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Immunogenicity and Smoking Cessation Outcomes for a Novel Nicotine Immunotherapeutic

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Abstract

NicVAX[®], a nicotine vaccine (3'AmNic-rEPA), has been clinically evaluated to determine if higher antibody concentrations are associated with higher smoking abstinence rates and if doses and frequency of administration are associated with increased antibody response. This randomized, double-blinded, placebo-controlled multicenter clinical trial (N=301 smokers) tested 200 and 400 µg doses administered 4 or 5 times over 6 months compared to placebo. 3'AmNic-rEPA recipients with the highest serum anti-nicotine antibody response (top 30% by AUC) were

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Conflict of interest/Disclosure

Ms. Akhavan, and Drs. Fahim, Kessler, Kalnik and Niknian are employees of Nabi Biopharmaceuticals and have received salary support, stock and options. All other authors were investigators on the clinical trial funded by NIDA and by Nabi Biopharmaceuticals and some served on an advisory panel. Other disclosures include the following: Dr. Douglas Jorenby has received research support from Pfizer. Dr. David Gonzales has received grant/research support from Pfizer, Addex Pharmaceuticals, Sanofi-Aventis and GlaxoSmithKline; consulting fees and honoraria from Pfizer, GlaxoSmithKline, and Evotech NeuroSciences; speakers fees from Pfizer; and owning five shares of Pfizer stock. Dr. Nancy Rigotti has received research grant support from Pfizer and is an unpaid consultant to Pfizer and Free & Clear. Dr. Elbert Glover has served as a speaker, consultant, grantee, provided advice or is on the advisory board/panel for Pfizer and served as a speaker for Nabi Biopharmaceuticals.

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significantly more likely to attain 8 weeks continuous abstinence from weeks 19 through 26 than the placebo recipients (24.6% vs. 12.0%, $p=0.024$, OR=2.69, 95% CI, 1.14–6.37). The 5 injection 400 µg dose regimen had the greatest antibody response and had significantly higher abstinence rates than placebo. This study demonstrates proof-of-concept that 3'AmNic-rEPA elicits antibodies to nicotine and is associated with higher continuous abstinence rates, justifying its further development as a treatment for nicotine dependence.

Keywords

3'AmNic-rEPA; NicVAX; nicotine immunotherapeutic; nicotine vaccine; cigarette; smoking cessation; antibody; *P. aeruginosa* r-Exoprotein A; aminomethyl nicotine; cotinine; CO

Introduction

Worldwide, smoking prevalence is 1.2 billion and approximately 5 million people die each year of smoking caused illnesses [1]. The global rate of smoking and smoking related deaths is anticipated to increase over the next 20 years unless significant public health measures are instituted. These include effective cessation interventions such as pharmacological treatments, which improve cessation rates by 1.5 to 3 fold over placebo intervention [2, 3]. Approved pharmacotherapies (e.g., nicotine replacements, bupropion SR, varenicline) for smoking cessation act on the central nervous system, each with a different mechanism of action. Other novel medications are being developed including immunotherapeutics targeting nicotine.

Nicotine conjugate vaccines stimulate the immune system to develop nicotine specific antibodies (Abs) using an immunogen comprised of nicotine covalently linked to a larger carrier protein. Conceptually, the mechanism of action is that anti-nicotine antibodies bind nicotine molecules and the resulting complex is too large to cross the blood-brain barrier. With increasing Ab levels more nicotine is captured and sequestered in the blood and prevented from entering the brain, leading to less reinforcing effects from nicotine. Animal studies have demonstrated that passive or active immunization results in approximately 30% to 90% less nicotine entering the brain compared to control rats [4–7] and attenuated locomotor [4, 5] and behavioral [8, 9] responses to nicotine. Furthermore, vaccination reduced nicotine elimination from the body in a study with rats [10, 11], which may also contribute to reduced smoking.

Although human studies are limited, published data evaluating different nicotine vaccines support the general concept that nicotine vaccines can be effective for smoking cessation in some smokers [12, 13]. Unfortunately, these studies either had small sample sizes [12], did not use an intent-to-treat population of smokers [13] or did not perform statistical analysis of the data [14].

The primary aim of the present study was to establish the proof-of-concept that (i) anti-nicotine antibodies are useful as an aid to smoking cessation and (ii) higher serum anti-nicotine antibody concentrations are associated with higher abstinence rates in an intent-to-treat population of smokers. One of the challenges with immunotherapeutics, such as

vaccines, is attainment of therapeutic levels of antibody in most people. Therefore, this study tested two different doses of 3'-aminomethylnicotine *P. aeruginosa* r-exoprotein A - NicVAX (3'AmNic-rEPA) to identify a dose and schedule for further development: 200 and 400 µg across two different schedules (4 or 5 injections) compared to placebo for immunogenicity, efficacy and safety.

Results

A total of 301 subjects were randomized. Figure 2 shows the disposition and number of subjects within each treatment group. No significant group differences were observed in the demographic or smoking history by treatment or antibody levels (see Table 1).

Compliance

All 301 subjects received injection 1, 96.7%, 84.1%, 72.4% and 61.2% received injections 2, 3, 4 and 5 (Schedule 2 subjects only), respectively. No significant differences were observed across treatment groups for subjects receiving injections 2 through 4 for Schedules 1 and 2. Mean in-study duration was 286 ± 121 days.

