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Oral Cancer Risk Assessment for Different Type of Smokeless Tobacco Products Sold Worldwide: A Review of Reviews and Meta-analyses

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Abstract

Smokeless tobacco (SLT) use is a significant cause of lip and oral cavity cancers. Globally, oral cancer (OC) prevalence is strongly linked to the types of tobacco products used, their chemical composition, and their pattern of use. Except snus, all SLT products sold in different WHO regions are strongly associated with OC incidence. Shammah showed the highest association odds ratio (OR) with 95% confidence intervals (CI) (OR 38.74, 95% CI 19.50-76.96), followed by oral snuff (OR 11.80, 95% CI 8.45-16.49), gutkha (OR 8.67, 95% CI 3.59-20.93), tobacco with betel quid (OR 7.74, 95% CI 5.38-11.13), toombak (OR 4.72, 95% CI 2.88-7.73) and unspecified chewing tobacco (OR 4.72, 95% CI 3.13-7.11). Most SLT products containing high levels of carcinogenic tobacco-specific nitrosamines (TSNAs) exhibit a high risk of oral cancer. There is an urgent need to frame and implement international policies for OC prevention through legal control of the TSNA levels in all SLT product types.

Prevention Relevance Statement

Most smokeless tobacco products sold worldwide, mainly shammah, toombak, gutkha, betel quid with tobacco, and dry snuff, are associated with a high risk of oral cancer. A high concentration of tobacco-specific nitrosamines in SLT products is the major causative factor for oral cancer development.

Introduction

Oral cancer (OC) is a highly lethal disease and one of the most debilitating and disfiguring malignancies globally. Head and neck cancers represent the sixth most common cancer worldwide and OC accounts for ~37% of head and neck cancers with more than 500,000 cases worldwide and are predicted to rise by 62% to 856,000 cases by 2035 (1). According to global cancer statistics, Globocan 2020, cancers of the lip and oral cavity pose an enormous global challenge, with 377,713 new cases and 177,757 deaths accounting for about 3.8% of all cancer cases and 3.6% of cancer deaths globally (2).

OC is most likely caused by a combination of extrinsic and intrinsic factors acting in concert over a period of time (3,4). Major risk factors implicated in the aetiology of OCs are tobacco use (5), areca nut use (6) alcohol consumption (7), ultraviolet radiation (UVR), and human papillomavirus (HPV) infection (8). Other factors include poor oral hygiene, low socioeconomic status and genetic factors, occupational exposure (9), weakened immune system, deficiencies in dietary intake, or lack of healthy eating (10). Gender, age, physical activity and environmental factors may also play a crucial role in the progression of the disease (11,12). Tobacco and alcohol use are two of the most common risk factors for oral cavity and oropharyngeal cancers (13). As dual use of tobacco products and alcohol act synergistically, and account for 3 out of 4 oral cavity cancer cases globally (14,15). Smokeless Tobacco (SLT) includes a large variety of commercial or non-commercial tobacco preparations used orally or nasally, without combustion. Chewing tobacco, moist snuff, and dry snuff are the three most common types of SLT products used worldwide. The chewing tobacco products mainly include betel quid with tobacco, khaini, zarda and gutkha. Non-chewing products include oral snuff, nasal snuff, and snus. Snuffed tobacco products are used in either wet or dry form. Use of wet snuff is more common in the Western world, while nasal snuff in dry powder form is used in the South East Asia and Eastern Mediterranean regions (16).

The WHO South-East Asia Region (SEAR), notably the Indian subcontinent, contains 90% of the world's 250 million SLT consumers and accounts for nearly one-third of all cancers (17,18). SLT use is culturally widely acceptable due to its association with socialisation and family tradition in various parts of the world (19). SLT products may be premade (ready-to-use) or custom-made. Premade products range from large factory manufactured products to small cottage industry products, while custom made are assembled by the user or a vendor in market stalls or shops according to one's preferences. Due to the vast heterogeneity and lack of standardization, the chemical formulation or composition of SLT products show great complexity. Factors for the high prevalence of SLT are its addictive properties, easy accessibility, low cost and lack of prohibitive legislation (20). This could be the reason that the US, Food and Drug Administration's (FDA) nicotine reduction strategy which

greatly improved the health consequences of tobacco dependence in smokers, could not be applied to SLT products.

SLT causes cancers of all parts of the oral cavity including the lip, tongue, palate, gum, cheek, buccal gingivae and floor of the mouth (21), along with oesophageal and pancreatic cancer, etc. (22). More than 180,000 cases of OC occur every year in SEAR with approximately 90% of which are due to tobacco use (23). The odds of developing OC in SEAR were more than four times higher among SLT users than non-tobacco users (24,25). India has one of the highest incidences of OC and accounts for about 30% of all new cases annually due to the high prevalence of SLT use and betel-quid chewing (26). Population-based studies from 13 countries showed that the OC incidence rate is increasing, especially among the younger population (27). Other than HPV, increased incidence of early-onset oral carcinoma in the United States (US) has been associated with SLT use, mainly chewing tobacco and snuff (28).

Due to increasing awareness about smoking-related harms and growing regulatory pressures on cigarettes, the global prevalence of smoking is showing a downward trend in the last two decades (29). A systematic analysis of the global burden of disease study results in 204 countries and territories between 1990–2019 indicated that the global age-standardised prevalence of smoking had decreased significantly during this period, while the use of SLT products continued unabated during this period (30). Such a trend could be one of the reasons that the incidence of nasopharyngeal cancers has decreased dramatically (estimated annual percentage change (EAPC) -1.5, 95% CI -1.7 to -1.3) from 1990 to 2017, while the global incidence for lip and oral cavity cancers has shown a substantial increase from 1990 to 2017 (EAPC 0.26, 95% CI 0.16–0.37). Globally, the absolute number of lip and oral cavity cancers incidence increased from around 186,000 in 1990 to 389,800 in 2017, which is about a 109% increase over 28 years (31).

Great diversity in the preparation and composition of SLT products makes their regulation a big challenge. For example, gutkha is chewing tobacco mixed with areca nut and slaked lime (32), often marketed as a mouth freshener due to added flavours (33). Shammah is a traditional form of fermented chewing tobacco popular in the Middle East (34) while toombak, a homemade oral snuff mainly used in Sudan, is prepared from the tobacco leaves of *Nicotiana rustica* species having high nicotine content (35). Weak enforcement of regulatory policies and aggressive marketing of SLT products by the tobacco industry worsens the situation (36,37).

Broadly, reports quantifying the promotion of all types of SLT, as a harm reduction strategy and as a safer alternative to cigarettes, have shown no apparent health benefits at a population level (38). On the other hand, this has caused an increase in the sale of SLT. Because nicotine content in a cigarette stick varies from 0.8 to 13.0 mg/g, while it ranges from 0.8 to 76.0 mg/g in SLT products (39), SLT

users absorb two to three times the amount of nicotine as those who smoke cigarettes (40). This is due to the high alkaline nature of most SLT products providing free nicotine at a high concentration in a short time. Excessive high nicotine concentration makes SLT products highly addictive, and nicotine is also a precursor of carcinogenic tobacco-specific N-nitrosamines (TSNAs) (41,42).

Nicotine and Tobacco Specific Nitrosamine (TSNA) levels

TSNAs are chemically stable compounds under physiological conditions and are found to be associated with carcinogenicity in humans and experimental animals (43). TSNAs mainly N'-nitrosonornicotine (NNN) and nicotine-derived nitrosamine ketone (NNK) are listed as group 1 human carcinogens by IARC (3). They are shown to disrupt DNA repair and molecular processes and are the prime cause of OC in SLT users (44–46).

Addictiveness and health hazards of SLT across the globe are largely dependent upon product's chemical composition and its use pattern (47). Globally, the magnitude of cancer risk due to SLT use shows disparity and is highly correlated with the variation in the levels of NNN and NNK present in diverse SLT products sold worldwide (48,49). Seeing the carcinogenicity of NNN and NNK in humans, the WHO Study Group on Tobacco Product Regulation in 2010 recommended a regulatory limit for maximal total concentration of NNN and NNK as less than 2 µg/g dry weight of tobacco (48). However, the levels of NNN and NNK, per unit dose, in SLT products are much higher as compared to cigarette smoke. While on an average mainstream cigarette smoke contains NNK and NNN in the range of 0.006–1.74 µg/g and 0.004–2.83 µg/g, respectively, SLT products sold across the world showed NNK levels between 0.019 to 7870 µg/g and NNN levels between 0.080 to 3080 µg/g against the WHO permissible limit of less than 2 µg/g.

Swedish Match, the principal manufacturer of Swedish moist snuff, adopted a voluntary standard for TSNAs levels, called the *GothiaTek* standard. (50). Table 1 represents comparative data on the type of SLT sold across the world, its preparation process and use, country/ WHO region, levels of nicotine, total TSNAs, NNN and NNK. SLT products viz. shammah gutkha, toombak, betel quid with tobacco, chewing tobacco (unspecified) along with dry snuff and moist snuff (snus) were found to contain high levels of carcinogenic TSNAs, mainly NNN and NNK in them.

Many research articles in the previous years have indicated the link between SLT and OC but the present systematic review, for the first time, describes the levels of risk estimates of OC associated with the major individual type of SLT products sold across the five world health organisation (WHO) regions. It also reports the WHO region-wise OC risk estimates associated with different SLT products and compiles data on the global pattern of different types of SLT product use and the concentration of nicotine, total TSNAs, NNN and NNK in them.

