



Culture-independent studies on bacterial dysbiosis in oral and oropharyngeal squamous cell carcinoma: A systematic review

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ABSTRACT

Imbalance within the resident bacterial community (dysbiosis), rather than the presence and activity of a single organism, has been proposed to be associated with, and to influence, the development and progression of various diseases; however, the existence and significance of dysbiosis in oral/oropharyngeal cancer is yet to be clearly established. A systematic search (conducted on 25/01/2018 and updated on 25/05/2018) was performed on three databases (Pubmed, Web of Science & Scopus) to identify studies employing culture-independent methods which investigated the bacterial community in oral/oropharyngeal cancer patients compared to control subjects. Of the 1546 texts screened, only fifteen publications met the pre-determined selection criteria. Data extracted from 731 cases and 809 controls overall, could not identify consistent enrichment of any particular taxon in oral/oropharyngeal cancers, although common taxa could be identified between studies. Six studies reported the enrichment of *Fusobacteria* in cancer at different taxonomic levels whereas four studies reported an increase in *Parvimonas*. Changes in microbial diversity remained inconclusive, with four studies showing a higher diversity in controls, three studies showing a higher diversity in tumors and three additional studies showing no difference between tumors and controls. Even though most studies identified a component of dysbiosis in oral/oropharyngeal cancer, methodological and analytical variations prevented a standardized summary, which highlights the necessity for studies of superior quality and magnitude employing standardized methodology and reporting. Indeed an holistic metagenomic approach is likely to be more meaningful, as is understanding of the overall metabolome, rather than a mere enumeration of the organisms present.

1. Introduction

Head and neck squamous cell carcinomas (HNSCC) are amongst the most common cancers in the world. Oral and Oropharyngeal cancers together constitute the major group of HNSCC and 51,540 new cases are expected to be diagnosed and 10,030 individuals are estimated to die in 2018 due to these cancers in US alone (Siegel et al., 2018). Numerous causative agents or factors have been described, notably tobacco-both smoked and smokeless, areca nut, alcohol, human papillomavirus (HPV), tooth-related factors/chronic mechanical irritation, dietary deficiencies and fungal infections, notably chronic candidiasis (Pollaers et al., 2017; Johnson, 2017). However, the exact role of each

factor remains uncertain despite the significant number of studies showing their association. HNSCC, like any other carcinoma, is a progressive chronic disease in which the healthy epithelium accumulates dysplastic changes and in some, but not all cases, finally transforms into invasive phenotypes. The events of carcinogenesis are complex and multifactorial and might not follow the same sequence in different individuals (Johnson, 2017).

An increasing body of evidence has been accumulating in recent years which lead experts to suggest a possible role of microorganisms at different stages of carcinogenesis (Althani et al., 2016; Meurman and Uittamo, 2008; Vogtmann et al., 2016). The connections between a consortium of bacteria and carcinogenesis have not been investigated

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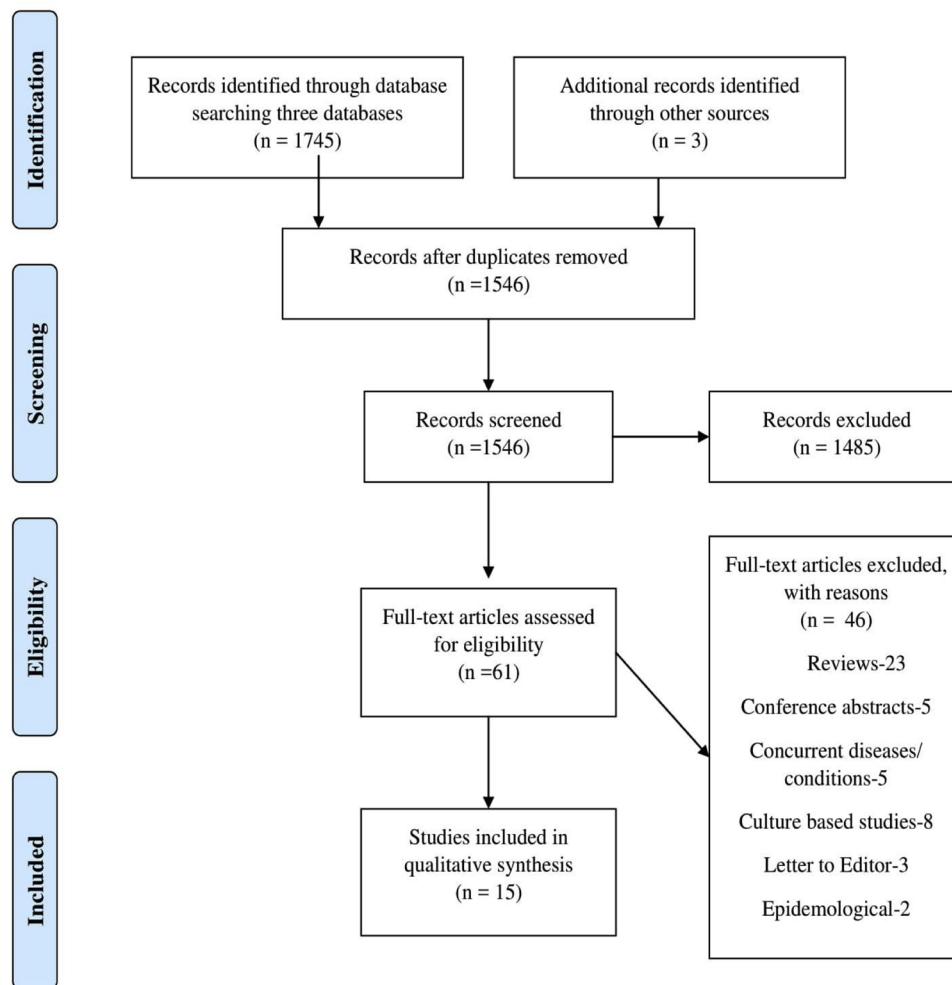


