Introduction

Smoking is one of the most important causes of mortality and morbidity in the world, especially in developing countries.1–4 According to the World Health Organization (WHO), there are around 1.1 billion smokers in the world, and half of them die every year (2015).1 There are many types of smoking, such as cigarette smoking, bidi smoking, hookah smoking, electronic cigarette (EC) smoking, passive smoking, and many more.2–7 All of these types of smoking are related to the risk factor and etiology of many health problems, especially in the respiratory tract. In the respiratory tract, smoking is related to upper respiratory tract infection, asthma, chronic obstructive pulmonary disease, nasopharyngeal cancer, and lung cancer.2,8–10 Smoking is a significant risk factor for respiratory diseases, considering its ability to lead to an alteration of nasal mucociliary clearance (NMC).2,11,12

Nasal mucociliary clearance is the primary innate defense mechanism of the nose and paranasal sinuses, and it consists of mucous layer, airway surface liquid layer, and ciliary epithelia.2,13–15 Mucus produced by the goblet cells of the
mucosa is required for binding of the airborne pathogens (such as inhaled microbes and irritants). Ciliated epithelial cells are expectorated or swallowed. Normal functioning of the NMC requires high frequency, coordinated, and directional ciliary beating (metachronal waves) as well as proper mucus secretion and airway surface liquid. Ciliary beat frequency (CBF) was shown to be the major determinant of NMC efficiency. Nasal mucociliary clearance is influenced by physiological factors, such as mucus production and CBF; anatomic factors, such as nasal airflow and patency of the sinus ostia in the prechambers; and biochemical factors, such as mucus composition.

Various factors, like aging, body temperature, drugs (like adrenaline, acetylcholine, corticosteroid, and intranasal drug), tobacco use and smoking, and environment factors (like pollutant, smoke, and dust) affect this system, besides pathological conditions such as rhinitis allergy, acute or chronic rhinosinusitis, and deviated nasal septum. Any dysfunction in this defense system increases the inflammatory process and stasis of airborne pathogens, and the respiratory system becomes prone to obstructive airway diseases and infections. Long-term smoking causes both structural and functional damage in the respiratory airways, leading to changes in NMC.

The adverse effects of smoking on NMC have been reported and explained in various studies, but we could not find any systematic review about NMC in smokers. The aim of the present study was to look systematically into the current literature and carefully collect and analyze results to explore NMC in smokers.

Review of the Literature

Scope of the Review: Inclusion and Exclusion Criteria

Inclusion criteria:
1. Publication type:
   - full-text articles discussing NMC in smokers
   - primary studies of every design (case study, case series, cross sectional, case control, cohort, clinical trial)
2. Language of publication: English
3. Time of publication: January 2000 to August 2019
4. Methodology: studies included must explain NMC in smokers
5. Population: human at any age

Exclusion criteria:
1. Population: in-vitro samples
2. Objective and outcome measures are not relevant (are not about NMC in smokers)
3. Confounding variables are related to outcome of NMC in smokers

Literature Search

The current systematic review was conducted in accordance with Cochrane handbook for systematic reviews and is reported by using the guidelines of the preferred reporting items for systematic review and meta-analysis (PRISMA). A systematic search strategy was conducted in the following electronic databases: Pubmed, Medline, Ebsco, Springer Link, Science Direct, Scopus, and Proquest. The search was conducted using the following keywords for title and abstract: nasal mucociliary clearance OR nasal mucociliary transport AND smoke OR smoker OR smoking OR cigarette. In the Pubmed database, the keywords were searched through the [tiab] and [MeSH] tags. No limitation was applied during the search. The reference lists of the retrieved papers were also examined to avoid missing any published data.

Data Collection and Analysis

Studies were selected for retrieval after two independent reviewers (A. P. and U. S.) had collected titles and abstracts identified in the electronic search. The results of the two reviewers were compared by a third independent reviewer (J. B.), and any differences of opinion were resolved by discussion. Full papers from potential studies were independently assessed by the investigators (A. P. and U. S.).