Proof-of-concept

Effects of high Ab on abstinence—High Ab responders to 3'AmNic-rEPA were defined as the top 30% of responders by AUC (0 to 26 weeks) and the low Ab group as the remaining bottom 70% of responders. 3'AmNic-rEPA recipients in the high Ab group were significantly more likely to attain 8 weeks of continuous abstinence from weeks 19 through 26 than placebo (24.6% vs. 12.0%, $p=0.024$, OR=2.69, 95% CI, 1.14–6.37). No significant differences were observed between the 3'AmNic-rEPA low Ab group vs. placebo (9.3% vs. 12.0%, $p=0.46$). As a secondary outcome, continuous abstinence rate (CAR) to 52 weeks were evaluated from weeks 19 to 52 and were significantly higher for the high Ab group vs. placebo (19.7% vs. 10.0%, $p=0.044$, OR=2.64, 95% CI, 1.03–6.79) with no significant difference between the low Ab group and placebo (7.1% vs. 10.0%, $p=0.43$). 7-day point prevalence abstinence results show that subjects with high Ab levels were significantly more likely to abstain compared to placebo at 26 weeks (36.1% vs. 16.0%, $p=0.0024$, OR=3.30, 95% CI, 1.53–7.13) and 52 weeks (31.1% vs. 12.0%, $p=0.0021$, OR=3.69, 95% CI, 0.42–2.14). No significant differences were observed in the point prevalence abstinence rates between the low Ab group and placebo at 26 and 52 weeks (12.9% vs. 16.0%, $p=0.51$ and 11.4% vs. 12.0%, $p=0.89$, respectively). As shown in Figure 3A, abstinence rates remained essentially the same following the target quit date (TQD) for the duration of the study.

To further validate the proof-of-concept, the relationship between abstinence during the final eight weeks of the study and anti-nicotine Ab concentrations (AUC) is shown in Figure 3B for all subjects receiving the vaccine. Continuous abstinence rates from week 45–52 are displayed for each 10th percentile increase in AUC. The proportion of abstinent subjects increased with increasing AUC percentile, and the ordered ranking maintained.

Effects of high Ab on time to continuous abstinence—An exploratory analysis examined the rate and time to continuous abstinence through study end (Figure 3C). Most

smokers quit soon after the TQD, with a clear divergence of the high Ab group from the placebo and low Ab group. Among the 18 high Ab continuous abstainers, 12 initiated abstinence prior to the primary endpoint while 6 initiated their abstinence after the start of the primary endpoint. Furthermore, 3 of the 15 subjects who abstained during the primary endpoint relapsed by the end of the study. Cox proportional hazards analysis demonstrated superiority of the high Ab group compared to placebo ($p=0.0069$, hazard ratio of 2.76).

Evaluation of long-term abstinence—As most subjects achieved abstinence shortly after their TQD, additional analyses were undertaken to evaluate prolonged abstinence to 6 and 12 months allowing for a 2 week grace period after the TQD [15]. (Prolonged abstinence is defined as not a single puff during the period from 2 weeks after the TQD for 20 weeks and 44 weeks, respectively).

Prolonged abstinence rates to 6 months were significantly higher in the high Ab group vs. placebo (19.7% vs. 6.0%, $p=0.0060$, OR=4.41, 95% CI, 1.53–12.71) with no significant differences between the placebo and low Ab groups (7.9% vs. 6.0%, $p=0.60$). Subjects with high Ab were also significantly more likely to be abstinent for 12 months compared to placebo (18.0% vs. 6.0%, $p=0.014$; OR of 3.84; 95% CI, 1.32–11.20). The low Ab group did not differ significantly from placebo (7.1% vs. 6.0%, $p=0.67$).

Cigarette smoking in non-quitters—Statistically significant differences were observed in reduction of daily cigarette consumption and cotinine between non-abstainers (weeks 19–52) with high Ab levels and non-abstainers in the placebo group ($p=0.0015$ and 0.019, respectively; see Figures 4A and 4C). The difference in the median reduction in cigarette consumption, following the TQD, between the high Ab non-abstainers and placebo non-abstainers was on average 4.6 cigarettes per day. Similarly, cotinine geometric mean concentrations (GMCs) were 19.0% lower on average following the target quit date in the high Ab non-abstainers as compared to placebo non-abstainers. Median cigarettes per day and cotinine GMC for placebo and low Ab group are nearly identical over the study period. Differences in mean CO were not observed across all three groups (see Figure 4B).

15 out of the 301 subjects smoked more than a 2-fold higher number of cigarettes following the TQD as compared to baseline with no significant difference between the placebo ($n=4/100$) and 3'AmNic-rEPA ($n=11/201$) groups. The highest smoking levels observed post-TQD were 5-fold baseline in the placebo and 4-fold baseline in the 3'AmNic-rEPA groups. Elevated smoking levels for 11 of these 15 subjects returned to below 2-fold baseline by study end. Subjects with remaining elevated levels were similar between the 3'AmNic-rEPA group ($n=3/201$) and placebo ($n=1/100$). Individual subjects with CO levels elevated by 2-fold or higher than baseline were also assessed and results were similar with no significant difference between the placebo ($n=5/100$) and 3'AmNic-rEPA ($n=13/201$) groups.

Withdrawal symptoms—Significant differences were not observed in overall withdrawal severity between placebo and the high and low Ab groups ($p > 0.22$).

Effects of dose and schedule

Immunogenicity and Efficacy by Study Group—Figure 5 depicts immune response by study group from baseline to week 52. Anti-nicotine Ab GMCs increased across all active treatment groups after each vaccination, with each subsequent dose resulting in a higher Ab response than the previous dose. Schedule 2 showed a higher initial increase in Ab concentrations. The 400 µg/Schedule 2 group demonstrated the highest Ab concentrations. However, no significant differences ($p > 0.05$) were observed in AUC, C_{\max} and C_{avg} across the treatment groups to 26 or 52 weeks likely because the study was not powered to detect such difference.

An intent-to-treat analysis demonstrated that the Schedule 2, 400 µg dose group had significantly higher prolonged abstinence to 6 months as compared to placebo (17.6% vs. 6.0%; $p=0.015$; OR of 4.14; 95% CI, 1.32–13.02), although not significant for the Schedule 2, 200 µg dose (14.0% vs. 6.0%, $p=0.054$; OR=3.23; 95% CI, 0.98–10.67) or between placebo and Schedule 1 for each of the doses ($p > 0.84$). The Schedule 2, 400 µg dose group also had the highest rates of prolonged abstinence to 12-months, significantly higher than placebo (15.7% vs. 6.0%, $p=0.038$; OR=3.44; 95% CI, 1.07–11.04), but not significant for the Schedule 2, 200 µg dose (14.0% vs. 6.0% for 200 ($p=0.056$; OR=3.21; 95% CI, 0.97–10.63) or between placebo and Schedule 1 for each dose ($p > 0.88$).

