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Early introduction of peanut reduces peanut allergy across risk groups in pooled and causal inference analyses

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Abstract

Background

The LEAP study has shown the effectiveness of early peanut introduction in prevention of peanut allergy (PA). In the EAT study, a statistically significant reduction in PA was present only in per-protocol (PP) analyses, which can be subject to bias.

Objective

To combine individual-level data from the LEAP and EAT trials and provide robust evidence on the bias-corrected, causal effect of early peanut introduction.

Method

As part of the European Union-funded iFAAM project, this pooled analysis of individual paediatric patient data combines and compares effectiveness and efficacy estimates of oral tolerance induction among different risk strata and analysis methods.

Results

An intention-to-treat (ITT) analysis of pooled data showed a 75% reduction in PA ($p < 0.0001$) among children randomized to consume peanut from early infancy. A protective effect was present across all eczema severity groups, irrespective of enrolment sensitization to peanut, and across different ethnicities. Earlier age of introduction was associated with improved effectiveness of the intervention. In the pooled PP analysis, peanut consumption reduced the risk of PA by 98% ($p < 0.0001$). A causal inference analysis confirmed the strong PP effect (89% average treatment effect relative risk reduction $p < 0.0001$). A multivariable causal inference analysis approach estimated a large (100%) reduction in PA in children without eczema ($p = 0.004$).

Conclusion

We demonstrate a significant reduction in PA with early peanut introduction in a large group of pooled, randomized participants. This significant reduction was demonstrated across all risk subgroups, including children with no eczema. Furthermore, our results point to increased efficacy of the intervention with earlier age of introduction.

Introduction

Despite strong findings favouring early peanut introduction, there are gaps in our knowledge base especially with respect to whether to target high risk or normal populations, and age of intervention. It is therefore not surprising that specialist organizations provide different guidelines regarding peanut introduction. Most recently, the EAACI guidelines gave a conditional recommendation for the introduction of peanut to the whole infant population as the strength of the evidence was considered to be moderate.¹ These recommendations were limited to countries with a high prevalence of peanut allergy. In order to increase our knowledge base for future guidelines, we conducted an integrated analysis of individual participant data from all available randomized controlled trials (RCT) of peanut introduction.

Until recently, UK and US guidelines recommended avoidance of peanut in infancy to prevent the development of peanut allergy.^{2,3} Despite these guidelines, the prevalence of peanut allergy continued to increase and subsequently a growing body of evidence emerged favouring early introduction of peanut rather than early avoidance.

An observational study comparing Israeli and UK children found that the early introduction and regular consumption of peanut in infancy was strongly protective against developing a peanut allergy (PA).⁴ A series of early introduction RCTs have subsequently taken place. There have been two independently powered and randomized UK-based cohorts investigating the early introduction of peanut.^{5,6} A recent systematic review and meta-analysis of these two early introduction RCTs was undertaken and the pooled risk ratio for peanut introduction was 0.29 (0.11-0.74).⁷ The review concluded that the current body of evidence supported the early introduction of peanut for the prevention of peanut allergy but was unable to investigate subgroups, such as infants with varying eczema severity, ethnicity, and sensitization.⁷

The LEAP (Learning Early About Peanut Allergy) study recruited infants at high risk of developing PA aged 4-10 months and showed an 81% relative reduction in PA prevalence between the peanut consumption and avoidance groups.⁵ The EAT (Enquiring About Tolerance) study enrolled children at 3 months of age and randomized them to either consume 6 allergenic foods including peanut (Early Introduction Group: EIG) or to avoid allergenic food consumption until 6 months (Standard Introduction Group: SIG). A 51% reduction in peanut allergy prevalence was observed but did not reach statistical significance in the intention-to-treat (ITT) analysis.⁶ Adherence to peanut consumption in the LEAP study was 92%. In comparison, peanut adherence was significantly lower in the EAT study (61% of 549 peanut adherence evaluable participants and 48% of all 652 EIG participants). The reasons for poor adherence in the EAT study have been extensively investigated previously where increasing maternal age, non-Caucasian ethnicity, and lower maternal quality of life were found to be important factors influencing adherence. In comparison to LEAP study participants, the EAT participants introduced 5 other allergenic foods alongside peanut making the intervention harder to follow.^{6,8}

Despite the LEAP study demonstrating that earlier age of introduction of peanut reduced PA in high-risk participants and subsequent changes to the American infant feeding guidelines,⁹ there remains uncertainty across clinical and policy spheres regarding the robustness of the results and their generalizability to lower-risk populations. The EAT study set out to address some of those concerns by enrolling a lower-risk cohort but ultimately did not demonstrate that the intervention had as strong of an effect in the ITT population. In particular, low protocol compliance in the EAT study led to complications in the interpretation of results.⁶

When adherence to an intervention is lacking, patients and investigators have been shown to prefer bias corrected estimates of an intervention's effectiveness across risk subgroups using valid per-protocol effect estimates.¹⁰ Since ITT analyses estimate the effect of treatment assignment, not the

actual treatment received, they can produce misleading causal estimates of an intervention when adherence is reduced. Therefore, effective interventions, diluted by non-adherence, can appear to be ineffective using ITT analyses. For these reasons and others, alternative analysis approaches have been proposed to adjust for post-randomization imbalances of treatment adherence.¹¹ In addition to ITT and PP analyses of the pooled data from both cohorts, we have implemented causal inference analysis approaches to estimate the effect of early introduction of peanut among the different risk strata and adherence populations, and to adjust per-protocol analyses for biases often present due to post-randomization imbalances of the intervention received rather than assigned. For example, a small number of participants in the intervention group had peanut allergy at baseline or developed peanut allergy while receiving the intervention and were subsequently instructed per the protocol not to begin or to discontinue consumption of peanut, respectively. In the per-protocol analysis of the original study reports, these participants were considered non-per-protocol and thus were removed from the analysis. However, under a causal inference framework, these participants are included as protocol adherent participants and analysed as receiving the intervention.¹¹

Based on evidence from the LEAP study, and supported by results of the EAT study, the National Institute of Allergy and Infectious Diseases (NIAID) and a number of other specialist national societies issued guidelines recommending the early introduction of peanut for peanut allergy prevention. The evidence cited to support the NIAID recommendations in different eczema subgroups is based on those LEAP infants recruited with egg allergy (who had varying levels of eczema) and also from the EAT study participants, most of whom had no or mild to moderate eczema at enrollment.⁹

The pooled analysis of data from the LEAP and EAT cohorts presented here provides a unique opportunity to look at the effect of the early introduction of peanut across a variety of risk strata (presence of eczema and its severity, enrolment sensitization status to peanut, and ethnicity), while

implementing regression adjustment methodologies to estimate the causal effect of oral tolerance induction when adhered to.

Accepted Article

Methods

Study-specific methods for the LEAP and EAT studies are published in full elsewhere and cohort demographics are summarized in Table S1.^{12, 13}

The European Union-funded iFAAM (Integrated Food Allergy and Allergen Management) project included a pooled analysis of individual-level data from the RCTs of early allergenic food introduction. This pooled analysis allows estimates of oral tolerance induction among a number of different risk strata. Pooled estimates are adjusted for study-specific and individual-specific factors. Pooling the data across studies provides a robust and powerful estimate of the effect of oral tolerance induction. Analysis of the individual participant data enables estimates to be derived for the effect of early introduction of allergenic foods across the different ethnicities, baseline eczema severities, and baseline IgE sensitization; a notable advantage in comparison to a meta-analysis approach of summarized results.

Detailed methods and levels of adherence for the LEAP and EAT studies have been reported elsewhere. Data collected in each study included allergy, sensitization and anthropometric endpoints, as well as demographics and family history of atopy. The challenges of harmonizing data from studies that have differences in design and methodology were extensively discussed and agreement was reached on what could feasibly be analysed in one dataset and a statistical analysis plan outlined. Using the iFAAM-funded Allerg-e-lab, each data point was annotated with descriptors, and individual datasets from each study were then recoded, renamed and relabelled using the agreed upon matched variables, thus creating individual study datasets containing identical names, formats, labels, and coding values. The harmonized datasets were merged together as a final integrated dataset and analysed using methods described below.