All studies selected for this systematic review were screened by two reviewers independently to validate the results (A. P. and U. S.). The data from all retrieved studies are presented in the summary table (Table 1) featuring key points of each study. The following data were collected: first author and year, study design, sample, sample characteristic (age and gender), smoking characteristic (type, years of smoking, cigarettes/day or packs/year), NMC measurement test, and result.

Table 1  Newcastle-Ottawa scale (prospective study)

<table>
<thead>
<tr>
<th>No.</th>
<th>First author, year</th>
<th>Selection</th>
<th>Comparability</th>
<th>Outcome</th>
<th>Total</th>
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<tbody>
<tr>
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<td>1 2 3 4</td>
<td>1 2 3</td>
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</tr>
<tr>
<td>1.</td>
<td>Dülger et al, 2018</td>
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<tr>
<td>2.</td>
<td>Utiyama et al, 2016</td>
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<td>3.</td>
<td>Yadav et al, 2014</td>
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<td>4.</td>
<td>Ramos et al, 2011</td>
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<td>7</td>
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</table>

*Maximum points for comparability were 2.

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Ottawa scale cohort version, and the Cochrane risk of bias were used to assess the methodological quality of the studies. The Newcastle-Ottawa scale adapted for cross-sectional studies was used to assess cross sectional studies; interpretation of the total score was: 9 to 10 points were considered very good studies, 7 to 8 points were considered good studies, 5 to 6 points were considered satisfactory studies, and 0 to 4 were considered unsatisfactory studies. The Newcastle-Ottawa scale cohort version was used to assess prospective studies; interpretation of the total score was: ≥ 7 points were considered good studies, 5 to 6 points were considered fair studies, and < 5 points were considered poor studies. The Cochrane risk of bias was used to assess randomized control trial studies, whose results were either high risk or some concerns or low risk.

Results

Selection of Articles for Review

Fig. 1 summarized the identified, screened, and included articles for review. Initially, 1,622 peer-reviewed articles were identified from electronic databases, and an additional 32 articles were identified through other sources (search engine). After removing duplicates, 384 articles remained for title and abstract screening. Articles that did not meet the inclusion and exclusion criteria were not further screened. Twenty-six articles were screened for eligibility, 16 of which met all the inclusion criteria.

Assessment of Study Validity (Risk of Bias)

All eligible studies were associated with NMC in smokers. Table 2 provides quality scores for cross-sectional studies; all studies that got 6 to 8 points were included in the satisfactory and good studies category. Table 1 provides quality scores for prospective studies; all studies that got 5 to 7 points were included in the fair and good studies category. Table 3 provides quality scores for randomized control trial studies; all studies presented results of some concerns or low risk.

Study Characteristic

Study characteristics for the included studies could be seen in Table 4. The majority of the studies followed the cross-sectional design (11 out of 16). Most of the samples were in productive age and reported cigarette smoking. Nasal mucociliary clearance measurement tests mostly used saccharin transfer/transit time test.

Most of the studies showed the impairment of NMC in smokers. Nasal mucociliary clearance in cigarette smokers was explained in 13 studies (11 studies reported significant impairment in smokers, 1 study reported insignificant impairment in smokers, 1 studies reported that the NMC value in smokers was significantly lower than in non-smokers). Nasal mucociliary clearance in passive smokers was explained in three studies, and all of the studies reported that the NMC value in passive smokers was significantly higher than in non-smokers. Nasal mucociliary clearance in bidi, hookah, and electronic smoking, respectively, was explained in only one study.

