Comparison of nicotine exposure during pregnancy when smoking and abstinent with nicotine replacement therapy: systematic review and meta-analysis

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ABSTRACT

Background and aims Smoking during pregnancy is strongly associated with negative pregnancy and perinatal outcomes. Some guidelines recommend nicotine replacement therapy (NRT) for smoking cessation during pregnancy, but adherence with NRT is generally poor and could be partially explained by nicotine-related safety concerns. We compared pregnant women’s cotinine and nicotine exposures from smoking with those when they were abstinent from smoking and using NRT.

Design Systematic review with meta-analysis and narrative reporting. Twelve studies were included: in most, only one type of NRT was used. Seven were quality-assessed and judged of variable quality. Setting Studies from any setting that reported nicotine or cotinine levels when smoking and later when abstinent and using NRT.

Participants Pregnant women who smoked and became abstinent but used NRT either in a cessation study or in a study investigating other impacts of NRT.

Measurements We quality-assessed longitudinal cohort studies using a modified version of the Newcastle–Ottawa scale. For meta-analysis, we used mean within-person differences in cotinine or nicotine levels when smoking and at later follow-up when abstinent and using NRT. Where such data were not available, we calculated differences in group mean levels and reported these narratively, indicating where data were not completely longitudinal.

Findings Of the 12 included studies, four cotinine-measuring studies (n = 83) were combined in a random effects meta-analysis; the pooled estimate for the mean difference (95% confidence intervals) in cotinine levels between when women were smoking and abstinent but using NRT was 75.3 (57.1 to 93.4) ng/ml (I² = 42.1%, P = 0.11). Of eight narratively-described studies, six reported lower cotinine and/or nicotine levels when abstinent and using NRT; two had mixed findings, with higher levels when abstinent but using NRT reported from at least one assay time-point.

Conclusions Pregnant women who use nicotine replacement therapy instead of smoking reduce their nicotine exposure.

Keywords Cotinine, nicotine, nicotine replacement therapy, pregnancy, smoking, smoking cessation.

INTRODUCTION

Smoking in pregnancy causes much morbidity and mortality [1] and rates are highest among younger, socially disadvantaged women [2]. Forty per cent of socio-economic inequalities in stillbirths and infant deaths are smoking-related [3], and smokers’ children are twice as likely to become smokers themselves [4]; however, this is all avoidable. Stopping smoking in pregnancy improves birth outcomes [5]; permanent cessation after pregnancy improves women’s health and may also improve their children’s health by diminishing second-hand smoke exposure and possibly by reducing penetration of smoking across the generations [6].

In the United Kingdom, when other cessation methods have been ineffective, pregnant women who want to stop can be recommended to use nicotine replacement therapy (NRT) [7] and guidance developed for use across the European Union (EU) takes the same approach [8]. All UK stop smoking services (SSS) offer NRT to pregnant smokers [9], and 11% of UK pregnant smokers receive NRT prescriptions [10]. Although NRT is effective outside pregnancy...
and the risk ratio (RR) [95% confidence interval (CI)] for cessation using NRT in non-pregnant smokers is 1.60 (1.53–1.68) [11], in pregnancy NRT has, at best, only borderline effectiveness for promoting smoking cessation. From all trials of NRT in pregnancy, the risk ratio (95% CI) for cessation with NRT in pregnancy is 1.43 (1.03–1.93), but when meta-analysis is restricted to include only least-biased, placebo randomized controlled trials (RCTs), there is less evidence that NRT works and the risk ratio (RR) is reduced further (RR 1.28, 95% CI = 0.99–1.66) [12]. One of the most plausible explanations for NRT appearing less effective in pregnancy is that pregnant women may not use NRT for long enough or in sufficient doses for it to be effective. For example, of UK pregnant smokers who are offered or prescribed NRT, 70% receive only a 2-week supply [10]. Similarly, in some trials which have enrolled pregnant smokers, only 7–30% of participants completed recommended courses of NRT [12]. In contrast, non-pregnant smokers enrolled into cessation trials adhere more strongly, using up to 94% of their intended NRT treatment courses [13]. Improving pregnant smokers’ adherence to NRT could result in this being more effective at helping them to stop smoking. In non-pregnant smokers, prescribing higher doses of NRT results in greater use of NRT [13], and this greater use of NRT is causally associated with successfully stopping smoking [13,14]. There is very little similar research in pregnancy; however, we know that the rate of nicotine metabolism is substantially accelerated in pregnancy [15,16]. This means that any given dose of NRT generates lower blood nicotine concentrations than the same dose used either before pregnancy or in the postpartum period. It is also known that, in pregnancy, faster nicotine metabolism is associated with lower cessation rates [17], possibly because pregnant NRT users have more rapid nicotine turnover and so will experience stronger nicotine withdrawal symptoms, be more likely to perceive NRT as unhelpful and stop using it. One would therefore only expect NRT to be as effective during pregnancy as it is either before or afterwards if pregnant women’s adherence levels were improved such that they obtained sufficient nicotine to ameliorate the impact of increased metabolism.

Pregnant women’s reluctance to use NRT seems to be partially explained by their worries about the safety of nicotine [18]. However, as NRT contains none of the harmful products of tobacco combustion there has long been consensus that, for pregnant women, NRT is probably safer than smoking [19]. Nevertheless, as we cannot be completely sure that nicotine is entirely safe in pregnancy, women probably need reassurance. Hence, to help pregnant women to decide about using NRT, clear information about nicotine exposures generated when smoking or using NRT could be useful. Such information could also assist health professionals who counsel pregnant women about using NRT. In this review, therefore, we aimed to identify and describe studies which report nicotine or cotinine levels in pregnant women when smoking and subsequently when abstinent from smoking and using NRT, comparing these to estimate any differences between body fluid concentrations. A secondary aim was to investigate how any differences might be influenced by type(s) of NRT used or health professionals’ instructions on how NRT should be used.

**METHOD**

This review followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) methods [20]. A review protocol has been published [21]. To be included, papers needed to study pregnant women who smoke and who subsequently became abstinent while using NRT. Studies had to report the same women’s nicotine or cotinine body fluid levels both when smoking and when using NRT. The design had to either be longitudinal or have a design which implied that longitudinal data might be available, even if such data were not reported in study publications (e.g. from NRT-allocated arms in RCTs of NRT).

