




# BMJ Open Time-varying association between cigarette and ENDS use on incident hypertension among US adults: a prospective longitudinal study

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## ABSTRACT

**Objective** Electronic nicotine delivery systems (ENDS) products have emerged as the most popular alternative to combustible cigarettes. However, ENDS products contain potentially dangerous toxicants and chemical compounds, and little is known about their health effects. The aim of the present study was to examine the prospective association between cigarette and ENDS use on self-reported incident hypertension.

**Design** Longitudinal cohort study.

**Setting** Nationally representative sample of the civilian, non-institutionalised population in the USA.

**Participants** 17 539 adults aged 18 or older who participated at follow-up and had no self-reported heart condition or previous diagnosis of hypertension or high cholesterol at baseline.

**Measures** We constructed a time-varying tobacco exposure, lagged by one wave, defined as no use, exclusive established use (every day or some days) of ENDS or cigarettes, and dual use. We controlled for demographics (age, sex, race/ethnicity and household income), clinical risk factors (family history of heart attack, obesity, diabetes and binge drinking) and smoking history (cigarette pack-years).

**Outcomes** Self-reported incident hypertension diagnosis.

**Results** The self-reported incidence of hypertension was 3.7% between wave 2 and wave 5. At baseline, 18.0% (n=5570) of respondents exclusively smoked cigarettes; 1.1% (n=336) exclusively used ENDS; and 1.7% (n=570) were dual users. In adjusted models, exclusive cigarette use was associated with an increased risk of self-reported incident hypertension compared with non-use (adjusted HR (aHR) 1.21, 95% CI 1.06 to 1.38), while exclusive ENDS use (aHR 1.00, 95% CI 0.68 to 1.47) and dual use (aHR 1.15, 95% CI 0.87 to 1.52) were not.

**Conclusions** We found that smoking increased the risk of self-reported hypertension, but ENDS use did not. These results highlight the importance of using prospective longitudinal data to examine the health effects of ENDS use.

## INTRODUCTION

Cigarette smoking is the leading cause of premature mortality in the USA,<sup>1 2</sup> and a

## STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ In this study, we examine the time-varying association between cigarette smoking and electronic nicotine delivery systems (ENDS) use on the incidence of hypertension among a nationally representative sample of US adults.
- ⇒ By examining the prospective incident cases of hypertension and using a lagged time-varying cigarette/ENDS exposure variable, our study does not have the same concerns with reverse causation that have been identified in cross-sectional studies.
- ⇒ We also controlled for the potential confounding of smoking history, measured as cigarette pack-years, which is important because most adults who use ENDS are either currently smoking cigarettes or have smoked cigarettes in the past.
- ⇒ Our study was limited by relying on self-reported hypertension, as systolic and diastolic blood pressure measures were not available.
- ⇒ Our non-randomised data mean that our results could be affected by unmeasured confounding, and the results should be interpreted with the same level of caution required in all prospective longitudinal studies.

significant proportion of smoking-attributable deaths are related to cardiovascular disease.<sup>3 4</sup> Smoking is known to cause an acute rise in blood pressure<sup>5</sup> and contribute to arterial stiffness,<sup>6</sup> and has been associated with an increased risk of developing hypertension.<sup>5 7–10</sup> Hypertension, in turn, is an important risk factor for most downstream cardiovascular diseases.<sup>11–13</sup> The health hazards of smoking on cardiovascular disease underscore the importance of further reducing smoking prevalence in the general population and the continued need to promote smoking cessation among adults who smoke.

Electronic nicotine delivery systems (ENDS) products became widely available



around 2010, and they refer to a broad range of devices that produce an aerosol from heating an e-liquid. ENDS products quickly emerged as the most popular alternative to combustible cigarettes in the USA, as their prevalence doubled among young adults between 2014 and 2018,<sup>14</sup> and more than 5.6million US adults reported ENDS use in 2018–2019.<sup>15</sup> Some adults use ENDS products as a way to help them quit smoking<sup>15 16</sup> and because they are generally believed to be less harmful than combustible cigarettes,<sup>17 18</sup> and it has been argued that their use should be encouraged as part of a harm minimisation strategy.<sup>17</sup> However, non-smoking youth are also using ENDS products,<sup>19</sup> raising concerns about tobacco use renormalisation. Furthermore, ENDS contain toxicants and chemical compounds that are potentially dangerous, including aldehydes, carbonyl, nicotine and flavouring additives.<sup>20 21</sup> Very little is known about the health consequences of ENDS product use,<sup>21</sup> and we need reliable and rigorous estimates of their health effects.

One potential consequence of ENDS product use may be an increased risk of hypertension. Evidence of a short-term elevation in both systolic blood pressure and diastolic blood pressure from ENDS product use has been found in experimental studies,<sup>22</sup> and a recent epidemiological study found evidence of a cross-sectional association between ENDS product use and self-reported hypertension among adults.<sup>23</sup> However, cross-sectional research on the cardiovascular risks of ENDS use has resulted in a contentious debate<sup>24–28</sup> largely centred around the issue of reverse causation.<sup>27</sup> Without information on the timing of both ENDS use and disease outcome, it is simply not possible to know whether ENDS use came before or after the disease outcome. The latter is likely common, given the use of ENDS by some smokers trying to quit after being diagnosed with a cardiovascular disease.<sup>29</sup> Therefore, the results from these cross-sectional studies need to be interpreted with caution. Researchers have highlighted the need for prospective longitudinal data to better understand the temporal ordering between ENDS use and cardiovascular disease endpoints.<sup>22 28</sup>

In this study, we use data from a nationally representative prospective cohort study to examine the time-varying association between cigarette and ENDS use on the incidence of self-reported hypertension, which limits potential concerns with reverse causation. In addition, we developed a composite exposure variable combining current cigarette and ENDS use to examine the relative contribution of exclusive cigarette use, exclusive ENDS use and dual cigarette/ENDS use, compared with no use. We also adjust for cigarette smoking history.

