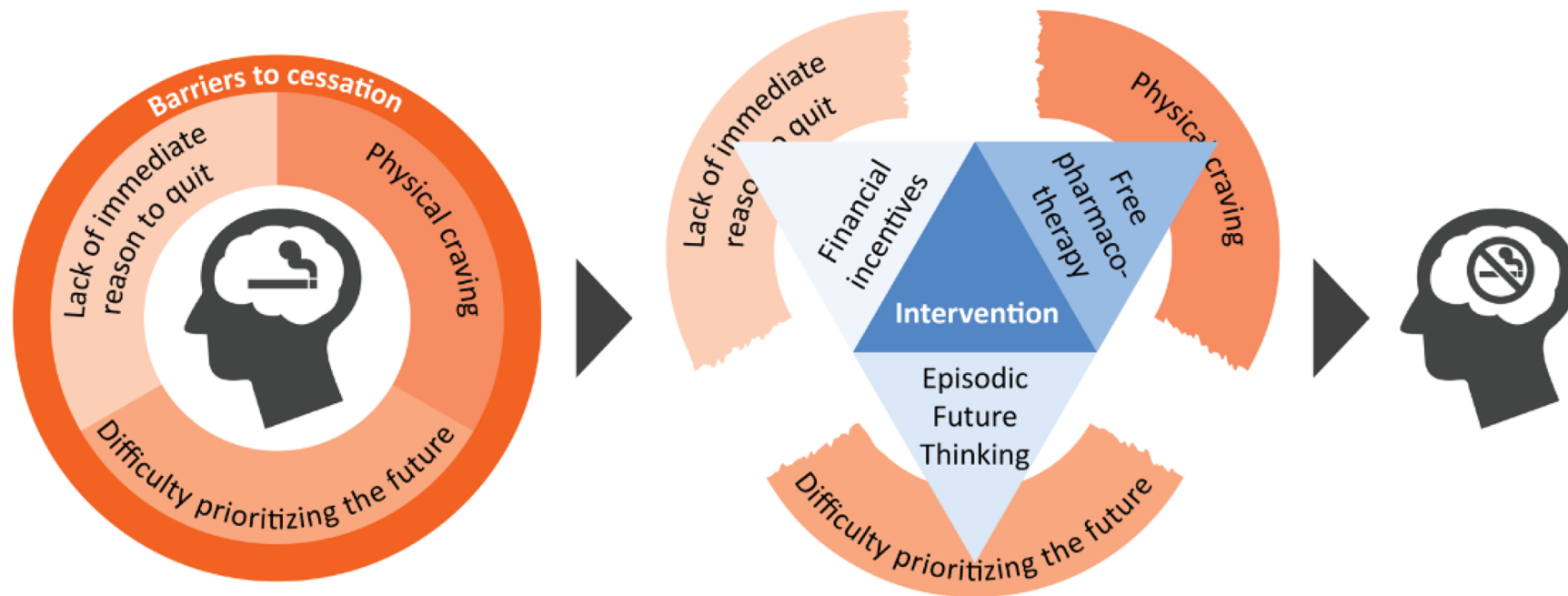