Discussion

Nasal Mucociliary Clearance in Smokers

Most of the studies (15 out of 16 studies) showed the impairment of NMC in smokers. One study (Nicola et al, 2014) reported that the NMC value in smokers was lower than non-smokers. Nicola et al (2014) speculated that young smokers may have a protective response to cigarette smoking...
Various chemicals present in cigarette smoke, including acrolein, formaldehyde, carbon monoxide, nicotine, cotinine, acetaldehyde, phenol, and potassium cyanide, have been identified as having high toxicity to NMC. The gaseous phase of cigarette smoke contains high concentrations of free radicals (\(\geq 10^{15}\) molecules per puff), resulting in increased oxidative stress that, in turn, causes changes in the structure and function of the NMC. The impairment of NMC in chronic exposure to cigarette smoke is caused by the ciliotoxic effect, hypersecretion and viscoelastic change of mucus (particularly of more viscous properties), airway surface liquid depletion, increased oxidative stress, and deteriorations in the inflammatory and immune systems (increased of inflammatory and immune systems (increased oxidative stress that, in turn, causes changes in the structure and function of the NMC)).

Macrophages, neutrophil, and proinflammatory cytokines, which cause elongation of the NMC time and stagnation of toxic substances. The ciliotoxic effect of cigarette smoking reduces cilia genesis, paralyzes ciliary beating activity (reduces CBF), is related to abnormality in the cilia ultrastructure, and decreases the number of cilia. Cigarette smoking also causes metaplastic changes of the respiratory mucosa, with increase in the number and size of the goblet cells that leads to increased production of respiratory airway secretions.

The im-...
Table 4: Study characteristics

<table>
<thead>
<tr>
<th>No.</th>
<th>First author, year</th>
<th>Study design</th>
<th>Sample (N)</th>
<th>Sample characteristic: age (year), gender (male, female)</th>
<th>Smoking characteristic: type, years of smoking, cigarettes/day and/or packs/year</th>
<th>NMC measurement test</th>
<th>Result</th>
</tr>
</thead>
</table>
| 1.  | Arıcil and Arbağ, 2018 | Cross-sectional | Non-smokers: 40 Smokers I (once every week): 20 Smokers II (more than once a week/2–5 session-week): 18 | Age: 18–41 years Non-smokers: 27.5 ± 6.4 Smokers I: 26.9 ± 6.8 Smokers II: 27.7 ± 6.3 Gender Non-smokers: 22, 18 Smokers I: 11, 9 Smokers II: 10, 8 | Hookah smoking | Saccharin transfer/transit time test | • NMC value (STT) in smokers was significantly higher than in non-smokers ($p < 0.001$).  
• NMC value (STT) in the smokers-II group was significantly higher than in the smokers-I group and non-smokers ($19.2 ± 2.5; 11.9 ± 2.8; 11.1 ± 3; p < 0.001$). |
| 2.  | Dulger et al, 2018 | Prospective study (2 years period) | Non-smokers: 35 Smokers: 50 | Age: 18–65 years Non-smokers: 35.8 Smokers: 31.4 Gender Non-smokers: 23, 12 Smokers: 30, 20 | Cigarette smoking Cigarettes/day: 20.6 Packs/year: ≤ 10 packs/year: 11 10–20 packs/year: 28 20–30 packs/year: 5 ≥ 30 packs/year: 6 | Saccharin transfer/transit time test | • NMC value (STT) in smokers was significantly higher than in non-smokers (12 minutes, 9 minutes; $p < 0.001$).  
• No statistically significant difference in nasal MCC value and packs/year ($p = 0.943$). |
| 3.  | Paul et al, 2018 | Cross-sectional | Non-smokers: 20 Cigarette smokers: 20 Bidi smokers: 20 | Age: 20–40 years Non-smokers: 29.3 ± 6.25 Cigarette smokers: 31.2 ± 5.42 Bidi smokers: 30.2 ± 6.77 Gender: male | Cigarette and Bidi smoking Packs/year: Cigarette smokers: 4.63 ± 2.74 Bidi smokers: 7.5 ± 5.1 | Methylene blue dye test | • NMC value was significantly decreased in bidi smokers as compared with cigarette smokers and non-smokers ($59.25 ± 12.38$ mm; $67.89 ± 4.10$ mm; $p < 0.05$).  
• Multivariate analysis revealed a significant association between NMC and bidi smoking, number of cigarettes or bidis smoked per day, and packs/year ($p < 0.05$). |
| 4.  | Solak et al, 2018 | Cross-sectional | Non-smokers: 74 Smokers: 123 | Age: 18–55 years Non-smokers: 38.79 ± 9.66 Smokers: 40.33 ± 8.94 Gender Non-smokers: 58, 16 Smokers: 23, 100 | Cigarette smoking Years of smoking: 24.31 ± 9.66 Cigarettes/day: 18.45 ± 8.78 Packs/year: 28.49 ± 15.68 | Saccharin transfer/transit time test | • NMC value (STT) in smokers was significantly higher than in non-smokers ($336.19 ± 254.81$ seconds $320.43 ± 184.98$ seconds $p < 0.001$).  
• Positive correlation between STT and number of cigarettes/day ($p = 0.012, r = 0.225$), STT and packs/year ($0.001, r = 0.296$), STT and years of smoking ($p = 0.002, r = 0.200$). |
| 5.  | Uzeloto et al, 2018 | Cross-sectional | Non-smokers: 69 Smokers: 70 | Age: 30–50 years Non-smokers: 39.5° Smokers: 41° Gender Non-smokers: 32, 37 Smokers: 33, 37 °: median | Cigarette smoking Years of smoking: 21.50° Cigarettes/day: 20.00° Packs/year: 22.00° °: median | Saccharin transfer/transit time test | NMC value (STT) in smokers was insignificantly higher than in non-smokers ($9.7$ minutes; $9.145$ minutes; $p > 0.05$). |
| 6.  | Kumral et al, 2016 | Prospective randomized | Non-smokers: 40 Smokers: 58 | Age Non-smokers: 38.0 ± 8.2 Smokers: 33.9 ± 7.9 | E-C smoking duration: 3 months | Saccharin transfer/transit time test | NMC value (STT) in electronic cigarette was significantly higher than in non-smokers ($p < 0.001$). |