## METHODS

### Data

We used data on adults from wave 1 to wave 5 (2013–2019) of the Population Assessment of Tobacco and Health (PATH) Study, a publicly available dataset. However, this analysis used the restricted-use files<sup>30</sup> in order to

use variables such as continuous age and cigarette pack-years. These variables are not available in the public-use files. Further details on how to access the restricted-use data are described in the PATH Study Restricted Use Files User Guide (available at <https://doi.org/10.3886/ICPSR36231.v33>).

The PATH study is an ongoing, nationally representative cohort study of the civilian, non-institutionalised population in the USA. A stratified area probability design was used to sample geographical segments from 156 geographical primary sampling units. An address-based sampling frame was then used to randomly select households based on residential addresses derived from the United States Postal Service. Once households were identified, an introductory letter and brochure were mailed to sampled addresses followed by an in-person field interview within 2 weeks. A two-phase sampling procedure was used to select adults within sampled households for the in-person interview.<sup>31</sup> African-Americans and tobacco users were oversampled in relation to population proportions, and weighting procedures were adjusted for over-sampling and non-response based on US Census Bureau Data. Data were collected from September 2013 to December 2014 for wave 1 (response rate among screened households, 74.0%); from October 2014 to October 2015 for wave 2 (response rate, 83.2%); from October 2015 to October 2016 for wave 3 (response rate, 78.4%); from December 2016 to January 2018 for wave 4 (response rate, 73.5%); and from December 2018 to November 2019 for wave 5 (response rate, 69.4%). All PATH survey interviews were completed in person using audio-assisted self-interviewing administrations, available in English or Spanish. Data collection protocols were used to ensure that follow-up interviews were close to the anniversary of their participation in the previous wave.<sup>32</sup> Further details about the design and methods of the PATH study have been published elsewhere.<sup>31–34</sup>

The analytical sample for the current study was restricted to adult respondents (18+) (wave 1, n=32 320) with no self-reported heart condition (eg, congestive heart failure, heart attack and stroke) or previous diagnosis of hypertension or high cholesterol at baseline (n=21 734). A total of 3203 respondents were excluded as they did not participate at any follow-up interview, and respondents who did not report a hypertension diagnosis were right censored at their last observation point. Respondents with missing variable information (n=992, 5.3%) were excluded from the analysis using listwise deletion. The final analytical sample consisted of 17 539 respondents. A flowchart summarising the analytical sample is provided in the supplemental material (online supplemental figure S1).

### Patient and public involvement

Patients and/or the public were not involved in the design, conduct, reporting or dissemination plans of this research.

### Self-reported hypertension

We examined the incidence of self-reported hypertension at follow-up among respondents who reported they had never been diagnosed with hypertension at baseline. The reliability and concurrent validity of self-reported hypertension have been established in a previous study using PATH study data.<sup>32</sup> In waves 2 and 3, *all* respondents were asked, 'In the past 12 months, has a doctor, nurse or other health professional told you that you had high blood pressure?' Due to a change in the skip pattern in waves 4 and 5, this question was *only* asked to respondents who reported they saw a 'medical doctor, nurse, or other health professional' during the past 12 months. We adopted an inclusive measurement strategy because self-reported hypertension is known to have low sensitivity (ie, it is underestimated) in epidemiological studies,<sup>35</sup> especially among women<sup>36</sup> and non-Hispanic (NH) black adults.<sup>37</sup> To minimise this bias, we classified respondents who answered 'yes' to the blood pressure question as having self-reported hypertension regardless of whether they reported seeing a doctor during the past year. In waves 4 and 5, we classified respondents who did not report seeing a doctor during the past year as not having self-reported hypertension.

### Cigarette/ENDS exposure variable

Our exposure variable was based on answers to questions about established cigarette (100 or more cigarettes smoked in lifetime) and ENDS (ever fairly regularly used ENDS) use, as well as everyday or someday use of cigarettes and ENDS (current use). Based on these variables, we developed a four-category exposure variable: non-current user (of either product, which included people who never used either product), exclusive cigarette smoker, exclusive ENDS user and dual user of cigarettes and ENDS. This variable was constructed at each wave and was included as a time-varying exposure. To minimise missing values for a given wave, we imputed missing tobacco exposure data borrowing information from a previous wave. To ensure that the tobacco product use exposure preceded the hypertension diagnosis, we lagged our time-varying exposure by one wave. The descriptive statistics of the time-varying tobacco use exposure can be found in the supplemental material (online supplemental table S1).

### Covariates

We included age (continuous ages 18–90), sex (0=female and 1=male), race/ethnicity (Hispanic, NH white, NH black, NH Asian or NH other) and household income (less than \$49 999, more than \$50 000 or missing) as baseline sociodemographic variables. Missing values for baseline sociodemographic variables were updated with data from other waves when available to reduce item non-response. We also included baseline risk factors to control for potential confounding, including familial history of heart attack/bypass surgery, obesity (body mass index >30), diabetes mellitus and regular binge drinking (five

or more drinks in one sitting on at least five separate days during the past month).

To account for the potential confounding effect of lifetime cigarette smoking, two additional covariates were included. First, we included a dichotomous predictor for former established smokers (smoked at least 100 cigarettes in a lifetime but reported no current use at baseline). Second, we included cigarette pack-years as a measure of lifetime cigarette smoking at baseline. Pack-years were calculated by multiplying the duration of cigarette smoking by the average number of packs of cigarettes smoked per day while individuals smoked. Respondents who reported smoking more than 200 cigarettes per day (10 packs per day) were considered implausible and were set to missing (n=99).