The outcomes evaluated were the point prevalence of peanut allergy by 3 years (EAT study primary endpoint) and by 5 years of age (LEAP study primary endpoint) and sensitization to peanut (measured as skin prick test [SPT] ≥ 1 mm or specific IgE ≥ 0.1 kU/L). Secondary outcome of baseline eczema, defined by objective Scoring Atopic Dermatitis (SCORAD) measurement, was also available from both studies.

The integrated dataset was validated to ensure published study results on primary and secondary outcomes from the individual trials could be replicated. All published data on peanut allergy and sensitization were replicated in the integrated dataset before combined analyses began.

Statistical analysis

The primary endpoint of this pooled analysis was peanut allergy prevalence – defined on the basis of a positive oral food challenge (OFC) or sensitization and symptom history where food challenge was not done. Ninety-one percent of peanut allergy diagnoses were made on the basis of OFC. The primary analysis methodology for the pooled endpoint was a logistic regression model. Univariate and multivariable adjustments to the risk of peanut allergy were made and shown as estimated probabilities of allergy, risk differences between the randomized groups, and as relative risk reductions calculated from the univariate and multivariable adjusted predicted probabilities from the logistic regression models. We pre-specified a number of subgroups for comparison and the raw-unadjusted proportions are shown according to the different pre-specified subgroups. All tests were two-sided at the $\alpha = 0.05$ level of significance. Pearson's chi-squared tests were also used to replicate individual study results. When one or more cells had expected counts less than 5, Fisher's exact test was used. Individual and combined study proportions were displayed as bar charts with frequencies, proportions, and p-values annotated. Relative risk reductions and risk differences with

95% confidence intervals were computed and displayed as forest plots among the analysis and study populations.

Causal inference methods were implemented to adjust per-protocol analyses for biases often present due to post-randomization imbalances of the intervention's uptake relative to assignment. The average treatment effect (ATE), which estimates the average treatment effect among the entire population, assumes each participant is able to receive the intervention. Doubly robust methods were used to estimate the average treatment effect (ATE) and 95% bootstrapped confidence intervals by combining inverse probability weighting and regression adjustment using the AIPW option and the bootstrap statement in SAS Proc Causaltrt.¹⁴⁻¹⁶ Estimates for the average treatment effect for the treated (ATT) and average treatment effect for the untreated (ATU) are reported in the supplementary appendix.

Causal inference per-protocol analyses were performed using SAS software version 9.4 (Proc Causaltrt).¹⁷ The estimates were adjusted for ethnicity, baseline eczema severity, and baseline peanut sensitization as covariates in each study separately and in the pooled study-adjusted analysis to produce the ATE, ATT, and ATU estimates. Complier average causal effect (CACE) was determined for each study and in a combined analysis adjusted for study using the R package ivpack version 1.2.¹⁸ The CACE analysis used an instrumental variable approach, with two-staged least squares regression and active participation in the intervention as the predictor and randomized treatment assignment as the instrument. This instrumental variable approach produces marginal estimates of the treatment effect without making conditional adjustments for the covariates used in the SAS Proc Causaltrt approach.

Results

Individual level participant data from the EAT and LEAP studies were combined to give a pooled estimate of peanut allergy prevalence in intervention and control groups based on data from 1943 children. Demographic characteristics, eczema severity, and baseline IgE sensitization were balanced between control and intervention groups in the pooled data (Table S1).

Peanut allergy

Peanut allergy status could be evaluated in 1796 of 1943 children in the LEAP and EAT cohorts, 86 of whom (4.8%) were allergic to peanut at 3-5 years of age (Table S1). The intention-to-treat analysis showed a 75% reduction in peanut allergy: 1.9% in the early introduction group versus 7.6% in the control group ($p < 0.0001$) (Figure 1a). Amongst those adherent to the early introduction or control protocols, the effect was strengthened with a 98% reduction in peanut allergy ($p < 0.0001$) (Figure 1b).

Eczema

Participants with eczema at baseline contributed disproportionately to the prevalence of peanut allergy: 0.9% (8/900) in those with no eczema versus 8.7% (78/895) in those with eczema. The effect of early introduction in these eczema subgroups is key to determining the potential impact of any early introduction strategy.

The protective effect of early introduction was evident across all eczema severities with significant intention-to-treat reductions in peanut allergy prevalence in mild (85% reduction), moderate (87% reduction) and severe (67% reduction) eczema severity groups (Figure 2a). Greater reductions in peanut allergy were seen in per-protocol analyses, with 100% reductions in children with mild and moderate eczema and a 96% reduction among the severe eczema group (Figure 2b). Among children with no eczema, there was a 36% intention-to-treat reduction ($p = 0.73$) and a 100% per-protocol reduction in peanut allergy ($p = 0.16$) (Figure 2).

Peanut sensitization

In both individual and pooled study results, skin prick test sensitization to peanut was significantly reduced at 12 months among those introducing peanut early in comparison to those avoiding or introducing after 6 months of age. IgE sensitization to peanut at 12 months of age remained similar between the randomized groups (Figure S1).

In the ITT population, children with specific IgE to peanut ≥ 0.1 k U/L at enrolment had a 75% reduction in peanut allergy prevalence at 3-5 years of age ($p < 0.0001$) (Figure 3a). There was also a statistically significant reduction in peanut allergy prevalence among children not sensitized to peanut at enrolment—72% reduction, $p = 0.003$ (Figure 3a). In per-protocol analyses, this reduction was greater, with 100% reduction in peanut allergy among sensitized children and a 94% reduction in non-sensitized children (Figure 3b).

Ethnicity

Non-Caucasian groups combined (approximately 20% of all participants) had an approximately 3-fold higher prevalence of peanut allergy compared to Caucasians (16.8% versus 5.5%, $p < 0.001$). However, among non-Caucasian ethnicities, peanut allergy rates were more similar ($p > 0.05$) (Figure S2). The intervention's effect size was similar in the different ethnicities, and a significant reduction in peanut allergy prevalence was seen in both Caucasian and non-Caucasian groups (72% and 82% reduction, respectively; $p < 0.0001$) (Figure 3c).

Causal effects

Due to lower protocol compliance in the EAT study, we implemented a causal inference approach to estimate the intervention's effect on peanut allergy. In pooled analyses adjusted for study, peanut consumption demonstrated statistically significant causal effects on peanut allergy prevalence. Under

all causal inference frameworks (ATE, ATT, ATU, CACE), the predicted mean allergy incidence was lower for the group of participants that consumed peanut; ATE analysis determined an absolute risk difference in peanut allergy of -7.18% [95% CI: (-9.01, -5.49); $p < 0.0001$], CACE analysis: -8.22% [95% CI: (-10.96, -5.47); $p < 0.0001$] (Table 1 and Figure 4; ATT and ATU reported in Table S2 and Figure S3). Furthermore, consumption of peanut contributed to a relative risk reduction (RRR) of greater than 88% in all causal effect analyses of pooled data (ATE RRR of 88.8%, CACE RRR of 88.1%) (Table 1, Figure 4; ATT and ATU reported in Table S2 and Figure S3). Causal effect subgroup analyses are provided in Figures S4 and S5. Lastly, a multivariable logistic regression analysis was fit to compare the probability of peanut allergy between the EAT and LEAP cohorts after risk factor adjustment. Specifically, adjustment was made for study, randomization assignment, SCORAD group, and egg allergy at baseline. Figure S6a displays the probability of peanut allergy between the EAT and LEAP cohorts according to each risk factor combination. The model-based estimates of the risk of peanut allergy were shown to be similar between the EAT and LEAP studies, demonstrating that combining these higher and lower risk cohorts using regression adjustment adequately controlled for the heterogeneity in the outcome between these cohorts. Moreover, the analysis demonstrates significantly lower risk of PA in the consumption arm in infants without eczema and with eczema (regardless of eczema severity) as well as in the presence or absence of egg allergy (Figures S4, S5, S6a). Figure S6b provides the raw data partitioned in a similar manner as S6a to show the unmodeled peanut allergy proportions, sample sizes, and number with peanut allergy in each of the risk strata, cohorts, and treatment groups. Using regression adjustment, the logistic model interpolates the allergy rate across all factors in the model to give average estimates in Figure S6a, which are less influenced by the small sample sizes and sparse cases of allergy in some of the risk strata shown in Figure S6b.

