Cohort study of electronic cigarette use: effectiveness and safety at 24 months

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INTRODUCTION
Despite their potential public health relevance, the current evidence on the safety and efficacy/effectiveness of e-cigarettes is scarce, and long-term data are urgently needed.1–7 The published prospective evidence on the efficacy/effectiveness for healthy participants consists of 2 randomised trials,8 9 2 single-arm small trials10–12 and 11 observational studies.13–23 However, these studies mostly included users of both tobacco and e-cigarettes followed for ≤12 months, used various assessment methods, and did not directly compare e-cigarette users and tobacco smokers.4 6 7 24 25 Moreover, the entire evidence on e-cigarette safety—at 6 months—is limited to 122 healthy participants, most of whom were also smoking tobacco for most of the follow-up.8 9 11 We previously reported the 12-month follow-up results26 of our 5-year study aimed at evaluating the long-term effects of e-cigarette use.27 We report the results of the 24-month follow-up, and include hospital discharge data.

METHODS
The protocol and 12-month results of this prospective cohort study are reported elsewhere,26 27 and registered in Clinicaltrials.gov (NCT01785537). In brief, we recruited adults (30–75 years) who were: (1) tobacco smokers of ≥1 tobacco cigarette daily for ≥6 months; (2) e-cigarette users of any type of e-cigarette for ≥6 months (3) dual users of tobacco and e-cigarettes for ≥6 months, via general practitioners, e-cigarette shops, internet advertisements and social networks.

Data were collected through a structured questionnaire, administered through phone interview and/or by internet, and follow-up will continue up to 60 months. Two investigators (MEF and LM) tested carbon monoxide levels in expired breath (Smokerlyzer piCO+, Bedfont Scientific) in a random sample of those declaring tobacco smoking abstinence (25% and 50% at 12 and 24 months, respectively).

The work was approved by Chieti University Ethics Committee; all participants provided written informed consent.

Outcome variables and data analysis
The primary outcome was the percentage of sustained (30 days) abstinence from tobacco smoking at 24 months. Other outcomes were the proportion of participants abstinent from both tobacco smoking and e-cigarette use, the number of tobacco cigarettes and/or e-cigarettes after 24 months, the difference in the number of tobacco cigarettes and/or e-cigarettes after 24 months, possibly related serious adverse events in 46.0% of the sample. The primary outcome was the percentage of sustained abstinence from tobacco cigarettes (25% and 50% at 12 and 24 months, respectively).

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ABSTRACT
Objective  To evaluate the safety and effectiveness of e-cigarettes, by comparing users of only e-cigarettes, smokers of only tobacco cigarettes and dual users.

Design  Prospective cohort study. We update previous 12-month findings and report the results of the 24-month follow-up.

Data sources  Direct contact and questionnaires by phone or via internet.

Methods  Adults (30–75 years) were classified as: (1) tobacco smokers, if they smoked ≥1 tobacco cigarette/day, (2) e-cigarette users, if they inhaled ≥50 puffs/week of any type of e-cigarette and (3) dual users, if they smoked tobacco cigarettes and also used e-cigarettes. Carbon monoxide levels were tested in 50% of those declaring tobacco smoking abstinence. Hospital discharge data were used to validate possibly related serious adverse events in 46.0% of the sample.

Main outcome measures  Sustained abstinence from tobacco cigarettes and/or e-cigarettes after 24 months, the difference in the number of tobacco cigarettes smoked daily between baseline and 24 months, possibly related serious adverse events.

Results  Data at 24 months were available for 229 e-cigarette users, 480 tobacco smokers and 223 dual users (overall response rate 68.8%). Of the e-cigarette users, 61.1% remained abstinent from tobacco (while 23.1% and 26.0% of tobacco-only smokers and dual users achieved tobacco abstinence). The rate (18.8%) of stopping use of either product (tobacco and/or e-cigarettes) was not higher for e-cigarette users achieving tobacco abstinence. The rate (18.8%) of stopping use of either product (tobacco and/or e-cigarettes) was not higher for e-cigarette users achieving tobacco abstinence. The rate (18.8%) of stopping use of either product (tobacco and/or e-cigarettes) was not higher for e-cigarette users achieving tobacco abstinence. The rate (18.8%) of stopping use of either product (tobacco and/or e-cigarettes) was not higher for e-cigarette users achieving tobacco abstinence. The rate (18.8%) of stopping use of either product (tobacco and/or e-cigarettes) was not higher for e-cigarette users achieving tobacco abstinence.
cigarettes smoked per day (and the proportion of participants reducing tobacco cigarette consumption by 50% or more between baseline and 24 months), self-reported health (assessed through the final item of the Italian version of the EuroQol EQ-5D-3L) and possibly related serious adverse events. Adverse events were both self-reported and gathered from data obtained from the regional hospital discharge administrative database (Italian Scheda di Dimissione Ospedaliera) for the residents in the Abruzzo Region (46.0% of the sample); and only self-reported for the rest of the sample. The details of the International Classification of Diseases, Ninth Revision, Clinical Modification (ICD-9-CM) codes used to extract adverse events are reported in the protocol and in the online supplementary material.