Fig. 1. PRISMA flow diagram for systematic review depicting phases of identification of studies.

thoroughly; perhaps because earlier research had identified individual pathogens such as *Helicobacter pylori*, linked to carcinogenesis (Wroblewski et al., 2010). Culture-dependent methods have examined the role of numerous individual or groups of bacteria and their association with cancers of oral cavity (Perera et al., 2016). However, it is now considered that 60–90% of the human microbiome is uncultivable which has prompted further research regarding the role of bacteria in carcinogenesis (D'Argenio and Salvatore, 2015; Dewhirst et al., 2010). Culture-independent techniques using molecular methods with recombinant DNA have endowed researchers with tools for recognizing the organisms that reside in microbial communities, reducing the potentially misleading results arising from the vagaries of cultivation. With the advent of next-generation sequencing, studies have focused on a more holistic approach for the characterization and identification of microbial communities and their possible involvement in various diseases, including different types of cancer (Khan et al., 2017; Zakariah et al., 2018; Alshamsana et al., 2019).

Microbial dysbiosis can be defined as any change to the composition of resident commensal communities, relative to the community found in healthy individuals (Petersen and Round, 2014). To date, very few studies have tried to address bacterial dysbiosis associated with oral/oropharyngeal cancers employing culture-independent approaches. Furthermore, clear evidence is still deficient on any association between this dysbiosis and oral/oropharyngeal cancers and whether the presence of specific bacterial taxa could be developed as biomarkers for diagnosis or risk assessment. Our aim in present article is to summarize current evidence on the association of bacterial dysbiosis and oral

cancer by performing a systematic and comprehensive review of published studies which have utilized culture-independent techniques to study the characteristics of the whole oral microbial community in association with oral/oropharyngeal cancers. Whilst recognizing that a microbiome constitutes fungi, bacteria and viruses, and occasionally other organisms, the present review is restricted to the 'bacteriome'. It is also important to differentiate the terms *microbiota*, which comprises all living microorganisms, and the *microbiome/bacteriome*, which includes all available genomes derived from dead or living microorganisms in its pure context.

2. Methods

Literature search and data extraction methods for this review were specified and documented as a protocol and registered in PROSPERO (CRD42017051160). This systematic review follows the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) criteria (Moher et al., 2009). The strategies for the screening and inclusion were decided by authors 1 and 2 in advance. A systematic search of the published literature was conducted on 25/01/2018 and updated on 25/05/2018 in PubMed, Scopus and Web of science to identify articles on oral cavity microbiome and oral cancer-without date limitations using specific search terms. The search strategy included the following words: oral, oral cavity, saliva, oral tissue, microbiome, microbiota, microbial diversity, micro flora, microorganisms, oral cancer, oral squamous cell carcinoma and head & neck squamous cell carcinoma in combination. Boolean operators such as "and" "or" were used to

stratify the search. After an initial screening of the titles and abstracts, two authors (DG and RKM) independently screened the full texts for eligibility. Moreover, manual searches of cross-references from relevant narrative reviews were also performed in order to exclude the possibility of omitting any important study. The search strategy was confined to English language comprising human studies.

Two authors independently assessed the risk of bias using an adapted Newcastle-Ottawa Scale for analytical non-randomized case-control studies similar to a recently published review on skin microbiome in atopic dermatitis (Bjerre et al., 2017).

2.1. Inclusion criteria

Original research articles investigating the ‘bacteriome’ community through culture-independent methods in tissue, saliva or swabs in patients with oral and/or oropharyngeal cancers and controls were included.

2.2. Exclusion criteria

Culture-based studies, conference abstracts, review articles, studies examining a cohort with oral or oropharyngeal cancer associated with other debilitating diseases and those studies without controls were excluded. Articles in languages other than English were excluded.

3. Results

1546 articles were identified in the initial search after removal of duplicates. After reviewing the title and the abstract, 61 articles were included for full-text screening. Based on our selection criteria, 15 studies were selected and examined in detail (Fig. 1).

3.1. Sample characteristics & methodology

Sample characteristics are described in detail in Table 1. There were a total of 731 cases and 809 controls represented in a total of fifteen publications. Various kinds of oral samples were used to study the ‘bacteriome’ including saliva (Guerrero-Preston et al., 2016; Pushalkar et al., 2011; Lee et al., 2017; Wolf et al., 2017), swabs from the surface of the tumor (Schmidt et al., 2014a; Zhao et al., 2017), the tumor tissue (Bebek et al., 2012; Hooper et al., 2007; Pushalkar et al., 2012; Wang et al., 2017; Al-Hebshi et al., 2017; Banerjee et al., 2017; Perera et al., 2018; Mukherjee et al., 2017) and oral rinses (Bornigen et al., 2017). Samples from normal healthy individuals as well as patient’s own mucosa were utilized as controls in the included studies. Those studies which used samples from normal healthy individuals as controls had varied matching criteria which included site (Pushalkar et al., 2011; Al-Hebshi et al., 2017; Perera et al., 2018), age (Lee et al., 2017; Al-Hebshi et al., 2017) and gender (Al-Hebshi et al., 2017; Perera et al., 2018). Other studies used tissues from contra lateral mucosa of the same patient (Schmidt et al., 2014a; Zhao et al., 2017; Banerjee et al., 2017) or mucosa adjacent to cancer tissue from the patient (Bebek et al., 2012; Hooper et al., 2007; Pushalkar et al., 2012; Wang et al., 2017; Mukherjee et al., 2017) as controls. The total sample size of the selected studies ranged from 5 to 363. However, none of the studies stated the rationale for their study population size: nonetheless, one of the studies claimed that their multivariate analysis remained well powered as the number of matched controls was twice that of cases (Bornigen et al., 2017).