### Statistical analysis

Descriptive statistics were first calculated for sociodemographic characteristics, cigarette/ENDS use and hypertension risk factors at baseline. The sample characteristics were then calculated according to respondent's cigarette/ENDS use at baseline.  $\chi^2$  tests or Fisher's exact test was used to test for statistically significant differences between groups. Life tables were then used to describe the distribution of the incident hypertension outcomes at follow-up (waves 2–5). The hazard estimates reflect the weighted conditional probability for self-reported hypertension for respondents in the risk set at each discrete-time interval.<sup>38</sup>

We used discrete-time survival models to analyse the incidence of self-reported hypertension across waves 2–5 of follow-up (approximately 5 years). Discrete-time survival models are appropriate when the exact timing until an event is not known.<sup>38</sup> The data were fit to an unbalanced person-period dataset where each individual contributed a number of rows equal to the time period until they were diagnosed with hypertension or were right censored.<sup>39</sup> As such, all 17539 respondents in the self-reported hypertension sample had a separate row of data for each period, with a maximum of four rows per respondent, resulting in a person-period dataset with 59367 observations. The structure of the reorganised person-period dataset allowed for an examination of the conditional probability of self-reported and medicated incident hypertension at each discrete-time interval. All discrete-time survival models were estimated using a complimentary log–log (cloglog) link function on the person-period dataset. Data were weighted using wave 1 weights, including full-sample and 100 replicate weights, to ensure that our respondents were representative of the non-institutionalised adult population in the USA at baseline.

Several sensitivity analyses were included as robustness checks. First, to assess the impact of attrition, we compared baseline characteristics for censored and non-censored respondents (online supplemental table S2). Second, because the censored respondents had a slightly different sociodemographic profile than the non-censored

respondents, as a sensitivity analysis, we estimated the discrete-time models using the ‘all waves weights’, which account for this type of attrition<sup>31</sup> and restrict the analysis to a longitudinal cohort of respondents who participated in all waves of the PATH study (online supplemental table S3). Third, to better approximate clinical hypertension and minimise potential false-positive errors in self-reported hypertension, we also included a measure of medicated hypertension as a sensitivity analysis. Respondents who self-reported hypertension and responded yes when asked ‘In the past 12 months, did you take heart or blood pressure medication regularly?’ were considered to have medicated hypertension (online supplemental table S4). Fourth, to examine whether more frequent cigarette/ENDS use was associated with incident hypertension, we included a more frequent cigarette/ENDS use exposure (measured as 10+ days in the past 30 days) as a sensitivity analysis (online supplemental table S5). Fifth, to more clearly distinguish between adults who never smoked cigarettes from former smokers, we created a revised exposure with adults who reported ‘never established smoking’ as the reference group, with the following use categories: (1) former cigarette, no ENDS; (2) current cigarette, no ENDS; (3) former cigarette, current ENDS; (4) current cigarette and ENDS; and (5) exclusive ENDS (see online supplemental table S6). Finally, we restricted our analysis to adults who reported they had never smoked 100 cigarettes in their lifetime at baseline and examined the association between ENDS use and hypertension among respondents who had never smoked (online supplemental table S7). For all analyses, variances were computed using the balanced repeated replication methods with Fay’s adjustment set to 0.3 as recommended by the PATH study.<sup>33 40</sup> All analyses were conducted using Stata V.16.1.<sup>41</sup>

## RESULTS

The weighted baseline sociodemographic characteristics, smoking behaviours and hypertensive risk factors for the self-reported hypertension (n=17539) analytical sample are outlined in [table 1](#). At baseline, respondents had a mean age of 39 years (SD=15.4) and were predominately female (53.9%) and NH white (63.0%) and reported a household income of less than \$50 000 (56.0%). Most respondents were not current cigarette or ENDS users at baseline (n=11 063, 79.2%), while a similar percentage of respondents were exclusive ENDS users (n=336, 1.1%) or dual users (n=570, 1.7%). Current cigarette use was the most common tobacco use status at baseline (n=5570, 18.0%). Of the respondents, 13.4% were former established smokers at baseline; among current or former established smokers, the average cigarette pack-years was 13.9 (SD=20.0). In terms of baseline hypertensive risk factors, approximately one-quarter of respondents reported a family history of heart attack (27.7%) and obesity (24.6%), while diabetes mellitus

**Table 1** Weighted sociodemographic characteristics, smoking behaviours and hypertensive risk factors for adult respondents (18+) at baseline, Population Assessment of Tobacco and Health study (Wave 1, 2013–2014)

	N	%*	95% CI
Age (years), mean (SD)	17 539	38.97 (15.42)	
Sex			
Female	9073	53.9	53.2 to 54.6
Male	8466	46.1	45.4 to 46.8
Race/ethnicity			
NH white	10 250	63	62.2 to 63.8
Hispanic	3446	17.6	17.0 to 18.2
NH black	2422	11	10.5 to 11.5
NH Asian	526	5.8	5.3 to 6.3
NH other	895	2.6	2.4 to 2.9
Household income			
<\$50 000	11 481	56	54.6 to 57.3
>\$50 000	5699	41.8	40.4 to 43.1
Missing	359	2.2	1.9 to 2.7
Cigarette/ecigarette baseline exposure			
Non-user	11 063	79.2	78.5 to 79.9
Cigarette only	5570	18	17.3 to 18.7
E-cigarette only	336	1.1	0.92 to 1.2
Dual user	570	1.7	1.6 to 2.0
Family history of heart attack			
No	12 852	72.3	71.2 to 73.3
Yes	4687	27.7	26.7 to 28.8
Obesity (BMI >30)			
No	13 318	75.4	74.3 to 76.5
Yes	4221	24.6	23.5 to 25.7
Diabetes diagnosis at baseline			
No	16 848	95.3	94.8 to 95.8
Yes	691	4.7	4.2 to 5.2
Regular binge drinking			
No	16 297	95.5	95.1 to 95.8
Yes	1242	4.5	4.2 to 4.9
Former established smoker at baseline			
No	15 618	86.6	85.8 to 87.5
Yes	1921	13.4	12.5 to 14.2
Pack-years among current/former smokers, mean (SD)†	8061	13.9 (20.0)	