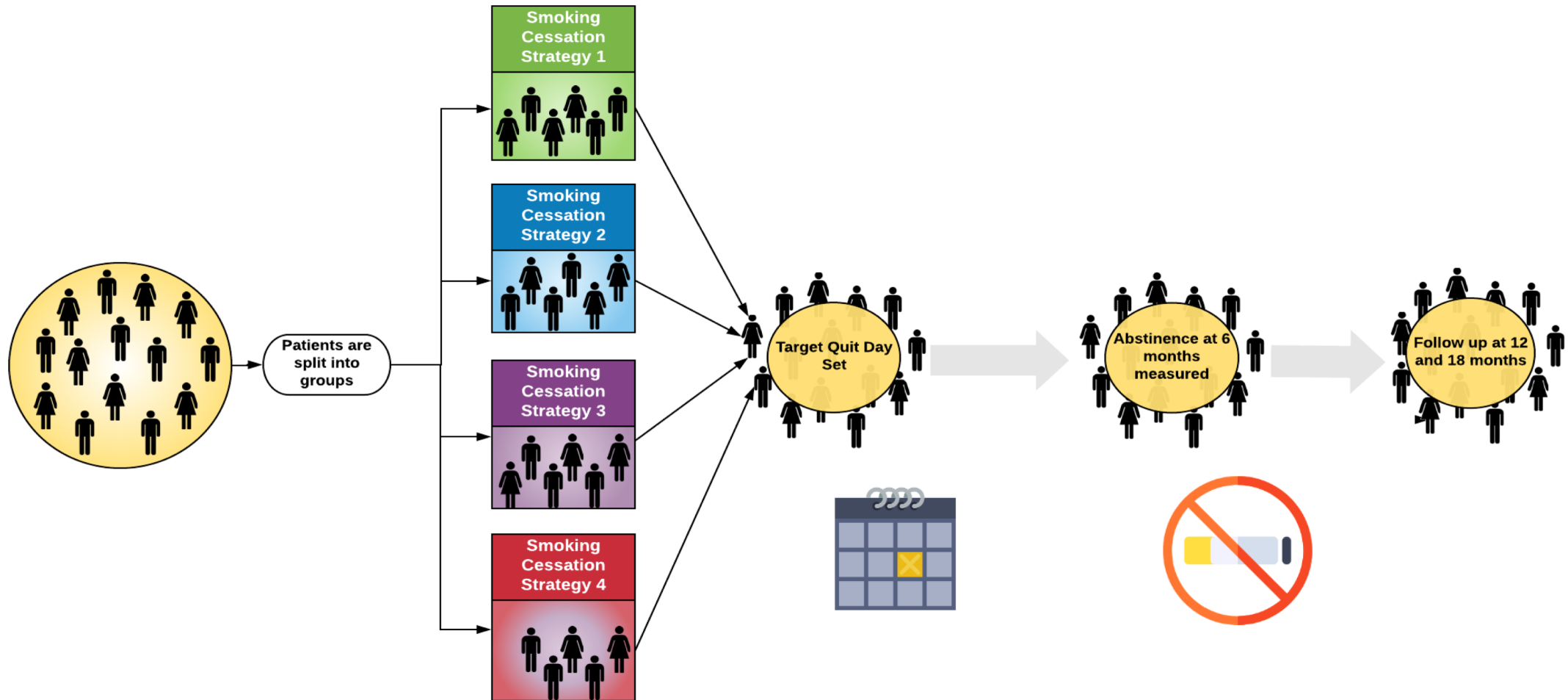