We found significant heterogeneity in the methodology of studies (Table 2). Numerous studies have used archival frozen specimens collected for other varied purposes and thus strict exclusion criteria were not properly defined. Accepted protocols of microbial wall lysis such as lysozyme treatment or bead beating were not reported in the selected studies except for three (Wang et al., 2017; Mukherjee et al., 2017; Bornigen et al., 2017). DNA extractions were performed using different

commercial DNA kits. The studies using 16S rRNA amplification approach targeted different hypervariable regions; however, significant overlap was observed in constant regions of the 16S gene. One of the earlier studies used three sets of primers, which covered bacteria of three main phyla found in *Spirochaetes*, *Bacteroides* & *domain Bacteria* (Zhao et al., 2017). Amount of PCR cycles ranged from 25 to 35, where as one study did not provide any information regarding the PCR cycles (Guerrero-Preston et al., 2016). Four studies utilized Topo TA cloning and Sanger sequencing for generating sequences with the number of colonies analyzed ranging from 30 to 95 per sample, whereas rest of the studies utilized 16S rRNA sequencing that allows deeper sequencing depths. One study utilized PathoChip™ array (a type of microarray) coupled with next generation sequencing for identification of the putative pathogens (Banerjee et al., 2017).

3.2. Risk of bias

The quality of the ‘bacteriome’ studies showed high variability (Table 3). None could be rated as very good. We qualified them as very good (total score = 9; N = 0), good (total score = 7–8; N = 8), fair (total score = 5–6; N = 5) and poor (< 5; N = 2). We could not identify any justification for sample size in the included studies. The main reason for downgrading the studies to fair was either lack of proper statistical analyses and lack of definable outcome measures.

3.3. Microbial diversity & relative abundance

Diversity and richness are two commonly used indices to report the diversity of the microbiome in human and environmental studies. Diversity can be defined as the number of phylotypes or operational taxonomic units found in a local habitat (Bornigen et al., 2017). Interestingly, among the fifteen studies, only nine reported the change in bacterial diversity of the microbiome when compared to controls. Three studies did not observe any differences in microbial diversity in tumors compared to controls. However, higher diversity in controls over cases was reported by Preston et al. (Guerrero-Preston et al., 2016) and Lee et al. (Lee et al., 2017) in saliva and Perera et al. (2018) as well as Mukherjee et al. (2017) in tissues. Contrary to this, Schmidt et al. (2014a) and Zhao et al. (2017) reported an increase in diversity in tumor swab specimens whereas Wolf et al. (2017) reported an increase in diversity in tumor salivary samples respectively. Another study reported a relative shift from gram-negative to gram-positive organisms in oral cancer tissue samples (Pushalkar et al., 2012).

Abundance is defined as the number of times each taxon is identified in the sample and often reported in terms of relative abundance, which helps in the comparison between the groups reported (Li et al., 2012). The studies varied in the reporting of abundance of taxa with a few concentrating on hierarchical differences from phyla all the way down to genera, whereas others only reported differences in genera.

The reported range of phyla varied from five to thirteen among the studies with many of them concentrating on the most abundant seven phyla in the oral cavity. *Firmicutes* is the most common phylum in the oral cavity (Petersen and Round, 2014) and all of the studies were in concordance regarding this finding except the study by Zhao et al which reported *Bacteroidetes* to be the most abundant (Zhao et al., 2017). However, the relative abundances of phylum *Firmicutes* varied between cases and controls among the studies. Three studies reported a reduction in *Firmicutes* in tumor tissue compared to controls (Wolf et al., 2017; Schmidt et al., 2014a; Zhao et al., 2017) whereas three other studies identified an increase in phylum *Firmicutes* in the saliva (Guerrero-Preston et al., 2016; Pushalkar et al., 2011) and tissue (Mukherjee et al., 2017) of cancer patients respectively when compared to controls. Reduction in abundances of *Actinobacteria* in the tumor samples was reported by three studies (Schmidt et al., 2014a; Zhao et al., 2017; Wang et al., 2017) and one among these studies observed a positive correlation between the reduction in the abundance of phyla

Table 1
Summary of sample characteristics and 'bacteriome'.