\*Percentages were calculated using W1 weights.  
†Mean pack-years value for ever established (both current and former) smokers.  
BMI, body mass index; NH, non-Hispanic; W, wave.

(4.7%) and regular binge drinking (4.5%) were reported less frequently.

[Table 2](#) presents the sample characteristics stratified by our tobacco exposure variable at baseline. Compared with

**Table 2** Sample characteristics by baseline cigarette/ENDS use, Population Assessment of Tobacco and Health study (wave 1, 2013–2014)

	Non-user % (95% CI)	Exclusive cigarette user % (95% CI)	Exclusive ENDS user % (95% CI)	Dual user % (95% CI)
Age (years), mean (SD)	39.6 (14.2)	37.1 (17.7)	33.2 (16.7)	34.2 (16.6)
Sex				
Female	55.9 (55.1 to 56.8)	45.9 (44.5 to 47.3)	45.9 (39.9 to 52.1)	47.5 (43.4 to 51.6)
Male	44.1 (43.2 to 44.9)	54.1 (52.7 to 55.5)	54.1 (47.9 to 60.1)	52.5 (48.4 to 56.6)
Race/ethnicity				
NH white	61.2 (60.1 to 62.4)	68.9 (67.3 to 70.5)	69.3 (63.0 to 75.0)	76.7 (72.7 to 80.4)
Hispanic	19 (18.2 to 19.7)	12.6 (11.7 to 13.6)	12.3 (9.1 to 16.5)	9.9 (7.5 to 13.0)
NH black	10.8 (10.2 to 11.4)	12.6 (11.5 to 13.7)	8.5 (5.6 to 12.5)	5.8 (3.8 to 8.7)
NH Asian	6.6 (6.1 to 7.3)	2.4 (1.8 to 3.2)	5.7 (2.7 to 11.5)	2.3 (1.0 to 4.9)
NH other	2.4 (2.1 to 2.7)	3.5 (3.1 to 3.9)	4.2 (2.4 to 7.1)	5.3 (3.8 to 7.2)
Household income				
<\$50 000	51.4 (49.9 to 52.9)	74.3 (72.7 to 75.9)	65.2 (59.3 to 70.7)	66.2 (61.0 to 70.9)
>\$50 000	46.2 (44.7 to 47.7)	23.8 (22.3 to 25.3)	33.2 (27.4 to 39.5)	32.2 (27.5 to 37.3)
Missing	2.4 (2.0 to 2.9)	1.9 (1.5 to 2.3)	1.6 (.65 to 3.7)	1.6 (.80 to 3.2)
Family history of heart attack				
No	72.8 (71.6 to 74.0)	70.6 (69.2 to 72.0)	68.3 (63.3 to 73.0)	65.8 (61.0 to 70.3)
Yes	27.2 (26.0 to 28.4)	29.4 (28.0 to 30.8)	31.7 (27.0 to 36.7)	34.2 (29.7 to 39.0)
Obesity (BMI >30)				
No	75.5 (74.1 to 76.8)	75.3 (73.8 to 76.7)	72 (65.7 to 77.5)	76.2 (72.2 to 79.8)
Yes	24.5 (23.2 to 25.9)	24.7 (23.3 to 26.2)	28 (22.5 to 34.3)	23.8 (20.2 to 27.8)
Diabetes diagnosis at baseline				
No	95.3 (94.6 to 95.8)	95.5 (94.9 to 96.0)	96.8 (94.3 to 98.2)	94.9 (92.3 to 96.6)
Yes	4.7 (4.2 to 5.4)	4.5 (4.0 to 5.1)	3.2 (1.8 to 5.7)	5.1 (3.4 to 7.7)
Regular binge drinking				
No	97.2 (96.8 to 97.5)	89 (88.0 to 89.9)	89.5 (85.1 to 92.7)	87.9 (84.6 to 90.6)
Yes	2.8 (2.5 to 3.2)	11 (10.1 to 12.0)	10.5 (7.3 to 14.9)	12.1 (9.4 to 15.4)
Former established smoker at baseline				
No	84 (82.9 to 85.0)	100	36.3 (30.3 to 42.9)	100
Yes	16 (15.0 to 17.1)	0	63.7 (57.1 to 69.7)	0
Pack-years smoking at baseline, mean (SD)*	13.9 (15.3)	14.1 (22.4)	17.9 (23.6)	11.1 (16.9)

\*Mean pack-years value for ever established (both current and former) smokers.