The differences by baseline cigarette use were evaluated using Kruskal-Wallis or one-way analysis of variance with Sidak correction for continuous variables, and χ² test for categorical variables. For the latter, when more than two categories were compared (such as for cigarette use), separate comparisons were made for one group versus the others, and different p values were computed. The difference in continuous variables (ie, number of cigarettes smoked per day) within groups between baseline and end of follow-up was evaluated through Wilcoxon matched-pairs signed-rank test.

Multivariable random-effect linear and logistic regressions, with geographical region as the cluster unit, were used to investigate potential predictors of continuous and categorical outcomes, respectively. We set eight multivariable models for the following outcomes: (A) tobacco smoking abstinence at 24 months; (B) abstinence from both tobacco smoking and e-cigarette use at 24 months; (C) possibly related serious adverse events at 24 months; (D) reduction by ≥50% of tobacco cigarette consumption from baseline to 24 months, by baseline cigarette use group (this model was fitted into two versions: the first included the whole sample of tobacco and dual users at baseline; the second included only the subsample of continuing smokers, excluding those who had quit and whose cigarette consumption was zero at follow-up); (E) reduction by ≥50% of tobacco cigarette consumption from baseline to 24 months, by 24-month cigarette use group (as most dual users switched to another group, this model and model G were needed to further investigate the potential predictors of tobacco smoking reduction among the switchers); (F) difference in the daily number of tobacco cigarettes smoked per day at 24 months compared with baseline, by baseline cigarette use group (this model was fitted into two versions: the first included the whole sample of tobacco and dual users at baseline; the second included only the subsample of continuing smokers, excluding those who had quit and whose cigarette consumption was zero at follow-up); (G) difference in the daily number of tobacco cigarettes smoked per day at 24 months compared with baseline, by 24-month cigarette use group and (H) difference in the self-rated health at 24 months compared with baseline. The users of e-cigarettes only at baseline were excluded from models D, E, F and G.

With the exceptions of models C, E and G, which included a limited number of successes and had to be fitted with a restricted set of covariates, for all other models the number of successes of categorical outcomes (excluding adverse events) approximated 10 successes for each recorded variable (thus avoiding overfitting requirements). It was thus decided a priori to include all recorded variables into all final models regardless of significance, unless inclusion would create multicollinearity or violate other assumptions. Each covariate was tested in its original form or transformed if needed: only the number of cigarettes smoked per day was transformed into its square root; because the results were similar to those with the covariate included in its untransformed version, it was thus kept in the model in its original form. Models A, B, D, F and H were adjusted for the following baseline characteristics: age, gender, body mass index, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-rated health and years of tobacco smoking (former smoking for e-cigarette-only users). We excluded physical activity from the final multivariate models due to substantial missing data (n=41) and its virtually null effect on any dependent variables. The number of tobacco cigarettes smoked per day was transformed to be included into models A, B, C and H: because no tobacco cigarettes were smoked at baseline by e-cigarette-only users, we recategorised the amount of smoking, using tertiles. Those smoking <10 tobacco cigarettes per day (or <50 puffs per day if e-cigarette-only smokers) at baseline were assigned to the lowest tertile of consumption; smokers of 10–19 tobacco cigarettes (or 50–100 puffs if e-cigarette-only users) per day at baseline were assigned to the intermediate tertile; smokers of 20 or more cigarettes (or 100 or more puffs if e-cigarette-only users) per day at baseline were classified into the highest tertile.