Materials and Methods

Electronic Searches

An electronic search was conducted on PubMed and Google Scholar for articles published between Jan 1, 2010, to Aug 5, 2021 using the key phrases “oral cancer”, “oral squamous cell carcinoma” “smokeless tobacco”, “chewing tobacco”, “betel quid”, “snuff”, “snus”, “gutkha/gutka”, “toombak” and “shammah”. The references of relevant articles were manually searched for additional eligible citations. This comprehensive review presents pooled data from the different studies.

Selection of Studies

Author, AKG extracted data through this literature search and identified studies. Duplicate records were removed, and the reference lists of the selected articles were screened for additional relevant articles. Titles and abstracts of papers identified through the search strategy were reviewed and relevant articles, potentially fulfilling the inclusion criteria, were retrieved in full text. A second reviewer (RM) screened the titles and abstracts of the retrieved articles to identify the relevance of the articles to the objectives of this review. Two authors, AKG and MK, independently assessed the eligibility of the selected data to assure quality and minimise biases. **Figure 1** provides the detailed strategy of the study selection process using PRISMA guidelines.

Inclusion Criteria -

- Oral cancer had to be one of the outcomes of smokeless tobacco use in the adult population.
- Articles presented only as reviews, systematic reviews and meta-analyses.
- Studies providing odds ratio (OR)/risk ratio (RR) estimates with corresponding 95% confidence intervals (CI).
- Articles published in English.

Exclusion Criteria

- Studies not designed to investigate SLT association with OC.
- Articles published before year 2010.
- Articles published in languages other than English.

Data Extraction

For articles meeting the eligibility criteria, the following information was extracted: the study authors with the date of publication, region of the study, the type of smokeless tobacco, period of study, OR/RR estimates and corresponding 95% CI. Information was extracted by one author AKG and checked by another author, MK. (Supplementary Table 1)

The region of the study was classified as global or as one of the WHO regions, namely, the American Region (AMR), Eastern Mediterranean Region (EMR) including Pakistan, European Region (EUR), African Region (AFR) and South-East Asian Region (SEAR). The type of tobacco was classified as: any type of smokeless tobacco, if not explicitly specified which type, shammah (Arabian chewing

tobacco), toombak (Sudanese dipping tobacco), gutkha (Indian chewing tobacco), betel quid with tobacco, chewing tobacco (unspecified), dry snuff and moist snuff (snus). If a review article had been updated, then the updated review estimates were used and if two reviews cite the same source, then the one reporting pooled estimates was used.

Data Analysis

We used forest plot graphs to represent the OR/RR estimates and 95% CI. Results were stratified by WHO region and by tobacco type. No overall pooled analysis was conducted. If a previous review reported individual studies without pooling the results, these were pooled if the estimates were provided together with 95% CI or other information to enable pooling the results. All studies were systematic reviews with meta-analysis except one study on toombak where the combined OR estimates were not reported and thus were calculated (see supplementary method).

Ethics Statement

Article does not contain any studies involving human or animal participants.

Data Availability Statement

The data generated in this study are available upon request from the first author AKG.

Note: Supplementary data for this article are available at Cancer Prevention Research Online (<http://cancerprevres.aacrjournals.org/>)

Results

Articles, published in the last decade, i.e., from 2010 to 2021 and reporting the OC risk estimates in the association of the SLT product, were selected for the present review. After removing duplicate records, titles and abstracts of 74 records were retrieved through the selected databases. The reference lists of the included articles were screened for 4 additional articles. All 78 articles were reviewed thoroughly. After removing 52 irrelevant articles, 26 were selected for the full-text study, of which, 17 which did not meet the selection criteria, were excluded. **Figure 1** demonstrates the flow-chart of the study selection process for smokeless tobacco use and oral cancer risk using PRISMA guidelines. Oral potentially malignant disorders are abbreviated as OPMD in fig 1).

Nine studies fulfilling all the eligibility criteria for inclusion were finally included in the current review. Of these, three reviews evaluated the risk of OC with the use of all types of SLT products combined (51–53). Three reported OR estimates for betel quid with tobacco (51,54,55). Dry snuff was evaluated for high risk of OC in three studies (51,56,57). Two studies mentioned chewing tobacco (without specifying the type) (51,56), while one study each was found on shammah (58), gutkha (51), toombak (59) and snus (51). All the selected studies are systematic reviews with meta-analysis and OR estimates were adjusted for confounding factors mainly smoking except for one study (59). (Supplementary Table 1)

Data analysis of all included studies together indicated that the individual product that showed the highest association (OR 38.74, 95% CI 19.50-76.96) was shammah, followed by oral snuff (OR 11.80, 95% CI 8.45-16.49), gutkha (OR 8.67, 95% CI 3.59-20.93), tobacco with betel quid (OR 7.74, 95% CI 5.3-11.13), toombak (calculated OR 4.72, 95% CI 2.88-7.73, please see supplementary method) and unspecified chewing tobacco (OR 4.72, 95% CI 3.13-7.11). Overall, all selected SLT product types, except snus, were found to have a strong association with OC incidence across the globe. **Figure 2** represents a forest plot of the included studies showing odds ratios and 95% confidence intervals (CI) for the association between the types of SLT products and the risk of OC. Region-wise analysis of SLT products showed that the overall global OR for OC for all SLT types combined, ranged from 3.53 (95% CI 2.76-4.52) to 3.94 (95% CI 2.70-5.75). In general, region-wise OC risk estimates, for all types combined, were highest for EMR with OR ranging from 1.28 (95% CI 1.05-1.57) to 14.52 (95% CI 7.69-27.41), followed by SEAR with OR 4.44 (95% CI 3.51-5.61) to 5.67 (95% CI 3.83-8.40) and for AMR, OR 0.95 (95% CI 0.71-1.25) to 4.72 (95% CI 0.66-33.69), while it was not statistically significant for EUR with OR 0.94 (95% CI 0.71-1.25). For further details, see **figure 3** which represents a forest plot of included studies by the WHO region.

A strong positive association of betel quid with tobacco and OC was seen globally OR 7.18 (95% CI 5.489-41) (51) while for Asian studies risk estimates for betel quid with tobacco range from OR 7.10 (95% CI 4.49–11.22) to 7.74 (5.38-11.13) (54,55), toombak and shammah use for EMR, showed highest OC risk estimate with OR 4.72 (95% CI 2.88-7.73) (56) and OR 38.74 (95% CI 19.50-76.96) respectively (58). Risk estimates for snuff-type products vary significantly among various WHO regions. In EUR and AMR, dry snuff and snus are more prevalent. Global OC risk estimates for oral snuff showed OR 4.18 (95% CI 2.37-7.38) (51) while for AMR, OR was 3.01 (95% CI 1.63-5.55) (56). Naswar, used in EMR was shown to have a high OR value of 11.80 (95% CI 8.45-16.49) (57). Globally, chewing tobacco, is shown to have a high OC risk with OR 4.37 (95% CI 3.27-5.84) as compared to non-chewing SLT products with OR 1.56 (95% CI 1.04- 2.35) (51). **Figure 3**

The level of TSNA in SLT products plays a significant role in carcinogen exposure levels. Thus the difference in the magnitude of OC risks can be correlated with the variation in the levels of NNN and NNK present in SLT products (49). TSNA levels varies from 0.08 µg/g to as high as 992 µg/g in the selected SLT products. **Figure 4** indicates that high levels of TSNA are present in SLT products with a high-risk ratio for OC. Fig 4 (a) presents TSNA values on the log scale while the original TSNA levels in µg/g are presented on the right-hand side of the y-axis. (b) OR and corresponding 95% CIs estimates are based on review studies from the same region that the SLT product TSNA values are based. The OR estimates for zarda and khaini are not product specific but those for all types of chewing tobacco from SEAR (54). For gutkha, dry snuff and snus the OR estimates are

based on global pooled estimates (51), whereas for naswar (a nasal snuff) these are based on EMR estimates only (57).

Discussion

Global Pattern of Oral Cancer Risk Estimates for different SLT Products

According to a recent study, published in Lancet Public Health, out of the total 273·9 million tobacco chewers (age 15 and above) in the world, about 228·2 million lived in SEAR (30). Over the past several decades, it has been seen that SLT use has increased by nearly 50% in low-and-middle-income countries (LMIC) while declining in high-income countries (60). Tobacco chewing and betel quid with tobacco are the two most prevalent forms of SLT use in Asia (61). In India, the majority of SLT users consume chewing tobacco (11.6% khaini, 8.2% gutkha preparations, 6.2% betel quid with tobacco, 4.7% oral snuff and 4.4% other SLT products) (51). Gutkha use has been gaining popularity in Europe and US in the last two decades due to its easy availability, low cost and extensive marketing (62). In the US, the sale of SLT products increased by 5.8% between 2011 and 2016, but declined by 3.9% from 2016 to 2019; however, the sale of snus consistently increased while the sale of chewing tobacco, dry snuff, and dissolvable decreased during this period (63).