**Searches**

We developed a search strategy in MEDLINE using a combination of MESH and plain text terms and adapted it to use in Web of Science and EBSCO (see Supporting information, Appendix S1); the strategy was optimized against its ability to find three studies which we knew should be included in the final review. Searches of these three platforms allowed access to six databases: MEDLINE, EMBASE (Excerpta Medica Database), PsycINFO, MIDRIS (Maternity and Infant Care Database), SSCI (Social Sciences Citation Index) and CINHAL (Cumulative Index to Nursing and Allied Health Literature), and were completed by 29 August 2017. We also searched GSK clinical trials (https://www.gsk-clinicalstudyregister.com/); World Health Organization International Clinical Trials Registry Platform (www.who.int/trialsearch); US National Library of Medicine Clinical Trials database (clinicaltrials.gov/); and the ISRCTN registry (http://www.isrctn.com/). Finally, we searched the Cochrane Library using the terms ‘smoking’, ‘pregnancy’ and ‘nicotine replacement’. Non-bibliographic database searches were completed by 7 September 2017. There were no language restrictions and literature was searched from 1980, as the first trials of NRT were reported after that. In tandem with electronic searches, we scanned the references of papers included in reviews identified by the searches, and which covered the topic of interest, but were not eligible for inclusion.
Study selection

Identified citations (titles and abstracts) were manipulated in an EndNote library. One reviewer (C.H.) screened these to assess whether or not articles should be included, and where there was uncertainty or papers were thought likely to be eligible, full texts were assessed by two reviewers with agreement on inclusion or exclusion being reached by consensus.

Data extraction

Data were extracted by one researcher (C.H.) and checked by a second (T.C.). The following study details were extracted: objectives, setting, inclusion and exclusion criteria, study design and analysis; and number and characteristics of participants providing data for this review, baseline information on nicotine addiction or heaviness of smoking. The following intervention details were extracted: completeness of follow-up for women in longitudinal analyses; reasons for dropout; biochemical confirmation of participant’s smoking abstinence or not; dose(s) and type(s) of NRT used; instructions given on how regularly and for how long NRT should be used. The following details on measurements were extracted: body fluids sampled; whether nicotine or cotinine was assayed; time-points at which samples were taken and timings of samples relative to smoking or NRT use; and relevant numerical findings (e.g. mean differences between concentrations of cotinine or nicotine concentrations at baseline and later time-points). For ongoing studies, we e-mailed the Principal Investigator enquiring whether data were available and we asked the same of corresponding authors for those papers which reported insufficient data for meta-analysis (see ‘Analysis’ below). For two studies [22,23] we converted graphical data to numerical using WebPlotDigitizer software [24].

Risk of bias assessment

We quality-assessed those studies which had been designed as longitudinal cohort studies and which stated, a priori, that a reason for the study was to take measurements when smoking and later abstinent and using NRT. These studies designs were, therefore, directly relevant to this review—any biases in methods used could be adjudged directly from published reports; this was performed using Wells’ modified version of the Newcastle–Ottawa Scale (NCOS) [25] (see Supporting information, Appendix S2). Papers were independently rated by two researchers, ratings were compared and disagreements resolved by discussion. We did not quality-assess studies which had not been designed as before–after longitudinal studies (e.g. RCTs or secondary analysis of RCTs). For these studies, as studies’ data were not being used in a manner consistent with their designs (e.g. data from RCT arms treated as cohorts), the quality of the original study would not necessarily be relevant to review analyses. Similarly, where authors provided additional, unpublished data, we did not attempt quality assessment.

Modifications to the Newcastle–Ottawa scale

Wells’ modified version of the NCOS allocates stars to reflect study quality on eight items grouped under three domains: selection or comparability of study group and ascertainment of exposure(outcome) [25]. We did not use three NCOS items and amended others, such that the maximum score was seven stars. Two items attracted up to two stars (‘representativeness of cohort’ and ‘adequacy of cohort follow-up’) and one star for the remaining three (ascertainment of exposures, method for confirming abstinence and appropriateness of sample timing). We did not use the item ‘Selection of the non-exposed cohort’, as included studies compared measurements from the same women at different times and did not have non-exposed controls. Demonstration that outcome of interest was not present at start of study was irrelevant, as all studies measured outcomes (e.g. cotinine) and ‘Comparability of cohorts on the basis of the design or analysis’ was not discriminatory, as all studies were longitudinal cohorts. All five items and scoring are fully described in Supporting information, Appendix S2.

Analysis

Longitudinal, within-person data, from the same women at baseline and at later time-points, were used to estimate the mean differences between body fluid levels of nicotine or cotinine when smoking and later when abstinent and using NRT. We aimed to provide a pooled estimate of this difference in body fluid levels and to investigate the impacts of the type and dose of NRT and gestational age, but anticipated that the meta-analysis undertaken would depend upon the available data and that a final decision on which studies (if any) to include in analyses would be taken once available literature were identified. For inclusion in meta-analyses, study manuscripts had to report such a mean difference and its standard error or to report sufficient other data from which these could be calculated. Where such data were not included in papers, we contacted authors requesting either aggregated data as mean differences and standard errors or as individual participants’ data. A saliva : blood cotinine ratio has been reported as 1.01 (95% CI = 0.99–1.04) [26], so blood and saliva cotinine levels were considered interchangeable; nicotine and cotinine values and also urinary and saliva cotinine readings are not interchangeable, so these data were not aggregated.

Meta-analysis was conducted in Stata version 15 using the Metan command employing random-effects models.
[27] to provide a pooled, weighted estimate for the mean difference in cotinine levels when smoking and later when abstinent and using NRT [28]. Two studies reported independent cohorts of women who had received different types or combinations of NRT [29,30]. As we anticipated that there would potentially be more variation between cohorts reported within one study, exposed to different types, doses or combinations of NRT than between cohorts reported in different studies, we treated such cohorts as independent studies in the random-effects meta-analysis. Heterogeneity was assessed using the $I^2$ statistic [31].

For studies which could not be included in meta-analysis, we calculated differences in group mean levels (of cotinine or nicotine) and report these narratively, indicating where data were not completely longitudinal. For studies which provided ‘within-participant’, longitudinal data with no loss to follow-up, percentage nicotine substitution was calculated by dividing follow-up mean cotinine (nicotine) levels by baseline ones and multiplying by 100. The percentage nicotine substitution measure indicates how completely NRT substitutes for nicotine from smoked tobacco.