Safety—Table 2 lists the number of subjects experiencing local and systemic reactogenicity. Reactogenicity events were aggregated over all injections. Overall, ache and tenderness were the most commonly reported local events, with at least one report by nearly all subjects (86% to 98%) in each treatment group. Myalgia, malaise, and headache were the most commonly reported systemic events (64% to 88% of subjects). Swelling, heat, burning, erythema, and nausea were reported by about half the subjects. Fever and vomiting were less common (4% to 16%).

A total of 1184 treatment-emergent adverse events (AEs), predominantly rated mild or moderate, were reported by 266 of 301 subjects; 87.1% of 3'AmNic-rEPA recipients and 91.0% of placebo recipients. On average, 3.7 and 4.3 events were observed per person in the vaccinated and placebo groups, respectively, including subjects with no events. The distribution of 161 physician-determined treatment-related AEs, according to severity and relationship to treatment, was similar for the 3'AmNic-rEPA and placebo arms. Seven 3'AmNic-rEPA recipients (3.5%) and 2 subjects in the placebo arm (2.0%) withdrew from the study due to adverse events.

18 Serious AEs were reported: 8 events in the 3'AmNic-rEPA treatment groups among 7 subjects (3.5% of the 3'AmNic-rEPA recipients) and 10 events in the placebo group among 5 subjects (5.0% of the placebo recipients). Only one of these SAEs (anaphylactic reaction in a 3'AmNic-rEPA 400 µg/Schedule 2 recipient) was considered by the Investigator to be treatment related. This subject, who had a history of urticaria to penicillin and seasonal allergies, experienced difficulty breathing, throat tightness, facial erythema and urticaria 70 minutes after the initial vaccination. The subject was treated with a single injection of subcutaneous epinephrine and diphenhydramine, which resolved the symptoms. Herpes Zoster was reported in 6 subjects; one occurring within 3–5 hours after the first vaccination

which would be impossible to link to vaccination. Of the remaining cases, 4 (2%) occurred in 3'AmNic-rEPA recipients; while 1 occurred in placebo (1%). In contrast, the related herpes virus, simplex infection was reported in 4 3'AmNic-rEPA recipients (1%) and 3 placebo recipients (2%).

Discussion

Results demonstrated the proof-of-concept that smokers who achieved higher anti-nicotine Ab concentrations were more likely to quit and remain abstinent from smoking. The high Ab group demonstrated the highest abstinence rates independent of the time period of ascertainment. Similarly, in a separate study conducted by Cornuz and coworkers [13] testing a different nicotine vaccine (nicotine derivative conjugated to a virus-like particle derived from bacteriophage Q β), *post hoc* analysis showed subjects stratified to the highest Ab group had a significantly higher quit rate than placebo. However, unlike the current study, which used the intent-to-treat (ITT) population to establish proof of concept, the reported finding by Cornuz et al. [13] was observed after eliminating about a third of the subjects who used nicotine replacement therapies during the course of the study or who had incomplete Ab titer values.

In the present 3'AmNic study, subjects in the high Ab group had observed odds ratios of 4.4 (95% CI 1.5–12.7) and 3.9 (95% CI 1.3–11.2) for prolonged abstinence rates to 6 and 12 months versus placebo. Although no direct comparisons can be made, these odds ratios are not unlike ones observed in the meta-analyses conducted for the U.S. Clinical Practice Guideline where the reported odds-ratios ranged from 1.5 (95% CI 1.2–1.7) for nicotine gum to 3.1 (95% CI 2.5–3.8) for varenicline at 6 months post-quit [3]. If current study findings are confirmed in larger studies, immunotherapeutics are likely to emerge as an important aid to smoking cessation.

No significant compensatory smoking in response to the presence of anti-nicotine antibodies, as determined by the number of cigarettes smoked per day, carbon monoxide levels or cotinine levels, was observed in this study. This result is consistent with observations from other studies [12, 13]. In the current study, subjects in the high Ab group who did not abstain smoked significantly lower number of cigarettes (median reduction of ~5 cigarettes/day) and experienced lower cotinine levels (~20%) than placebo subjects along with no differences observed between the low Ab and placebo groups. However, a small number of individual subjects (N=15/301) smoked more than 2-fold the number of cigarettes at baseline with no significant differences between the active treatment and placebo groups.

A major challenge for immunotherapeutics is to stimulate high Ab levels in the vast majority of smokers trying to quit. Vaccine dose and frequency of vaccination have an impact on the Ab levels attained. The 5 injection/400 μ g dose was associated with the highest Ab response, although not statistically significant, possibly due to the small sample size. Importantly, this dose and schedule demonstrated statistically significant improved 44-week CAR compared to placebo. Because the 4 injection/400 μ g dose was not associated with higher abstinence rates, this result demonstrates that consideration of both dose and schedule of injection are critical. In an independent, follow-up immunogenicity study to examine if peak Ab levels

could be elevated, a total of 74 subjects received 6 injections of 400 µg 3'AmNic-rEPA at week 0, 4, 8, 12, 16, and 26. More than 80% of subjects receiving the 6-dose immunization regimen exceeded a target level of Ab (25 µg/ml) by week 14. In contrast, only 50% of subjects receiving 5 injections of the 400 µg dose achieved this level by week 14 in the current study and, only 7% of subjects attained this level by the target quit date. This finding suggests that more frequent injections and a later quit date may increase treatment success.

In general, 3'AmNic-rEPA was well-tolerated. The frequencies of local and systemic vaccine reactogenicity events were similar in the vaccine and placebo groups and similar to licensed adult vaccines containing Alum adjuvant [16]. The slight increase in cases of *herpes zoster* observed in this study may be spurious, but continued monitoring is necessary to determine if a causal relationship exists. The occurrence of a single anaphylactic reaction would suggest a need for continued monitoring and follow-up, even considering the subjects' history of prior drug allergy to penicillin.¹

In summary, results from this study support the concept that high anti-nicotine Ab levels are associated with higher rates of abstinence. These findings suggest that vaccines attaining high Ab levels by the target quit date may be more effective. Other future strategies may include examining additional ways to increase Ab levels across all individuals. Nonetheless, this study demonstrates that 3'AmNic-rEPA has significant potential as a smoking cessation, and perhaps a relapse prevention aid.