A recent CDC report indicated that the incidence of cancers of the oral cavity and pharynx (all sites), not associated with HPV, increased in the US during 2007–2016 (64). In 2018, an estimated 120,000 new OC patients were diagnosed with 72,000 deaths in India alone (65). Studies revealed that a higher risk of OC was observed for SLT products sold before 1990 (OR 6.6, 95% CI 5.3-8.2) as compared to that sold after 1990 (OR 3.0, 95% CI 2.3-3.9) (17). Dry snuff sold in the US and Western Europe, before 2000, was shown to have higher relative risks for OCs (RR 8, 95% CI 2.7-20.0) (66). This is due to improvement in the quality of manufactured tobacco products. Most SLT products sold in the US after 1990, achieved TSNA levels below 20 ppm as compared to generally high TSNA levels (above 100 ppm) in earlier SLT products, sold before 1990 (67). Previous studies showed that snus had an association with an increased risk of oral or pancreatic cancer as compared to non-tobacco users (68,69). However, the current prevalence statistics and epidemiological data on snus use, in the European population, do not indicate an increased risk of OC compared to cigarettes (70).

More than 50% of OCs are attributable to using SLT products in Sudan and India compared to about 4% in US men (65). Literature studies show that toombak has a major role in the aetiology of oral/oropharyngeal cancer in Sudan (71,72) and sub-Saharan Africa (73). OC occurrence is about 3 to 6 times higher in North-East Nigeria than reported for the US and Europe -mainly due to the use of dry snuff (OR 10, 95% CI 4.1-4.3) (74,75). Oral cancer is the third most common malignancy in Saudi Arabia mainly due to the use of shammah, the traditional form of chewing tobacco prevalent in

the Middle East, Yemen and Sudan (76). A review of studies by Awan and Patil showed that in the SEAR, the OC risk estimates (OR) for betel quid varied from 3.1 to 15.7 (95% CI 11.0-22.1) and from 1.2 (95% CI 1.0-1.4) to 12.9 (95% CI 7.5-22.3) for chewing tobacco (43).

The frequency of SLT use was also seen to vary substantially across countries and by sex, age, ethnic origin, and socioeconomic characteristics within a country (77). A linear dose-response association was observed between OC and chewing tobacco regarding age at initiation, duration, and frequency of chewing per day (78).

Most SLT users have limited awareness of its association with OC due to a lack of knowledge of its harmful constituents and high use due to cultural traditions/ religious norms (79). According to the Global Adult Tobacco Survey in India (GATS, 2016-17), the prevalence of SLT use is very high, especially in females, which could be due to a lack of awareness and knowledge about the health hazards of the SLT product used (80). In the Indian subcontinent, betel quid chewing, with added tobacco has a much higher risk ratio in women (OR 14.6, 95% CI 7.6-27.8) (55). Globally, gender-wise sub-group analysis showed a higher risk for females with (OR 5.8, 95% CI 2.9-11.6), as compared to males (OR 2.7, 95% CI 1.7-4.3) (51).

High Levels of Nicotine and Tobacco-specific Nitrosamines (TSNAs) in Smokeless Tobacco Products and Oral Cancer

High nicotine content in SLT products is responsible for the increased levels of TSNAs which are primarily formed during tobacco fermentation and storage, especially at elevated temperature and moisture (81). A global surveillance study across 113 countries from five WHO regions over the past 10 years, indicated that diverse SLT products sold worldwide seem to contain high levels of carcinogenic TSNAs (52). Maximum concentrations of NNN and NNK content for toombak products from Sudan were found to be 3085 and 7870 µg/g respectively which were remarkably higher than most of the products sold worldwide (82). Average levels of NNN, in a brand of khaini, marketed as snus, were 22.9 and 2.6 µg/g tobacco respectively (83). Khaini, sold in South Asia, contains alarmingly high levels of NNN (39.4-76.9 µg/g) and NNK (2.34-28.4 µg/g) (84). Snuff sold in America was shown to have TSNAs levels as high as 76.5 µg/g, while NNN (0.37-42.6 µg/g) and NNK (0.38-9.9 µg/g) (85). The literature did not report levels of TSNAs in shammah, showing the highest OR. On average, NNN and NNK levels showed an almost 70-fold variation with NNN concentrations ranging from 0.09 to 76.9 µg/g while NNK levels ranged from 0.04 to 28.4 µg/g in all selected SLT products (6). Fermented SLT products, like toombak, shammah, dry snuff, khaini, gutkha, have been found to contain higher levels of TSNA than pasteurised products like snus (84). Shammah, a highly fermented product with high nicotine content (86), is made under long anaerobic conditions so more nitrite is generated which increases TSNA concentration. However, no study was

found reporting the TSNA levels in shammah (34). The OR of developing OC, for shammah users was 38.7 (95% CI 19.5-77.0) which was nearly 39 times higher than non-shammah users (58). Studies showed that NNN and NNK levels for toombak were about 100 folds higher than most of the products sold worldwide (87,88). OC risk estimate for toombak use was significantly high among users in comparison with controls (OR 3.8, 95% CI 1.7-8.6) (89). A report showed that US snus had high TSNA levels with NNN and NNK as high as 42.55 and 9.95 µg/g, respectively (90). Dry snuff, the major factor for tongue carcinoma in the US, is shown to contain high TSNA levels (91). On the other hand, Swedish snus made with improved manufacturing techniques has low OC risks due to low levels of NNN and NNK (92). Thus, the levels of nicotine and TSNA showed several hundred-fold variations across different product types and substantial vendor-to-vendor variation within some product categories (93).

Thus, SLT products with higher NNN concentration pose higher cancer risks, so reducing the levels of carcinogenic nitrosamines in finished SLT products could prove a beneficial strategy to reduce OR risk for OC (94,95).

For the protection of public health, FDA has proposed a tobacco product standard rule, which states that the mean level of NNN in any batch of finished SLT product should not exceed 1.0 µg/g of tobacco (on a dry weight basis) at any time through the product's labelled expiration date (96). However, constituent regulation and control of SLT products lag far behind cigarettes, mainly due to non-standardised production and storage methods, greater heterogeneity and the lack of strict legal policies for SLT (39).

Conclusions

The current review is to bring attention to the prevention community to the risks of individual smokeless tobacco product for risk of oral cancer. Most carcinogenic SLT types sold across the various geographic regions worldwide, mainly shammah, toombak, gutkha, betel quid with tobacco, dry snuff were found to be associated with high OC risks. Data analysis indicated that the shammah showed the highest association (OR 38.7, 95% CI 19.5-77.0), followed by oral snuff (OR 11.8, 95% CI 8.4-16.4), gutkha (OR 8.7, 95% CI 3.6-20.9), tobacco with betel quid (OR 7.7, 95% CI 5.3-11.1), toombak (OR 4.7, 95% CI 2.9-7.7) and unspecified chewing tobacco (OR 4.7, 95% CI 3.0-7.1). The difference in the magnitude of OC risks has been found to correlate highly with regional variation in the SLT product type which showed great diversity and heterogeneity in its composition, usage and manufacturing process. A decrease in smoking and the prevalence of lung cancer in the US shows the effectiveness of decades of public education and tobacco control policies (97). However, the rising incidence of OC across the world, primarily associated with SLT use, indicates that the tobacco control policies do not have a more prominent effect on SLT usage. The huge variation in the levels

of carcinogenic TSNA's, especially NNN and NNK, in diverse types of SLT products, hinders the comparability of results from evaluating the global risks estimate of SLT to human health across the globe. It is imperative to develop and effectively implement strategies for monitoring TSNA levels in SLT products. There is a critical need for systematic surveillance of all types of SLT products through legal control of the permissible TSNA levels. Global standards for testing and measuring TSNA's levels in all types of SLT products, with effective measures to minimise the levels of TSNA, can significantly help reduce OC risk associated with individual SLT products.

Road Ahead

The high concentration of TSNA's, mainly NNN and NNK, in diverse types of SLT products is the major causative factor for the development of OC. Applying a grassroots approach to lower the levels of carcinogenic TSNA's at various stages of SLT production, right from its growth, processing, manufacturing, and storage, could prove to be a beneficial strategy. This includes the use of tobacco plant varieties having low levels of nitrate and TSNA's precursors, decreasing the use of nitrate fertilisers and chemical pesticides while growing tobacco, avoiding microbial contamination during tobacco processing, air-curing of leaves instead of fire curing under controlled conditions, use of newer technologies like heat treatment, pasteurisation for tobacco processing and avoiding tobacco fermentation etc. can significantly lower the concentration of carcinogenic TSNA's in the finished SLT products (39).

As the majority of OC are preventable through risk factors intervention, creating awareness about their carcinogenicity among consumers, constituent's disclosure along with their health hazard information on all SLT products may play a key factor in reducing oral cancer incidence in the future. Strict regulatory measures are to be taken for the additives and flavouring agents in SLT products, which make them palatable and more appealing especially amongst youth (98).

For the first time, the World Health Assembly, in 2007, passed a resolution on oral health and oral cancer prevention to be an integral part of national cancer control programs. The WHO global oral health program was launched to work for the capacity building in OC prevention in different countries, inter-country exchange and the development of global surveillance systems for OC and risk factors. With the establishment of more cancer registries across the globe and their secondary data analysis, the surveillance of SLT products should become easier.

Conflict of Interest statement:

The authors declare no potential conflicts of interest.

References

1. Shield KD, Ferlay J, Jemal A, Sankaranarayanan R, Chaturvedi AK, Bray F, et al. The global incidence of lip, oral cavity, and pharyngeal cancers by subsite in 2012. CA Cancer J Clin.