RESULTS

After removing duplicates, 3576 potentially relevant citations were found from library databases (131 from other sources, Fig. 1), 30 full texts were reviewed, one study was ongoing [32]. 12 studies were included in the review and four were meta-analysed. Table 1 gives the studies’ characteristics, including the numbers of participants providing longitudinal data and hence which could potentially be aggregated in a meta-analysis. This was not always the total number of study participants; for example, from RCTs, only women randomized receiving NRT could provide such data. Two study reports contained sufficient data for inclusion in meta-analysis [29,30]. For another two, authors re-analysed their data to provide sufficient information [27,39,40]. Eight studies were reported narratively; for one of these, the authors provided sufficient extra data for a ‘within-person’ mean difference in urinary cotinine values to be calculated; this could not be combined with values obtained from saliva assays, however [38]. For the seven remaining narratively reported studies, mean differences were calculated by subtracting published group mean cotinine or nicotine levels when abstinent and using NRT from those measured when smoking, ignoring between-participant variability. In two of these seven studies, only some follow-up women were abstinent and using NRT and these women could not be identified from other study participants [22,37].

Characteristics of included studies

Seven studies were set in the United States [22,23,30,34–36,38], two recruited from Denmark [29,37] and one each from France [39], United Kingdom [40] and Sweden [33]. Seven were longitudinal cohorts designed to investigate the impacts of smoking and then NRT use in the same women [23,29,30,33–36], and five were either RCTs [22,37–40] or presented secondary analyses of RCT data [40]. Studies tested 2 mg nicotine gum [29,33,38]; 4 mg gum [33]; 7 mg/24-hour nicotine patch [22]; 14 mg/24-hour nicotine patch [22]; 1.5 mg/16-hour nicotine patch [29,30,37,40]; 15 mg/24-hour nicotine patch plus 2 mg gum (often called ‘dual NRT’) [29]; 21 mg/24-hour nicotine patch [23,35]; 22 mg/24-hour patch [36]; nicotine nasal spray [30]; and also individualized nicotine dosing based on saliva cotinine levels [39]. Three papers reported studies recruiting relatively early in pregnancy; two RCTs reported women’s mean gestational age at enrolment as 17 weeks [38,39] and another as 14 weeks [40]. All except one of the remaining studies (22 weeks) [29] reported mean gestations at recruitment of 27 weeks or later. Timings for body fluids sampling while using NRT varied greatly; in laboratory (or in-patient) studies sampling occurred within 1 hour [33]; at 30 minutes and 5 days [34]; at up to 8 hours [23,35]; during a 4-day period [36]; and at both 8 hours and 4 days [30]. In non-laboratory studies (mainly RCTs), sampling occurred at 1–2 weeks [29], 8–11 weeks and also 4 weeks before delivery [37]; 6 weeks [38]; 2–4 weeks [22,40]; and at both 2 and 8 weeks after starting NRT [39].

Quality assessment

Quality assessments are reported in Table 2. The seven longitudinal cohort studies were of variable quality; six were awarded three or more stars out of seven. Studies used appropriate biochemical validation methods and generally scored well on follow-up completeness, but they scored less strongly with regard to the timing of samples when smoking or using NRT or in how abstinence was confirmed before or while using NRT, usually due to lack of detail in study descriptions.

Studies’ findings

Meta-analysis

Data obtained from 83 participants in four saliva cotinine-measuring studies were included in a meta-analysis (Fig. 2; Table 3, rows 1–4) [29,30,38,40]. The pooled estimate for the mean difference (95% CIs) between saliva cotinine levels when smoking and when using NRT and abstinent from smoking was 75.3 (57.1–93.4) ng/ml ($I^2 = 42.1\%$).
Within these studies, percentage nicotine substitution varied between 26.5% (2 mg gum) [29] and 60.0% (15 mg/16-hour patch).

Narratively reported studies
In six of the eight narratively reported studies, irrespective of body fluid (or substance assayed), exposure levels were higher when smoking than when abstinent and using NRT. In the remaining two studies, findings were mixed and details follow: Table 3 shows mean differences and explains which data were used to derive these and reasons for exclusion from meta-analysis. Also, numbers of participants for whom longitudinal data were available are given and, as relevant, how these related to total study samples.

The study summarized in row 5 reported higher urinal cotinine levels when smoking [35]. Although longitudinal data were available, findings from this study could not be used for the meta-analysis as other studies in this analysis reported saliva cotinine.

In rows 6–9, four longitudinal cohort studies are described [23,33–35]: in three, exposure measured as nicotine or cotinine was higher when smoking [23,33,34].