Potential interactions with the dependent variable and/or quadratic/cubic terms were investigated for all covariates. In logistic regression analyses, the outlier analysis was based on Pearson calculation and standardised residuals, the change in Pearson χ², and Dbeta influence statistics and leverage (hat diagonal matrix). The validity of the final linear regression models was assessed as follows. The assumption of constant error variance was checked graphically, plotting Pearson residuals versus fitted values, and formally, using the Cook-Weisberg test for heteroscedasticity. High leverage observations were identified by computing Pearson, standardised and studentised residuals, Cook’s D influence, Welsch distance and the hat diagonal matrix (LC Hamilton. Statistics with Stata: Version 12, Eighth Edition. Boston: Cengage 2013). We found <30 influential or high-leverage observations in almost all models. In all cases, we repeated the analyses excluding these observations, with no substantial changes, and we thus kept all observations in the models.

We had very few missing data for all outcome variables (<5) except self-rated health, which was not answered by 56 participants at baseline (but only 30 of them were kept in the 24-month assessment), and by 3 participants at the 24-month follow-up. Models A, B, C and H were rerun without self-rated health at baseline (~30 participants), with no appreciable variation, therefore it was retained. Model H was inevitably fitted with 33 missing observations. Given that the participants lost in model H were relatively few (3.3% of the sample) and balanced across baseline groups, no missing data imputation technique was adopted. Finally, the distribution of the difference in self-rated health was relatively skewed (Shapiro-Wilk p<0.01) and model G was set also using its cubic form. However, again the estimates of p values of cigarette use covariates were similar and the dependent variable was maintained in its original form to facilitate results interpretation.

The results of the logistic regression analyses are presented as ORs and the corresponding 95% CIs whereas the results of the linear regression analyses are presented as β-coefficients and 95% CIs. A two-tailed p value of 0.05 was considered significant for all analyses, which were performed using Stata V13.1 (Stata Corp, College Station, Texas, USA, 2014).
RESULTS
The flow of the participants is shown in figure 1 and baseline characteristics are reported in online supplementary table S1. After 24 months, 61.1% of the 229 baseline e-cigarette-only users were still abstinent from tobacco smoking; 23.1% of the 480 baseline tobacco smokers and 26.0% of the 223 baseline dual users achieved tobacco abstinence (p<0.001 for e-cigarette-only users vs tobacco smokers or dual users; table 1). The proportion of participants who achieved complete abstinence (who were using neither tobacco cigarettes nor e-cigarettes) did not significantly differ by baseline use group: 18.8%, 17.5% and 14.3% among e-cigarette users, tobacco smokers and dual users, respectively (all p>0.05).

The proportion of participants reducing tobacco cigarette consumption by 50% or more, or by ≥5 tobacco cigarettes per day, and the average daily number of cigarettes, did not change by baseline group (table 1; all p>0.05). Importantly, the majority (83.4%) of dual users at baseline abandoned e-cigarettes and continued to smoke only tobacco (57.4%), or quit tobacco (11.7%) or both tobacco cigarettes and e-cigarettes (14.3%) during follow-up (table 1). Therefore, a second analysis was conducted stratifying by baseline and 24-month cigarette use, in order to assess the variation of the pattern of consumption among the switchers as well. Among the 603 tobacco-only smokers or dual users at baseline, 21 of the 40 (52.5%) participants who started or continued dual use during the follow-up reduced tobacco cigarette consumption by ≥50%, while only 67 of the 489 (13.7%) participants who started or continued only tobacco smoking showed a ≥50% cigarette reduction (p<0.001, table 1).

No significant differences in average self-rated health were noted by baseline group (all p>0.05). A substantial improvement, however, was observed for tobacco smokers or dual users who switched to e-cigarettes only (+1.1 or 1.0 in EuroQol mean score, respectively; p<0.05; table 1).

We recorded an identical number of mouth irritations (n=38) and possibly related serious adverse events (n=38; table 1). Potential adverse events were reported by 4.4%, 2.9% and 6.3% of baseline e-cigarette users, tobacco smokers and dual users, respectively (p<0.05 for the comparison of tobacco smokers vs dual users). Mouth irritation rates were 6.1%, 4.2% and 1.8% in the above groups (p<0.05 for the comparison of e-cigarette users vs dual users). The distribution of adverse events by baseline and 24-month group is reported in figure 1. The characteristics of the participants reporting a serious adverse event, and its type, are reported in online supplementary table S2. Also, the cigarette use throughout the follow-up of the 38 participants experiencing a possibly related serious adverse event is shown in online supplementary figure S1. Notably, most tobacco-only smokers at baseline, who experienced an adverse event, remained tobacco-only smokers or quit both tobacco smoking and e-cigarette use. In contrast, most of the e-cigarette-only and dual users at baseline who experienced a serious adverse event (n=24), switched group during the 24-month follow-up: 13 switched to tobacco smoking only, and 5 quit both tobacco cigarettes and e-cigarettes.