- American Cancer Society; 2017;67(1):51–64.
2. WHO-IARC. Globocan 2020: New Global Cancer Data,UICC. Available from:
<https://gco.iarc.fr/today/data/factsheets/cancers/1-Lip-oral-cavity-fact-sheet.pdf>
3. International Agency for Research on Cancer. A Review of Human Carcinogens. E. Personal Habits and Indoor Combustions. IARC Working Group on the Evaluation of Carcinogenic Risks to Humans. 2012;100E:585.
4. Conway DI, Purkayastha M, Chestnutt IG. The changing epidemiology of oral cancer: Definitions, trends, and risk factors. Br Dent J. Nature Publishing Group; 2018;225(9):867-73.
5. Petti S. Lifestyle risk factors for oral cancer. Oral Oncol. 2009;45(4-5):340–50.
6. Gupta AK, Tulsyan S, Thakur N, Sharma V, Sinha DN, Mehrotra R. Chemistry, metabolism and pharmacology of carcinogenic alkaloids present in areca nut and factors affecting their concentration. Regul Toxicol Pharmacol. 2020;110:104548.
7. Akinkugbe AA, Garcia DT, Brickhouse TH, Mosavel M. Lifestyle risk factor related disparities in oral cancer examination in the U.S: A population-based cross-sectional study. BMC Public Health. BioMed Central Ltd.; 2020;20(1):153.
8. Tanaka TI, Alawi F. Human Papillomavirus and Oropharyngeal Cancer. Dent. Clin. North Am. W.B. Saunders; 2018;62(1):111–20.
9. Zini A, Czerninski R, Sgan-Cohen HD. Oral cancer over four decades: epidemiology, trends, histology, and survival by anatomical sites. J Oral Pathol Med. John Wiley & Sons, Ltd; 2009;39:299–05.
10. Freedman N, Park Y, Subar A, Hollenbeck A, Leitzmann M, Schatzkin A, et al. Fruit and vegetable intake and head and neck cancer in a large United States prospective cohort study. Cancer Res. 2007;67:849.
11. Elshahat S, Treanor C, Donnelly M. Factors influencing physical activity participation among people living with or beyond cancer: a systematic scoping review. Int. J. Behav. Nutr. Phys. Act. BioMed Central Ltd; 2021;18(1):50.
12. Curtis DC, Eckhart SC, Morrow AC, Sikes LC, Mridha T. Demographic and behavioral risk factors for oral cancer among Florida residents. J Int Soc Prev Community Dent. Wolters Kluwer (UK) Ltd.; 2020;10:255–61.
13. Goldenberg D, Lee J, Koch WM, Kim MM, Trink B, Sidransky D, et al. Habitual risk factors for head and neck cancer. Otolaryngol. - Head Neck Surg. Otolaryngol Head Neck Surg; 2004; 131(6) 986–93.
14. Blot WJ, McLaughlin JK, Winn DM, Austin DF, Greenberg RS, Preston-Martin S, et al. Smoking and Drinking in Relation to Oral and Pharyngeal Cancer. Cancer Res.

1988;48(11):3282-7.

15. Pelucchi C, Gallus S, Garavello W, Bosetti C, La Vecchia C. Cancer risk associated with alcohol and tobacco use: Focus on upper aero-digestive tract and liver. *Alcohol Res. Heal.* National Institute on Alcohol Abuse and Alcoholism; 2006;29(3):193–8.
16. National Cancer Institute and Centers for Disease Control and Prevention. *Smokeless Tobacco and Public Health: A Global Perspective*. Bethesda, MD: U.S. Department of Health and Human Services, Centers for Disease Control and Prevention and National Institutes of Health, National Cancer Institute. NIH Publication No. 14-7983 2014;79-83
17. Warnakulasuriya S. Global epidemiology of oral and oropharyngeal cancer. *Oral Oncol.* Pergamon; 2009;45(4-5):309–16.
18. Datta S, Chaturvedi P, Mishra A, Pawar P. A review of Indian literature for association of smokeless tobacco with malignant and premalignant diseases of head and neck region. *Indian J Cancer.* Medknow Publications; 2014;51(3):200-8.
19. Mishra S, Mishra MB. Tobacco: Its historical, cultural, oral, and periodontal health association. *J of Internat Soc. of Preventive and Comm. Dentistry* 2013;3(1):12-18.
20. Yadav A, Singh PK, Yadav N, Kaushik R, Chandan K, Chandra A, et al. Smokeless tobacco control in India: policy review and lessons for high-burden countries. *BMJ Glob Heal.* BMJ Publishing Group. 2020;5:2367.
21. McGuire S. *World Cancer Report 2014*. Geneva, Switzerland: World Health Organization, International Agency for Research on Cancer, WHO Press, 2016. *Adv Nutr.* 2016;7(2):418–9.
22. Gupta S, Gupta R, Sinha DN, Mehrotra R. Relationship between type of smokeless tobacco & risk of cancer: A systematic review. *Indian J. Med. Res. Suppl.* Indian Council of Medical Research; 2018;148(1)56–76.
23. Jiang X, Wu J, Wang J, Huang R. Tobacco and oral squamous cell carcinoma: A review of carcinogenic pathways. *Tob Induc Dis.* The International Society for the Prevention of Tobacco Induced Diseases. 2019;17:29.
24. Sinha DN, Suliankatchi RA, Gupta PC, Thamarangsi T, Agarwal N, Parascandola M, et al. Global burden of all-cause and cause-specific mortality due to smokeless tobacco use: systematic review and meta-analysis. *Tob Control.* 2018;27:35–42.
25. Miranda-Filho A, Bray F. Global patterns and trends in cancers of the lip, tongue and mouth. *Oral Oncol.* Elsevier Ltd; 2020;102:104551.
26. IARC Monographs on the Evaluation of Carcinogenic Risks to Humans VOLUME 85, *Betel-quid and Areca-nut Chewing and Some Areca-nut-derived Nitrosamines*; 2004; 80-112.
27. Hussein AA, Helder MN, de Visscher JG, Leemans CR, Braakhuis BJ, de Vet HCW, et al.

Global incidence of oral and oropharynx cancer in patients younger than 45 years versus older patients: A systematic review. *Eur. J. Cancer*. Elsevier Ltd. 2017; 82:115–27.

28. Campbell BR, Sanders CB, Netterville JL, Sinard RJ, Rohde SL, Langerman A, et al. Early onset oral tongue squamous cell carcinoma: Associated factors and patient outcomes. *Head Neck*. John Wiley and Sons Inc.; 2019;41(6):1952–60.
29. Wang TW, Kenemer B, Tynan MA, Singh T, King B. Consumption of Combustible and Smokeless Tobacco — United States, 2000–2015. *MMWR Morb Mortal Wkly Rep*. Centers for Disease Control MMWR Office. 2016;65:1357–63.
30. Kendrick PJ, Reitsma MB, Abbasi-Kangevari M, Abdoli A, Abdollahi M, Abedi A, et al. Spatial, temporal, and demographic patterns in prevalence of chewing tobacco use in 204 countries and territories, 1990–2019: a systematic analysis from the Global Burden of Disease Study 2019. *Lancet Public Heal*. Elsevier. 2021;6(7):E482–99.
31. Du M, Nair R, Jamieson L, Liu Z, Bi P. Incidence Trends of Lip, Oral Cavity, and Pharyngeal Cancers: Global Burden of Disease 1990–2017. *J. Dent. Res*. SAGE Publications Inc. 2020; 99(2):143–51.
32. Niaz K, Maqbool F, Khan F, Bahadar H, Ismail Hassan F, Abdollahi M. Smokeless tobacco (paan and gutkha) consumption, prevalence, and contribution to oral cancer. *Epidemiol Health*. Korean Society of Epidemiology. 2017;39:e2017009.
33. Sankhla B, Kachhwaha K, Hussain SY, Saxena S, Sireesha SK, Bhargava A. Genotoxic and Carcinogenic Effect of Gutkha: A Fast-growing Smokeless Tobacco. *Addict Heal*. Addict Health; 2018;10(1):52–63.
34. Alsanosy RM. Smokeless tobacco (shammah) in Saudi Arabia: A review of its pattern of use, prevalence, and potential role in oral cancer. *Asian Pacific J. Cancer Prev*. Asian Pacific Organization for Cancer Prevention; 2014;15(16):6477–83.
35. Idris AM, Prokopczyk B, Hoffmann D. Toombak: A major risk factor for cancer of the oral cavity in Sudan. *Prev Med (Baltim)*. *Prev Med*; 1994;23:832–9.
36. Kaur J, Thamarangsi T, Rinkoo AV. Regulating smokeless tobacco and processed areca nut in South-East Asia region: The journey so far and the road ahead. *NLM (Medline)*; 2017;61:S3-S6.
37. Kumar A, Bhartiya D, Kaur J, Kumari S, Singh H, Saraf D, et al. Regulation of toxic contents of smokeless tobacco products. *Indian J. Med. Res. Suppl*. Indian Council of Medical Research; 2018;148(1):14–24.
38. Mejia AB, Ling PM, Glantz SA. Quantifying the effects of promoting smokeless tobacco as a harm reduction strategy in the USA. *Tob Control*. BMJ Publishing Group Ltd;

2010;19(4):297–305.