Figure 1 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) diagram: study selection. From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(7): e1000097. https://doi.org/10.1371/journal.pmed1000097
| Study                          | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures | Body fluid used and assay timings | Assay results |
|-------------------------------|-----------------------------------------------------------|---------------------------------|-----------------------------|---------------------------------|---------------|
| **Cohort studies**            |                                                            |                                 |                             |                                 |               |
| Gennser, Maršál & Brantmark 1975 [33] | 12 Values are ranges                                       | Sweden                           | Blood nicotine              | Mean ± SEM (ng/ml)               |               |
|                               | Maternal age 20–31 years<sup>a</sup>                       | Laboratory-based longitudinal cohort study: not a smoking cessation intervention study | 5 and 30 minutes after starting to smoke a cigarette | Smoking (n = 12)                |               |
|                               | Gestational age 33–39 weeks<sup>b</sup>                    | One of each below exposure was used on consecutive days | 30 and 60 minutes after starting to chew 1 piece of gum | 5 minutes 41.3 ± 3.5, 30 minutes 19.6 ± 1.4 |               |
|                               | Daily cigarette consumption 7–20<sup>a</sup>                | • 1 ‘standard’ cigarette: talking not permitted and deep inhalation encouraged; average smoking time was 5 minutes | 2 mg gum (n = 6) | 2 minutes 10.4 ± 0.6, 60 minutes 10.0 ± 0.7 |               |
|                               |                                                            | • 2 mg gum chewed for 30 minutes | Also presented trough and peak nicotine levels after smoking one cigarette and chewing 1 piece 2 mg gum | 4 mg gum (n = 6) | 30 minutes 17.5 ± 1.3, 60 minutes 14.7 ± 1.3 |               |
|                               |                                                            | • 4 mg gum chewed for 30 minutes |                                      |                                 |               |
| Oncken et al. 1996 [34]       | 15 Values are means ± SD                                  | United States                    | Blood cotinine              | Mean ± SD (ng/ml)               |               |
|                               | Maternal age 28 ± 6<sup>b</sup>                           | Longitudinal cohort study: not a smoking cessation intervention study | 30 minutes after last cigarette | Smoking (n = 15)               | 153 ± 18      |
|                               | Weight (lb) 169 ± 29<sup>b</sup>                          | Women allocated to receive a 5-day course of 2 mg NRT gum and advised to use ≥ 6 pieces/day but < 30 pieces/day and ≤ 2 pieces/hour | After 5 days using nicotine gum while remaining abstinent | 5 days gum (n = 15) | 33 ± 8        |
|                               | Gestational age 28.1 ± 3.2<sup>b</sup>                    |                                    | Also presented trough and peak nicotine levels after smoking one cigarette and chewing 1 piece 2 mg gum |                                      |               |
|                               | Daily cigarette consumption 19 ± 6<sup>b</sup>             | Observed chewing to ensure proper technique on first use |                                      | 1 cigarette | Trough 6.7 ± 0.8, Peak 19.7 ± 1.5 |               |
|                               | Fagerström value 5.7 ± 2<sup>b</sup>                     |                                    |                                      |                                 |               |
|                               |                                                            | Reported daily adherence (mean ± SD) |                                      |                                 |               |
|                               |                                                            | Day 1, 6 ± 3 pieces, day 2, 9 ± 5, day 3, 8 ± 3, day 4, 8 ± 3, day 5, 5 ± 2 (last day was only a half-day) |                                      |                                 |               |

(Continues)
| Study                  | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures | Body fluid used and assay timings | Assay results                      |
|-----------------------|----------------------------------------------------------|--------------------------------|-----------------------------|----------------------------------|-----------------------------------|
| Oncken et al. 1997 [35] | 15                                                       | Values are means ± SD          | United stated               | Blood nicotine                   | Mean ± SD (ng/ml)                 |
|                       |                                                          | Maternal age 28 ± 5.4          | Laboratory-based longitudinal cross-over study; not a smoking cessation intervention study | Hourly assays when smoking       | Maximal plasma level smoking 19.7 ± 8.09 Patch 16.0 ± 3.5 |
|                       |                                                          | Gestational age 28 weeks, 3 days ± 20 days | Two groups of women used | Assays at 2, 3, 4, 6 & 8 hours after placement of nicotine patch | Time to reach max Smoking 5 hours ± 2.4 Patch 3.2 hours ± 1.7 |
|                       |                                                          | Height 161 ± 5.9 cm            | Smoked ad libitum for 8 hours | Maximum mean nicotine concentration, time to reach max, and the mean area plasma nicotine concentration-time curve were reported in the text of this paper |
|                       |                                                          | Weight 63.3 ± 9.2 kg           | Abstinent for 13 hours before placement of a 21-mg transdermal nicotine patch for 8 hours | Mean, range (μg/l) Cotinine values (n = 6) Smoking 100 (40–155) Patch 55 (20–100) |
|                       |                                                          | Daily cigarette consumption 20.2 ± 5.2 | They were randomized to either smoking or patch use in session 1 and then a week later crossed over to the other experimental condition | Mean difference ± SEM 4.8 ± 10.3 ng hour/ml |
|                       |                                                          | Values are ns                  |                             |                                  |                                   |
|                       |                                                          | Ethnicity data                 |                             |                                  |                                   |
|                       |                                                          | White 10                       |                             |                                  |                                   |
|                       |                                                          | Black 2                        |                             |                                  |                                   |
|                       |                                                          | Hispanic 3                     |                             |                                  |                                   |
| Wright et al. 1997 [23] | 6                                                        | Values are means & ranges      | United States               | Saliva nicotine and cotinine     | Mean, range (μg/l) Cotinine values (n = 6) Smoking 100 (40–155) Patch 55 (20–100) |
|                       |                                                          | Maternal age 25.7 (21–31)      | Laboratory-based cohort study; not a smoking cessation intervention study | Minimum 1 hour after last cigarette |
|                       |                                                          | Weight 82.05 kg (66.1–87.5 kg) | Smoking as normal prior to starting study | 8 hours after of started patch |
|                       |                                                          | Gestational age 34.2 (28.1–37) | Abstinent for 11 hours prior to patch placement. Transdermal patch 21 mg worn for 8 hours | Nicotine values (n = 6) smoking 19 (7–48) Patch 19 (6–41) |
| Study                        | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures                                                                                     | Body fluid used and assay timings | Assay results |
|------------------------------|----------------------------------------------------------|---------------------------------|---------------------------------------------------------------------------------------------------------------|----------------------------------|----------------|
| **Ogburn et al. 1999 [36]**  | 21                                                       | Values are means ± SD           | United States                                                                                                 | Blood nicotine and cotinine      | Mean ± SD (ng/ml) |
|                              |                                                          | Maternal age 26.5 ± 5.7         | Laboratory-based cohort study; not a smoking cessation intervention study                                     | At 2 p.m. after a normal morning’s smoking and at least 10 minutes after last cigarette | Cotinine values |
|                              |                                                          | Gestation at enrolment 27.4 ± 2.7| Smoking as usual prior to day of admission                                                                   | On all of days 1–4, 8 hours after patch placement on each day | Smoking 116 ± 54 (n = 21) |
|                              |                                                          | Daily cigarette consumption 20.5 ± 8.7 | Abstinence started on admission and continued for in-patient period Transdermal patch 22 mg/24 hours worn each day of in-patient stay |                                    | Day 1 patch 142 ± 47 (n = 21) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 2 patch 128 ± 38 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 3 patch 123 ± 42 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 4 patch 117 ± 38 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Nicotine values |
|                              |                                                          |                                 |                                                                                                              |                                   | Smoking 14.4 ± 9.7 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 1 patch 12.7 ± 4.2 (n = 21) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 2 patch 12.8 ± 4.5 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 3 patch 13.7 ± 6.0 (n = 20) |
|                              |                                                          |                                 |                                                                                                              |                                   | Day 4 patch 11.8 ± 3.9 (n = 20) |