Multivariate analyses substantially confirmed univariate results (table 2): when several potential confounders were adjusted for, tobacco smoking abstinence remained significantly more likely among e-cigarette users (adjusted OR 5.56; 95% CI 3.89 to 7.95; p<0.001); the likelihood of abstinence from tobacco smoking and e-cigarette use, as well as self-reported health, did not significantly vary by baseline group (p>0.05); the probability of halving or reducing the average number of daily tobacco cigarettes smoked did not change by baseline group, but tobacco smokers who started dual use or dual users who switched to e-cigarettes only (+1.1 or 1.0 in EuroQol mean score, respectively; p<0.05; table 1).

No significant differences in the following adverse events were noted by baseline group (all p>0.05): mouth irritation (6.1%, 4.2% and 1.8% in the above groups), serious adverse events related to e-cigarettes use (6.3%, 4.4% and 2.9% in the above groups), serious adverse events related to tobacco smoking use (6.1%, 4.2% and 1.8% in the above groups), serious adverse events related to both e-cigarettes and tobacco smoking use (6.3%, 4.4% and 2.9% in the above groups), serious adverse events related to tobacco smoking and e-cigarettes use (6.1%, 4.2% and 1.8% in the above groups), serious adverse events related to both tobacco smoking and e-cigarettes use (6.1%, 4.2% and 1.8% in the above groups), and serious adverse events related to tobacco smoking, e-cigarettes and other factors (6.1%, 4.2% and 1.8% in the above groups).

Figure 1 Flow of the participants, number of quitters and possibly related serious adverse events.

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Figure 1 Flow of the participants, number of quitters and possibly related serious adverse events.
### Table 1  Main outcomes at 24 months