(Continued)
<table>
<thead>
<tr>
<th>No.</th>
<th>First author, year</th>
<th>Study design</th>
<th>Sample (N)</th>
<th>Sample characteristic: age (year), gender (male, female)</th>
<th>Smoking characteristic: type, years of smoking, cigarettes/day and/or packs/year</th>
<th>NMC measurement test</th>
<th>Result</th>
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<tbody>
<tr>
<td></td>
<td>Prospective study (12 months duration)</td>
<td>Quitters: 20 Smokers: 13</td>
<td>Age Quitters: 51 ± 9 Smokers: 52 ± 10 Gender Quitters: 9, 11 Smokers: 6, 7</td>
<td>Cigarette smoking Packs/year: Quitters: 40 ± 27 Smokers: 45 ± 28</td>
<td>Saccharin transfer/transit time test</td>
<td>• NMC value (STT) in smokers showed increases of impairment after 12 months observation (±14 minutes; ±15 minutes). &lt;br&gt;• NMC value (STT) in quitter showed decreases of impairment after 12 months observation (±15 minutes, ±10 minutes).</td>
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<td>Cross-sectional Group I (control): 18 Group II (living with at least one adult smoker outside the house): 15 Group III (living with at least one adult smoker inside the house): 17</td>
<td>Age 6–14 years Group I: 10.22 ± 2.39 Group II: 11.2 ± 1.97 Group III: 10.65 ± 2.09 Gender Group I: 9.9 Group II: 7.8 Group III: 9.8</td>
<td>Passive smoking</td>
<td>Saccharin transfer/transit time test</td>
<td>• NMC value (STT) in group II was insignificantly higher than in group I (p = 0.067). &lt;br&gt;• NMC value (STT) in group III was significantly higher than in group I (p &lt; 0.001). &lt;br&gt;• NMC value (STT) in group III was insignificantly higher than in group II (p = 0.173). &lt;br&gt;• NMC value (STT) in groups I, II, and III: 7.33 ± 2.91 minutes, 10.00 ± 4.78 minutes, 12.41 ± 3.44 minutes.</td>
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<td>Cross-sectional Non-smokers: 30 Ex-smokers: 30 Smokers: 30</td>
<td>Age Non-smokers: 53.17 ± 5.53 Ex-smokers: 50.73 ± 6.51 Smokers: 51.97 ± 6.02 Gender Non-smokers: 19, 11 Ex-smokers: 23, 7 Smokers: 24, 6</td>
<td>Cigarette smoking Cigarettes/day: Ex-smokers: 25 ± 7.76 smokers: 24.67 ± 6.3</td>
<td>Saccharin transfer/transit time test</td>
<td>NMC value (STT) in smokers was significantly higher than in others (smokers: 11.77 minutes, ex-smokers: 11.71 minutes, non-smokers: 11.71 minutes, p &lt; 0.0001).</td>
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<td></td>
<td>Cross-sectional Non-smokers: 30 Smokers: 30</td>
<td>Age: 21–40 years Non-smokers: 26.8 ± 1.2 Smokers: 24.96 ± 1 Gender: male</td>
<td>Cigarette smoking</td>
<td>Saccharin transfer/transit time test</td>
<td>• NMC value (STT) in smokers was significantly higher than in the non-smokers group (481.2 ± 29.83 seconds 300.32 ± 17.42 seconds; p &lt; 0.01). &lt;br&gt;• A statistically significant increase in the NMC value was observed with an increase in duration of smoking habit (NMC in smoking &lt; 1 year: 492.25 ± 79.93 seconds; 1–5 years: 516.7 ± 34.01 seconds &gt; 5 years: 637.5 ± 28.49 seconds p = 0.0000).</td>
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### Table 4 (Continued)