39. Gupta AK, Tulsyan S, Bharadwaj M, Mehrotra R. Grass roots approach to control levels of carcinogenic nitrosamines, NNN and NNK in smokeless tobacco products. *Food Chem Toxicol.* Elsevier Ltd; 2019;124:359–66.
40. Wollina U, Verma SB, Ali FM, Patil K. Oral submucous fibrosis: an update. *Clin Cosmet Investig Dermatol.* Dove Press; 2015;8:193–204.
41. Hecht SS. Tobacco carcinogens, their biomarkers and tobacco-induced cancer. *Nat Rev Cancer.* Nature Publishing Group; 2003;3:733–44.
42. Hecht SS, Carmella SG, Murphy SE, Riley WT, Le C, Luo X, et al. Similar Exposure to a Tobacco-Specific Carcinogen in Smokeless Tobacco Users and Cigarette Smokers. *Cancer Epidemiol Biomarkers Prev.* 2007;16(8):1567–72.
43. Awan KH, Patil S. Association of smokeless tobacco with oral cancer - evidence from the South Asian studies: A systematic review. *J. Coll. Physicians Surg. Pakistan. College of Physicians and Surgeons Pakistan;* 2016;26(9):775–80.
44. De Geu JL, Wambier L-MLM, Loguercio ADA-DAD, Reis A, de Geus J-L, Wambier L-MLM, et al. The smokeless tobacco habit and DNA damage: A systematic review and meta-analysis. *Med Oral Patol Oral Cir Bucal. Medicina Oral S.L.;* 2019;24:e145–55.
45. Critchley JA, Unal B. Health effects associated with smokeless tobacco: A systematic review. *Thorax.* Thorax; 2003;58(5):435–43.
46. Xue J, Yang S, Seng S. Mechanisms of Cancer Induction by Tobacco-Specific NNK and NNN. *Cancers (Basel).* Multidisciplinary Digital Publishing Institute (MDPI); 2014;6:1138–56.
47. Giovino GA, Mirza SA, Samet JM, Gupta PC, Jarvis MJ, Bhala N, et al. Tobacco use in 3 billion individuals from 16 countries: An analysis of nationally representative cross-sectional household surveys. *Lancet.* Lancet Publishing Group; 2012;380:668–79.
48. World Health Organization Study Group on Tobacco Product Regulation. WHO Study Group on Tobacco Product Regulation. Report on the scientific basic of tobacco product regulation: fourth report of a WHO study group. *World Health Organ Tech Rep Ser.* 2012;(967):1-83
49. Hatsukami DK, Stepanov I, Severson H, Jensen JA, Lindgren BR, Horn K, et al. Evidence Supporting Product Standards for Carcinogens in Smokeless Tobacco Products. *Cancer Prev Res.* 2015;8(1):20–6.
50. Bates C, Fagerström K, Jarvis MJ, Kunze M, McNeill A, Ramström L. European Union policy on smokeless tobacco: a statement in favour of evidence based regulation for public health. *Tob Control.* Tob Control; 2003;12(4):360–7.

51. Asthana S, Labani S, Kailash U, Sinha DN, Mehrotra R. Association of Smokeless Tobacco Use and Oral Cancer: A Systematic Global Review and Meta-Analysis. *Nicotine Tob Res.* NLM (Medline); 2019;21(9):1162–71.
52. Siddiqi K, Husain S, Vidyasagaran A, Readshaw A, Mishu MP, Sheikh A. Global burden of disease due to smokeless tobacco consumption in adults: an updated analysis of data from 127 countries. *BMC Med. BioMed Central*; 2020;18(1):222.
53. Sinha DN, Abdulkader RS, Gupta PC. Smokeless tobacco-associated cancers: A systematic review and meta-analysis of Indian studies. *Int J Cancer.* Wiley-Liss Inc.; 2016;138(6):1368–79.
54. Khan Z, Tönnies J, Müller S. Smokeless tobacco and oral cancer in South Asia: a systematic review with meta-analysis. *J Cancer Epidemiol.* Hindawi Publishing Corporation; 2014;2014:394696.
55. Guha N, Warnakulasuriya S, Vlaanderen J, Straif K. Betel quid chewing and the risk of oral and oropharyngeal cancers: A meta-analysis with implications for cancer control. *Int J Cancer.* Wiley-Liss Inc.; 2014;135(6):1433–43.
56. Wyss AB, Hashibe M, Lee Y-CCA, Chuang S-CC, Muscat J, Chen C, et al. Smokeless tobacco use and the risk of head and neck cancer: Pooled analysis of US studies in the inance consortium. *Am J Epidemiol.* Oxford University Press; 2016;184(10):703–16.
57. Khan Z, Suliankatchi RA, Heise TL, Dreger S. NASWAR (smokeless tobacco) use and the risk of oral cancer in Pakistan: A systematic review with meta-analysis. *Nicotine Tob. Res.* Oxford University Press; 2019;21(1):32–40.
58. Quadri MFA, Tadakamadla SK, John T. Smokeless tobacco and oral cancer in the Middle East and North Africa: A systematic review and meta-analysis. *Tob. Induc. Dis. International Society for the Prevention of Tob Induc Dis*; 2019;17:56
59. Patil S, Arakeri G, Alamir AWH, Patil S, Awan KH, Baeshen H, et al. Is toombak a risk factor for oral leukoplakia and oral squamous cell carcinoma ? A systematic review. *J. Oral Pathol. Med.* Blackwell Publishing Ltd; 2020;49(2):103–9.
60. Mummudi N, Agarwal JP, Chatterjee S, Mallick I, Ghosh-Laskar S. Oral Cavity Cancer in the Indian Subcontinent – Challenges and Opportunities. *Clin Oncol.* Elsevier Ltd; 2019;31(8):520–8.
61. Sinha DN, Bajracharya B, Khadka BB, Rinchen S, Bhattad VB, Singh PK. Smokeless tobacco use in Nepal. *Indian J. Cancer.* Indian J Cancer; 2012;49(4): 352–6.
62. Changrani J, Cruz G, Kerr R, Katz R, Gany FM. Paan and Gutka Use in the United States. *J Immigr Refug Stud.* 2006;4(1):99–110.

- 580 63. Delnevo CD, Hrywna M, Miller Lo EJ, Wackowski OA. Examining Market Trends in
581 Smokeless Tobacco Sales in the United States: 2011-2019. *Nicotine Tob Res* Oxford
582 University Press 2021;23(8):1420–4.
- 583 64. Ellington TD, Henley SJ, Senkomago V, O’Neil ME, Wilson RJ, Singh S, et al. Trends in
584 Incidence of Cancers of the Oral Cavity and Pharynx — United States 2007–2016. *MMWR*
585 *Morb Mortal Wkly Rep. Centers for Disease Control MMWR Office*; 2020;69(15):433–8.
- 586 65. Boffetta P, Hecht S, Gray N, Gupta P, Straif K, Paolo Boffetta, Stephen Hecht, Nigel Gray,
587 Prakash Gupta KS. Smokeless tobacco and cancer. *Lancet Oncol. Lancet Publishing Group*;
588 2008;9(7):667–75.
- 589 66. Lee PN, Hamling J. The relation between smokeless tobacco and cancer in Northern Europe
590 and North America. A commentary on differences between the conclusions reached by two
591 recent reviews. *BMC Cancer. BioMed Central*; 2009;9:256.
- 592 67. Rodu B, Jansson C. Smokeless tobacco and oral cancer: A review of the risks and
593 determinants. *Crit. Rev. Oral Biol. Med. Crit Rev Oral Biol Med*; 2004;15(5):252–63.
- 594 68. Roosaar A, Johansson ALV, Sandborgh-Englund G, Axéll T, Nyrén O. Cancer and mortality
595 among users and nonusers of snus. *Int J Cancer. 2008*;123(1):168–73.
- 596 69. Luo J, Ye W, Zendehdel K, Adami J, Adami HO, Boffetta P, et al. Oral use of Swedish moist
597 snuff (snus) and risk for cancer of the mouth, lung, and pancreas in male construction workers:
598 a retrospective cohort study. *Lancet. Lancet*; 2007;369(9578):2015–20.
- 599 70. Clarke E, Thompson K, Weaver S, Thompson J, O’Connell G. Snus: A compelling harm
600 reduction alternative to cigarettes. *Harm Reduct. J. BioMed Central Ltd.*; 2019;16:62.
- 601 71. Mustafa MB, Hassan MO, Alhussein A, Mamoun E, El Sheikh M, Suleiman AM. Oral
602 leukoplakia in the Sudan: clinicopathological features and risk factors. *Int Dent J. Wiley-*
603 *Blackwell Publishing Ltd*; 2019;69(6):428–35.
- 604 72. Ahmed HG. Aetiology of oral cancer in the Sudan. *J oral Maxillofac Res. Stilus Optimus*;
605 2013;4(2):e3.
- 606 73. Faggons CE, Mabedi C, Shores CG, Gopal S. Review: Head and neck squamous cell
607 carcinoma in sub-Saharan Africa. *Malawi Med J. Malawi Medical Journal*; 2015;27(3):79–87.
- 608 74. Otoh EC, Johnson NW, Olasoji HO, Danfillo IS, Adeleke OA. Intra-oral carcinomas in
609 Maiduguri, north-eastern Nigeria. *Oral Dis. 2005*;11(6):379–85.
- 610 75. Onoh I, Owopetu O, Olorukooba AA, Umeokonkwo CD, Dahiru T, Balogun MS. Prevalence,
611 patterns and correlates of smokeless tobacco use in Nigerian adults: An analysis of the Global
612 Adult Tobacco Survey. Glantz SA, editor. *PLoS One. Public Library of Science (PLoS)*;
613 2021;16(1):e0245114.