BMI, body mass index; ENDS, electronic nicotine delivery systems; NH, non-Hispanic.

all the other groups, respondents who exclusively smoked cigarettes were the most likely to be NH black (12.6%), most likely to report household incomes under \$50 000 (74.3%). Compared with exclusive cigarette users, exclusive ENDS users at baseline were younger (33.2 (SD=16.7) years vs 37.1 (SD=17.7) years), reported higher household incomes (33.2% vs 23.8%) and were more likely to report a family history of heart attack (31.7% vs 29.4%) and obesity (33.2% vs 23.8%). Importantly, nearly two-thirds of exclusive ENDS users were former established smokers at baseline (63.7%). The average pack-year value for exclusive ENDS users who were former established smokers (17.9, SD=23.6) was higher than for current

exclusive cigarette users (14.1, SD=22.4) at baseline. Dual users shared similar sociodemographic characteristics with exclusive ENDS users, except dual users were more likely to be NH white (76.7% vs 69.3%), to have diabetes mellitus (5.1% vs 3.2%) and reported more regular binge drinking (12.1% vs 10.5%–10.3%). The average pack-year value for dual users (11.1, SD=16.9), on the other hand, was lower than exclusive cigarette users (14.1, SD=22.4), and for former smokers who were non-current users (13.9, SD=15.3) or exclusive ENDS users (17.9, SD=23.61) at baseline.

Life tables describing the conditional probability for self-reported incident hypertension are displayed in

**Table 3** Life tables describing the incidence of self-reported hypertension among adults (18+), Population Assessment of Tobacco and Health study (W1–W5, 2013–2019)

Interval	Total	Diagnosis	Censored	Hazard estimate*
Period 1 (W1–W2)	17 539	652	1230	0.039
Period 2 (W2–W3)	15 660	464	1137	0.033
Period 3 (W3–W4)	14 067	334	1632	0.029
Period 4 (W4–W5)	12 101	480	11 612	0.046

\*Hazard estimates were calculated using W1 weights. W, wave.

**table 3.** Hypertension was self-reported by 1930 respondents in the analytical sample, with an annual incidence hazard of 3.7% (range 2.9% to 4.6% between wave 2 and wave 5). The hazard estimates were similar across all discrete-time intervals, with slight increases between wave

4 and wave 5, reflecting a 2-year time interval between waves.

**Table 4** presents discrete-time hazard models examining the risk of self-reported incident hypertension. In the unadjusted model, respondents who exclusively

**Table 4** Discrete-time survival analysis predicting incidence of self-reported hypertension among adults, Population Assessment of Tobacco and Health study (waves 1–5, 2013–2019)

	Unadjusted		Adjusted	
	Hazard	95% CI	Hazard	95% CI
Time-varying cigarettes/ENDS use				
Non-use	Ref	Ref	Ref	Ref
Exclusive cigarette use	1.28	1.15 to 1.42	1.21	1.06 to 1.38
Exclusive ENDS use	0.84	0.58 to 1.21	1	0.68 to 1.47
Dual use	1	0.77 to 1.30	1.15	0.87 to 1.52
Sociodemographic risk factors				
Age (mean)*	1.03	1.03 to 1.04	1.03	1.03 to 1.04
Sex (male=1)	1.28	1.11 to 1.48	1.33	1.16 to 1.53
Race/ethnicity				
NH white	Ref	Ref	Ref	Ref
Hispanic	0.83	0.71 to 0.98	0.99	0.84 to 1.17
NH black	1.44	1.24 to 1.68	1.62	1.38 to 1.90
NH Asian	0.38	0.23 to 0.64	0.55	0.33 to 0.94
NH Other	1.03	0.73 to 1.44	1.06	0.76 to 1.49
Household income				
<\$50 000	Ref	Ref	Ref	Ref
>\$50 000	0.8	0.70 to 0.92	0.83	0.72 to 0.96
Missing	0.67	0.32 to 1.39	0.58	0.27 to 1.22
Baseline risk factors				
Family history of heart attack	1.43	1.24 to 1.66	1.27	1.08 to 1.49
Obesity (BMI >30)	1.89	1.66 to 2.15	1.71	1.50 to 1.96
Diabetes diagnosis	2.48	2.0 to 3.06	1.74	1.37 to 2.21
Binge drinking	1.22	0.99 to 1.50	1.25	1.01 to 1.56
Smoking history variables				
Former established smoker	1.42	1.18 to 1.72	1.03	0.83 to 1.27
Pack-years (intervals of 10)*	1.17	1.13 to 1.21	1.03	0.98 to 1.08

Person n=17 539, risk period n=59 367.

\*For interpretation, pack-years were rescaled to intervals of 10 pack-years.

BMI, body mass index; ENDS, electronic nicotine delivery systems; NH, non-Hispanic.

smoked cigarettes had a significantly higher risk of self-reported incident hypertension compared with those who did not currently use cigarettes or ENDS products (HR 1.28, 95% CI 1.15 to 1.42). The risk did not statistically differ for respondents who used ENDS, either exclusively (HR 0.84, 95% CI 0.68 to 1.47) or with cigarettes (HR 1.00, 95% CI 0.77 to 1.30), from respondents who did not use either product. After adjusting for sociodemographic risk factors, baseline risk factors and smoking history variables, the results were very similar as exclusive cigarette use was associated with a 21% higher risk of self-reported incident hypertension (95% CI 1.06 to 1.38), while exclusive ENDS use (adjusted HR (aHR) 1.0, 95% CI 0.68 to 1.47) and dual use (aHR 1.15, 95% CI 0.87 to 1.52) were not. Other hypertensive risk factors associated with an increased risk of self-reported hypertension included older age, male sex, NH black (vs NH white) race/ethnicity, lower (vs higher) household income, family history of heart attack, obesity, diabetes diagnosis and regular binge drinking at baseline in adjusted (multi-variable) models.