| Hegard, Kjærgaard, Møller, Wachmann & Ottesen 2004 [29] | 75 | Values are means ± SD | Denmark | Longitudinal cohort study; smoking cessation intervention study | Baseline measurement taken before starting NRT, while still smoking but no time from last cigarette given | Mean ± SD (ng/ml) |
|----------------------------------------------------------|----|-----------------------|---------|---------------------------------------------------------------|----------------------------------------------------------------------------------------------------------|----------------|
|                                                          |    | Gestation at enrolment 21.5 ± 8.4 |         | Baseline measurement taken before starting NRT, while still smoking but no time from last cigarette given | Baseline measurement taken before starting NRT, while still smoking but no time from last cigarette given | Smoking 132 ± 95 |
|                                                          |    | Daily cigarette consumption 12.5 ± 5.2 |         | Baseline measurement taken before starting NRT, while still smoking but no time from last cigarette given | Baseline measurement taken before starting NRT, while still smoking but no time from last cigarette given | NRT 35 ± 28 |
|                                                          |    |                                 |         | Mean difference, 95% CI = –97 (–6–200)                         | Mean difference, 95% CI = –97 (–6–200)                                                                   | (Continues |
|                                                          |    |                                 |         |                                                                 |                                                                                                          |     |
|                                                          |    | Fagerström value n             |         |                                                                 |                                                                                                          |     |
|                                                          |    | 2–4                            | 6       | Women with Fagerström score of 2–4 used up to 12 pieces of 2 mg gum daily | Women with Fagerström score of 2–4 used up to 12 pieces of 2 mg gum daily | Smoking 173 ± 41 |
|                                                          |    | 4–7                            | 7       | For scores of 4–7, 15 mg/16-hour nicotine patches were used   | For scores of 4–7, 15 mg/16-hour nicotine patches were used   | NRT 70 ± 33 |
|                                                          |    | 7–10                           | 5       | For scores of 7–10, 15 mg/16-hour nicotine patches plus up to 8 daily pieces of 2 mg gum were used | For scores of 7–10, 15 mg/16-hour nicotine patches plus up to 8 daily pieces of 2 mg gum were used | Mean difference, 95% CI = –103 (60–146) |
|                                                          |    |                                 |         |                                                                 |                                                                                                          |     |

(Continues)
| Study | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures | Body fluid used and assay timings | Assay results |
|-------|----------------------------------------------------------|---------------------------------|-----------------------------|----------------------------------|---------------|
| Oncken, Campbell, Chan, Hatsukami & Kranzler 2009 [30] | 14 | Values are means ± SD | United States Longitudinal cohort study; smoking cessation intervention study | Serum nicotine & cotinine (cotinine used in analysis) | Mean ± SD (ng/ml) |
| | | Maternal age | | Patch (n = 7) | 1st laboratory-based smoking session; smoking 138 ± 55 |
| | | Patch 29.86 ± 6.52 | Women smoked 7 cigarettes over 7 hours (1 per hour) in a monitoring session | smoking 38 ± 55 | Spray 75 |
| | | Spray 30.29 ± 5.09 | | | Mean difference ± SD = 63 ± 33 |
| | | Gestation at enrolment | Then they were allocated to one of two types of NRT for 4 days, or the placebo equivalent. They were not monitored during the 4 days and then returned for a 2nd monitoring session | 2nd laboratory session: on the 5th treatment day, after overnight NRT abstinence. In both patch and spray groups, samples were taken at equivalent times to the smoking session; average of these samples reported | Spray (n = 7) |
| | | Patch 32.06 ± 2.64 | | Spray 130 ± 57 |
| | | Spray 31.70 ± 3.2 | | NRT session spray 39 |
| | | Daily consumption of cigarettes | | Mean difference ± SD = 91 ± 38 |
| | | Patch 19.64 ± 3.66 | | |
| | | Spray 16.71 ± 5.90 | | |
| | | Fagerström value | | |
| | | Patch 6.71 ± 1.60 | | |
| | | Spray 6.29 ± 1.11 | | |
| | | Ethnicity data | | |
| | | Caucasian patch 85.7%; spray 71.4% | | |
| | | Nasal spray (1 mg/dose) dose = 1 spray to each nostril; instructed to use up to 24 daily doses. During monitoring session nasal spray was used twice at the equivalent times to cigarette 1, 4 and 7, and once in place of the 2nd, 3rd, 5th and 6th cigarette | | |
| Study                              | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures | Body fluid used and assay timings | Assay results |
|-----------------------------------|----------------------------------------------------------|---------------------------------|---------------------------|----------------------------------|---------------|
| **Randomized controlled trial data** |                                                          |                                 |                           |                                  |               |
| Wisborg, Henriksen, Jespersen & Secher 2000 [37] | 124 randomized to nicotine | Values are means ± SD | Denmark | Saliva cotinine | Mean ± SD (ng/ml) |
|                                    |                                                           | Maternal age 28.2 ± 4.9 \(^b\) | Randomized controlled trial study: smoking cessation intervention study; recruited from Aarhus Hospital | While still smoking usual amount: no time from last cigarette reported | Smoking (baseline) \(n = 124\) 231 ± 125 |
|                                    |                                                           | Daily cigarette consumption 13.4 ± 4.0 \(^b\) | | | 8 weeks \(^d\) patch \(n = 90\) 153 |
|                                    |                                                           | Women randomized to NRT were issued with 8 week's 15 mg/16 hours nicotine patches, followed by a further 3 weeks of 10 mg/16 hours nicotine patches | | | 11 weeks \(^d\) patch \(n = 83\) 121 |
| Oncken et al. 2008 [38]           | 100 randomized to nicotine | Values are means ± SD | United States | Urine cotinine | Mean ± SD (ng/ml) |
|                                    |                                                           | Maternal age 25.5 ± 6.8 \(^b\) | Randomized controlled trial: smoking cessation intervention study; recruited from Hartford, New Britain, Springfield medical sites | While still smoking usual amount: no time from last cigarette reported | Smoking (baseline) \(n = 93\) 672 ± 438 |
|                                    |                                                           | Gestation at enrollment 17.1 ± 5.6 \(^b\) | | | 6 weeks \(^d\) of 2 mg gum \(n = 51\) 542 ± 454 |
|                                    |                                                           | Daily consumption of cigarettes before pregnancy 17.5 ± 9.6 \(^b\) | | | 32–34 weeks gestation |
|                                    |                                                           | last 7 days before enrollment 10.2 ± 6.6 | | | 32–34 weeks \(^d\) gestation after NRT treatment finished \(n = 54\) 492.45 ± 443 |
|                                    |                                                           | Fagerström value 3.83 ± 1.91 \(^b\) | | | |
|                                    |                                                           | Values are \(n\) | | | |
|                                    |                                                           | Ethnicity data \(^b\) | | | |
| Hispanic                           | 53                            | | | | |
| Non-Hispanic white                 | 38                            | | | | |
| Non-Hispanic black                 | 8                             | | | | |
| Other                              | 1                             | | | | |