Methods

Study Population

Smokers were recruited via advertisement across nine geographically diverse U.S. sites. Interested subjects were screened over the telephone and more extensively screened at the first Screening Visit. Subject informed consent was obtained prior to screening. Subjects were 18 years of age or older, smoked ≥15 cigarettes/day, had exhaled CO ≥10 ppm; wanted to quit smoking, and were in good general physical and mental health. Exclusion criteria included recent use of any medications or drugs that might interfere with immune response, interact with the vaccine, and pharmacotherapies or other treatments for smoking cessation. For females, a negative urine pregnancy test at enrollment and active use of acceptable birth control or documentation of surgical sterility was required.

Study Design

This Phase 2 study was a randomized double-blind, placebo-controlled, parallel-arm trial design (see Figure 1). Four treatment groups varied the dose and/or schedule of intramuscular vaccination: 200 or 400 µg of 3'AmNic-rEPA or placebo according to Schedule 1 (weeks 0, 6, 12, and 26) or Schedule 2 (weeks 0, 4, 8, 16, and 26). Subjects (N=150) were first randomized within Schedule 1 groups in a 1:1:1 ratio (200 µg: 400 µg: placebo) and then 151 subjects randomized within Schedule 2 groups in the same ratio. The

¹In subsequent completed and on-going clinical studies comparing 3'-AmNic-rEPA and placebo, over 1800 subjects have received 1–6 vaccinations with 400 µg 3'AmNic-rEPA or placebo. Only two additional cases of *herpes zoster* have been observed. Moreover, no additional anaphylactic/anaphylactoid-type SAEs have been reported to date.

TQD was 1 week after the second injection (end of week 7 for Schedule 1 and end of week 5 for Schedule 2). If the subjects relapsed (seven consecutive days of smoking) after the quit date, a second quit date coinciding with a future clinic visit was allowed between time of relapse and week 18. Cessation counseling (based on the USDHHS Clinical Practice Guidelines [17]) for the first quit attempt involved 5 standardized face-to-face sessions (10 minutes) and for the second quit attempt, a face-to-face session plus 2 post-quit telephone counseling sessions.

Subjects were followed for 52 weeks after randomization and the first injection on Day 0 for a total of 21 visits. During the injection day, subjects remained at the study site for 30 to 60 minutes for observation and attended a visit 24 hours after each injection to assess side effects. Otherwise, visits ranged from weekly to bi-weekly and were less frequent after later injections.

Subjects who failed to quit smoking on their TQDs were encouraged to remain in the study and continue to attempt to achieve abstinence. Subjects who terminated from the study were not replaced and presumed to be smokers.

Institutional Review Board approval was obtained from all institutions involved in the study. A Data and Safety Monitoring Board (DSMB) met four times during the study.

Investigational Product

The active investigational product was purified 3'-aminomethylnicotine conjugated to *P. aeruginosa* r-exoprotein A. For the 200 and 400 µg/mL dose, each single-use syringe contained 3'-aminomethylnicotine conjugated to 200 or 400 µg rEPA, respectively, adsorbed to 1.2 mg aluminum hydroxide adjuvant (Alhydrogel 85) in 1 mL phosphate buffered saline (0.15 M NaCl, 0.002 M NaPO₄, pH 7.2, 0.01% polysorbate 80; PBS). For the placebo dose, PBS with 1.2 mg Alhydrogel 85 was included in a 1 mL single-use syringe.

Measures

Cigarette use was recorded daily on an electronic diary for 182 days, and then weekly for the remainder of the study. Exhaled CO and urine cotinine were measured at each study visit, except for visits within 24 hours of vaccination. Questionnaires were collected via electronic diary: Fagerström Test for Nicotine Dependence [18] (administered days 0, 210, 364), Minnesota Nicotine Withdrawal Scale [MNWS, 19] (administered weekly until Month 6) and data on other tobacco usage.

Sera were collected for immunogenicity measurements at 16–17 time points (schedule-dependent) from baseline to week 52. Anti-nicotine antibody concentrations were measured using ELISA (Enzyme-linked Immunosorbent Assay) [12]. Subjects recorded standard local and systemic reactogenicity events for 7 days after each injection. All reactogenicity events were followed until resolution or study completion. Treatment emergent AEs were recorded until 4 weeks after the last dose, with the exception of SAEs, which were collected through week 52. Subjects were also periodically monitored at clinic visits for vital signs, weight, hematology, chemistry and urinalysis.

Statistical Analysis

The Intent-to-Treat (ITT) population was used for evaluation of efficacy, safety and immunogenicity. The ITT population was defined as all subjects who were randomized to treatment.

The primary endpoint was continuous smoking abstinence for a total of 8 weeks measured from the beginning of week 19 to the end of week 26 (determined from subject diaries confirmed by exhaled CO \leq 8 ppm). The analysis for proof-of-concept stratified the active-treatment recipients into high (top 30% AUC from weeks 0 to 26) and low (bottom 70% AUC from weeks 0 to 26) Ab responders, regardless of treatment group. An a priori decision was made to establish the antibody cutoff between 50% and 25%. Top 30% by AUC group was selected as the largest group of high antibody responders between the 25% and 50% cutoffs that demonstrated statistical significance as compared to placebo. Smoking outcomes were compared between subjects with high Ab and pooled placebo recipients using logistic regression.

Secondary aims of this study were to assess a) 7-day point prevalence abstinence at various times, CARs during 52 weeks and time to sustained abstinence defined as attaining 8 weeks of continuous abstinence at anytime prior to week 46 *and* maintaining continuous abstinence through 52 weeks; b) impact on compensatory smoking amongst non-abstainers; c) withdrawal symptoms and d) immunogenicity, efficacy and safety of administration of either 4 or 5 of the 200 and 400 μ g doses.