Age of introduction

In an exploratory analysis we examined the association between age of introduction of peanut and the prevalence of peanut allergy at 36 months in the EAT study (Figure 5). This analysis looked at age of introduction irrespective of the intervention group to which the subject was randomized. All but one participant introducing peanut before 6 months of age were randomized to the Early Introduction Group (EIG); however, seventy-one (13%) of EIG participants introduced peanut after the key early introduction period, at 6 months or later. Similarly, one SIG participant introduced peanut before 6 months of age; however, there was a large amount of variability in age of introduction within the SIG, which introduced peanut, at parental discretion, at 6 months and beyond. The analysis presented in Figure 5 uses the variability in the combined randomized groups from EAT to investigate the association between age of introduction and peanut allergy. The overlaid regression line shows an increase in prevalence of peanut allergy at 3 years of age with increasing age of introduction.

Discussion

In order to maximize the use of data from the EAT and LEAP studies and assess the efficacy of intervention in important subgroups we undertook a patient-level meta-analysis. The EAT study included participants with varying risk, while LEAP was limited to high-risk participants. To account for these differences, we included ethnicity, baseline eczema, egg allergy, and baseline IgE in our models. A recent systematic review and meta-analysis concluded that there was moderate evidence to support the early introduction of peanut as a method of reducing peanut allergy prevalence.⁷ However, this approach has limitations, for example, a meta-analysis cannot adjust for the varying severity profiles of the participants in the two studies. This European Union-funded iFAAM project has successfully harmonized data from two large randomized controlled trials on early peanut introduction and analysed pooled, individual-level data on 1943 children from normal and high-risk populations. While the combined population does not completely reflect the general population, the pooled analysis increases the power to analyse subgroups (e.g., eczema severity levels, sensitization groups, and ethnic groups) and allows for risk factor adjustment within the integrated dataset. This approach diminishes the need for an impractically large trial in low-risk infants.

In summary, the ITT analysis of pooled data showed a 75% reduction in peanut allergy prevalence among children randomized to consume peanut from an early age; moreover, significant reductions were demonstrated in peanut allergy across all eczema severity levels, sensitization groups, and ethnic groups. Importantly, a causal inference analysis approach (ATE) demonstrated a significant benefit of the intervention among those without eczema, who make up a majority of children at risk for peanut allergy across the whole population. Moreover, a non-randomized analysis of age of introduction in the EAT cohort demonstrates an association between earlier age of introduction and increased efficacy (Figure 5). For the first time these results show, using integrated participant-level data, that early peanut introduction is successful and generalizable to a wider, multi-ethnic population, and provides a strong basis for a broad public health measure. This is now supported by the recently

published PreventADALL randomised controlled trial in a general population in Sweden and Norway which demonstrated an odds ratio of 0.4 (95% confidence interval 0.2 to 0.8) for peanut allergy with early introduction of peanut products.¹⁹

Specialist guidelines often base their recommendations purely on the ‘real world effectiveness’ that intention-to-treat analyses provide, whilst ignoring per-protocol analyses. For example, while the EAT per-protocol analysis showed 100% efficacy ($p=0.003$), this analysis was not taken into account by the EAACI Guidelines committee, as their selection criteria was limited to consider only ITT analyses.¹ We argue that the primary ITT EAT study results are ‘diluted by non-adherence,’ and the intervention’s actual efficacy is evident using a per-protocol analysis.

The reluctance to accept per-protocol analyses as evidence for specialist guidelines has been justified by the concern of introducing bias, since randomization is not always preserved (e.g., more atopic participants could be disproportionately dropped from the intervention arm post-randomization and thus excluded from per-protocol analysis, resulting in bias). However, the use of newer causal inference methods have the ability to mitigate this bias with the use of propensity scores, regression adjustment, and instrumental variable analysis. It is remarkable that the 2021 Nobel Prize for economics was awarded to David Card, Joshua Angrist, and Guido Imbens for, “Answering causal questions using observational data.” In this award, Section 1.3 “Causal effects in a world with imperfect compliance and individual heterogeneity,” the committee comments that, “Imperfect compliance with treatment assignment makes it more difficult to identify the average effect of treatment, in particular when causal effects vary in the population under study. . . . An ITT analysis thus provides an unbiased estimate of the effect of the treatment assignment in the study population, but not the actual causal effect of the treatment itself.”²⁰ Using the methods pioneered by these scholars, we have attempted to address the discordance between the ITT results observed in EAT, where

compliance was clearly imperfect (48%), with the results observed in LEAP, where compliance was very high (92%).

A comparison of naïve per-protocol analyses with the newer causal inference approaches demonstrated consistency of the intervention's efficacy when adhered to. Peanut consumption reduced the risk of PA by 98% ($p < 0.0001$) in the pooled per-protocol analysis. The causal inference multivariable analysis showed an 89% average treatment effect (ATE) relative risk reduction, $p < 0.0001$. Lower adherence (49%) in the EAT ITT population with no eczema at enrolment may have diluted the effectiveness of treatment in this underpowered, lower-risk subgroup (prevalence of PA 1%).^{21,22} However, a multivariable causal inference analysis approach (ATE) estimated a complete (100%) and statistically significant ($p = 0.004$) reduction in PA in children without eczema (Figure S4A and S5A). The consistency of the different causal effect estimates, and the similarity of these estimates to that reported for the per-protocol analyses, provides evidence that over 85% of allergy can be prevented with early introduction of peanut.

The main strength of our study was the integration and validation of individual-level data from two large randomized controlled trials (Figure 1), enabling us for the first time to determine efficacy of the intervention across the whole range of risk groups that comprise a normal population. Furthermore, the large number of diagnoses based on food challenge (91%) validates the estimates of allergy prevalence, and the harmonization of criteria used to diagnose food allergy and sensitization adds consistency to the data presented.

These findings are not without limitations. Firstly, the LEAP and EAT study populations were selected very differently (high and normal risk populations respectively) and followed up to aged 5 and 3 years respectively. However, the lower risk EAT study is a heterogeneous population with both low and high-risk participants. A causal inference analysis with adjustment for risk factors such as eczema and egg

allergy controls for these differences and shows the intervention to be effective. Secondly, the discrepancy between the per-protocol and ITT results, especially among infants at lower risk of allergy, points to a difference between the intervention's 'idealized' efficacy (per-protocol) and its 'real world' effectiveness (ITT).²³ An intervention may be very efficacious (e.g., folate supplementation to prevent neural tube defects) but if its uptake is low, it will not be effective. While cultural and social norms are likely to play an important role in early feeding of peanuts, these preferences can change following the publication and dissemination of trial results showing efficacy of peanut introduction.⁴ Soriano (2019) showed a 3-fold increase in peanut introduction in Australia by age 1 year in 2018 compared with 2007-2011 (from 30% to 90%), which coincided with changes to national infant feeding guidelines following publication of the LEAP study.^{24,25} Good adherence to early dietary introduction of peanut products is essential for a successful peanut allergy prevention strategy.

Another weakness of our findings is that the association between age of peanut introduction and prevalence of allergy in the EAT study is, at least partially, an un-randomized (i.e. observational) comparison. Therefore, this analysis may be confounded by other, unaccounted for factors. A similar analysis was previously performed in the LEAP population.^{26,27} However, this analysis showed no association between age of introduction and prevention of allergy. This apparent discrepancy in the age dependent effect in EAT and the age independent effect in LEAP on the interventional efficacy can be explained for two reasons. Firstly, in the LEAP study the intervention was adhered to and prevented peanut allergy across the entire age range;^{26,27} whereas in EAT, less peanut was consumed, and protocol adherence was low (48%). Secondly, at the LEAP screening visit 76 out of the 899 participants did not meet the inclusion criteria as they were considered already peanut allergic (peanut SPT >4mm). These excluded infants were significantly older than the rest of the LEAP screening cohort (mean age 8.3 months, SD 1.88),²⁸ and it was therefore not possible to prevent peanut allergy in this older group of infants.