<table>
<thead>
<tr>
<th>No.</th>
<th>First author, year</th>
<th>Study design</th>
<th>Sample (N)</th>
<th>Sample characteristic: age (year), gender (male, female)</th>
<th>Smoking characteristic: type, years of smoking, cigarettes/day and/or packs/year</th>
<th>NMC measurement test</th>
<th>Result</th>
</tr>
</thead>
</table>
Smokers < 2.5 packs/year: 20  
Smokers > 2.5 packs/year: 20 | Age: 18–35 years  
Non-smokers: 21  
Smokers < 2.5 packs/year: 19  
Smokers > 2.5 packs/year: 24  
Gender  
Non-smokers: 29, 3  
Smokers < 2.5 packs/year: 20, 0  
Smokers > 2.5 packs/year: 17, 3 | Cigarette smoking  
Years of smoking  
Smokers < 2.5 packs/year: 3  
Smokers > 2.5 packs/year: 7  
Pack/year index: 2.9  
: median | Saccharin transfer transit time test | NMC value (STT) in smokers was significantly lower than in non-smokers (5.9 ± 3.1 minutes; 7.7 ± 4.1 minutes; p = 0.033). |
| 12. | Yadav et al, 2014<sup>3</sup> | Prospective study (five years duration) | Non-smokers: 50  
Active smokers: 50  
Passive smokers: 50 | Age: 20–50 years  
Non-smokers: 38.7  
Active smokers: 39.1  
Passive smokers: 35.1  
Gender  
Non-smokers: 44.6  
Active smokers: 42.8  
Passive smokers: 24.26 | Cigarette smoking (active and passive) | Saccharin transfer transit time test | NMC value (STT) in smokers (active and passive) was significantly higher than in light smokers and non-smokers (23.59 ± 12.41 minutes; 12.6 ± 4.67 minutes; p < 0.0003).  
NMC value (STT) in active smokers was insignificantly higher than in passive smokers (p = 0.03).  
A positive correlation was observed between STT and cigarettes/day (r = 0.3; p = 0.02). |
| 13. | Xavier et al, 2013<sup>20</sup> | Cross-sectional | Non-smokers: 24  
Light smokers: 14  
Moderate smokers: 34  
Heavy smokers: 27 | Age: 20–35 years  
Non-smokers: 50 ± 11  
Light smokers: 51 ± 15  
Moderate smokers: 49 ± 7  
Heavy smokers: 46 ± 8  
Gender  
Non-smokers: 7.17  
Light smokers: 5.9  
Moderate smokers: 14, 20  
Heavy smokers: 13, 14 | Cigarette smoking  
Years of smoking  
Light smokers: 32 ± 17  
Moderate smokers: 29 ± 9  
Heavy smokers: 32 ± 8  
Cigarettes/day: Light smokers: 9 ± 1  
Moderate smokers: 18 ± 3  
Heavy smokers: 39 ± 11  
Pack/year index: 31 ± 10  
: median  
Light smokers: 27 ± 11  
Heavy smokers: 59 ± 44 | Saccharin transfer transit time test | NMC value (STT) in moderate and heavy smokers was significantly higher (p = 0.0001) than in light smokers and non-smokers.  
A positive correlation was observed between STT and cigarettes/day (r = 0.3; p = 0.02). |
Passive smokers: 15  
Active smokers: 17 | Age: 17–47 years  
Active smokers: 28.12 ± 10.79  
Passive smokers: 29.17 ± 12.18  
Non-smokers: 27.92 ± 11.29  
Gender  
Active smokers: 11.6  
Passive smokers: 8.7  
Non-smokers: 6.9 | Cigarette smoking (active and passive) | Saccharin transfer transit time test | NMC value (STT) in active smokers was significantly higher than in passive smokers and non-smokers (23.59 ± 12.41 minutes; 12.6 ± 4.67 minutes; 6 ± 1.55 minutes; p < 0.01).  
NMC value (STT) in the passive group was significantly lower than in the non-smokers group (p < 0.01). |
| 15. | Proença et al, 2011<sup>35</sup> | Cross-sectional | Non-smokers: 19  
Smokers: 19 | Age  
Non-smokers: 47 ± 11  
Smokers: 51 ± 16  
Gender:  
Non-smokers: 8.7  
Smokers: 6.9 | Cigarette smoking  
Years of smoking: 33 ± 11  
Cigarettes/day: 27 ± 11  
Pack/year index: 44 ± 25 | Saccharin transfer transit time test | NMC value (STT) in smokers was significantly higher than in non-smokers 8 hours after smoking (16 ± 6 minutes; 10 ± 4 minutes, p = 0.005) and insignificantly higher immediately after smoking. |
Nasal Mucociliary Clearance in Smokers
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Table 4 (Continued)