- 614 76. Subapriya R, Thangavelu A, Mathavan B, Ramachandran CR, Nagini S. Assessment of risk
615 factors for oral squamous cell carcinoma in Chidambaram, southern India: A case-control
616 study. *Eur J Cancer Prev. Eur J Cancer Prev*; 2007;16(3):251–6.
- 617 77. Maria E. Leon, Alessandra Lugo, Paolo Boffetta, Anna Gilmore, Hana Ross, Joachim Schüz,
618 et al., Smokeless tobacco use in Sweden and other 17 European countries, *Eur J Pub Health*;
619 2016;26(5)817–21.
- 620 78. Gupta B, Bray F, Kumar N, Johnson NW. Associations between oral hygiene habits, diet,
621 tobacco and alcohol and risk of oral cancer: A case–control study from India. *Cancer*
622 *Epidemiol. Elsevier Ltd*; 2017;51:7–14.
- 623 79. Kakde S, Bhopal RS, Jones CM. A systematic review on the social context of smokeless
624 tobacco use in the South Asian population: Implications for public health. *Public Health.*
625 *Public Health*; 2012;126(8):635–45.
- 626 80. Tata Institute of Social Sciences (TISS), Mumbai and Ministry of Health and Family Welfare,
627 Government of India. Global Adult Tobacco Survey GATS 2 India 2016-17.T | Report.
628 2018;161-165 SBN : 978-81-937917-0-7
- 629 81. IARC Working Group on the Evaluation of Carcinogenic Risks to Humans. Smokeless
630 tobacco and some tobacco-specific N-nitrosamines. *IARC Monogr Eval Carcinog Risks Hum.*
631 2007;89:1-592. PMID: 18335640; PMCID: PMC4781254.82.
- 632 82. Ahmed HG, Mahgoob RM. Impact of Toombak dipping in the etiology of oral cancer: gender-
633 exclusive hazard in the Sudan. *J Cancer Res Ther. Medknow Publications and Media Pvt. Ltd*;
634 2007;3(2):127–30.
- 635 83. Stepanov I, Gupta PC, Dhumal G, Yershova K, Toscano W, Hatsukami D, et al. High levels of
636 tobacco-specific N-nitrosamines and nicotine in Chaini Khaini, a product marketed as snus.
637 *Tob Control. NIH Public Access*; 2015;24(e4):e271–4.
- 638 84. Stanfill SB, Connolly GN, Zhang L, Jia LT, Henningfield JE, Richter P, et al. Global
639 surveillance of oral tobacco products: total nicotine, unionised nicotine and tobacco-specific
640 N-nitrosamines. *Tob Control.* 2011;20(3):e2.
- 641 85. Mehrotra R, Sinha DN, Szilagyi T. WHO FCTC Global Knowledge Hub on Smokeless
642 Tobacco. 2017. 29-40.
- 643 86. Allard WF, DeVol EB, Te OB. Smokeless tobacco (shamma) and oral cancer in Saudi Arabia.
644 *Community Dent Oral Epidemiol. Blackwell Munksgaard*; 1999;27(6):398–405.
- 645 87. Hassanin AA, Idris AM. Attribution of oral cancer in the Sudan to Toombak dipping. *Transl*
646 *Res Oral Oncol. SAGE Publications*; 2017;2:1-5.
- 647 88. Idris AM, Nair J, Friesen M, Ohshima H, Brouet I, Faustman EM, et al. Carcinogenic tobacc-

- specific nitrosamines are present at unusually high levels in the saliva of oral snuff users in sudan. *Carcinogenesis*. Oxford University Press; 1992;13(6):1001–5.
89. Sami A, Stanton C, Ross P, Ryan T, Elimairi I. Ultra- structure of Toombak; smokeless tobacco of Sudan and its effects on oral and systemic health. *Access Microbiol. Microbiology Society*; 2020;2(7A):836.
 90. Richter P, Hodge K, Stanfill S, Zhang L, Watson C. Surveillance of moist snuff: total nicotine, moisture, pH, un-ionized nicotine, and tobacco-specific nitrosamines. *Nicotine Tob Res*. 2008;10(11):1645–52.
 91. Stepanov I, Biener L, Yershova K, Nyman AL, Bliss R, Parascandola M, et al. Monitoring Tobacco-Specific N-Nitrosamines and Nicotine in Novel Smokeless Tobacco Products: Findings From Round II of the New Product Watch. *Nicotine Tob Res*. 2014;16(8):1070–8.
 92. Araghi M, Galanti MR, Lundberg M, Liu Z, Ye W, Lager A, et al. No association between moist oral snuff (snus) use and oral cancer: pooled analysis of nine prospective observational studies. *Scand J Public Health*. Sweden: SAGE Publications Ltd; 2021;49(8):833–40.
 93. Stepanov I, Gupta PC, Parascandola M, Yershova K, Jain V, Dhumal G, et al. Constituent Variations in Smokeless Tobacco Purchased in Mumbai, India. *Tob Regul Sci*. 2017;3(3):305–14.
 94. Bennett JE, Fowler EA. Federal Register, FDA Proposed Rules, Tobacco Product Standard for N-Nitrosonornicotine Level in Finished Smokeless Tobacco Products. 2017;82(13):8004-8030.
 95. Appleton S, Olegario RM, Lipowicz PJ. TSNA levels in machine-generated mainstream cigarette smoke: 35 years of data. *Regul Toxicol Pharmacol*. Academic Press; 2013;66(2):197–207.
 96. Berman ML, Hatsukami DK. Reducing tobacco-related harm: FDA’s proposed product standard for smokeless tobacco. *Tob Control*. BMJ Publishing Group Ltd; 2018;27(3):352–4.
 97. de Groot PM, Wu CC, Carter BW, Munden RF. The epidemiology of lung cancer. *Transl Lung Cancer Res*. AME Publications; 2018;7(3):220-33.
 98. Gupta AK, Mehrotra R. Increasing use of flavoured tobacco products amongst youth. *Indian J Tuberc*. Elsevier; 2021.68:S105-S107.

S no	SLT type	Preparation and Use	Region	Countries with major consumption	Nicotine* mg/g	Total TSNA* µg/g	NNN*µg/g	NNK* µg/g
1	Snuff	Finely cut or ground air-cured flavoured tobacco dry or moist, placed in the mouth and sucked.	America	USA, Canada, Mexico,	3.9–40.1	0.3–76.5	0.37-42.6	0.38-9.9
2	Snus (Swedish)	Pasteurized finely ground moist tobacco, moisturizers, sodium carbonate, salt, sweeteners	Europe	Sweden, Denmark, Finland, Iceland, Norway,	7.8–15.2	0.6–0.7	0.42-3.28	0.13-1.1
3	Nass (Naswar)	Sun-dried and powdered tobacco; ash, oil, placed in the mouth and sucked	Parts of Europe and Eastern Mediterranean	Uzbekistan, Kyrgyzstan, Tajikistan, Afghanistan, Pakistan, Iran	8.9–14.2	0.5–1.4	0.59-1.3	0.07-0.21
4	Toombak	Fermented and grounded Tobacco, baking soda and water. Oral and nasal use	Parts of Eastern Mediterranean and Africa	Sudan, Chad	9.6–28.2	295–992	115-3085	147-7870
5	Dry Snuff	Finely ground powder, inhaled	Africa	Nigeria, Ghana, Algeria, Cameroon, Chad, South Africa	1.2–17.2	1.7–20.5	2.4-18.1	0.58-6.4
7	Gutkha (Chewing tobacco)	Commercial preparation, finely chopped tobacco with flavourings and sweeteners, Sucked and chewed	SEAR	India, Pakistan, Bangladesh, Nepal, Myanmar, Sri Lanka, UK	0.2–4.2	0.1–23.9	0.1-1.1	0.04-0.43
8	Khaini (Chewing tobacco)	Coarsely cut tobacco leaves mixed with slaked lime, Sun-dried or fermented.	South East Asia, Western Pacific and Eastern Mediterranean Europe	India, Bangladesh, Nepal, Bhutan	2.5–4.8	21.6–23.9	13.2-76.9	0.11-28.4
9	Zarda (Chewing Tobacco)	Shredded tobacco leaves are boiled with lime and saffron; often used with betel quid	SEAR	Bangladesh, India, Pakistan, Myanmar, Thailand, Indonesia, Nepal, Maldives, Sri Lanka, UK	9.5–30.4	5.5–53.7	4.79-19.9	0.22-24.1
10	Betel quid with tobacco	Mixture of betel quid with areca nut, with or without tobacco. May also be mixed with slaked lime and tobacco. be mixed with slaked lime, or sweeteners	SEAR	India, Pakistan, Bangladesh, Nepal, Myanmar	6.7-8.4	0.17-2.1	1.2-48.6	0-14.3
11	Shammah (Chewing tobacco)	Powdered tobacco used with slaked lime, oil, flavouring, kept in the mouth and sucked	Middle East	Saudi Arabia, Yemen, Algeria.	37.82-87.56	DNA**	DNA**	DNA**

682 Note: List of products is not exhaustive. *Figures are adapted from refs (26, 37, 52, 93 and 99); **DNA: Data not available.