The results of this pooled analysis provide new evidence for the efficacy of early introduction of peanut in children with all degrees of eczema severity and moreover demonstrates efficacy in those without eczema. Moreover, this efficacy is demonstrated irrespective of ethnic group, peanut sensitization status, and presence of egg allergy. These results suggest that recommending early consumption of peanut as a prevention strategy be broadly applied to the entire population, rather than targeting selected higher risk groups. Moreover, our results point to increased efficacy of the intervention with an age of introduction below six months, calling into question recommendations for exclusive breast feeding during the first six months of life in resource rich regions. Utilizing individual-level data from all RCTs of peanut introduction to date, these new analyses thus strengthen the evidence that underlie the EAACI guidelines recommending early introduction of peanut to the general population.

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Tables

Table 1 Summary of causal effect of intervention on peanut allergy outcome in EAT and LEAP and combined study-adjusted analyses. *Not adjusted for study.

Analysis Population	Study	n randomized control	n randomized intervention	n intervention received	Relative risk reduction	Risk difference % (95% CI)	p-value, risk difference
ITT	EAT	597	571	315	51.2	-1.29 (0.26, -2.83)	0.106
PP	EAT	525	310	315	100	-2.48 (-1.15, -3.81)	0.003
CACE	EAT	597	571	315	100	-2.33 (-5.16, 0.5)	0.105
ATE	EAT	597	571	315	100	-2.68 (-3.92, -1.66)	<.0001
ITT	LEAP	314	314	305	81.5	-14.01 (-9.41, -18.62)	<.0001
PP	LEAP	295	294	305	98.0	-16.95 (-12.58, -21.31)	<.0001
CACE	LEAP	314	314	305	86.2	-14.43 (-19.15, -9.69)	<.0001
ATE	LEAP	314	314	305	85.9	-14.77 (-19.01, -10.49)	<.0001
ITT	Combined	911	885	620	74.6	-5.65 (-3.71, -7.59)	<.0001
PP	Combined	820	604	620	97.9	-7.64 (-5.77, -9.5)	<.0001
CACE	Combined	911	885	620	88.1 [‡]	-8.22 (-10.96, -5.47)	<.0001
ATE	Combined	911	885	620	88.8	-7.18 (-9.01, -5.49)	<.0001

Average treatment effect (ATE) was estimated through regression and propensity score adjustment methods. The ATE estimates the average treatment effect among the entire population, assuming each participant is able to receive the intervention. Complier average causal effect (CACE) analysis used an instrumental variable approach, with two-staged least squares regression and active participation in the intervention as the predictor and randomized treatment assignment as the instrument.

[‡]Two-staged least squares regression produces a risk difference estimate only; relative risks were computed by determining a theoretical peanut allergy rate among control participants, under the assumption of the CACE model that participants of the control group have the same probability of being non-compliant as participants of the intervention group and being offered the intervention has no effect on the outcome. Thus, the combined relative risk under the CACE approach does not adjust for study.

Figure Legends

Figure 1 Peanut allergy prevalence at 3-5 years of age – a) Intention-to-treat and b) per-protocol populations.

Prevalence of peanut allergy in each randomized treatment group is shown within individual and combined studies in a) ITT analysis and b) PP analysis. Number of peanut allergic (PA) and total number of participants (N) are annotated at the top. Each bar is annotated with the prevalence of peanut allergy as determined by OFC at 3 years (EAT) or 5 years of age (LEAP); relative risk reduction in individual and combined studies is annotated in red. Pearson's chi-squared was used to determine annotated p-values; when expected values were less than 5, Fisher's two-tailed exact test was used. Fisher's p-value is reported for the EAT PP analysis.

Figure 2 Peanut allergy prevalence by eczema severity – a) Intention-to-treat and b) Per-protocol populations.

Prevalence of peanut allergy in each randomized treatment group is shown by baseline eczema subgroups when data from the EAT and LEAP studies are combined in a) ITT analysis and b) PP analysis. Number of peanut allergic (PA) and total number of participants (N) are annotated at the top. Each bar is annotated with the prevalence of peanut allergy as determined by OFC at 3 years (EAT) or 5 years of age (LEAP); relative risk reduction in individual and combined studies is annotated in red. Pearson's chi-squared was used to determine annotated p-values; when expected values were less than 5, Fisher's two-tailed exact test was used. *Fisher's p-value is reported for the ITT and PP analyses in the subgroup of participants that did not have eczema at baseline as per the original statistical analysis plan. However, this analysis is expanded in Supplementary Figure 4a using a model based causal inference approach producing a p-value of 0.004.

Figure 3 Peanut Allergy Prevalence by specific IgE sensitization status at enrolment – a) Intention-to-treat and b) Per-protocol population; in Caucasian and non-Caucasian children – c) Intention-to-treat and d) Per-protocol population.

Prevalence of peanut allergy in each randomized treatment group is shown in baseline sensitized (peanut-specific IgE ≥ 0.1 kU/L) and unsensitized (peanut-specific IgE < 0.1 kU/L) subgroups for individual and combined studies in a) ITT analysis and b) PP analysis; in Caucasian and non-Caucasian subgroups within individual and combined studies in c) ITT analysis and d) PP analysis. Number of peanut allergic (PA) and total number of participants (N) are annotated at the top. Each bar is annotated with the prevalence of peanut allergy as determined by OFC at 3 years (EAT) or 5 years of

age (LEAP); relative risk reduction in individual and combined studies is annotated in red. Pearson's chi-squared was used to determine annotated p-values; when expected values were less than 5, Fisher's two-tailed exact test was used. Fisher's p-value is reported for the ITT analysis among sensitized participants in the EAT study, PP analysis in both sensitized and unsensitized subgroups in the EAT study, ITT analysis among non-Caucasian participants in the EAT study, and PP analysis in both Caucasian and non-Caucasian subgroups in the EAT study.

Figure 4 Forest plot of peanut allergy prevention effect sizes among the different study populations and analysis methods.

This forest plot shows the percentage relative risk reductions in peanut allergy (PA) annotated below the marker and risk differences (x-axis) in PA are annotated above the marker among the different cohorts (EAT, LEAP, and combined studies). Relative risk reductions and risk differences are relative to the control (peanut avoidance) group and determined by intention-to-treat (ITT), per-protocol (PP), and causal inference methods: complier average causal effect (CACE) and average treatment effect (ATE). Additionally, average treatment effect for the treated (ATT) and average treatment effect for the untreated (ATU) are shown in supplementary figure S2.

*Not adjusted for study; two-staged least squares regression produces a risk difference estimate only; relative risks were computed by determining a theoretical peanut allergy rate among control participants, under the assumption of the CACE model that participants of the control group have the same probability of being non-compliant as participants of the intervention group and being offered the intervention has no effect on the outcome. Thus, the combined relative risk under the CACE approach does not adjust for study.

Figure 5 Prevalence of peanut allergy by age of introduction to peanut in the EAT study.

The prevalence of peanut allergy at 36 months is shown according to the month of peanut introduction in the EAT cohort with available data (n=1,168) irrespective of randomization group. Those starting to consume peanut at 9 months of age or older or who never introduced peanut were grouped into a single 9+ category. The smoothed regression line and bootstrapped confidence intervals are created with a cubic spline to visualize the prevalence of peanut allergy conditional on when peanut was introduced into the diet. The adherent EAT participants randomized to the early introduction group (EIG) are shown in the 3-5 month bins whereas the non-adherent participants are dispersed along with the standard introduction participants in the 6 to 9+ bins. The number of subjects with peanut allergy (PA), the total sample size (N), and the proportion (%) with peanut allergy are annotated above each bar. The large risk of peanut allergy apparent in the 6-month

category may be an artefact of the study design where participants were asked to consume before the 6-month time point resulting in an artificially low sample size.