Secondary smoking cessation analyses used logistic regression for binary outcomes and Cox proportional hazards regression models and log-rank tests for time-to-sustained abstinence analyses. Mixed-effects repeated-measures analyses of the number of cigarettes smoked, CO and cotinine adjusted for baseline were utilized in assessing compensatory smoking among non-abstainers or in assessing withdrawal symptoms.

Anti-nicotine Ab responses were summarized as GMC with 95% confidence intervals. Safety was assessed throughout this study primarily using reactogenicity and adverse events (AEs). Reactogenicity data for 7 days after each injection were tabulated, and the proportions of subjects with any post-vaccination reactogenicity, aggregated over all injections, among the five treatment groups were compared using the Generalized Cochran-Mantel-Haenszel test.

When a subject dropped out of the study, they were assigned presumed to be smokers. Otherwise, all missing diary data related to cigarette use were imputed utilizing the Last Observation Carried Forward principle. No imputation was carried out for CO level. The missing serology data were imputed by first defining a set of injection windows for each schedule. The missing serology was imputed by using the next available measured serology in its corresponding window; if the next value was not available the value of the nearest previous time point in that window was used. The AUC for anti-nicotine Ab was calculated based on imputed data. For Ab values one week after the target quit date, the value at week 9 was used for Schedule 1 as sera were not collected at week 8.

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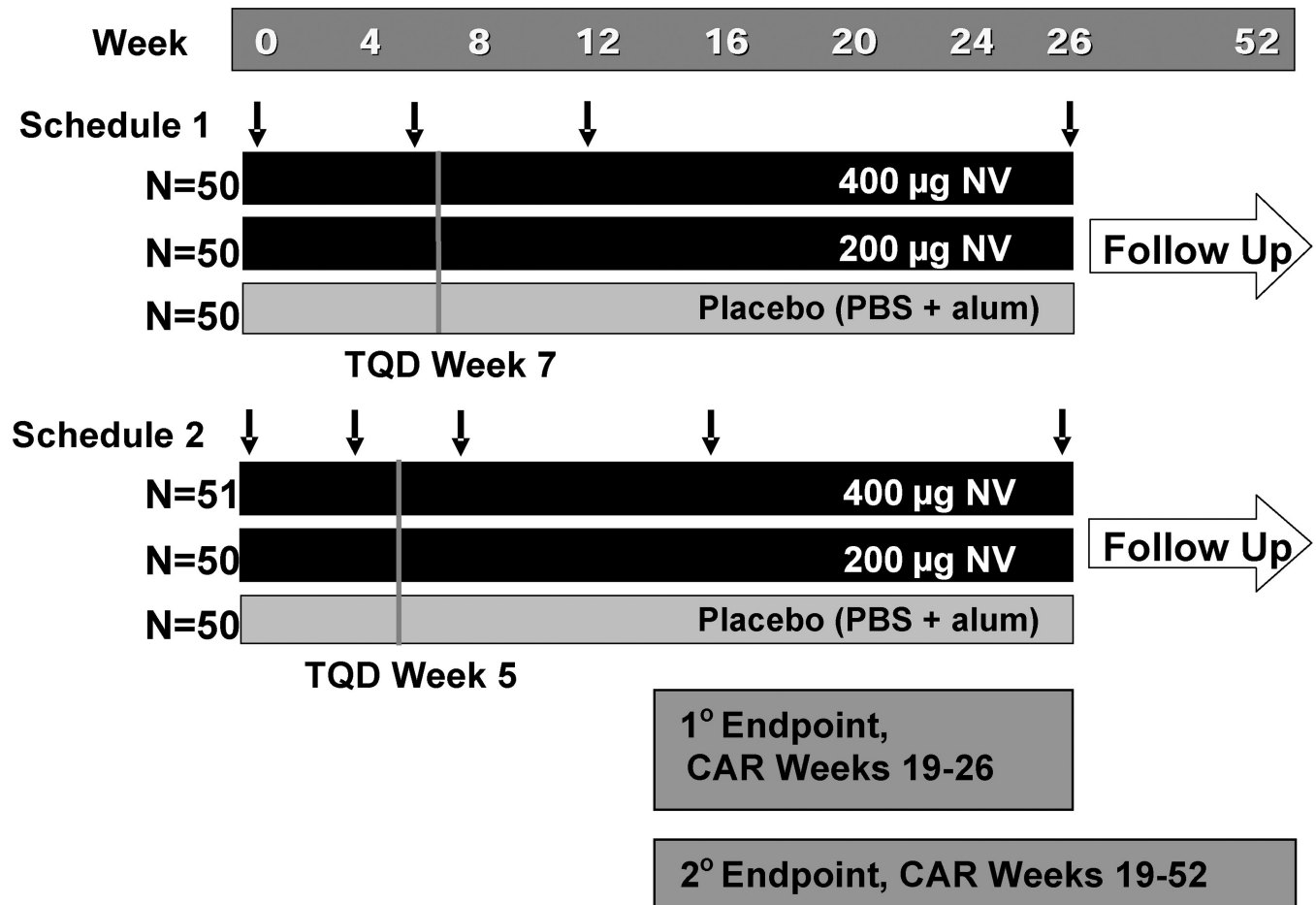


Figure 1.

Study design. Arrows denote timing of vaccinations for Schedule 1 (Week 0, 6, 12 and 26) and Schedule 2 (Week 0, 4, 8, 16, and 26). Primary endpoint (percent of subjects abstinent Week 19–26) and secondary endpoint (percent of subjects abstinent Week 19–52) are shown. NV (3'AmNic-rEPA); PBS (Phosphate buffered saline); Alum (aluminum hydroxide adjuvant); TQD (Target quit date).