No	Author year	Sample	Age (Mean) Sex	Site	Study setting	Nature of control	Study population size		Other clinical factors studied	RESULTS	Diversity and Richness	Abundance of taxa ^a
							Cases	Controls				
1	Hooper et al., 2007	Fresh Tissue	65.3 Both	Oral	UK	Normal mucosa from same subject	10	10	NA	NA		<ul style="list-style-type: none"> • <i>Clavibacter michiganensis</i> subsp. <i>tellurii</i>, <i>Fusobacterium naviforme</i>, <i>Ralstonia insidiosa</i> detected in > 30% more of tumor • <i>Granulicatella adiacens</i>, <i>Porphyromonas gingivalis</i>, <i>Sphingomonas</i> sp. PC5.28 and <i>Streptococcus mitis/oralis</i> detected in > 30% more of non tumor tissue • Most prevalent in OSCC: <i>Streptococcus</i>, <i>Gemella</i>, <i>Rothia</i>, <i>Peptostreptococcus</i>, <i>Lactobacillus</i>, <i>Porphyromonas</i> and <i>Lactobacillus</i> (genus) • Most prevalent in controls: <i>Prevotella</i>, <i>Neisseria</i>, <i>Lepiostichia</i>, <i>Capnocytophaga</i>, <i>Actinobacillus</i>, <i>Oribacterium</i>, SR1 and TM7 (genus) • <i>Enterobacteriaceae</i> (family) & <i>Terrivibrio</i> (phylum) found to be significantly enriched in tumor tissue
2	Pushalkar et al., 2011	Saliva	> 50 M	Oral	USA	Saliva of controls	3	2	NA	NA		
3	Bebek et al., 2012	Fresh frozen and paraffin embedded tissue	62 M/F	HN-SCC	USA	Adjacent normal tissue	42	42	Age, Sex, Site, Metastases, HPV, Tobacco, Alcohol, Node, Methylation	NA		
4	Pushalkar et al., 2012	Tissue	59 M/F	Oral	USA	Adjacent normal tissue	10	10	NA	Richness higher in tumor, No difference in diversity		<ul style="list-style-type: none"> • Taxa enriched in tumor - <i>Streptococcus</i> sp. oral taxon 058, <i>Peptostreptococcus stomatis</i>, <i>Streptococcus salivarius</i>, <i>Streptococcus gordonii</i>, <i>Gemella haemolysans</i>, <i>Gemella morbillorum</i>, <i>Johnsonella ignava</i> and <i>Streptococcus parasanguinis</i> • Taxa enriched in non tumor - <i>Granulicatella adiacens</i> • Tumor-Significant reduction in <i>Firmicutes</i> and <i>Actinobacteria</i> (phyla) • Increase in <i>Fusobacteria</i> • Tumor-Significant increase in <i>Streptococcus</i> and <i>Lactobacillus</i> (genus) • Non tumor-Significant increase in <i>Haemophilus</i>, <i>Neisseria</i>, <i>Lepiostichia</i>, <i>Aggregatibacter</i>, <i>Lautropia</i> (genus) • Tumor-Significant reduction in <i>Actinobacteria</i>, Significant increase in <i>Parvimonas</i> (genus) up to parent family <i>Tissierellaceae</i> • Increase in <i>Fusobacteria</i> • Tumor-<i>F. nucleatum</i> subsp. <i>polymorphum</i>, <i>Pseudomonas aeruginosa</i> and <i>Campylobacter</i> sp. Oral taxon 44 (significantly abundant) tumors • Controls-S. <i>mitis</i>, <i>R. mucilaginosa</i> and <i>H. parainfluenzae</i> • Highly abundant in SSC patients - Phylum <i>Actinomyces</i> (<i>Actinobacteria</i>), <i>Schwartzia</i> (<i>Firmicutes</i>), <i>Treponema</i> (<i>Spirochaetes</i>) and <i>Selenomonas</i> (<i>Firmicutes</i>). • Highly abundant in Controls-Phylum <i>Bacteroidetes</i> (<i>Prevotella</i>), <i>Proteobacteria</i> (e.g. <i>Haemophilus</i> and <i>Neisseria</i>) and <i>Firmicutes</i> (e.g. <i>Streptococcus</i> and <i>Veillonella</i>)
5	Schmidt et al., 2014a	Swabs	62.4 M/F	Oral	USA	Contra lateral normal tissue	15	15	Smoking, Nodal status	Diversity higher in tumor samples		
6	Guerrero-Preston et al., 2016	Saliva	NS	Oral/Oropharyngeal	USA	Saliva of healthy controls	17	25	HPV status, Tumor site	Significantly higher diversity and richness in controls		
7	Wang et al., 2017	Archival Frozen Tissue	63 M/F	HN-SCC	USA	Adjacent normal tissue	121	121	Clinical stage of cancer	No difference in diversity and richness		
8	Al-hebshi et al., 2017	Archival Frozen Tissue	56.9 M/F	Oral	Yemen	Deep epithelial swabs from gender and age matched healthy controls	20	20	Nil	No difference in diversity and richness		
9	Wolf et al., 2017	Saliva	61.6 M/F	Oral/Oropharyngeal	Australia	Saliva from healthy controls	11	11	Age, Alcohol Consumption, Tumor size, Lymph node status, Smoking, HPV status	Higher diversity in saliva of cancer patients		

(continued on next page)

Table 1 (continued)