Figure Legends

Figure 1: Search strategy flow-chart of study selection process for smokeless tobacco use and oral cancer risk using PRISMA guidelines.

Figure 2: Forest plot of studies showing oral cancer risks associated with various types of SLT product. Data presented also include: the SLT type, the study reference, region, the odds ratio and corresponding 95% confidence interval, in addition, where available the number of estimates (No. Est) that the pooled estimate is based on are provided.

Figure 3: Forest plot of studies showing WHO region-wise oral cancer risks associated with various SLT products. Data presented also include: the SLT type, the study reference, the odds ratio and corresponding 95% confidence interval, in addition, where available the number of estimates (No. Est) that the pooled estimate is based on are provided.

Figure 4. Tobacco specific nitrosamines (TSNAs) levels and odds ratio for oral cancer in diverse SLT products. (BQ+ denotes betel quid with tobacco) (a) TSNA values are presented on the log scale; the original TSNAs levels in $\mu\text{g/g}$ are presented on the right hand side y-axis. (b) OR and corresponding 95% CIs estimates are based on review studies from the same region that the SLT product TSNAs values are based. The OR estimates for zarda and khaini are not product specific but those for all types of chewing tobacco from SEAR (ref 54). For gutkha, dry snuff and snus, the OR estimates are based on global pooled estimates (ref 51), whereas for naswar these are based on EMR estimates only (ref 57).