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Fig1a

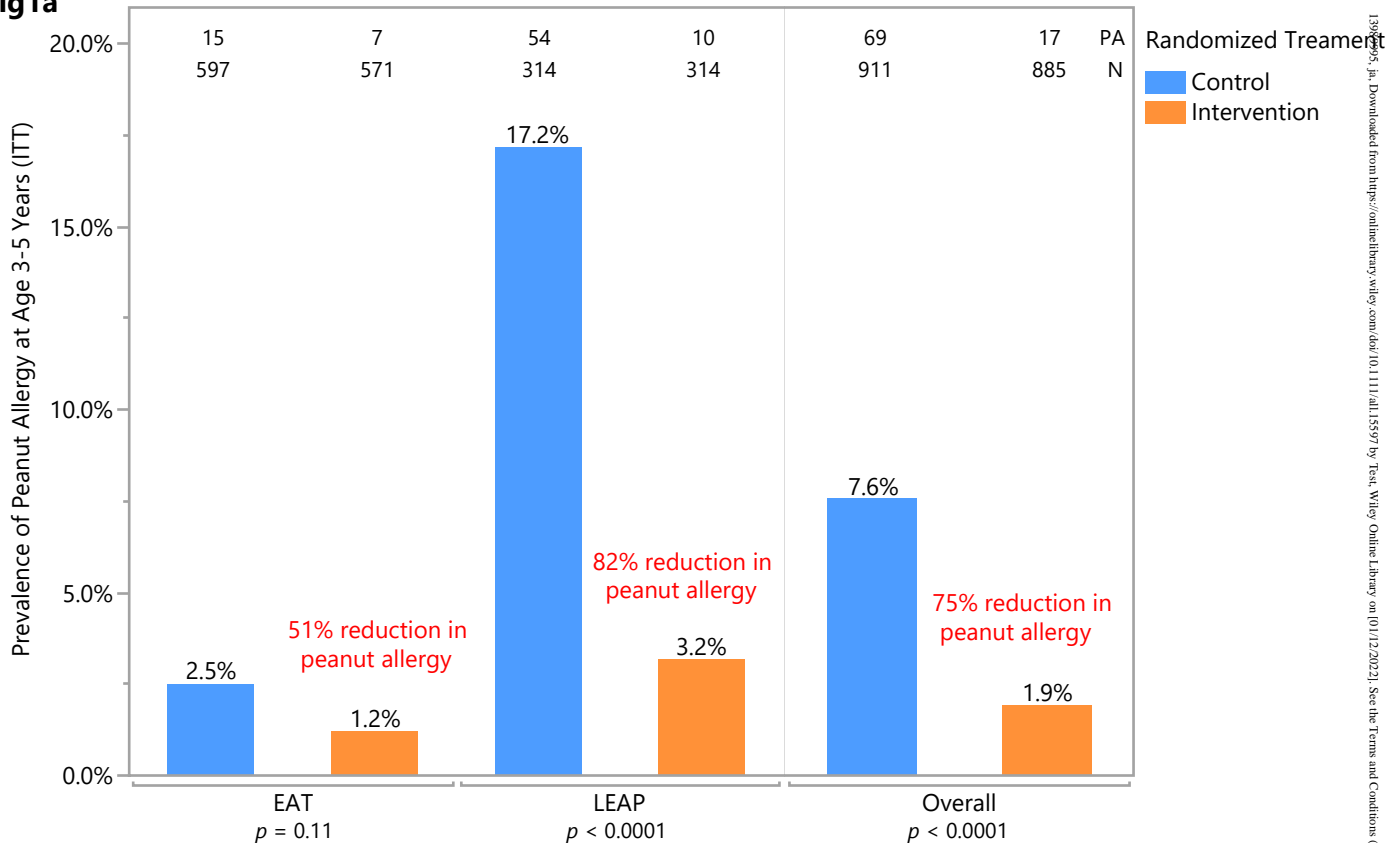


Fig1b

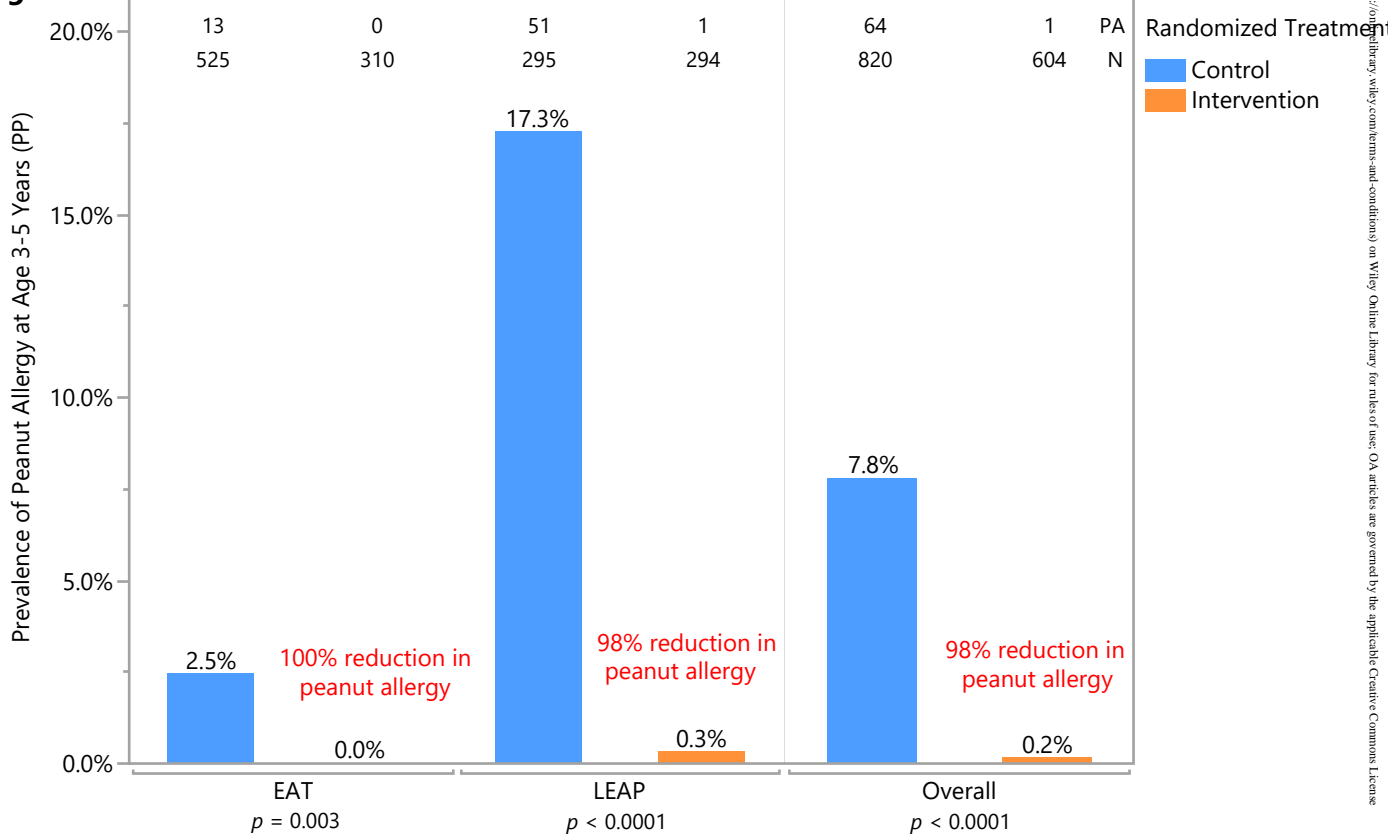