No	Author year	Sample	Age (Mean)	Site	Study setting	Nature of control	Study population size		Other clinical factors studied	RESULTS	Abundance of taxa*
							Cases	Controls			
10	Lee et al., 2017	Saliva	53 M/F	Oral	Taiwan	Saliva from healthy controls	127	125	Smoking, Chewing	Higher diversity in saliva of normal controls	<ul style="list-style-type: none"> • <i>Bacteroides</i>, <i>Escherichia</i> and <i>Bifidobacteria</i> were present in over of 40% in cancer patients • <i>Cloacibacillus</i>, <i>Gemmiger</i>, <i>Oscillospora</i> and <i>Roseburia</i> were 20 times abundant in cancer patients • Genus <i>Dialister</i> were relatively highly abundant in cases when compared to controls. • Orders <i>Actinomycetales</i> and <i>Lactobacillales</i> were significantly underrepresented in oral cancer. • Spirochaetes, Fusobacteria and Bacteroidetes were significantly enriched in cases • <i>Firmicutes</i> and <i>Actinobacteria</i> were significantly decreased in cases • 17 taxa were enriched at genus level in cancers • Phylum <i>Proteobacteria</i> was more pronounced in cases • Phylum <i>Bacteroides</i> were more pronounced in healthy controls • 13 genera were specifically associated with tumor samples
11	Bornigen et al., 2017	Oral rinse	NS M/F	Oral/Oropharyngeal	USA	Oral rinse from healthy controls	121	242	Smoking, Alcohol, Tooth status, HPV status, Periodontitis	NA	
12	Zhao et al., 2017	Swabs	62 M/F	Oral	China	Normal mucosa from same subject	80	80	Nil	Higher diversity in cancer patients	<ul style="list-style-type: none"> • <i>Firmicutes</i> and <i>Actinobacteria</i> were significantly decreased in cases • 17 taxa were enriched at genus level in cancers • Phylum <i>Proteobacteria</i> was more pronounced in cases • Phylum <i>Bacteroides</i> were more pronounced in healthy controls • 13 genera were specifically associated with tumor samples
13	Banerjee et al., 2017	Formalin fixed paraffin embedded tissue	NS	Oral/Oropharyngeal	USA	Adjacent normal tissue as well as healthy control tissue	100	20 + 20	Nil	NA	
14	Mukherjee et al., 2017	Archival Frozen tissue	60.5 M/F	Oral (Tongue)	USA	Adjacent normal mucosa	39	39	Gender, Race, Age, Smoking	Lower diversity & richness in cancer tissues	<ul style="list-style-type: none"> • Genus <i>Streptococcus</i>, <i>Actinomyces</i>, <i>Rothia</i>, <i>Corynebacterium</i>, <i>Enterococcus</i> and <i>Micrococcus</i> significantly increased in tumor group • Genus <i>Campylobacter</i>, <i>Pseudomonas</i>, and <i>Atopobium</i> were associated with tumor • <i>Lautropia</i>, <i>Staphylococcus</i>, <i>Propionibacterium</i>, and <i>Sphingomonas</i> were the most significantly abundant in controls
15	Perera et al., 2018	Deep Tissue	61 M	Oral	Sri Lanka	Fibroepithelial polyp from age and gender matched controls	25	27	Nil	Lower diversity and richness in cancer tissues	

* Described in terms used by authors in the respective articles as no standardized way of reporting was observed among the articles.

** Not statistically significant #NA-Not available #NS- Not specified.

Table 2
Summary of techniques of DNA extraction, amplification & sequencing.

No	Author year	Sample	Method of DNA extraction	DNA amplification	Sequencing	Reference Data base
1	Hooper et al., 2007	Tissue	Pure gene DNA isolation kit	3 sets of primers targeting 16S rRNA genes of <i>Spirochaetes, Bacteroides</i> & domain <i>Bacteria</i>	Topo TA cloning and sequencing of 30 clones samples	Gene bank
2	Pushalkar et al., 2011	Saliva	Incubation in Proteinase K & DNA purification kit (Epicenter)	V4-V5 region	454 parallel DNA sequencing	RDP ii
3	Bebek et al., 2012	Tissue	Tissue homogenized in trisodium citrate enzymatic lysis & DNA extraction by QIAamp DNA mini kit	V1-V4 region	Topo TA cloning and sequencing of 90 clones from each sample	RDP*
4	Pushalkar et al., 2012	Tissue	Incubation in Proteinase K & DNA purification kit (Epicenter)	V4-V5 region	Topo TA cloning and sequencing of 96-96 clones from each individual	HOMD**
5	Schmidt et al., 2014a	Swabs	Incubation in Proteinase K & DNA easy kit (Qiagen)	V4 region	454 pyrosequencing	Greengenes
6	Guerrero-Preston et al., 2016	Saliva	Saliva pellets were digested with SDS and Proteinase K & DNA extraction by phenol-chloroform method	V3-V5 region	Roche/454 GS pyrosequencing	RDP
7	Wang et al., 2017	Tissue	Bead homogenization of tissue + enzymatic lysis & DNA extraction by QIAamp DNA mini kit (Qiagen)	V1-V4 region	Strataclone psc vector cloning and sequencing of 95 clones per sample	RDP
8	Al-hebshi et al., 2017	Caeser-Tissue /Controls-swab	DDK DNA isolation kit	V1-V3 region	MiSeq platform (Illumina, USA) (300 base pair chemistry)	HOMD
9	Wolf et al., 2017	Saliva	MagNA pure LC DNA Isolation Kit III	V4-V5 region	MiSeq platform (Illumina, USA)	Greengenes
10	Lee et al., 2017	Saliva	QIAamp DNA Blood Kit	V4 region	MiSeq platform (Illumina, USA)	SILVA
11	Mukherjee et al., 2017	Tissue	QIAamp DNA Stool mini Kit	V4 region	Ion Torrent Thermofisher (USA)	UNITE 5.8 s
12	Bornigen et al., 2017	Oral rinse	QIAasympfony kit	V4 region	MiSeq platform (Illumina, USA)	Greengenes
13	Zhao et al., 2017	Swabs	QIAamp DNA Mini Kit	V4-V5 region	MiSeq platform (Illumina, USA)	RDP
14	Banerjee et al., 2017	Tissue	All Prep DNA/RNA FFPE kit	NA	MiSeq platform (Illumina, USA)	RDP
15	Perera et al., 2018	Tissue	Centra Puregene Tissue kit	V1 - V3 region	MiSeq platform (Illumina, USA)	Species-level taxonomy assignment algorithm (23)

*RDP-Ribosomal Database Project** HOMD-Human Oral Microbiome Database.

Table 3
Risk of Bias Assessment by adjusted Newcastle-Ottawa Scale for Case Control studies.