(Continues)
| Study                                      | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures                                                                 | Body fluid used and assay timings | Assay results |
|-------------------------------------------|----------------------------------------------------------|---------------------------------|------------------------------------------------------------------------------------------------|----------------------------------|---------------|
| El-Mohandes et al. 2013 [22]              | 26 randomized to nicotine                                | Values are means ± SD           | United States                                                                                   | Saliva cotinine                  | Mean ± SD (ng/ml) |
|                                           |                                                          | Maternal age 27.5 ± 5           | Randomized clinical trial; smoking cessation intervention study; recruited from prenatal care sites in the DC metropolitan area | While still smoking usual amount: no time from last cigarette reported | Smoking (baseline) (n = 26) |
|                                           |                                                          | Gestation at enrolment 19.6 ± 5.1 |                                                                                                 | After using a 14 mg patch for approx. 2–4 weeks (and having been using NRT for average 3.7 weeks) | 171 ± 14.3 |
|                                           |                                                          | Ethnicity data                 |                                                                                                 | After using a 7 mg patch for approx. 2 weeks (and having been using NRT for average 7.8 weeks) | 14 mg patch (n = 26) 142 |
|                                           |                                                          | BMI 28.1 ± 10.1                |                                                                                                 |                                  | 7 mg patch (n = 26) 129 |
|                                           |                                                          | Daily cigarette consumption 7 ± 7 |                                                                                                 |                                  |               |
| Berlin, Grangé, Jacob & Tanguy 2014 [39]  | 203 randomized to nicotine                               | Values are median & interquartile range |                                                                                                 | Saliva cotinine                  | Mean ± SE (μg/l) |
|                                           |                                                          | Maternal age 29.1 (25–34)       | France                                                                                          | While still smoking usual amount: no time from last cigarette reported | Smoking (baseline) 119 ± 1.09 |
|                                           |                                                          | Gestation at randomization 17 (15–20) |                                                                                                 |                                  | 2 weeks patch 108 ± 1.1 |
|                                           |                                                          | BMI pre-pregnancy 23 (20–27)     |                                                                                                 |                                  | 8 weeks patch 80 ± 1.1 |

(Continues)
Table 1. (Continued)

| Study | Participants potentially providing longitudinal data (n) | Characteristics of participants | Setting, design & exposures | Body fluid used and assay timings | Assay results |
|-------|----------------------------------------------------------|---------------------------------|-----------------------------|----------------------------------|--------------|
| Bowker, Lewis, Coleman, Vaz & Cooper 2014 [40] | 33 Values are median & interquartile range | Values are median & interquartile range | United Kingdom Secondary analysis of data from intervention arm of randomized controlled trial of NRT; smoking cessation intervention study; recruited from antenatal clinics | Saliva cotinine | Median & interquartile range (ng/ml) |
| | | Maternal age 26.12 (22.29–32.35) | | | Smoking (n = 33) 98.5 (71.3–177.8) |
| | | Gestation at baseline 14.4 (13.3–17.8) | | | NRT (abstinent) (n = 33) 62.8 (33.3–82.7) |
| | | Ethnicity data | | | Median cotinine difference with baseline cotinine: >150 ng/ml −134.8 < 150 ng/ml−27.9 |
| | | All white except one Asian/other | | | Reported difference in cotinine from smoking to NRT use when starting Cotinine level > 150 ng/ml and when <150 ng/ml |
| | | BMI 25.6 (22.7–29.3) | | | |
| | | Women with partners who smoke 23 (70%) | | | |
| | | Daily cigarette consumption 11 (8–15)b | | | |
| | | Fagerström value 5 (3–6)b | | | |
| | | Values are ns | | | |
| | | Ethnicity datab | | | |
| | | European 194 | | | |
| | | African 6 | | | |
| | | Asian 1 | | | |
| | | Other 2 | | | |
| | | Time to first cigaretteb | | | |
| | | > 60 minutes 31 | | | Unclear how many women sampled at each visit |
| | | 31–60 minutes 36 | | | |
| | | 6–30 minutes 74 | | | |
| | | ≤ 5 minutes 62 | | | |
The fourth [35] had inconsistent findings; peak exposure (mean maximal plasma nicotine) was higher but total exposure (area under a nicotine concentration versus time graph) was lower after smoking.

Row 10 describes a longitudinal cohort study in which women were followed-up daily for 4 days when abstinent after starting NRT [36]; cotinine levels (Table 3) were higher and nicotine levels (not shown) were lower at all follow-up points, with the day 1 cotinine difference reaching statistical significance. For three follow-up comparisons, a participant (from 21) was lost to follow-up (Table 1).

Rows 11 and 12 describe women in NRT arms of RCTs [22,37]; in both studies, exposures (group mean cotinine levels) were higher in smokers at baseline, but it was not possible to identify separately those using NRT and abstinent.

DISCUSSION

A meta-analysis comparing cotinine exposures when pregnant women smoke with those when they use NRT found that levels were, on average, 75.3 ng/ml lower when abstinent and using NRT than when the same women smoked. Similarly, lower exposures after NRT occurred in six of the remaining eight studies.