Fig2a

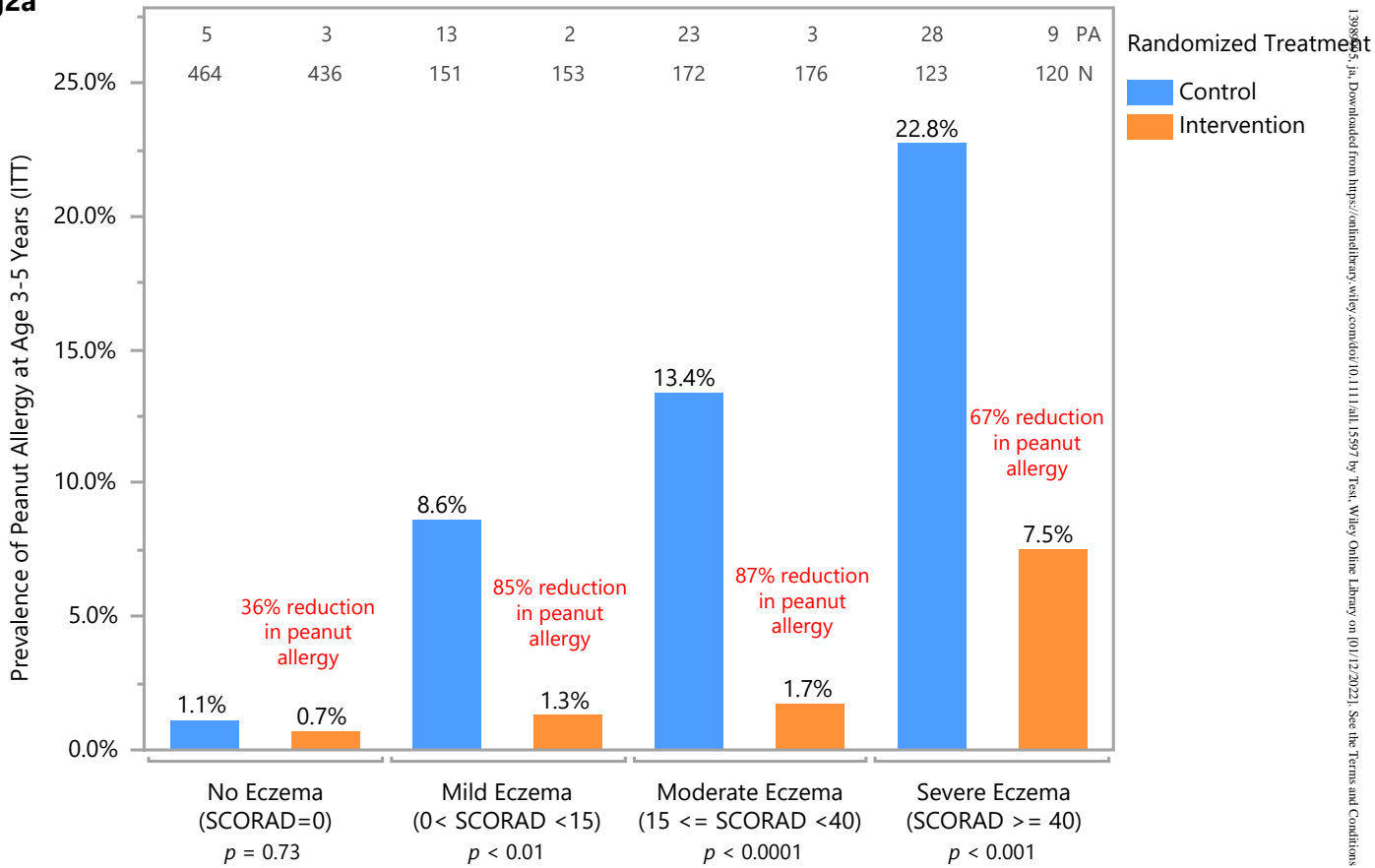


Fig2b

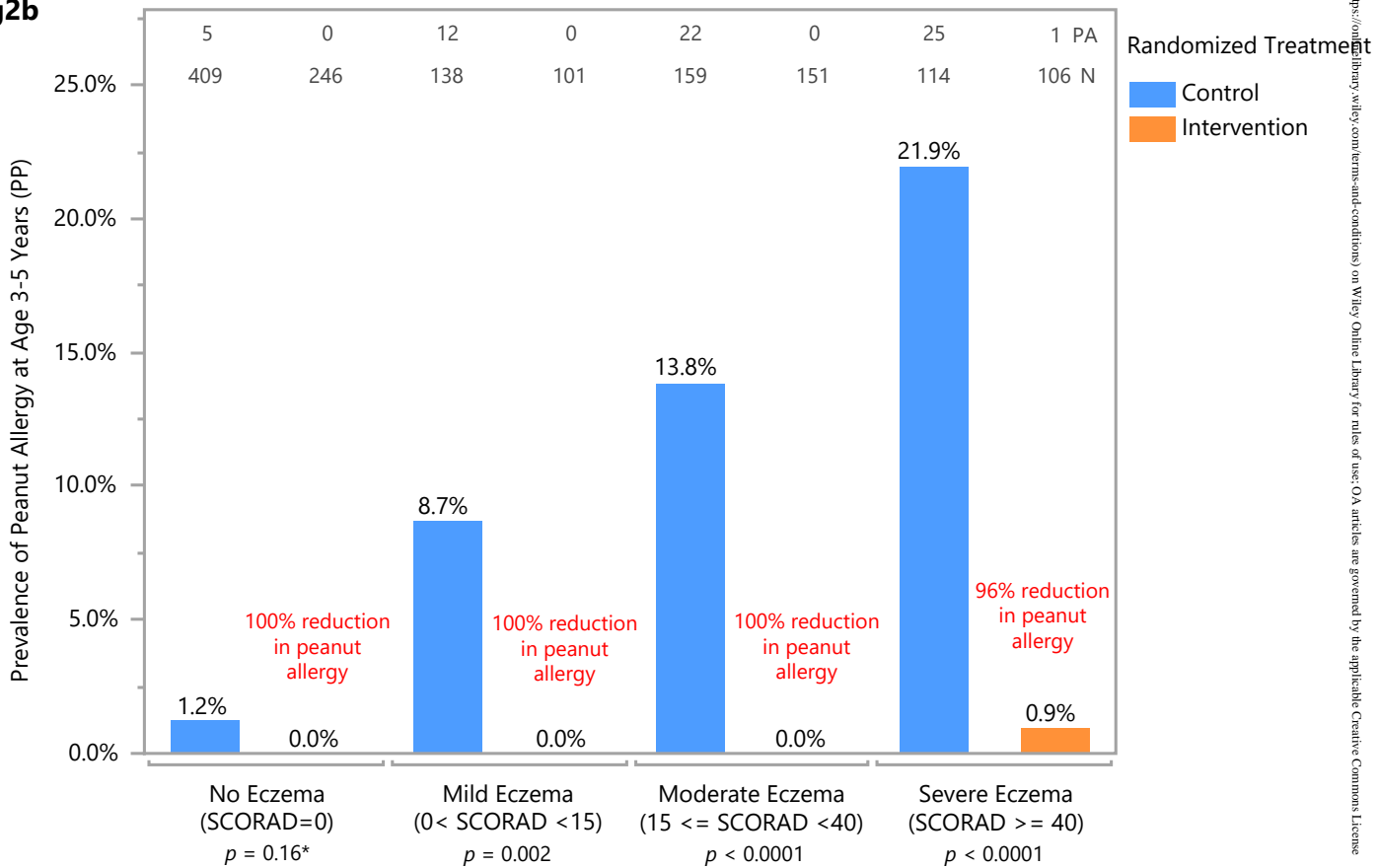


Figure 3

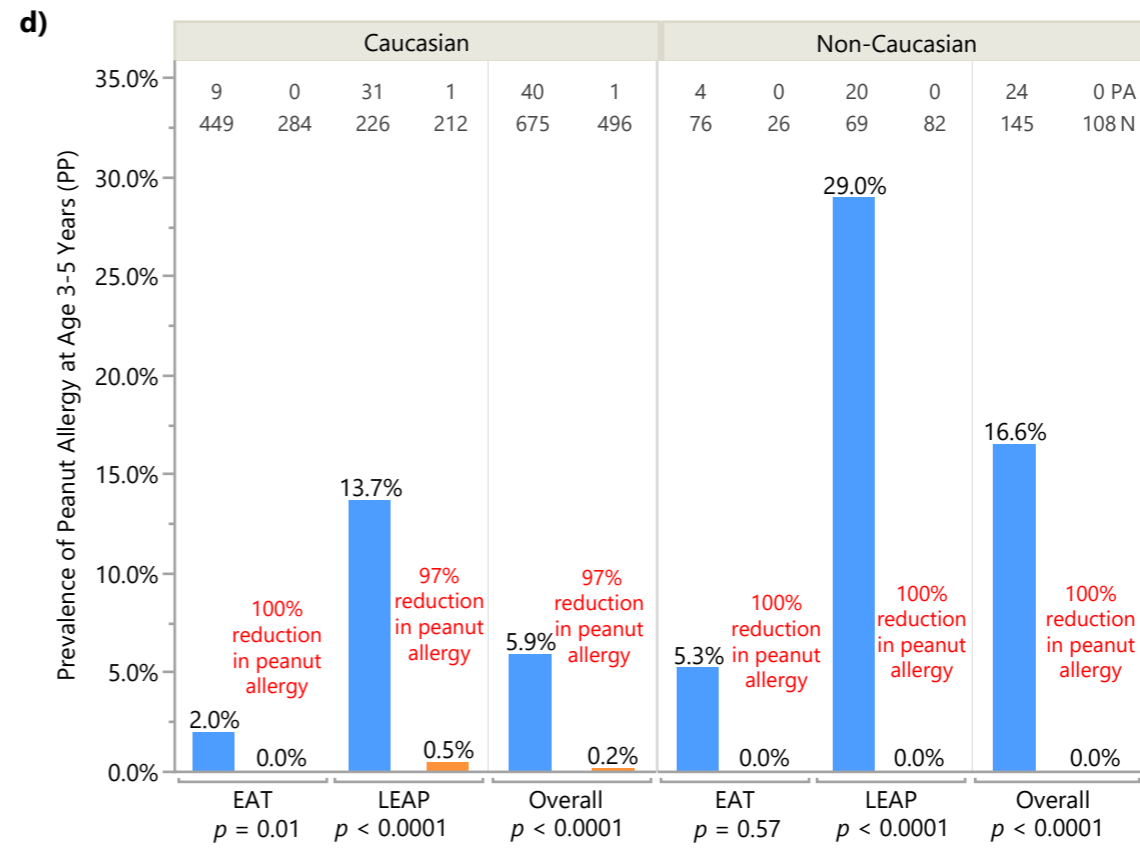