No	Studies	Selection (Definition and selection of cases and controls :max = 4*)	Comparability (cases and controls :max = 2*)	Outcome (outcomes measured as relative abundances of bacteria, Same method for cases and controls, statistical analysis :max = 3*)	Total (max = 9)
1	Hooper et al., 2007	**	*	*	4
2	Pushalkar et al., 2011	**	*	*	4
3	Bebek et al., 2012	**	**	**	6
4	Pushalkar et al., 2012	***	**	*	6
5	Schmidt et al., 2014a	***	**	**	8
6	Guerrero-Preston et al., 2016	***	*	***	7
7	Wang et al., 2017	**	**	***	7
8	Al-Hebshi et al., 2017	**	*	***	6
9	Wolf et al., 2017	**	**	***	7
10	Lee et al., 2017	***	**	***	8
11	Bornigen et al., 2017	***	**	***	6
12	Zhao et al., 2017	***	**	***	8
13	Banerjee et al., 2017	***	**	***	6
14	Mukherjee et al., 2017	***	**	***	8
15	Perera et al., 2018	***	**	***	8

Actinobacteria and the severity of the disease (increase in T stage) (Wang et al., 2017). Two studies reported significant enrichment (Zhao et al., 2017; Al-Hebshi et al., 2017) where as one study reported reduction (Mukherjee et al., 2017) of *Fusobacteria* at phylum level in cancer samples compared to controls. Apart from these three changes at the phylum level, significant enrichment in phylum *Tenericutes* was reported by Bebek et al. (2012) and *Spirochaetes* and *Bacteroidetes* by (Zhao et al., 2017) in cancer samples.

Numerous changes were reported at the genus level (Table 1). A study conducted on a US population reported a significant reduction in the relative abundance of genus *Lactobacillus* and *Streptococcus* and enrichment of *Staphylococcus* and *Parvimonas* in the saliva of oral cancer patients when compared to controls (Guerrero-Preston et al., 2016). Two other reports (Pushalkar et al., 2012; Wang et al., 2017) also identified an increase in *Parvimonas* in tumors relative to controls. Tumor samples also had loss of abundance of different taxa such as *Neisseria*, *Aggregatibacter*, *Haemophilus*, and *Leptotrichia* at genus level compared to controls as reported by Preston et al. (Guerrero-Preston et al., 2016). Another study from USA reported the difference in abundance of 22 bacteria at genus level in tumor and non-tumor groups. Among them, *Streptococcus* was the most abundant genera to be significantly increased in tumor group compared to controls along with *Actinomyces*, *Corynebacterium*, *Rothia*, *Enterococcus* and *Micrococcus* (Mukherjee et al., 2017). Bornigen et al. reported higher relative abundances of genus *Dialister* and depletion of *Scardovia* in oral rinse from oral cancer patients (Bornigen et al., 2017). A study from Yemen reported strong association of genus *Fusobacterium*, *Campylobacter* and *Pseudomonas* with OSCC tissues, while *Streptococcus*, *Haemophilus* and *Rothia* were the most common genera in the controls (Al-Hebshi et al., 2017). Statistically significant differences in abundances of seven genera namely, *Lachnoanaerobaculum*, *Filifactor*, *Mycoplasma*, *Fretibacterium*, *Megasphaera*, *Tannerella* and *Peptostreptococcus* between the saliva of normal and oral cancer groups were reported in a Taiwanese population (Lee et al., 2017). However, they reported a significant presence of *Escherichia coli* in normal controls and cancer tissue: this pathogen is not a common inhabitant of the oral cavity and is often considered as a contaminant in microbiome studies. Perera et al. reported the association of *Fusobacterium*, *Capnocytophaga*, *Pseudomonas*, and *Atopobium* with OSCC tissues from Sri Lankan males and *Lautropia*, *Staphylococcus*, *Propionibacterium*, and *Sphingomonas* with controls (Perera et al., 2018). Wolf identified high abundance of genera, notably *Actinomyces* (Actinobacteria), *Schwartzia* and *Selenomonas* (Firmicutes), and *Treponema* (Spirochaetes) in SSC patients whereas *Prevotella* (Bacteroidetes), *Streptococcus* and *Veillonella* (Firmicutes) & *Haemophilus* and *Neisseria* (Proteobacteria) were more in controls in an European population (Wolf et al., 2017). Significantly higher abundances of *Campylobacter*, *Eikenella*, *Mycoplasma*, *Fusobacterium*, *Selenomonas*, *Treponema*, *Dialister*, *Centipeda*, *Lachnospiraceae_G_7*, *Alloprevotella*, *Peptostreptococcus*, *Filifactor*, *Peptococcus*, *Catonella*, *Parvimonas*, and *Peptostreptococcaceae_XI_G_7*, & *Capnocytophaga* in cancer samples were reported in a Chinese population (Zhao et al., 2017). On the contrary, *Stomatobaculum*, *Lautropia*, *Veillonella*, *Megasphaera*, *Streptococcus*, *Scardovia*, *Rothia*, *Granulicatella*, and *Actinomyces* were more abundant in controls.

Apart from phylum and genus levels, two studies reported an increased abundance of families *Enterobacteriaceae* (Bebek et al., 2012) and *Tissierellaceae* (Wang et al., 2017) in tumors compared to adjacent normal control tissue whereas another study reported a decrease in orders *Actinomycetales* and *Lactobacillales* in oral rinses of cancer patients relative to controls (Bornigen et al., 2017).

3.4. Association with clinical variables

Associations of microbiome with other variables commonly associated with HNSCC were also investigated by some of these studies (Table 1). Bornigen et al. observed a loss of bacterial diversity in oral

rinse of smokers when compared to non smokers though not statistically significant (Bornigen et al., 2017). However Schmidt et al could not find any overall changes in the microbiome that might be attributed to the use of tobacco owing to their limited sample size (Schmidt et al., 2014a). Bebek et al reported that microbiome subpopulations differentiated tobacco users from non-tobacco users with tobacco users showing a significant decrease in *Flavobacteria* (class) and *Tenericutes* (phylum) compared to the other group (Bebek et al., 2012). However, they could not find any association between specific bacterial subpopulations with alcohol use, nor with HPV status of the neoplasm. Wolf et al. observed that patients who were HPV + exhibited a ‘bacteriome’ composition that resembled healthy controls (Wolf et al., 2017) whereas Guerrero-Preston et al. reported a significant abundance of the genus *Gemellaceae* and *Leuconostoc* in HPV + HNSCC when compared to HPV–HNSCC patients (Guerrero-Preston et al., 2016).