| Cigarette use at baseline | E-cigarettes only (n=229) | Tobacco cigarettes only (n=480) | Dual use (n=223) | p Value<sup>a</sup> | p Value<sup>b</sup> | p Value<sup>c</sup> |
|--------------------------|---------------------------|-------------------------------|------------------|--------------------|--------------------|--------------------|
| 1. Cigarette use at 24 months | Tobacco cigarettes only, per cent (n) | Continuous tobacco abstinence from baseline or cessation from tobacco smoking during follow-up<sup>+</sup> | 61.1 (140) | 23.1 (111) | 26.0 (58) | <0.001 | 0.4 | <0.001 |
| | | | 38.9 (89) | 76.9 (369) | 74.0 (165) | <0.001 | 0.4 | <0.001 |
| | All product use (cigarettes and/or e-cigarettes), per cent (n) | Quit using any product (either tobacco and/or e-cigarette) | 18.8 (43) | 17.5 (84) | 14.3 (32) | 0.7 | 0.3 | 0.20 |
| | | Using e-cigarettes only | 42.4 (97) | 5.6 (27) | 11.7 (26) | <0.001 | 0.004 | <0.001 |
| | | Dual use (tobacco cigarettes and e-cigarettes) | 8.3 (19) | 1.3 (6) | 16.6 (37) | <0.001 | <0.001 | 0.007 |
| | | Smoking tobacco cigarettes only | 30.6 (70) | 75.6 (363) | 57.4 (128) | <0.001 | <0.001 | <0.001 |
| 2. Number of tobacco cigarettes | Mean number of tobacco cigarettes daily at 24 months (SD) | Stratified by baseline group | 10.0 (8.4) | 11.2 (9.8) | 0.09 | |
| | | Stratified by cigarette use at 24 months | Started or continued tobacco cigarettes only | 0.0 (0.0) | 14.8 (8.1) | 15.8 (9.1) | – | 0.2 | – |
| | | | Started or continued dual use only | 0.0 (0.0) | 19.0 (6.3) | 14.0 (9.5) | – | 0.2 | – |
| | | | (p=0.2)<sup>†</sup> | (p=0.3)<sup>†</sup> | |
| | Percentage of participants reducing tobacco cigarettes of 50% or more from baseline to 24 months | Stratified by baseline group | 34.5 | 39.7 | – | 0.2 | – |
| | | Stratified by product use at 24 months | Started or continued tobacco cigarettes only | – | 14.1 (n=51) | 9.4 (n=12) | – | 0.2 | – |
| | | | Started or continued dual use only | – | 66.7 (n=4) | 50.0 (n=17) | – | 0.5 | – |
| | | | (p=0.001)<sup>†</sup> | (p=0.001)<sup>†</sup> | |
| | Percentage of participants who smoked ≥5 tobacco cigarettes less between baseline and 24 months | Stratified by baseline group | – | 38.6 | 42.5 | – | 0.3 | – |
| | | Stratified by product use at 24 months | Started or continued tobacco cigarettes only | – | 23.5 (n=85) | 22.1 (n=28) | – | 0.7 | – |
| | | | Started or continued dual use only | – | 66.7 (n=4) | 55.9 (n=19) | – | 0.6 | – |
| | Mean difference in the daily number of tobacco cigarettes between 24 months and baseline (SD) | Stratified by baseline group | – | −4.1 (8.1) | −4.0 (11.8) | – | 0.9 | – |
| | | Stratified by product use at 24 months | Started or continued tobacco cigarettes only | – | −1.6 (6.4) | −0.8 (9.6) | – | 0.2 | – |
| | | | Started or continued dual use only | – | −5.3 (3.3) | −6.5 (10.7) | – | 0.8 | – |
| | | | (p=0.2)<sup>†</sup> | (p=0.003)<sup>†</sup> | |
| 3. Self-rated health—Mean difference between 24 months and baseline (SD) | Stratified by baseline group | 0.0 (1.7) | 0.0 (1.4) | −0.1 (1.7) | 0.9 | 0.4 | 0.5 |
| | Stratified by product use at 24 months | Quit using any product (either tobacco and/or e-cigarette) | +0.2 (1.6) | +0.3 (1.4) | −0.2 (2.1) | 0.7 | 0.14 | 0.4 |
| | | Using e-cigarettes only | +0.3 (1.4) | +1.1 (1.7) | +1.0 (1.6) | 0.014 | 0.8 | 0.03 |
| | | Dual use (tobacco cigarettes and e-cigarettes) | −0.3 (1.7) | +0.3 (1.5) | +0.2 (1.7) | 0.4 | 0.9 | 0.3 |
| | | Smoking tobacco cigarettes only | −0.5 (1.9) | −0.1 (1.4) | −0.4 (1.4) | 0.041 | 0.039 | 0.7 |
| 4. Safety—possibly related sAEs, per cent (n) | Stratified by baseline group | Mouth irritation | 6.1 (14) | 4.2 (20) | 1.8 (4) | 0.27 | 0.10 | 0.019 |

<sup>a</sup> Cigarette smoking (continued or relapsed)†<sup>b</sup> Quitting smoking only during follow-up†<sup>c</sup> Smoking tobacco cigarettes only during follow-up†<sup>d</sup> Smoking only e-cigarettes during follow-up†<sup>e</sup> Smoking tobacco cigarettes and e-cigarettes during follow-up†<sup>f</sup> Smoking e-cigarettes only during follow-up†<sup>g</sup> Smoking at baseline or cessation from smoking during follow-up†<sup>h</sup>
### Table 1

Cigarette use at baseline

|                     | E-cigarettes only (n=229) | Tobacco cigarettes only (n=480) | Dual use (n=223) |
|---------------------|---------------------------|--------------------------------|-----------------|
| **p Value**         |                           |                                |                 |
| Any sAE¶            | 4.4 (10)                  | 2.9 (14)                       | 6.3 (14)        |
| Any cancer           | 2.6 (6)                   | 1.0 (5)                        | 1.4 (3)         |
| Quit using any product (either tobacco and/or e-cigarette) | 4.6 (2) | 6.0 (5) | 9.4 (3) |
| Using e-cigarettes only | 2.1 (2) | 3.7 (1) | 11.5 (3) |
| Dual use (tobacco cigarette and e-cigarettes) | 5.3 (1) | 0.0 | 0.0 (0) |
| Smoking tobacco cigarettes only | 7.1 (5) | 2.2 (8) | 6.3 (8) |

*p Value for the following groups at baseline: A tobacco cigarettes only versus e-cigarettes only; B tobacco cigarettes only versus dual use; C e-cigarettes only versus dual use.