Table 3: Pragmatism of the proposed trial on PRECIS-2 criteria*

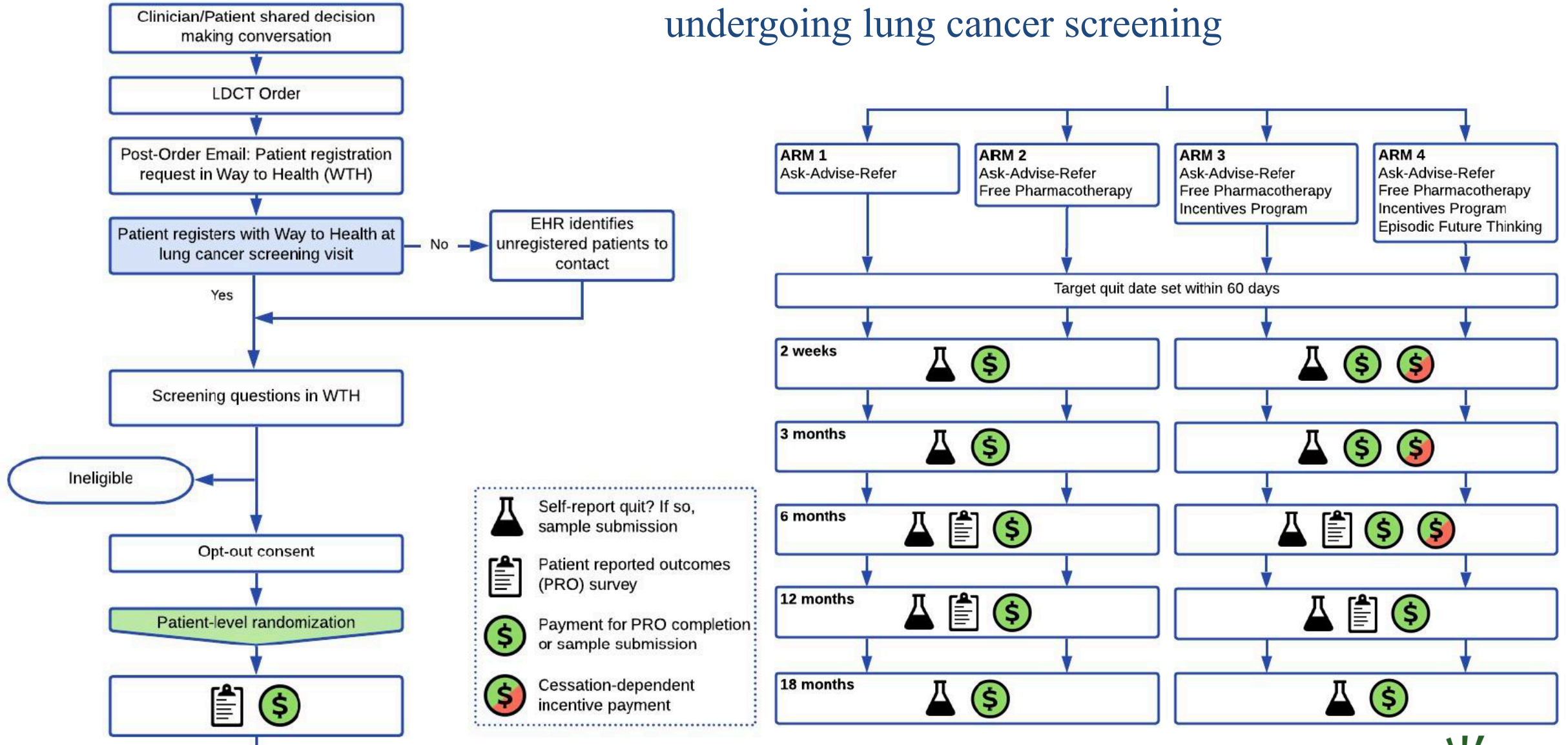
| <i>Domain</i> | <i>Relevant trial features</i> |
|---------------------------------|--|
| <i>Eligibility</i> | All underserved smokers getting lung-cancer screening |
| <i>Recruitment</i> | Built into routine workflow; use of opt-out consent |
| <i>Setting</i> | More than 30 screening centers within 4 large health systems |
| <i>Organization</i> | No clinician training required, minimal onsite research staff |
| <i>Flexibility of delivery</i> | Interventions delivered in ways compatible with usual care |
| <i>Flexibility of adherence</i> | Automated prompting of clinical staff to distribute iPads to promote enrollment and access to interventions |
| <i>Follow-up</i> | Outcomes data collected through automated, web-based research portal, including uploading of laboratory test results |
| <i>Primary outcome</i> | Most widely used outcome, of importance to all stakeholders |
| <i>Primary analysis</i> | Data available for all participants, intention-to-treat analyses |

*Criteria from Loudon et al. *The PRECIS-2 tool: designing trials that are fit for purpose. BMJ 2015*