### Sensitivity analyses

As sensitivity analyses, discrete-time models were estimated using the longitudinal cohort who participated in all waves of follow-up (online supplemental table S3), with a medicated hypertension outcome (online supplemental table S4) and with cigarette/ENDS use measured as 10+ days in the past 30 days rather than everyday or someday use (online supplemental table S5). Across these sensitivity analyses, the substantive results remained robust as exclusive cigarette use was associated with an increased risk of incident hypertension compared with non-use in both unadjusted and fully adjusted models. In contrast, compared with non-use, exclusive ENDS use and dual use were not associated with increased hypertension risk in unadjusted or fully adjusted models in any of these analyses. Discrete-time models were also estimated with an expanded cigarette/ENDS exposure incorporating never and former smoking as a sensitivity analysis (online supplemental table S6). Compared with never smoking, current cigarette smoking and non-ENDS use (aHR 1.20, 95% CI 1.04 to 1.38) was associated with an increased risk of incident hypertension, while current ENDS use among respondents who had formerly smoked (aHR 1.01, 95% CI 0.64 to 1.60) and dual ENDS and cigarette smoking (aHR 1.13, 95% CI 0.84 to 1.52) were not associated with increased hypertension risk. Finally, respondents with established cigarette use patterns were removed from the analytical sample, and the association between ENDS use and hypertension was examined among respondents who never smoked as an additional sensitivity analysis (online supplemental table S7). Time-varying ENDS use was not associated with an increased risk of incident hypertension compared with non-ENDS use in either unadjusted (HR 0.56, 95% CI 0.28, 1.13) or adjusted models (aHR 0.75, 95% CI 0.37 to 1.52).

### DISCUSSION

This study examined the time-varying association between cigarette smoking and ENDS use on the incidence of self-reported hypertension among a nationally representative sample of US adults. We found that exclusive cigarette use was associated with an increased risk of incident hypertension in both unadjusted and fully adjusted models. While the association between chronic cigarette use and hypertension is complex,<sup>42</sup> and the causal link is still debated,<sup>42 43</sup> this finding aligns with previous research indicating a modest association between current cigarette smoking and the risk of incident hypertension.<sup>5 8 10 44 45</sup> Moreover, this finding is consistent with hypertension risk prediction models that include current cigarette smoking as a covariate<sup>7</sup> and with the findings from the 2014 Surgeon General's report, which concluded that cigarette smoking is directly associated with coronary heart disease, including hypertension.<sup>9</sup> In contrast, studies examining the effects of ENDS use on hypertension have only recently been published,<sup>22</sup> and in a longitudinal follow-up of approximately 5 years, we found no evidence that short-term and time-varying ENDS use was associated with an increased risk of incident hypertension.

Dual use of cigarettes and ENDS was not associated with incidence of hypertension, although the direction of the hazard estimates was positive in fully adjusted models for both self-reported and medicated hypertension outcomes. However, it is important to note that dual users were different from exclusive cigarette smokers, and the non-significant association between dual use and incident hypertension may be partially explained by residual confounding by sociodemographic characteristics and tobacco use histories of dual users. In our study, dual users were younger and more likely to be NH white and reported higher household incomes than exclusive cigarette smokers. These characteristics are all correlated with lower risk of hypertension.<sup>8 46 47</sup> In addition, dual users had lower pack-year values than exclusive cigarette users, with pack-year values very similar to those of exclusive ENDS users. The different smoking histories between exclusive cigarette and dual users are consistent with other research finding that dual use is associated with reduced cigarette consumption<sup>48–50</sup> and may represent part of a transitional state as smokers move away from smoking cigarettes.<sup>50 51</sup> It is possible that dual users may have a different risk profile than exclusive cigarette users, which may then translate into a lower risk of disease relative to exclusive cigarette users. Studies with a larger number of ENDS users are needed to better understand the risk of incident hypertension among dual users.

Taken together, the results from this study do not support an association between ENDS use and self-reported incident hypertension. By examining the prospective incident cases of hypertension and using a lagged time-varying cigarette/ENDS exposure variable, our study does not have the same concerns with reverse



causation that have been identified in cross-sectional studies.<sup>27</sup> This is the most likely reason why our findings differ from a recent cross-sectional examination of the lifetime prevalence of hypertension using PATH data,<sup>23</sup> where the authors did not account for the relative timing of the ENDS exposure and hypertension. In addition, we also controlled for the potential confounding of cigarette smoking history, measured as pack-years, which is important, given that 64% of exclusive adult ENDS users at baseline were former established cigarette smokers. The substantial history of cigarette use among the majority of exclusive ENDS users further highlights the importance of controlling for their cigarette smoking history when trying to estimate the independent effect of ENDS use on hypertension and other health outcomes.

### Limitations

Our study has several important limitations that need to be considered. First, the results from this study are based on observational data from a prospective longitudinal study, and the results should be interpreted with the same level of caution required in all self-reported studies. Our non-randomised data mean that our results could be affected by unmeasured confounding, and while we included a measure of medicated hypertension as a sensitivity analysis, both our hypertensive outcomes are self-reported. Since systolic and diastolic blood pressure measures are not available in the PATH study, the reported incidence may underestimate the true incidence of hypertension,<sup>35 36</sup> particularly for some sociodemographic groups.<sup>35</sup> Future research would benefit from including measured hypertension instead of self-reported hypertension where possible. Second, while the PATH study was representative of the US population at baseline, the loss to follow-up was significant, and respondent attrition may not have been random. While we examined differences between censored and uncensored cases and conducted a sensitivity analysis with weights meant to adjust for attrition, this problem cannot be fully eliminated, as is true of most longitudinal studies. The discrete-time survival approach, which allows us to include all available information from respondents at each time interval, is a way to maximise information on the longitudinal sample. Third, while PATH has the biggest representative sample of longitudinal tobacco use and health in the US, ENDS use was only reported by a relatively small number of participants, limiting the power to detect statistical associations between ENDS use and incident hypertension. Fourth, if some respondents used ENDS to quit smoking cigarettes, it is possible that these respondents also made other lifestyle changes that may have concomitantly reduced the impact of ENDS use on incident hypertension. Similarly, some might have decided to switch in response to symptoms or health issues. Future research is needed to better understand the characteristics of respondents who