Only 12 empirical studies were included: five had not been designed as longitudinal cohorts and most did not publish sufficient details to be included in a meta-analysis. Nevertheless, longitudinal, within-participant data were available from 10 studies and so only two were of limited use for answering review questions [22,37]. Participants were recruited to either hospital in-patient/laboratory studies with intensive protocols or into clinical trials, but the consistency of outcomes from studies in very different settings suggests the principal finding that using NRT exposes pregnant women who are fully abstinent from smoking to less nicotine than smoking is valid. Although the amount of useable data from studies was small, by focusing on ‘within-individual’ differences in cotinine levels, study women effectively acted as their own controls and external impacts on cotinine levels, apart from NRT doses used, were eliminated. Only factors which changed within individual women between baseline and follow-up could be expected to affect the pooled estimate for mean difference in cotinine levels. One such factor is the rate of nicotine metabolism, which is significantly accelerated by the second trimester [15]. Adjusting findings for increasing rates of nicotine metabolism as pregnancy progressed could have helped us to understand how much lower cotinine levels on NRT might be attributable to faster metabolism; however, this was beyond the scope of the review. Nevertheless, there are two reasons to suspect that increased nicotine metabolism had little overall impact on

| Study | Setting, design & exposures | Body fluid used and assay timings | Assay results | Characteristics of participants | Cotinine levels in RCTs |
|-------|-----------------------------|----------------------------------|---------------|-------------------------------|-------------------------|
|       |                             |                                  |               | Heaviness of smoking index 3 (2–3) |                          |
|       |                             |                                  |               | Values are ns                 |                          |
|       |                             |                                  |               | Number of cigarettes smoked   |                          |
|       |                             |                                  |               | 5–10 cigarettes (23) 11–15 (8) > 20 |
|       |                             |                                  |               |                              |                          |
|       |                             |                                  |               |                              |                          |

Table 1. (Continued)
findings. First, the mean differences from studies which measured these only hours after stopping smoking [23,34,35] were comparable to those in whom cotinine (nicotine) levels on NRT were measured weeks afterwards or even later in pregnancy [22,37–40]. Secondly, findings from those studies which recruited more women who were under 18 weeks gestation [38–40] appeared similar to remaining studies which recruited later in pregnancy.

We believe this study is original, and the systematic approach used combined with the rigorous contact made with authors should have sourced all available data within identified studies. Despite substantial variation in the types of NRT issued and in how participants were instructed to use this, and also in the timings of sample measurement across studies, the low level of heterogeneity in the pooled mean difference estimate indicates that

| Study           | Representativeness | Ascertainment of exposures | Method for confirming abstinence | Appropriateness sample timing | Completeness of follow-up | Stars (out of 7) |
|-----------------|--------------------|-----------------------------|---------------------------------|-----------------------------|---------------------------|-----------------|
| Gennser 1975 [33] | –                  | *                           | *                               | **                          |                           | 5*              |
| Oncken 1996 [34] | **                 | *                           | *                               |                             |                           | 5*              |
| Oncken 1997 [35] | **                 | –                           | *                               |                             |                           | 5*              |
| Wright 1997 [21] | –                  | *                           |                                 |                             |                           | 3*              |
| Ogburn 1999 [36] | *                  | *                           |                                 |                             |                           | 3*              |
| Hegard 2004 [29] |                     |                             |                                 |                             |                           | 2*              |
| Oncken 2009 [30] | **                 | *                           |                                 |                             |                           | 7*              |

Figure 2. Forest plot showing meta-analysis of mean difference in saliva cotinine levels when smoking and when abstinent but using nicotine replacement therapy (NRT). Hegard 1, Hegard2 and Hegard3 2004 represent cohorts of women given different forms of NRT and reported in Hegard 2004 [29]; ditto for Oncken1 and Oncken2 2009 and Oncken 2009 [30] [Colour figure can be viewed at wileyonlinelibrary.com]
| Row | Study | Participants abstinent on NRT, providing longitudinal data and loss to follow-up (n) | Body fluid & assay times (i.e. after starting NRT) | Type and dose of NRT | Outcome(s) | Percentage substitution (%) | Derivation of outcome data |
|-----|-------|-----------------------------------------------------------------|-----------------------------|------------------|-------------|-----------------------------|-----------------------------|
| 1   | Hegard 2004 [29] | 18 analysed Saliva cotinine, 2 weeks | 2 mg gum (n = 6) | 97 (51.5) | 26.5 | Mean differences taken directly from paper SEMs calculated from CI presented in paper |
|     |       | Of 75 enrolled 40 were excluded: 16 dropped out 13 stopped NRT 11 smoked the day before second sample 17 were not analysed; 15 samples not collected or not treated properly or went missing 2 used a 10-mg patch | 15 mg/16-hour patch (n = 7) | 103 (21.5) | 40.5 |
|     |       |                                                                 | 15 mg/16-hour patch and 2 mg gum (n = 5) | 141 (47.5) | 42.7 |
| 2   | Oncken 2009 [30] | 14 Serum cotinine, 4 days | 15 mg/16-hour patch (n = 7) | 63 (12.47) | 54.3 | Mean differences taken directly from paper |
|     |       |                                                                 | Spray 24 dose/day = 24 mg/24 hours (n = 7) | 91 (14.36) | 30.0 | SEM calculated from SD of difference |
| 3   | Bowker 2014 [40] | 33 Saliva cotinine, 4 weeks | 15 mg/16-hour patch | 51.2 (10.77) | 60.0 | Mean difference and SEM calculated using original study data |
| 4   | Berlin 2014 [39] | 18 Saliva cotinine, 2 weeks | Patch, variable strength | 70.22 (20.07) | 49.3 | Mean difference and SEM of abstinent women only calculated by study author using original study data |

**Narratively reported studies**

| Row | Study | Participants abstinent on NRT, providing longitudinal data and loss to follow-up (n) | Body fluid & assay times (i.e. after starting NRT) | Type & dose of NRT | Outcome(s) | Percentage substitution (%) | Derivation of outcome data |
|-----|-------|-----------------------------------------------------------------|-----------------------------|-------------------|-------------|-----------------------------|-----------------------------|
| 5   | Oncken 2008 [38] | 4 Urine cotinine, 6 weeks | 2 mg gum (n = 4) | 130.00 (245.718) | 65.2 | Mean difference and SD of abstinent women using gum at 6 weeks calculated by author from original data Not included in meta-analysis as urine rather than saliva cotinine value |