Identification

Screening

Eligibility

Included

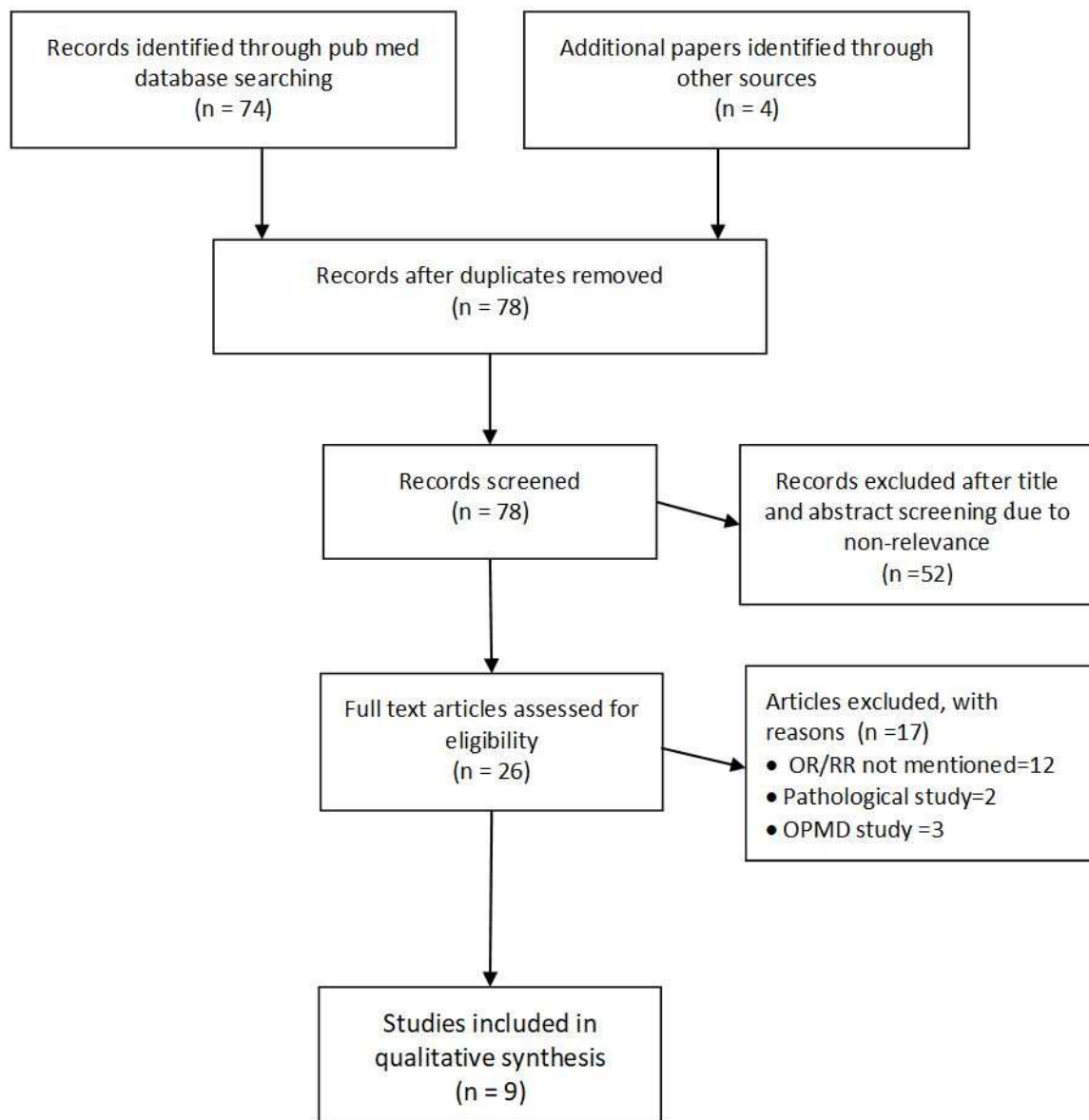


Figure 1

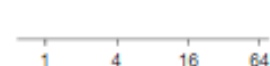
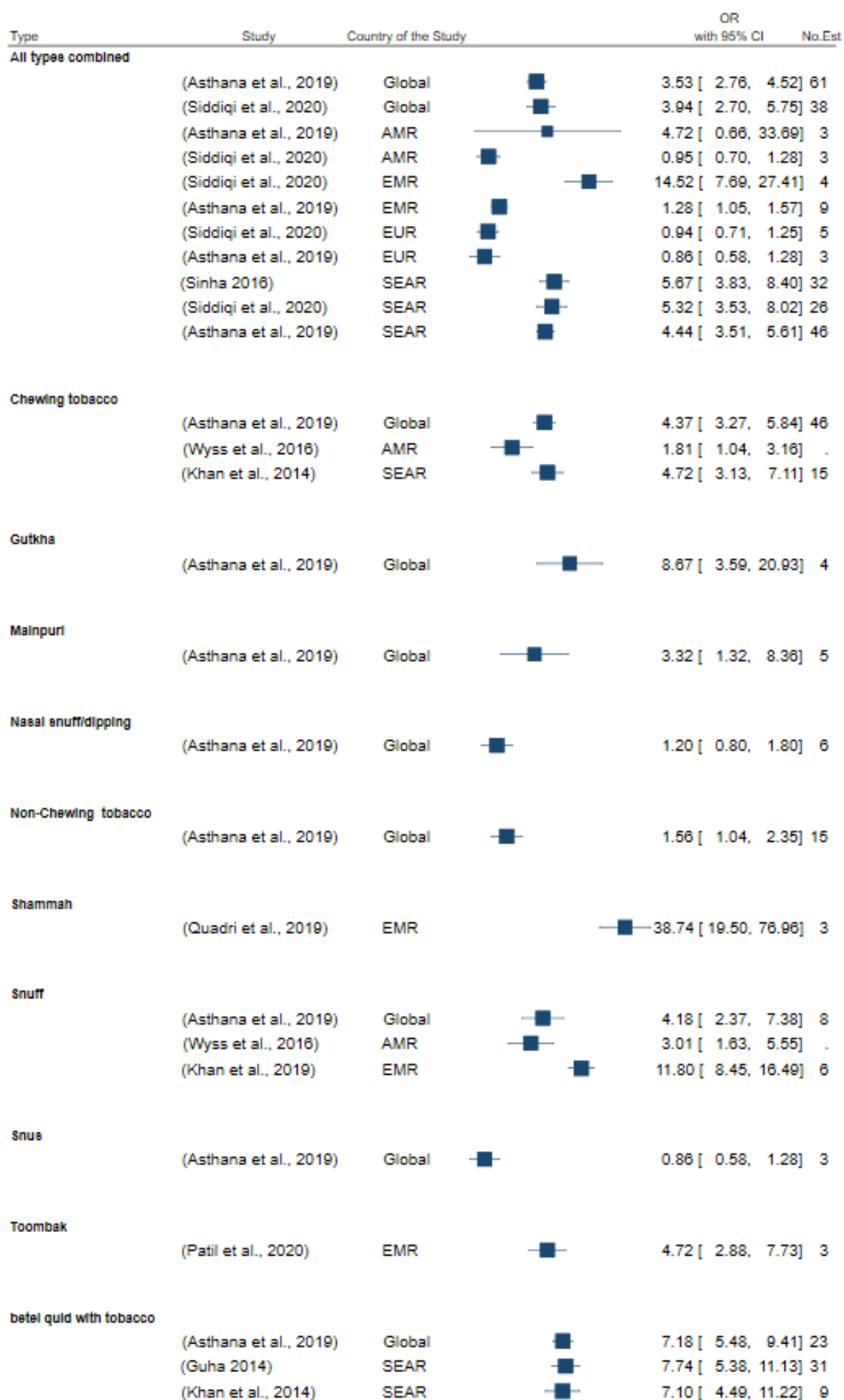
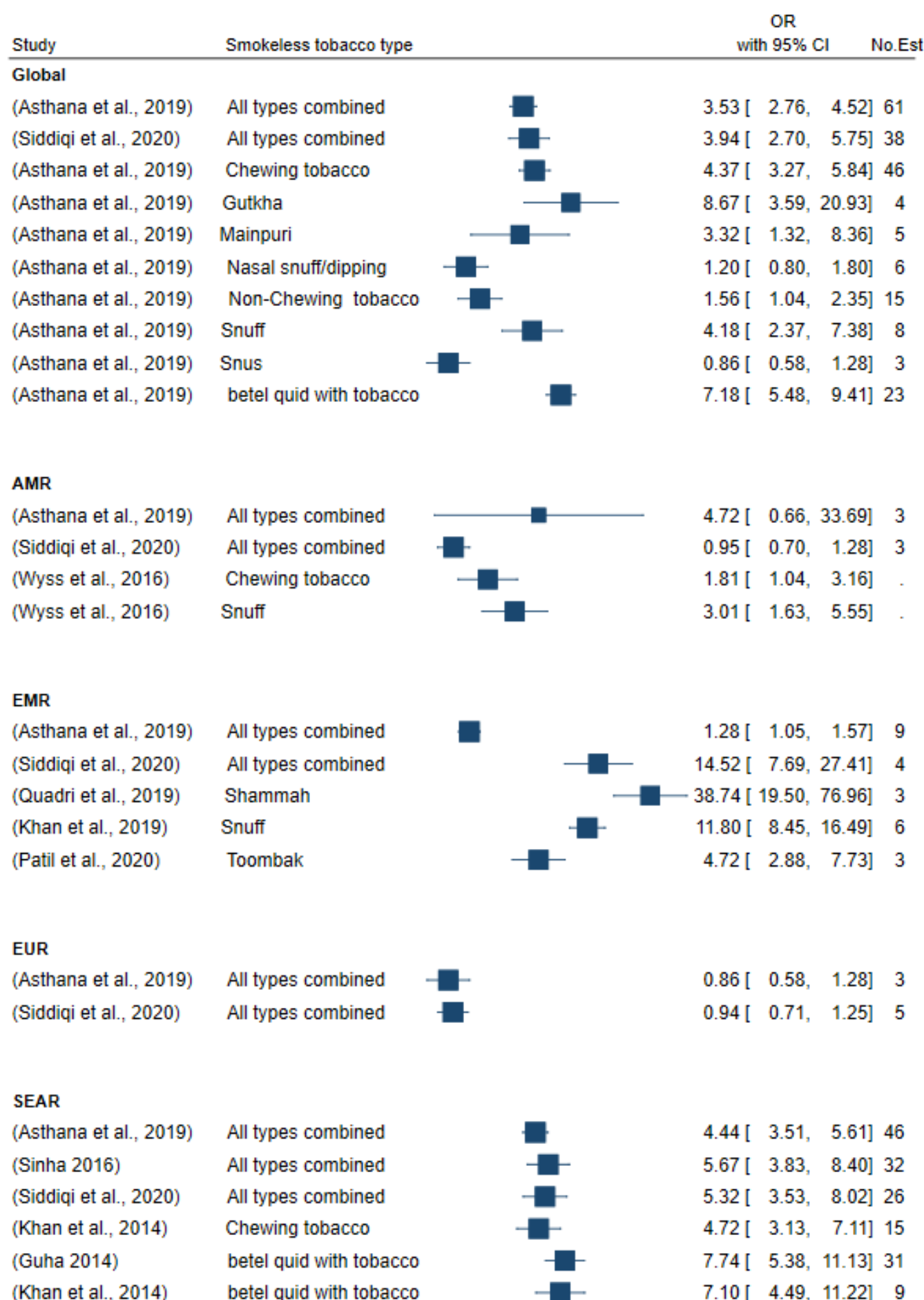


Figure 2



1 4 16 64

Figure 3

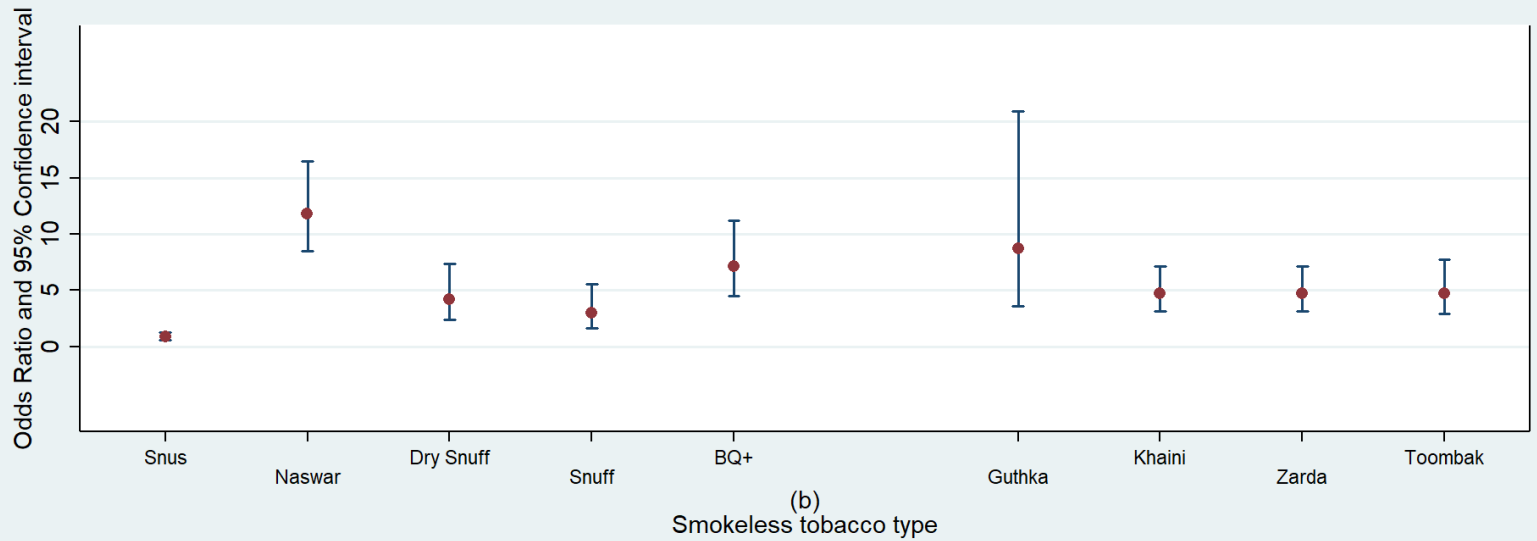
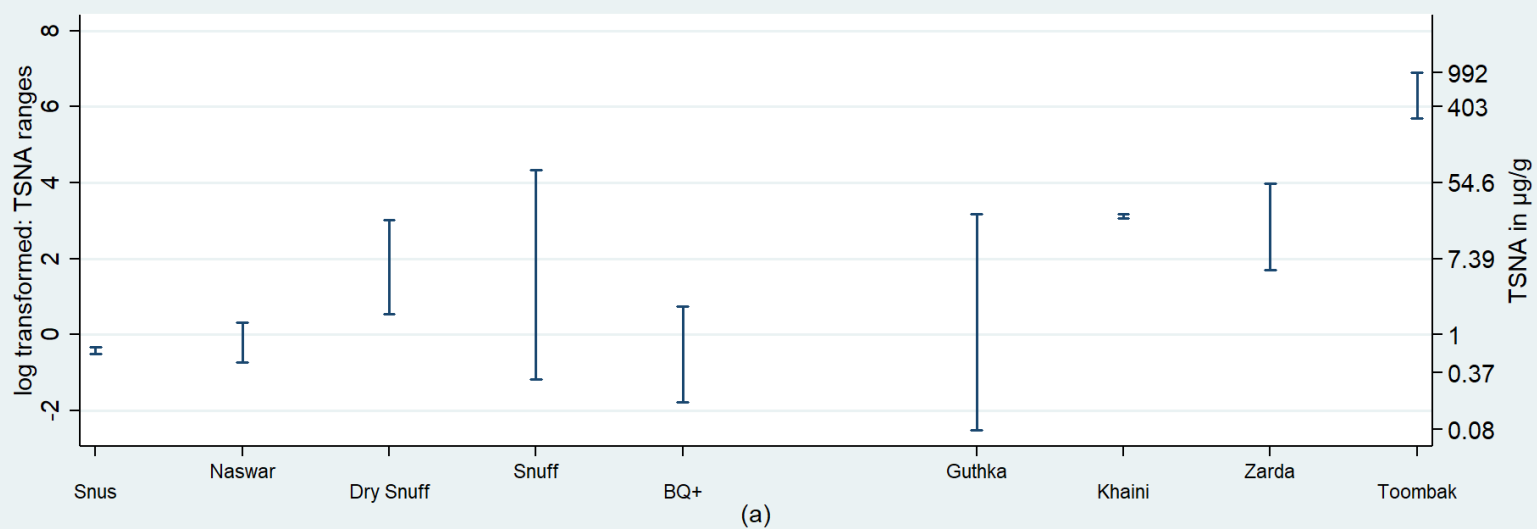


Figure 4