transition from cigarettes to ENDS use, their reasons for doing so and the future health outcomes of these transitions. Finally, ENDS products have only been widely available in the USA for a little more than a decade.<sup>52</sup> The findings from our study are based on approximately 5 years of longitudinal follow-up, and longer exposure to ENDS products may be required to more fully understand the role of ENDS use on the risk of hypertension. Moreover, ENDS products continue to evolve, and more recent generations of ENDS products have more efficient nicotine delivery. This study did not adjust for cumulative exposure to ENDS or for nicotine level by product type. Future studies should seek to develop valid methods for better understanding exposure to ENDS products, and this analysis will need to be updated as more longitudinal data on longer-term ENDS use becomes available.

### CONCLUSIONS

Using nationally representative prospective longitudinal data among US adults, we found that time-varying cigarette smoking increased the risk of self-reported incident hypertension, but time-varying ENDS use did not. These results highlight the importance of using prospective longitudinal data to disentangle the temporal ordering between cigarette and ENDS use and the need to control for the potential confounding effect of cigarette smoking histories among ENDS users. This type of longitudinal analysis can be extended in future research examining the cardiovascular health effects of ENDS use as longer-term data become available.

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**Contributors** SC conducted the data analysis and drafted and revised the manuscript. J LH and NLF initiated the research project in collaboration with RM and DL. IB and RM provided statistical consultation. GB and DA provided medical expertise and helped interpret the findings. EJM, AP and JJ created the measures used in the analysis. All coauthors revised the draft of the paper, and NLF revised the final draft prior to submission. SC is the author acting as guarantor.

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## REFERENCES

- Carter BD, Freedman ND, Jacobs EJ. Smoking and mortality--beyond established causes. *N Engl J Med* 2015;372:2170.
- Jha P, Ramasundarahettige C, Landsman V, et al. 21st-century hazards of smoking and benefits of cessation in the United States. *N Engl J Med* 2013;368:341–50.
- Lariscy JT, Hummer RA, Rogers RG. Cigarette smoking and all-cause and cause-specific adult mortality in the United States. *Demography* 2018;55:1855–85.
- Rostron B. Smoking-attributable mortality by cause in the United States: revisiting the CDC's data and estimates. *Nicotine & Tobacco Research* 2013;15:238–46.
- Niskanen L, Laaksonen DE, Nyssönen K, et al. Inflammation, abdominal obesity, and smoking as predictors of hypertension. *Hypertension* 2004;44:859–65.
- Doonan RJ, Hausvater A, Scallan C, et al. The effect of smoking on arterial stiffness. *Hypertens Res* 2010;33:398–410.
- Echouffo-Tcheugui JB, Batty GD, Kivimäki M, et al. Risk models to predict hypertension: a systematic review. *PLoS One* 2013;8:e67370.
- Gao K, Shi X, Wang W. The life-course impact of smoking on hypertension, myocardial infarction and respiratory diseases. *Sci Rep* 2017;7:1–7.
- Services UDoHaH. *The health consequences of smoking—50 years of progress: a report of the surgeon general*. 2014.
- Dikalov S, Itani H, Richmond B, et al. Tobacco smoking induces cardiovascular mitochondrial oxidative stress, promotes endothelial dysfunction, and enhances hypertension. *Am J Physiol Heart Circ Physiol* 2019;316:H639–46.
- Dubow J, Fink ME. Impact of hypertension on stroke. *Curr Atheroscler Rep* 2011;13:298–305.
- Kjeldsen SE. Hypertension and cardiovascular risk: general aspects. *Pharmacol Res* 2018;129:95–9.
- Fuchs FD, Whelton PK. High blood pressure and cardiovascular disease. *Hypertension* 2020;75:285–92.
- Dai H, Leventhal AM. Prevalence of e-cigarette use among adults in the United States, 2014–2018. *JAMA* 2019;322:1824–7.
- Mayer M, Reyes-Guzman C, Grana R, et al. Demographic characteristics, cigarette smoking, and e-cigarette use among US adults. *JAMA Netw Open* 2020;3:e2020694.
- Levy DT, Yuan Z, Luo Y, et al. The relationship of e-cigarette use to cigarette quit attempts and cessation: insights from a large, nationally representative U.S. survey. *Nicotine Tob Res* 2018;20:931–9.
- Abrams DB, Glasser AM, Pearson JL, et al. Harm minimization and tobacco control: reframing societal views of nicotine use to rapidly save lives. *Annu Rev Public Health* 2018;39:193–213.
- Shahandeh N, Chowdhary H, Middlekauff HR. Vaping and cardiac disease. *Heart* 2021;107:1530–5.
- Tam J, Brouwer AF. Comparison of e-cigarette use prevalence and frequency by smoking status among youth in the United States, 2014–19. *Addiction* 2021;116:2486–97.
- Cheng T. Chemical evaluation of electronic cigarettes. *Tob Control* 2014;23 Suppl 2:ii11–7.
- Tarran R, Barr RG, Benowitz NL, et al. E-cigarettes and cardiopulmonary health. *Function (Oxf)* 2021;2:zqab004.
- Martinez-Morata I, Sanchez TR, Shimbo D, et al. Electronic cigarette use and blood pressure endpoints: a systematic review. *Curr Hypertens Rep* 2021;23:1–10.
- Miller CR, Shi H, Li D, et al. Cross-sectional associations of smoking and E-cigarette use with self-reported diagnosed hypertension: findings from wave 3 of the population assessment of tobacco and health study. *Toxics* 2021;9:52.
- Alzahrani T, Glantz SA. Adding data from 2015 strengthens the association between e-cigarette use and myocardial infarction. *Am J Prev Med* 2019;57:569–71.
- Alzahrani T, Glantz SA. The association between e-cigarette use and myocardial infarction is what one would expect based on the biological and clinical evidence. *Am J Prev Med* 2019;56:627.
- Bhatta DN, Glantz SA. Electronic cigarette use and myocardial infarction among adults in the US population assessment of tobacco and health. *J Am Heart Assoc* 2019;8:e012317.
- Farsalinos K, Niaura R. E-cigarette use and myocardial infarction: association versus causal inference. *Am J Prev Med* 2019;56:626–7.
- Farsalinos KE, Polosa R, Cibella F, et al. Is e-cigarette use associated with coronary heart disease and myocardial infarction? Insights from the 2016 and 2017 National Health Interview Surveys. *Ther Adv Chronic Dis* 2019;10:2040622319877741.
- Stokes A, Collins JM, Berry KM, et al. Electronic cigarette prevalence and patterns of use in adults with a history of cardiovascular disease in the United States. *JAMA* 2018;7.
- United States Department of H, Human Services. National Institutes of Health. National Institute on Drug A, United States Department of H, Human Services F, Drug Administration. Center for Tobacco P. Population assessment of tobacco and health (PATH) study [United States] restricted-use files. In: *Inter-university Consortium for Political and Social Research [distributor]*. 2021.
- Hyland A, Ambrose BK, Conway KP, et al. Design and methods of the population assessment of tobacco and health (PATH) study. *Tob Control* 2017;26:371–8.
- Mahoney MC, Rivard C, Hammad HT, et al. Cardiovascular risk factor and disease measures from the population assessment of tobacco and health (PATH) study. *Int J Environ Res Public Health* 2021;18:7692.
- Piessse A, Opsomer J, Dohrmann S, et al. Longitudinal uses of the population assessment of tobacco and health study. *Tob Regul Sci* 2021;7:3–16.
- Tourangeau R, Yan T, Sun H, et al. Population assessment of tobacco and health (PATH) reliability and validity study: selected reliability and validity estimates. *Tob Control* 2019;28:663–8.
- Gonçalves VSS, Andrade KRC, Carvalho KMB, et al. Accuracy of self-reported hypertension: a systematic review and meta-analysis. *J Hypertens* 2018;36:970–8.
- Wellman JL, Holmes B, Hill SY. Accuracy of self-reported hypertension: effect of age, gender, and history of alcohol dependence. *J Clin Hypertens (Greenwich)* 2020;22:842–9.
- Mentz G, Schulz AJ, Mukherjee B, et al. Hypertension: development of a prediction model to adjust self-reported hypertension prevalence at the community level. *BMC Health Serv Res* 2012;12:312.
- Singer JD, Willett JB, Willett JB. *Applied longitudinal data analysis: modeling change and event occurrence*. Oxford university press, 2003.
- Jenkins SP. *Introduction to the analysis of spell duration data*. ISER, University of Essex, 2004.
- Judkins DR. Fay's method for variance estimation. *J Off Stat* 1990;6:223–39.