*p Value for the comparisons of the following groups at baseline: A tobacco cigarettes only versus e-cigarettes only at baseline who quit e-cigarette use and did not start tobacco smoking; B smokers of tobacco cigarettes only at baseline who quit tobacco cigarette use and did not start e-cigarette use; C dual users at baseline who quit both tobacco smoking and e-cigarette use; D e-cigarette users at baseline who continued to use only e-cigarettes; E tobacco cigarette smokers at baseline who switched to e-cigarettes; F dual users at baseline who quit smoking and continued to use e-cigarettes only.

‡p Value for the comparison of the tobacco-only smokers at baseline who started or continued tobacco cigarettes only versus tobacco-only smokers at baseline who started or continued dual use only.

§p Value for the comparison of the dual users at baseline who started or continued tobacco cigarettes only versus dual users at baseline who continued dual use only.

¶Mouth irritation excluded; chronic obstructive pulmonary disease, stroke, heart failure, myocardial infarction, angina, pneumonia and any type of cancer included.

sAE, serious adverse event.

### DISCUSSION

Switching or quitting cigarette use (n=276, 29.6%) or stopped using either product (n=159; 17.1%) using either product, of 24 months. Overall, 435 participants switched (n=276; 29.6%) or stopped using either product (n=159; 17.1%) using either product, 29 to tobacco smoking only, 4 to e-cigarettes only and 6 were lost to follow-up, 91 remained abstinent and 34 relapsed (27.2%).

To date, this is the only study to directly compare smokers of tobacco cigarettes and e-cigarettes. Other secondary findings on CO levels, the other predictors of tobacco smoking and adverse events, are reported in the online supplementary material.
# Table 2  Tobacco smoking and/or e-cigarette use abstinence or cessation, possibly related adverse events, difference in daily tobacco cigarette consumption and self-reported health: results of the multivariate analyses

| Outcomes at 24 months | Adjusted OR (95% CI) | p Value | Crude OR (95% CI) | p Value |
|-----------------------|----------------------|---------|-------------------|---------|
| Continuous tobacco abstinence from baseline or quit tobacco smoking during follow-up*,† | Tobacco cigarettes only at baseline 1 (ref cat) – – 1 (ref cat) – – | E-cigarettes only at baseline 5.56 (3.89 to 7.95) <0.001 5.23 (3.72 to 7.35) <0.001 | Both tobacco and e-cigarettes at baseline 1.25 (0.85 to 1.84) 0.2 1.17 (0.81 to 1.69) 0.4 |
| Quit all cigarettes (electronic and/or traditional)* | Tobacco cigarettes only at baseline 1 (ref cat) – – 1 (ref cat) – – | E-cigarettes only at baseline 1.12 (0.73 to 1.72) 0.6 1.09 (0.73 to 1.64) 0.7 | Both tobacco and e-cigarettes at baseline 0.94 (0.59 to 1.49) 0.8 0.79 (0.51 to 1.23) 0.3 |
| Safety—possibly related sAEs‡ | Tobacco cigarettes only at baseline 1 (ref cat) – – 1 (ref cat) – – | E-cigarettes only at baseline 1.48 (0.63 to 3.47) 0.4 1.52 (0.66 to 3.48) 0.3 | Both tobacco and e-cigarettes at baseline 2.40 (1.09 to 5.26) 0.029 2.23 (1.04 to 4.76) 0.038 |
| Reduction of tobacco cigarette consumption by 50% or more from baseline to 24 months§ | Tobacco cigarettes only at baseline 1 (ref cat) – – 1 (ref cat) – – | Both tobacco and e-cigarettes at baseline 1.28 (0.90 to 1.82)¶ 0.2 1.25 (0.89 to 1.76)¶ 0.2 | 0.99 (0.57 to 1.71)** 0.9 1.25 (0.76 to 2.05)** 0.4 |
| Reduction of tobacco cigarette consumption by 50% or more from baseline to 24 months†† | Tobacco or dual users at baseline who switched to or continued with tobacco smoking only 1 (ref cat) – – 1 (ref cat) – – | Tobacco or dual users at baseline who started or continued dual use 8.48 (4.05 to 17.8) <0.001 7.47 (3.81 to 14.7) <0.001 | 0.99 (0.57 to 1.71)** 0.9 1.25 (0.76 to 2.05)** 0.4 |