<table>
<thead>
<tr>
<th>Study design</th>
<th>Sample size (N)</th>
<th>Sample characteristic: age (years)/gender</th>
<th>Sample characteristic: smoking characteristic (year)/type, cigarettes/day and/or packs/year</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prospective study (1 year period)</td>
<td>Non-smokers: 33; smokers: 33</td>
<td>Age: 18-28; gender: non-smokers: 18, 15; smokers: 18, 15</td>
<td>Cigarette smoking (C6/6 minutes): non-smokers: 10, 9; smokers: 11, 8</td>
</tr>
</tbody>
</table>

Abbreviations: min, minutes; mm, millimeters; NMC, nasal mucociliary clearance; s, seconds; STT, saccharin transfer time.

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...and defense mechanism) returns to its “normal” (impaired) state.35

The improvement of NMC due to smoking cessation is caused by normalization of the cilia structure, genesis, and also ciliary beating function; changes of the cytomorphic features, such as fewer columnar cells, less mucus and reduced epithelial-cell metaplasia; restoration of the ionic transport function and mucous hydration at the basolateral membrane (increased mucus clearability); and detoxification of hydrogen peroxide through the activity of the glutathione peroxidase (antioxidant enzyme).11,31

The finding of increased NMC value in passive smokers indicates that passive smokers present impairment of the NMC just like active smokers.7,29,38 It has been observed that chronic passive exposure to cigarette smoke in the household has slightly higher serum continine, which is a metabolite of nicotine that decreases ciliary activity.7 Other studies also reported that passive smoking has a negative impact on cilia regeneration, as it impairs the structure of the nasal mucosa (patchy loss of cilia, generalized loss of cilia, squamous metaplasia, vascular congestion, hyperplasia of goblet cells and seromucinous acini), and impairs epithelial ion transport (inhibit chloride secretion and basolateral K+ conductance), which can negatively affect NMC.7,29,38