4. Discussion

Characterization of the microbiome as an entire community has broadened our views of microbial ecology, including community-level functions that integrate with the normal functioning of the human ecosystem. Results from the overall comparison of diversity and richness in tumor versus normal tissue in oral/oropharyngeal cancers are conflicting. Similarity in overall taxonomical composition of the microbiota was found when paired samples from the same individual were compared. As suggested by Wang et al, similarity in the resident microbiome from paired samples in the same patient is unsurprising: oral fluids are common to both, and the situation may be due to the histological analogy of adjoining ‘normal’ tissue to the tumor tissue: inappropriate due to the likelihood of ‘field cancerization’ (Wang et al., 2017). In contrast, the studies that have compared cases with normal healthy controls have identified a distinct pattern of the microbiome to be associated with cancer.

Our review demonstrates that the predominant phyla present in cases and controls are essentially the same. Nevertheless, there were notable discrepancies in relative abundances of three predominant phyla namely *Firmicutes*, *Actinobacteria* and *Fusobacteria*. Three of the published studies reported a significant decrease in the relative abundance of phylum *Actinobacteria* in tumors. It has been reported that *Actinomyces* spp. which is a genus under phylum *Actinobacteria*, exerts a shielding effect on the mucosa through the release of protease-inhibitors which are capable of inhibiting tumorigenesis (Hozumi et al., 1972). It has also been postulated that *Actinomyces* spp. could have been outnumbered by oral commensal bacteria which grow faster in the relatively acidic tumor microenvironment, aided by hypoxia and glucose-deprivation (Wang et al., 2017; Schmidt et al., 2014b; Li et al., 2012; Hozumi et al., 1972; Schulze and Harris, 2012).

Taxa “*Fusobacterium*” was reported to be significantly more abundant in cancer samples: these bacteria have a special property to cohere with other species and therefore can play a critical role in helping other pathogens to colonize (Bradshaw et al., 1998). *F. nucleatum* is known to be associated with colorectal carcinoma (CRC) and has been shown to promote cellular proliferation and invasion in human CRC cell lines *in vitro* as well as to boost the evolution of OSCC and CRC in animal experiments (Binder Gallimidi et al., 2015; Kostic et al., 2013; Nakatsu et al., 2015; Rubinstein et al., 2013; Chen et al., 2012). However, this needs to be investigated in detailed prospective studies. Four studies reported the enrichment of *Parvimonas* at the genus level (Guerrero-Preston et al., 2016; Zhao et al., 2017; Pushkar et al., 2012; Wang et al., 2017). *Parvimonas* are frequently seen to be enriched in colorectal cancer as well (Nakatsu et al., 2015; Rubinstein et al., 2013); however, the association has not been characterized in detail. The enrichment of these reported taxa in these studies is notable, despite numerous methodological variances between the studies, such as sample type, collection techniques and methods of sequencing.

We cannot discriminate whether the observed changes in the

microbial community are cancer-associated, cancer-promoting or involved in cancer-pathogenesis. Two hypotheses have been proposed on the association between microbiome and cancers in the gut and extrapolation of these models to oral cancer can be convincing (Rubinstein et al., 2013; Chen et al., 2012; Sears and Pardoll, 2011; So et al., 2016; Tjalsma et al., 2012; Yang et al., 2014). In the “alpha bug model”, it is assumed that certain bacterial species might induce alterations in the mucosa that can support carcinogenesis such as epithelial cell hyperproliferation and inflammation (Rubinstein et al., 2013). Moreover, these changes may also lead to the reduction of health-promoting bacteria. *Fusobacterium* can be considered as a plausible alpha bug in oral carcinogenesis, as it has been shown to initiate secondary colonization and induce cellular proliferation in carcinoma cells *in vitro* (Al-Hebshi et al., 2017). The “driver-passenger model” for colorectal cancer hypothesizes that the putative pathogenic microorganisms grow easily in the gut of those people prone to develop cancer, and exert their effect via a considerable reduction in the fraction of health-promoting microorganisms (Tjalsma et al., 2012). These pathogenic bacteria can be considered as “bacterial drivers”, driving out the health-associated organisms, and inducing inflammation in the gut which can promote gene mutations within colonic epithelia which can lead to its proliferation. Inflammation increases intestinal permeability, modifying the microenvironment so as to allow substitution of the driver bacteria by opportunistic bacteria which, although considered as “bacterial passengers”, may aid tumorigenesis as well (Tjalsma et al., 2012; Yang et al., 2014; Wang and Jia, 2016). The literature highlights that there is an insurgence of bacteria long regarded as periodontal pathogens in oral cancer samples, which can also act as bacterial “drivers”. However, a “snapshot” of the community composition of tumor-associated microbiome captured in cross-sectional evaluations may highlight “passenger” bacteria which are present at that time and may mask the true drivers of carcinogenesis (Wang and Jia, 2016). In other words, a temporal change in the oral microbiome might be involved in progression of cancer, and such perturbations may not be visible when samples are analyzed at a single time point.