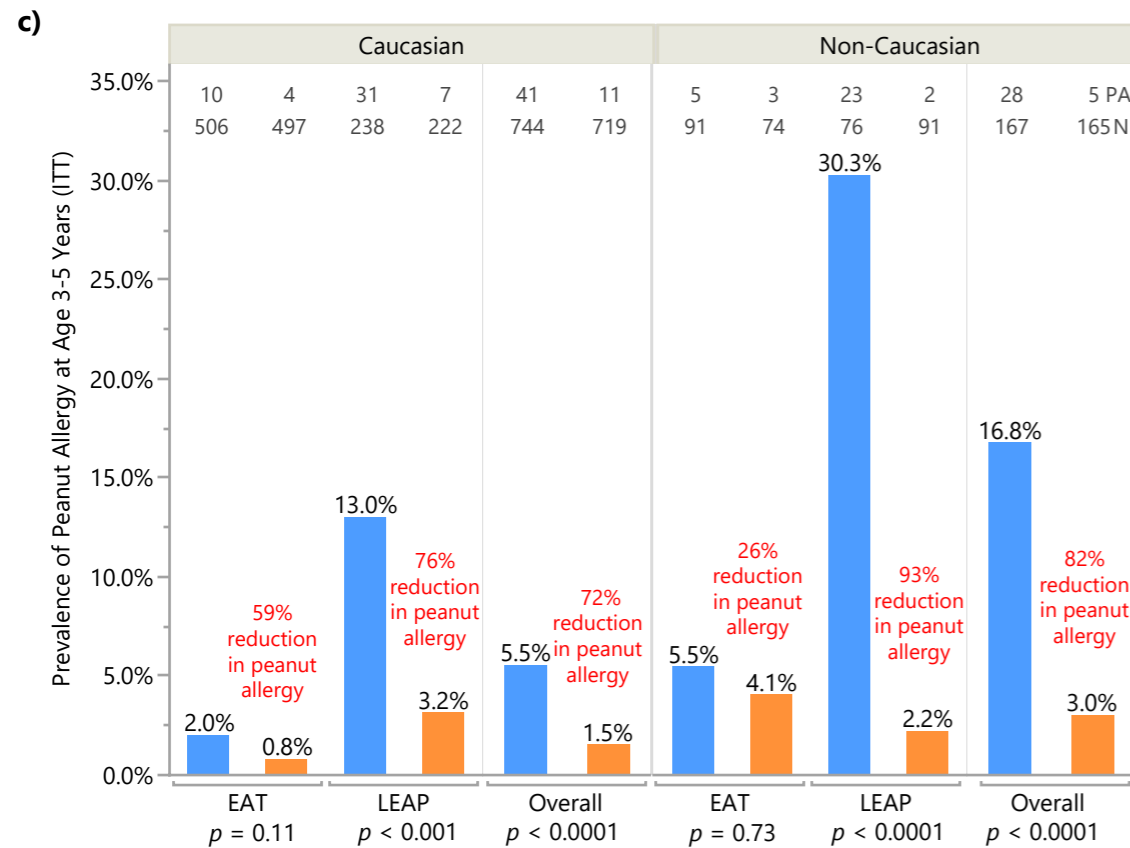
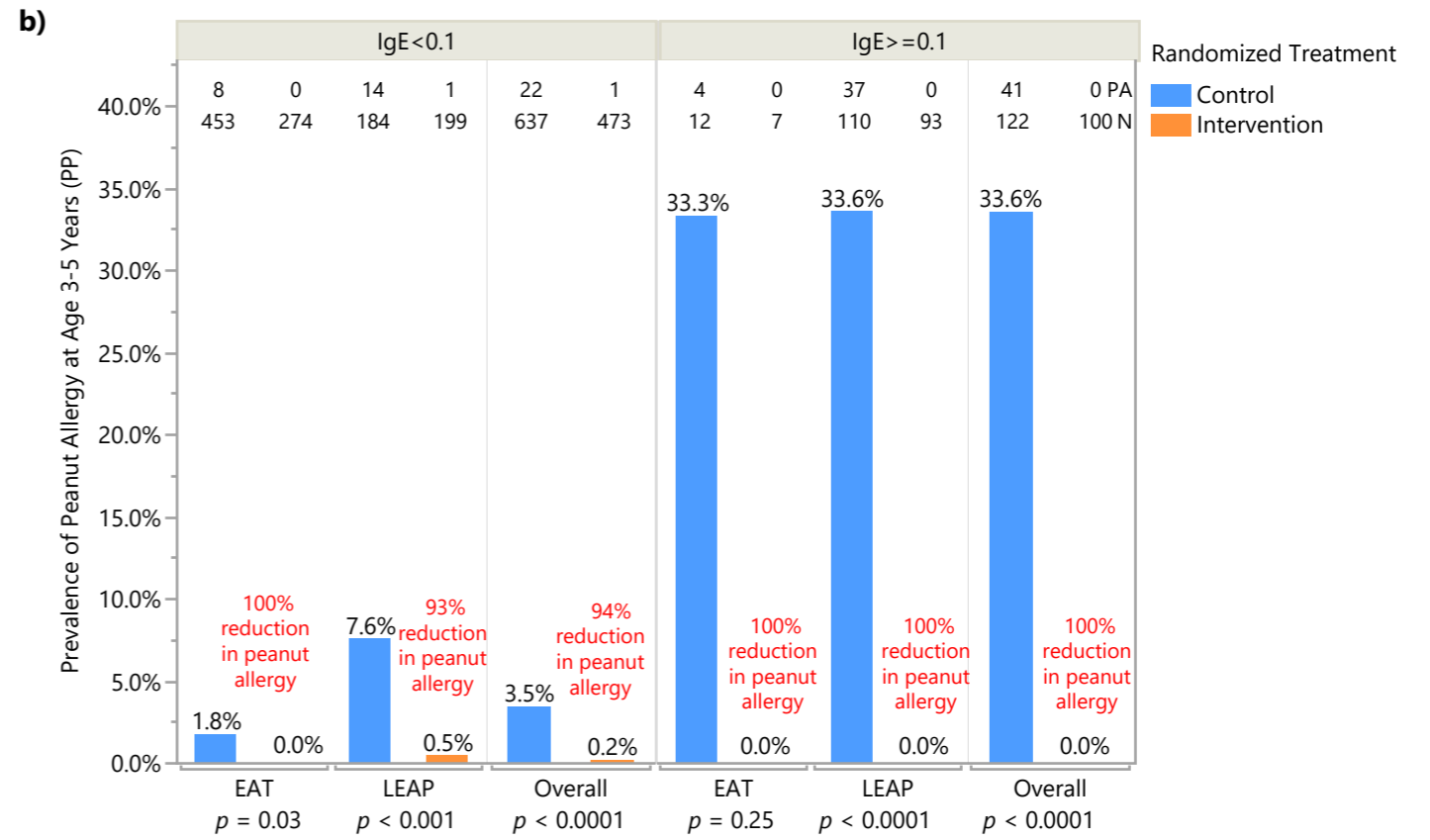
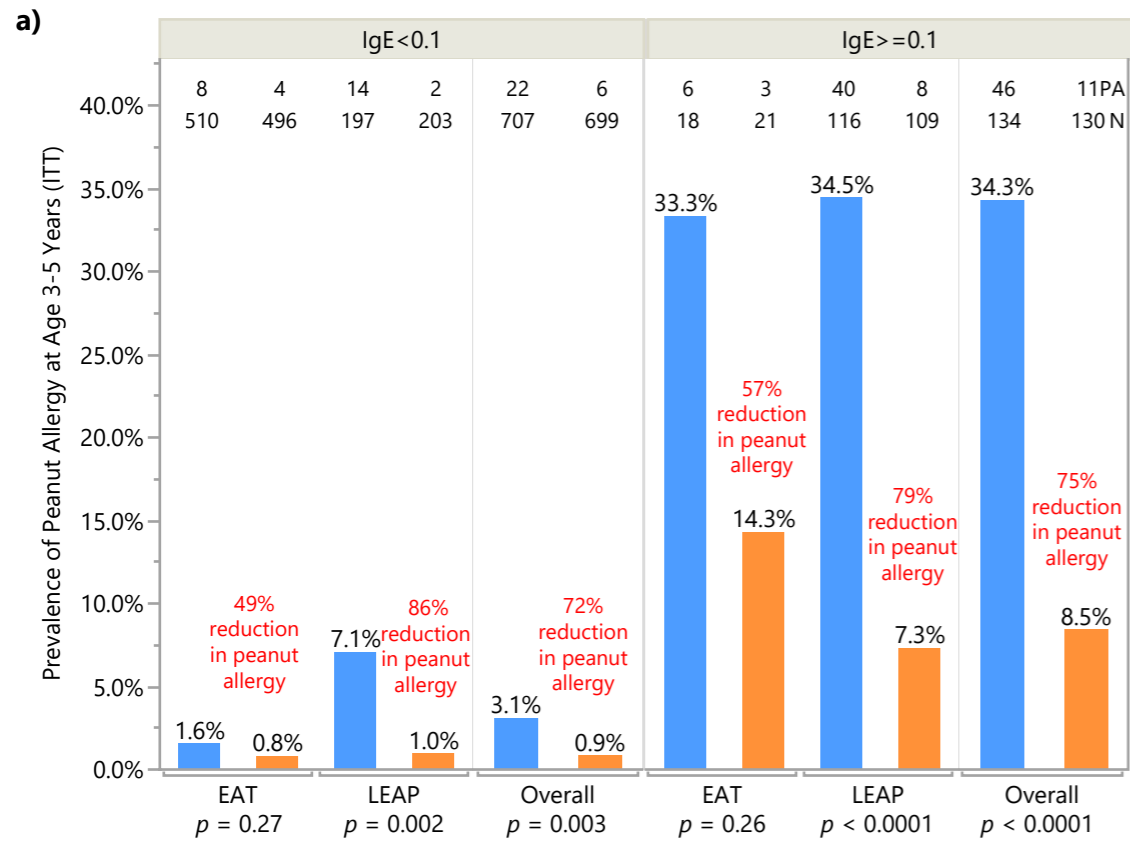