| Adjusted coefficient (95% CI) | Raw coefficient (95% CI) |
|-----------------------------|--------------------------|
| Difference in the daily number of tobacco cigarettes from 24 months to baseline‡‡ | Tobacco cigarettes only at baseline 0 (ref cat) – – 0 (ref cat) – – | Both tobacco and e-cigarettes at baseline 0.44 (–0.88 to 1.75)¶ 0.5 0.10 (–1.41 to 1.61)¶ 0.9 | 0.75 (–1.13 to 2.63)** 0.4 0.87 (–0.42 to 2.15)** 0.2 |
| Difference in the self-reported health score from 24 months to baseline¶¶ | Tobacco cigarettes only at baseline 0 (ref cat) – – 0 (ref cat) – – | E-cigarettes only at baseline 0.14 (–0.08 to 0.35) 0.2 –0.03 (–0.27 to 0.22) 0.8 | Continued |
Table 2 Continued

| Both tobacco cigarettes and e-cigarettes at baseline | Adjusted coefficient (95% CI) | Raw coefficient (95% CI) |
|---------------------------------------------------|-------------------------------|--------------------------|
| **Continued**                                     |                               |                          |

*Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking (former smoking for e-cigarette users), number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 902 participants were included in the final model due to 30 missing items in the self-reported health item at baseline.

†Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day.

§Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 682 participants were included in the final model due to 21 missing items in the self-reported health item at baseline.

¶Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 899 participants were included in the final model due to 30 missing items in the self-reported health item at baseline and 3 missing items in the self-reported health item at baseline.

**Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 682 participants were included in the final model due to 21 missing items in the self-reported health item at baseline.

††Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 529 participants were included in the final model due to 21 missing items in the self-reported health item at baseline.

‡‡Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, BMI, marital status, educational level, occupation, alcohol use, hypertension, hypercholesterolaemia, diabetes, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day (or puffs per day for e-cigarette only smokers). A total of 529 participants were included in the final model due to 21 missing items in the self-reported health item at baseline.

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**Computed from a model that included only the subsample of baseline smokers who continued smoking at 24 months (excluding those who had quit and whose cigarette consumption was zero at follow-up).

†††Restricted to tobacco only smokers or dual users at baseline who did not quit or switch to e-cigarettes only (n=529). Random-effect logistic regression with region as the cluster level, adjusting for the following baseline characteristics: age, gender, self-reported health, years of tobacco smoking, number of tobacco cigarettes smoked per day.

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Information on dual users. However, when baseline and 24-month use data were considered together, we found that baseline and 24-month use data were more likely to reduce the number of daily cigarettes smoked compared to those who returned to or continued to use e-cigarettes only, but it may facilitate the reduction of smoking, particularly for those who returned to only smoking after quitting during the 24-month follow-up.

CONCLUSIONS

The first 2 years of the study confirmed that switching completely to e-cigarettes may help tobacco smokers quit smoking, but it may facilitate the likelihood of quitting tobacco use of e-cigarettes and not using e-cigarettes. The results were similar to those of previous studies, but not using e-cigarettes. Adverse events were more prevalent in participants who had used e-cigarettes before as compared to those who had never used them. The results were similar to those of previous studies, but they were not always consistent. The results were similar to those of previous studies, but they were not always consistent. The results were similar to those of previous studies, but they were not always consistent. The results were similar to those of previous studies, but they were not always consistent. The results were similar to those of previous studies, but they were not always consistent.
next years of follow-up will help clarify safety concerns, which remain the most important issue to support policies on e-cigarettes use.

What this paper adds

- Despite the potential public health relevance, the current evidence on long-term safety and efficacy/effectiveness of e-cigarettes is scarce and conflicting.
- After 24 months of a prospective follow-up, most users of e-cigarettes alone were able to remain abstinent from tobacco smoking.
- Dual use of e-cigarettes with tobacco cigarettes did not encourage quitting tobacco or e-cigarette use, but may be helpful to reduce tobacco consumption.

Collaborators

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Contributors

All the authors and the collaborators participated in the design, analysis and interpretation of the study. LM, MEF, CLV, WR, MFi and PV were involved in all phases of the study. MFe and CM collected baseline and follow-up data, and participated in data-analysis. LM and MEF tested CO levels, linked hospital admissions and wrote the manuscript. LM is the guarantor for all data.

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Competing interests

None declared.

Patient consent

Obtained.

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Data sharing statement

The raw data set is available from the corresponding author on request.

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