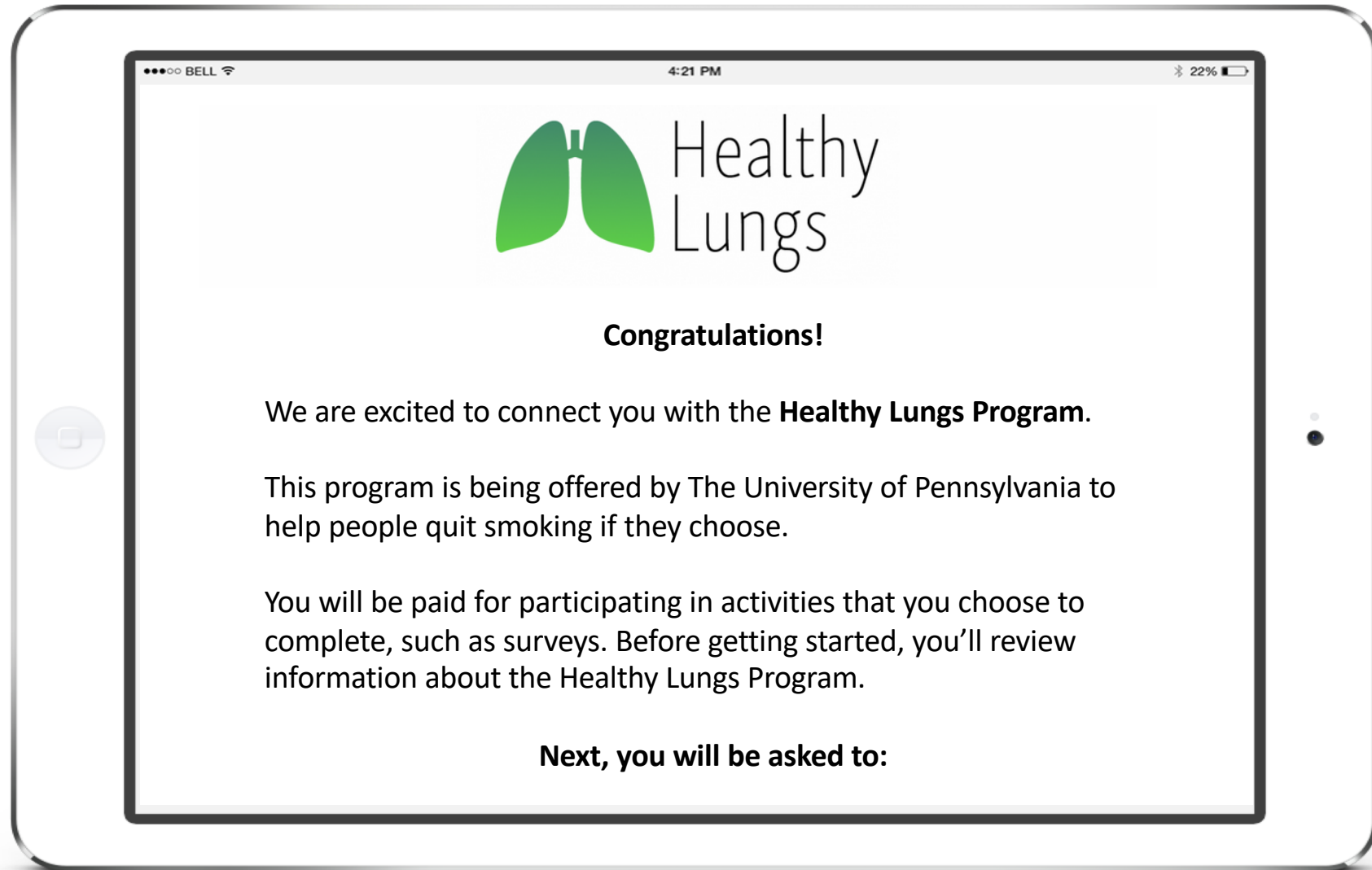
Simplified Patient Flow



Target sample size: 3,200 underserved smokers undergoing lung cancer screening



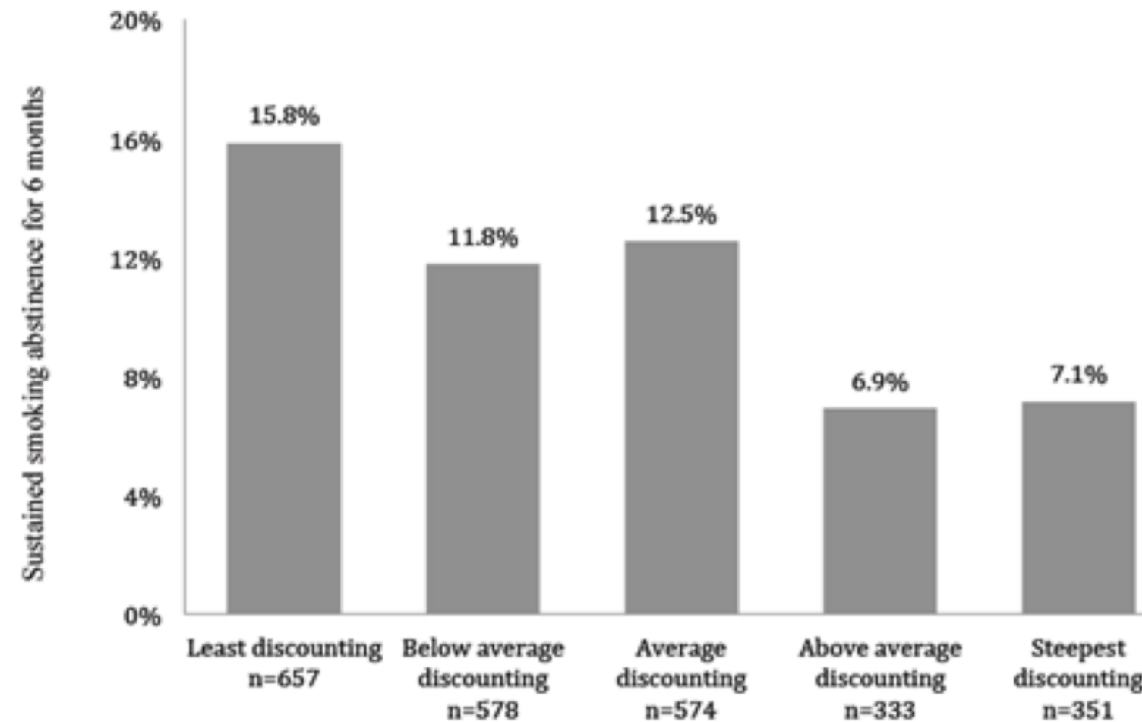
Patient Enrollment via iPad



Heterogeneity in the Effects of Reward- and Deposit-based Financial Incentives on Smoking Cessation