Figure 4

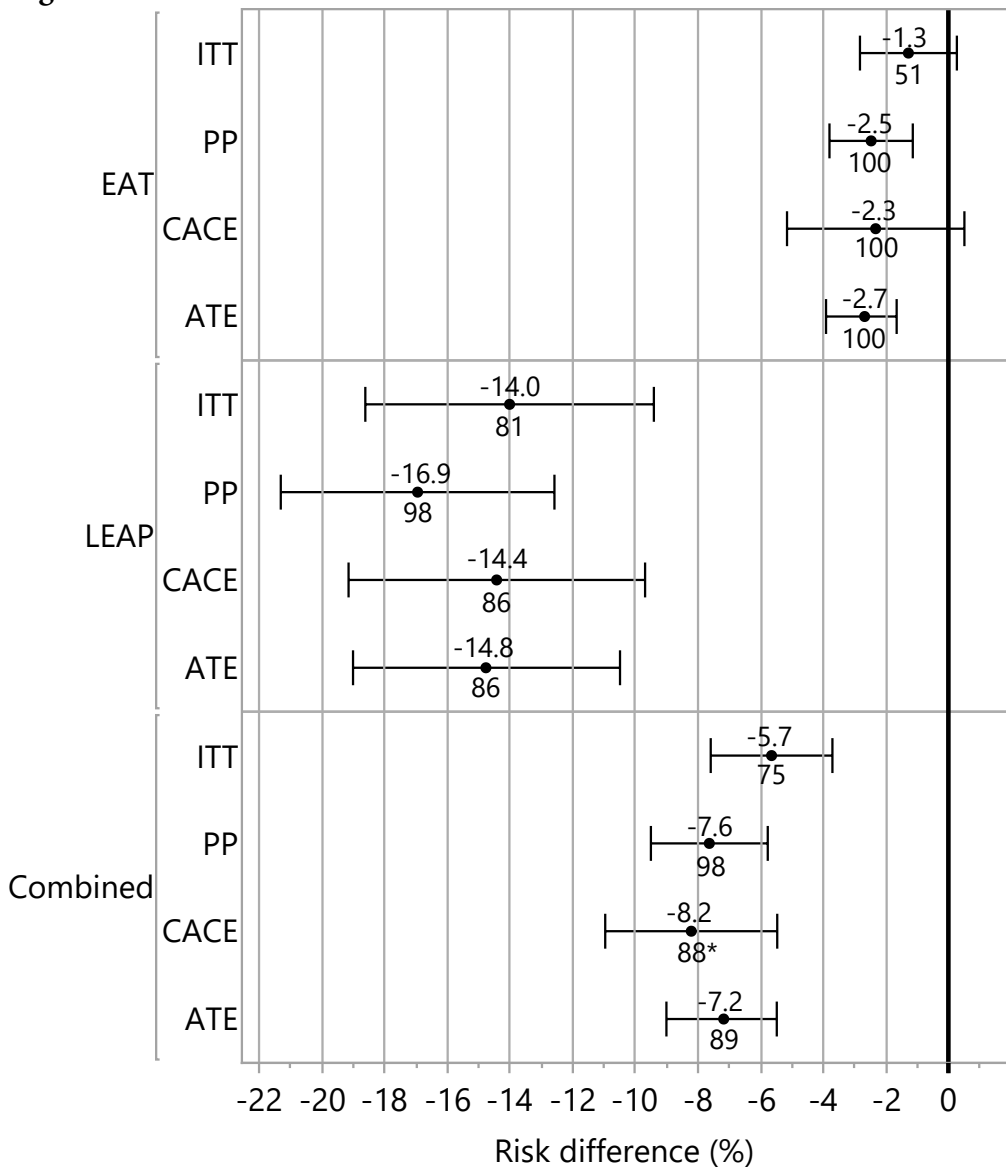


Figure 5