Bidi is a hand-rolled tobacco product, and, unlike cigarettes, it is devoid of a filter (the filter of a cigarette functions as a barrier, by preventing the noxious smokes from reaching the nasal cavity).3,39 Although a bidi contains a lower amount of tobacco compared with a cigarette, it produces higher levels of noxious substances like nicotine, carbon monoxide, tar, phenols, and ammonia and results in a higher degree of impairment and addiction.3,40 It has been reported that bidi produces three times the amount of carbon monoxide and nicotine, and five times the amount of tar than cigarettes. Given the low combustibility of the tendu leaf wrapper, bidi smokers must take more frequent and deeper puffs, resulting in the inhalation of more smoke, which is disseminated deeper into the lungs. The decreased NMC detected in bidi smokers may be due to reduced CBF, decline in the number of cilia, and alterations in the viscoelastic properties of mucus.3

The hookah, also known as a shisha or water pipe, is a traditional method of smoking tobacco.5,41 Both hookah and cigarette contain nicotine, harmful gases such as carbon monoxide and volatile aldehydes, ultrafine particles, and carcinogenic polycyclic aromatic hydrocarbons (PAHs).5,42,43 After comparing a 45-minute hookah session to smoking a single cigarette, it was found that hookah smokers had higher concentrations of nicotine, carbon monoxide, and PAHs than cigarette smokers.5,44 Hookah smoking resulted in increased airway resistance, inflammation, immune cells (neutrophils and lymphocytes), oxidative stress, nitric oxide, and catalase activity in the lungs and NMC of animals.5,45,46 In the study by Ancgul, the group that used hookahs more than once a week was shown to have impaired NMC. This predisposes them to upper respiratory tract inflammation and injury.5

The electronic cigarette (EC) is a device that carries aerosolized nicotine to the respiratory tract.6 The EC is marketed as a safer alternative to conventional cigarettes.
due to its defined composition and noncombustible nature. A short-term study investigating the possible side-effects of EC use revealed that EC was safer than cigarettes but had more side-effects than nicotine replacement therapy. Propylene glycol is the primary ingredient in the majority of EC cartridges on the marketplace today. Nasal mucociliary clearance in EC smokers was impaired by oxidative stress induced by nicotine exposure via transient receptor potential ankyrin 1 (TRPA 1) receptors. These receptors induce airway surface liquid volume loss and decrease mucus density. Other studies also reported that formaldehyde in ECIG-generated aerosols related to DNA strand breaks and cell death and propylene glycol thicken the respiratory epithelium by increasing the number of goblet cells and increasing the content of mucin within the goblet cell.

**Strength and Limitation of the Study**

The present systematic review included studies that reported NMC in many types of smoking (cigarette, passive, bidi, hookah, and electronic smoking). In addition, a comprehensive literature search was followed as well as bias protection methods, such as using three independent reviewers.

The limitation of the study was related to the minimal sample of each study and the fact that most of the studies were cross sectional. There were only one randomized controlled trial (RCT) and three prospective studies.

**Future Implication**

The current systematic review is expected to be a scientific consideration to clinician-related NMC in smokers and general information related the dangers of smoking for the public society. Further research is needed on each component of the NMC for development upon this systematic review.

**Final Comments**

Our findings suggest that there is an impairment of NMC in smokers. The impairment is not only seen in cigarette smokers but also in passive, bidi, electronic, and hookah smokers. The impairment of NMC in chronic exposure to smoking is caused by the ciliotoxic effect, hypersecretion and viscoelastic change of mucus, airway surface liquid depletion, increased oxidative stress, and deteriorations in the inflammatory and immune systems.

**Conflict of Interests**

The authors have no conflict of interests to declare.

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