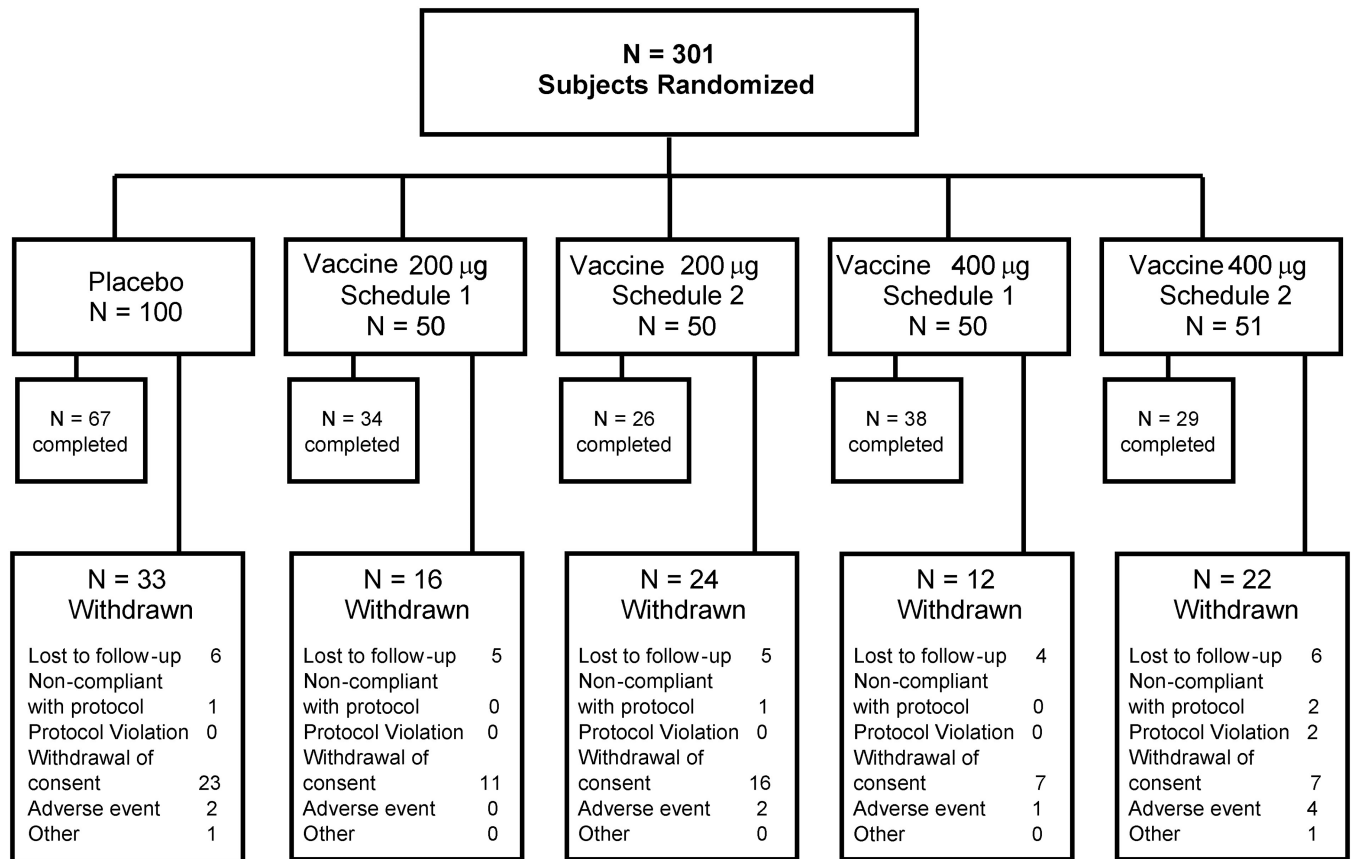
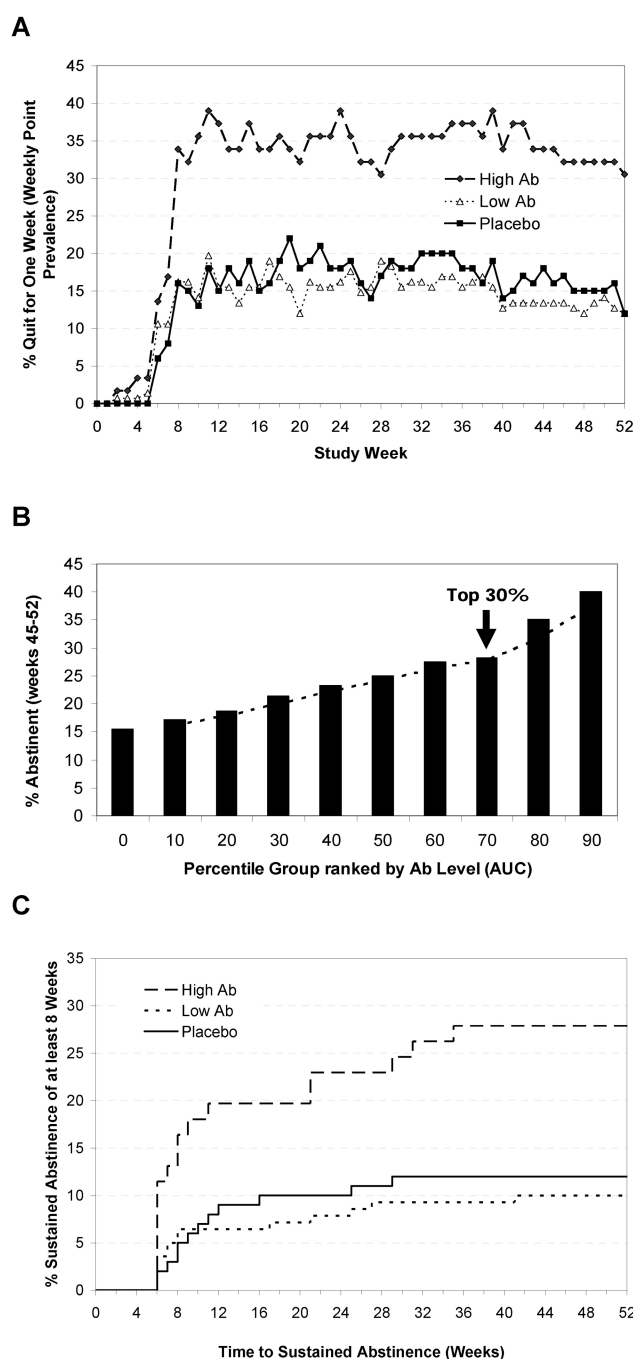


Figure 2.
Subject disposition.