**Studies where only published data used**

| Row | Study | Participants abstinent on NRT, providing longitudinal data and loss to follow-up (n) | Type & dose of NRT body fluid & assay times (i.e. after starting NRT) | Outcome(s) (ng/ml) | Reasons for non-inclusion in meta-analysis |
|-----|-------|-----------------------------------------------------------------|-----------------------------|-------------------|------------------------------------------|
| 6   | Gennser 1975 [13] | Unclear; 12 participants smoked at baseline, 6 were later using NRT but it is not clear which of the original 12 these were | Gum, 2 & 4 mg Blood nicotine, 30 minutes | Mean ± SEM Cig. 19.6 ± 1.4 (n = 12) 2 mg gum 10.0 ± 0.7 (n = 6) Difference in means = 9.6 Cig. 19.6 ± 1.4 (n = 12) 4 mg gum 14.7 ± 1.3 (n = 6) Difference in means = 4.9 | Only group means and their standard errors presented in paper; standard error of mean difference could not be calculated Difference in means calculated by review team as (group mean value for smoking) – (group mean value for gum use) |
| 7   | Oncken 1996 [34] | Of 19 enrolled 4 dropped out: 2 due to non-substance 2 suffered severe nausea | Gum, 2 mg blood cotinine, 5 days | Mean ± SD Smoking 153 ± 18 5 days gum 33 ± 8 Difference in means = 120 | Only group means and their standard errors presented in paper; standard error of mean difference could not be calculated Difference in means calculated by review team as (group mean value for smoking) – (group mean value for gum use) |

(Continues)
the data synthesis undertaken was valid and the estimate is robust.

It was not possible to combine studies’ findings to investigate the impacts of different NRT doses or regimens on cotinine levels. However, consideration of individual studies’ findings does not suggest that different NRT doses or giving different instructions about using NRT has substantial impact. For example, the mean differences in cotinine levels obtained when smoking and later from women who were abstinent and used NRT, and so were adherent, were
similar in two major RCTs investigating NRT in which participants were told to use this treatment in different ways [39,41]. In one trial [41], a single nicotine patch dose was provided for only an 8-week treatment course and participants were instructed to remove patches during smoking lapses. In the other trial, however, nicotine patch doses were personalized, and there was potential for higher doses to be delivered to women who were told that they could continue using NRT during brief smoking lapses and even for the whole of pregnancy, if desired [39]. The meta-analysis showed that cotinine levels when abstinent and on NRT were reduced, on average, by 70.3 ng/ml compared to smoking, and throughout the four meta-analysis studies cotinine levels when smoking varied between 99 and 246 ng/ml, suggesting that reductions in nicotine exposure while using NRT are clinically meaningful. Review studies, SNIPP excepted [39], used standard rather than higher doses of nicotine patches and these delivered no more than 15 mg cotinine in 16 hours or the 24-hour equivalent. An important, unequivocal message is, therefore, that when pregnant smokers become abstinent and adhere with to ‘standard’ doses of NRT they are, on average, exposed to less nicotine than from smoking. One arm of one study delivered both 15 mg/16-hour nicotine patches and 2 mg gum to five women [29] who had high baseline cotinine levels when smoking [mean (SD)] 246 (91) ng/ml, and the mean difference (95% CI) between this and cotinine levels on NRT was large [mean difference (95% CI)] –141 (47–236). This estimate lacks precision, however, and provides no evidence that higher-dose NRT might expose women to more nicotine; nevertheless, more studies are needed.

A key reason for this study was to determine whether pregnant smokers who have concerns about the safety of nicotine in pregnancy and which might deter them from using NRT regularly enough and in sufficiently high doses to help them stop smoking could be reassured about its use [18,40]. The review demonstrates clearly that NRT exposes pregnant women to much smaller nicotine doses than smoking and, clearly, pregnant women considering NRT use in pregnancy can be strongly reassured on this point. It was not an aim of this paper to determine whether or not nicotine is harmful to the developing baby; however, the accruing literature suggests that this is not the case. Although rodent studies have suggested that fetal nicotine exposure may cause infant behavioural problems [42], the only RCT of NRT for smoking cessation in pregnancy found that NRT group infants had better developmental outcomes [43]. Additionally, large studies of NRT used in routine health care have found no consistent relationship between NRT use in pregnancy and stillbirth [44,45], congenital abnormalities [46,47], preterm birth [48], low birth weight [49] or strabismus [50]. It seems most probable that most, if not all, the fetal harms caused by smoking in pregnancy are due to other tobacco smoke toxins. Pregnant women should avoid unnecessary toxin exposure and, compared to smoking, NRT both eliminates exposure to numerous tobacco smoke toxins and reduces nicotine exposure. However, NRT also has great potential for improving fetal health and averting adverse pregnancy outcomes by helping some pregnant women to stop smoking. Review findings could, therefore, help to reassure pregnant women about the probable safety of using NRT to maintain smoking abstinence and also about the use of higher-dose NRT. Although using ‘dual NRT’, an NRT patch and a short-acting NRT together would generate higher nicotine exposures, ‘standard NRT dose’-generated nicotine exposure in abstinent pregnant women is so much lower than that from smoking that dual NRT could well also deliver lower nicotine doses than cigarettes. However, ‘dual NRT’ would be more likely to alleviate withdrawal symptoms and so women would probably use this for longer; this may explain why an observational analysis of UK Stop Smoking Services’ routine data found dual NRT but not standard-dose NRT associated with smoking cessation in pregnancy [51].

CONCLUSIONS

Among pregnant women who quit smoking, standard-dose NRT generates lower nicotine exposure than smoking. This lower exposure, combined with the very strong likelihood that nicotine is not responsible for the majority of fetal harms caused by tobacco smoke, makes it very likely that relative to smoking, NRT is safer for the fetus than smoking. Additionally, when NRT promotes maternal smoking cessation this is very likely to improve fetal health by reducing adverse pregnancy outcomes.

Propero protocol

Propero protocol registration number: CRD42017081914

Declaration of interests

I.B. has received honoraria from Pfizer Ltd for talks and participation in advisory board. C.O. has received study medication (nicotine inhaler and placebo) from Pfizer Ltd for an NIH-funded study of a nicotine inhaler for smoking cessation during pregnancy. C.H., S.L., K.C., R.C., S.C., T.C/-H. and T.C.: none to declare.

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**Supporting Information**

Additional supporting information may be found online in the Supporting Information section at the end of the article.

**Appendix S1** Example search run in MEDLINE.

**Appendix S2** Amended Newcastle-Ottawa Scale for Cohort Studies: item definitions.