Scott D. Halpern^{1,2,3,4}, Benjamin French^{2,3}, Dylan S. Small^{2,5}, Kathryn Saulsgiver^{2,3}, Michael O. Harhay³, Janet Audrain-McGovern^{2,6}, George Loewenstein^{2,7}, David A. Asch^{1,2,4,8,9,10}, and Kevin G. Volpp^{1,2,4,8,9,10}

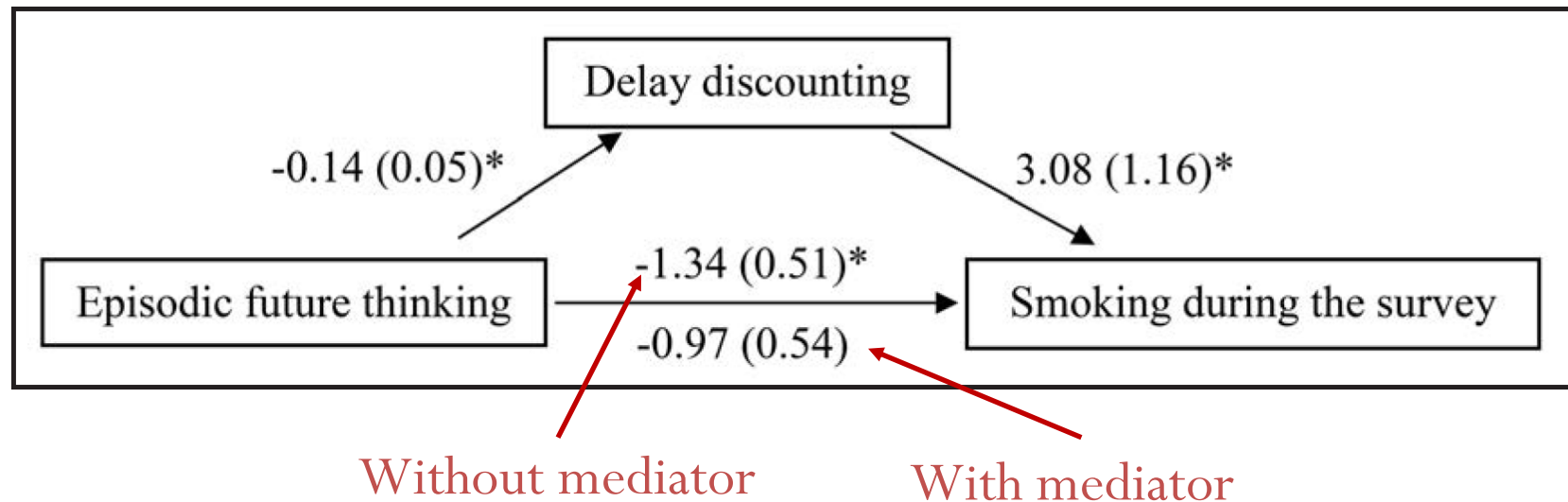
Steep temporal discounting reduces incentive effectiveness