- 41 StataCorp. *Stata statistical software: release 17*. College Station, TX: Stata Corp LLC, 2021.
- 42 Viridis A, Giannarelli C, Neves MF, *et al*. Cigarette smoking and hypertension. *Curr Pharm Des* 2010;16:2518–25.
- 43 Mills KT, Stefanescu A, He J. The global epidemiology of hypertension. *Nat Rev Nephrol* 2020;16:223–37.
- 44 Bowman TS, Gaziano JM, Buring JE, *et al*. A prospective study of cigarette smoking and risk of incident hypertension in women. *J Am Coll Cardiol* 2007;50:2085–92.
- 45 Halperin RO, Gaziano JM, Sesso HD. Smoking and the risk of incident hypertension in middle-aged and older men. *Am J Hypertens* 2008;21:148–52.
- 46 Deere BP, Ferdinand KC. Hypertension and race/ethnicity. *Curr Opin Cardiol* 2020;35:342–50.
- 47 Grotto I, Huerta M, Sharabi Y. Hypertension and socioeconomic status. *Curr Opin Cardiol* 2008;23:335–9.
- 48 Farsalinos KE, Romagna G, Voudris V. Factors associated with dual use of tobacco and electronic cigarettes: a case control study. *Int J Drug Policy* 2015;26:595–600.
- 49 Lee PN, Fry JS, Forey BA, *et al*. Cigarette consumption in adult dual users of cigarettes and e-cigarettes: a review of the evidence, including new results from the path study. *F1000Res* 2021;9:630.
- 50 Selya AS, Shiffman S, Greenberg M, *et al*. Dual use of cigarettes and JUUL: trajectory and cigarette consumption. *Am J Health Behav* 2021;45:464–85.
- 51 Brouwer AF, Jeon J, Hirschtick JL, *et al*. Transitions between cigarette, ends and dual use in adults in the path study (waves 1–4): multistate transition modelling accounting for complex survey design. *Tob Control* 2020. 10.1136/tobaccocontrol-2020-055967 [Epub ahead of print 16 Nov 2022].
- 52 Cahn Z, Drope J, Douglas CE, *et al*. Applying the population health standard to the regulation of electronic nicotine delivery systems. *Nicotine Tob Res* 2021;23:780–9.