**Figure 3.**

For the intent-to-treat population: (A) 7-day point prevalence abstinence rates for high antibody (top 30% AUC), low antibody (bottom 70% AUC) groups and placebo over the course of 52 weeks; (B) Percent abstinent for week 45–52 (8 week continuous abstinence) by AUC. AUC is displayed in 10th percentile point increments; (C) Time to 8-week of sustained abstinence prior to week 46 and continuous abstinence maintained through week 52, stratified by group with high antibody (top 30% AUC), low antibody (bottom 70% AUC) and placebo (dropouts censored at week 52).

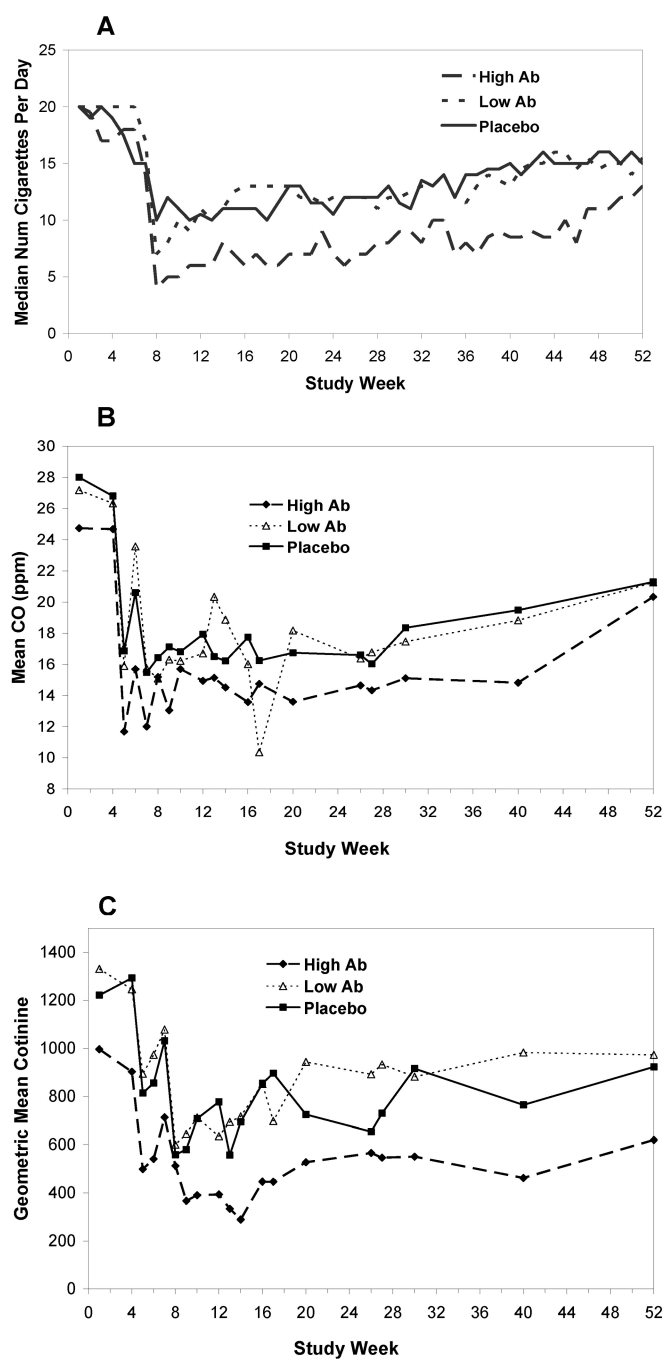


Figure 4.

(A) Median number of cigarettes per day, (B) CO levels and (C) geometric mean cotinine levels among those subjects not abstaining from cigarettes across those subjects with high antibody (top 30% AUC), low antibody (bottom 70% AUC) and placebo. Time on the x-axis is adjusted to align the target quit date between Schedule 1 and Schedule 2.