Episodic Future Thinking Involving the Nonsmoking Self Can Induce Lower Discounting and Cigarette Consumption

WEN-BIN CHIOU, PH.D.,^{a,*} & WEN-HSIUNG WU, PH.D.^b

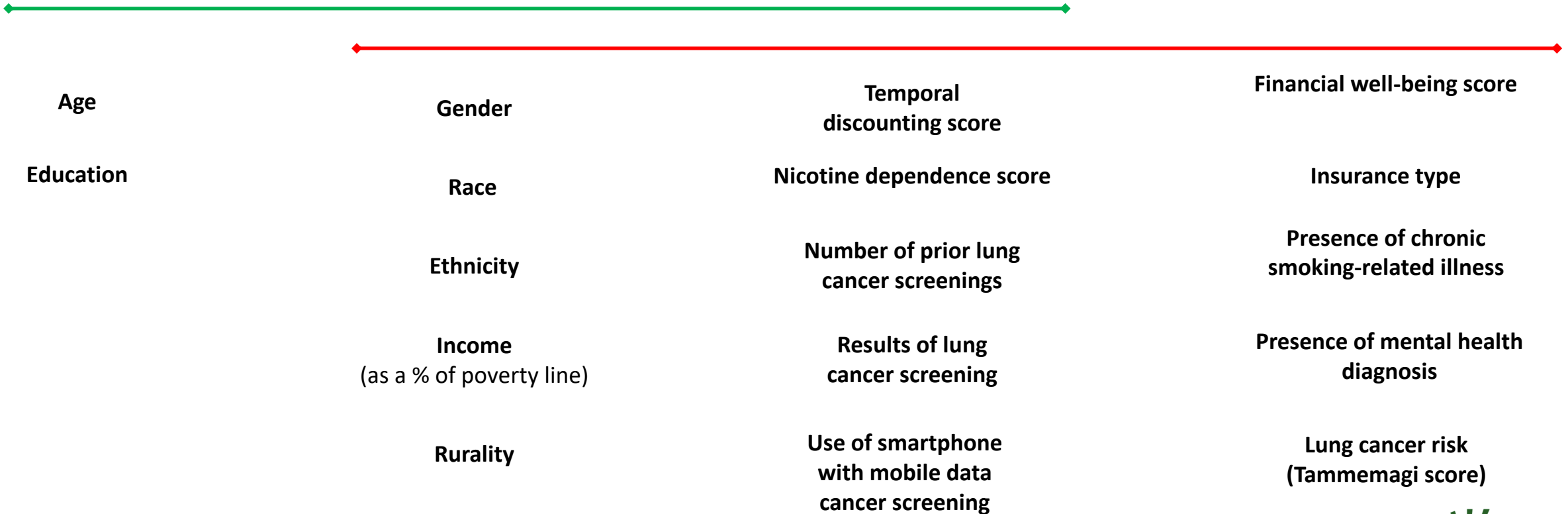
Delay discounting partially mediates relation between EFT and smoking



Proposed precision variables and effect modifiers

Precision variables

Test for effect modification



Discussion