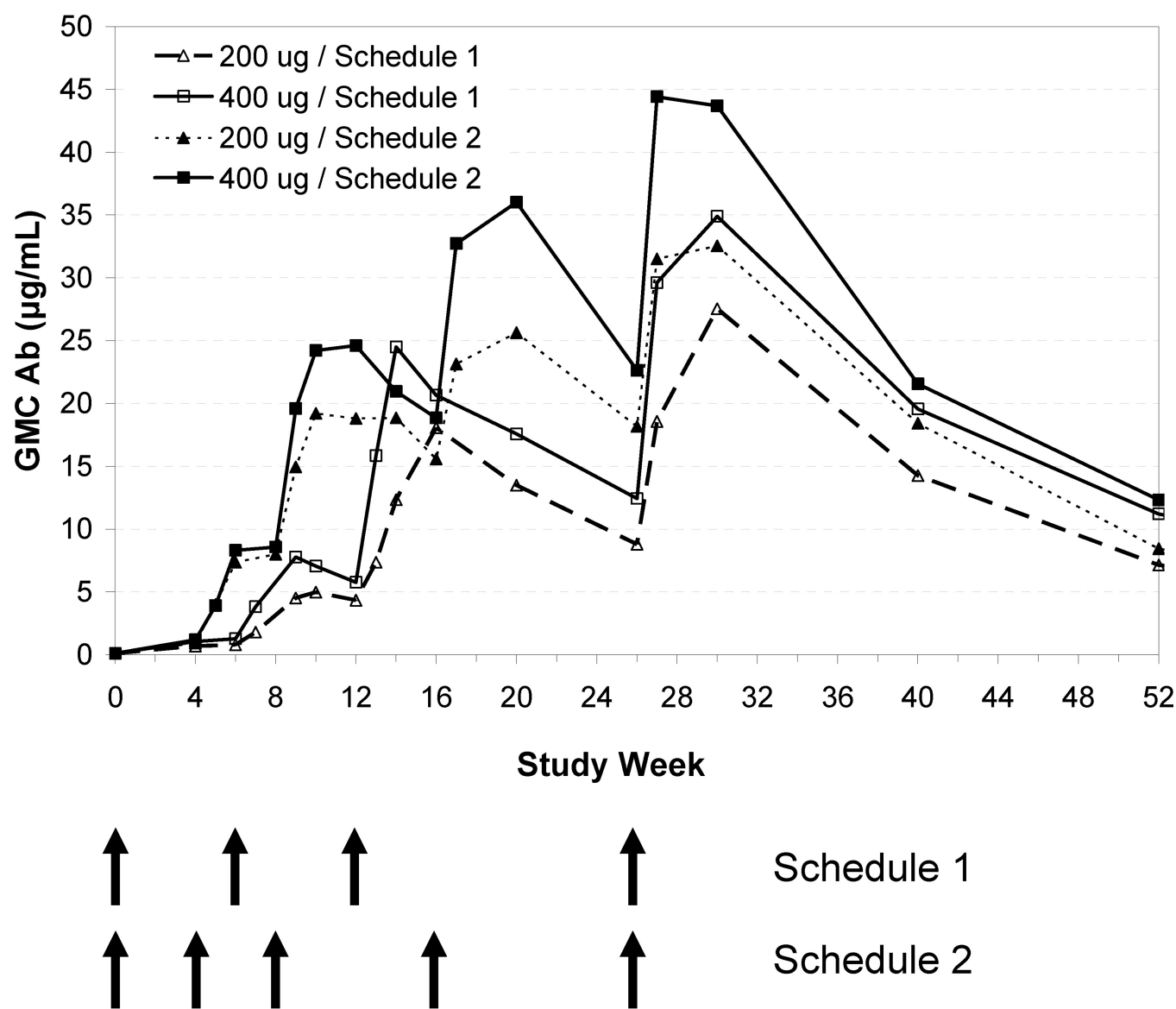


Figure 5.
Geometric mean antibody concentrations ($\mu\text{g/mL}$) by treatment group.

Table 1

Demographics and Smoking History at Baseline

Variables	Placebo N=100	Vaccine 200 µg Schedule 1 N=49-50	Vaccine 200 µg Schedule 2 N=49-50	Vaccine 400 µg Schedule 1 N=50	Vaccine 400 µg Schedule 2 N=51	Total N=301
Age (years)						
Mean±SD	47±11	50±10	46±11	48±11	48±12	48±11
Gender						
	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	N (%) ^b
Female	50 (50.0)	26 (52.0)	29 (58.0)	29 (58.0)	24 (47.1)	158 (52.5)
Male	50 (50.0)	24 (48.0)	21 (42.0)	21 (42.0)	27 (52.9)	143 (47.5)
Race						
	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	N (%) ^b
White	88 (88.0)	47 (94.0)	42 (84.0)	47 (94.0)	47 (92.0)	271 (90.0)
Other	12 (12.0)	3 (6.0)	8 (16.0)	3 (6.0)	4 (8.0)	30 (10.0)
Fagerström Total^c						
Mean±SD	6.1±1.9	6.3±2.1	5.8±2.0	5.8±2.0	6.6±1.7	6.0±2.0
Median (range)	6 (1-10)	7 (1-10)	6 (1-10)	6 (1-10)	6 (3-10)	6 (1-10)
Cigarettes Per Day						
Mean±SD	24.7±8.8	24.8±9.1	22.6±7.0	24.3±9.4	25.6±10.5	24.0±9.0
Median (range)	20 (15-50)	20 (15-50)	20 (15-40)	20 (15-60)	20 (15-70)	20 (15-70)
Previous Quit Attempts						
	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	n (%) ^a	N (%) ^b
Yes	96 (96.0)	48 (96.0)	47 (94.0)	48 (96.0)	49 (96.1)	288 (95.7)

^a Percentages based on total number of subjects within treatment group.

^b Percentages based on total number of subjects who received treatment.

^c Fagerström score assesses the severity of nicotine addiction ranging from 0 (minimum) to 10 (maximum).

Table 2

Comparison of the Presence of Reactogenicity Events by Treatment Group

	Number (Percent) of Subjects by Treatment					p-value
	Placebo (N=100)	200 µg Schedule 1 (N=50)	200 µg Schedule 2 (N=50)	400 µg Schedule 1 (N=50)	400 µg Schedule 2 (N=51)	
Local						
Ache	96 (96.0)	48 (96.0)	43 (86.0)	47 (94.0)	49 (96.1)	0.129
Burning	42 (42.0)	23 (46.0)	22 (44.0)	21 (42.0)	23 (45.1)	0.988
Erythema	39 (39.0)	29 (58.0)	23 (46.0)	27 (54.0)	22 (43.1)	0.180
Heat	42 (42.0)	23 (46.0)	25 (50.0)	27 (54.0)	22 (43.1)	0.661
Swelling/Induration	60 (60.0)	33 (66.0)	32 (64.0)	33 (66.0)	29 (56.9)	0.827
Tenderness	95 (95.0)	49 (98.0)	44 (88.0)	48 (96.0)	50 (98.0)	0.126
Systemic						
Fever	10 (10.0)	5 (10.0)	2 (4.0)	8 (16.0)	6 (11.8)	0.403
General Discomfort/Malaise	79 (79.0)	38 (76.0)	38 (76.0)	42 (84.0)	42 (82.4)	0.803
Headache	67 (67.0)	35 (70.0)	32 (64.0)	35 (70.0)	36 (70.6)	0.945
Myalgia	86 (86.0)	39 (78.0)	41 (82.0)	46 (92.0)	45 (88.2)	0.315
Nausea	44 (44.0)	17 (34.0)	25 (50.0)	21 (42.0)	25 (49.0)	0.501
Vomiting	6 (6.0)	3 (6.0)	2 (4.0)	2 (4.0)	8 (15.7)	0.111