Results from microbiome data will vary depending on the molecular methods adopted. thirteen out of the fifteen studies included in this review have analyzed the microbiome after amplification of 16S rRNA genes. Amplification of 16S gene regions, which are significant for structural configuration of ribosome structure and function, using universal primers, can help to identify bacterial taxa (Isenbarger et al., 2008). Nevertheless, a significant limitation of this technique is that precision of the classification is dependent on the variable regions chosen to be amplified (Liang et al., 2011). Moreover, some bacterial species have multiple copies of 16S rRNA genes, which may lead to their artificial over-representation (Liang et al., 2011). Some early studies used Topo TA cloning and Sanger sequencing, which is labor-intensive compared to the recent methods. The small quantity of clones (90–150) that are usually picked and sequenced in this way is unlikely to represent adequately the genetic makeup of the amplified material in a patient sample as compared to NGS (which accounts for 10,000–50,000 reads on average) (Liang et al., 2011): conclusions thus drawn may bias by underestimation of important organisms (Palatinszky et al., 2011; Schulze-Schweifing et al., 2014). Newer whole metagenomic sequencing approaches, which allow investigating the entire gene content of a microbial population, have been utilized to study other types of cancer-associated microbiome (Dewhirst et al., 2010; Zhang et al., 2015); however, to our knowledge, there are no similar studies yet on oral cancer. An advantage of metagenomic sequencing over 16S rRNA-based sequencing is that we can understand the functional implications of the dysbiosis, which may be invaluable in identifying the role of the microbiome in carcinogenesis.

Microbiome studies in general including studies that have investigated the “bacteriome” of oral/oropharyngeal cancers face an ardent task of eliminating compositional and phylogenetic variations as a result of different variables like geographic location and factors that

affect the lifestyle of an individual such as diet, habits and socioeconomic status. Bacteria commonly found in the intestines of swine and fowls have been reported to be present in oral cancer specimens from Taiwan. Constant exposure of the oral cavity to external environment and distinct dietary habits in Taiwan could have contributed to the presence of bacteria like *Cloacibacillus*, *Gemmiger*, *Oscillospira*, and *Roseburia* in samples procured from the oral cavity (Ding et al., 2017; Looft et al., 2013). Further, the published data on the oral and oropharyngeal cancer microbiome is, so far, limited to few geographic locations currently (Table 1), with sparse representation elsewhere in the globe, which highlights the need for more studies with a wider geographical coverage. A subject's recent use of an antibiotic is a common exclusion criterion in microbiome studies: antibiotics can influence the oral microbiome for as long as 3–36 weeks (Lazarevic et al., 2013; Abeles et al., 2016) depending on the type of antibiotic and the personalized response of an individual to a particular antibiotic (Dethlefsen et al., 2008). However, exclusion of patients with recent antibiotic use has not been considered in a number of studies.

Sample collection is a challenging yet important aspect of microbiome research which might critically influence the data generated (Sinha et al., 2016; Kim et al., 2017). Data generated from archival frozen specimens may differ from DNA retrieved from fresh or paraffin-embedded tissue: all risk contamination in the clinic or laboratory.

Methodologic variations in the laboratory workflow can influence the microbiome composition. Chemical and mechanical methods for sample homogenization have been shown to break bacterial cell walls and consequently improve species representation (Gill et al., 2016). It is noteworthy that only three of the included studies have reported the use of enzymatic lysis or bead beating (Wang et al., 2017; Mukherjee et al., 2017; Bornigen et al., 2017) even though they are the recommended approaches of sample treatment for microbiome studies. One study included in the current review has reported the use of reagent controls to confirm the absence of contaminating bacteria which can influence data interpretation (Wang et al., 2017). This is surprising, considering that the use of controls in order to identify contaminant taxa from reagents is highly recommended in microbial studies (Adams et al., 2015; Adams et al., 2014; Salter et al., 2014). Biological variability prevails in all ecological communities and human microbiome can show evident inherent biological diversity. So it is imperative for the microbiome studies to be powered statistically in order to estimate the magnitude of microbial divergence between health and disease (Kim et al., 2017). Nevertheless sample size estimation was not detailed in any of the studies included in the review. To summarize, inconsistencies and non-uniformity in data reporting and methodological approaches can be observed in oral microbiome studies.

A key undesirable outcome of all the aforementioned inconsistencies and variations in oral/oropharyngeal cancer microbiome studies is the consequent inability to perform a meta-analysis. Even though the raw data from some of the studies were publicly deposited and obtainable to everyone, a meta-analysis of datasets with clone libraries and next generation sequencing datasets could not be performed because of the differences in scale and magnitude of sequence counts and the laboratory procedures, which are fundamentally different (Adams et al., 2015). Collective datasets generated with uniform methodology can bring us closer to deciphering microbial signatures specific to oral/oropharyngeal cancers. All these caveats highlight the emerging need for a standardized guidelines for analyzing and reporting microbiome data (Menon and Gopinath, 2018).

5. Conclusion

To our knowledge, this is the first systematic and comprehensive review that summarizes relevant epidemiological studies regarding microbial dysbiosis in association with oral/oropharyngeal cancer, and discusses discrepancies among them. Preliminary microbiome studies on oral cancer have demonstrated alteration in bacteriome which

include pathogens of known significance. This could point towards the role of bacteriome associated inflammatory changes as a contributory event in oral cancer. Equally, bacterial changes could also be a reflection of the response to the changes in oral environment due to malignancy. Nevertheless identification of specific microbial changes is a positive stride towards developing salivary based microbial biomarkers for the clinical assessment of oral cancer progression. We highlight issues which are needed to be considered for future research. Despite accumulating evidence on the metagenomic characterization of the microbiome in several human cancers, few studies have so far used NGS in the study of cancers of oral cavity. Ethnicity, dietary patterns, behavioral risk factors, and timing of collection, have rarely been considered in any of these studies. Anatomical site and sub-site of the neoplasm, sample site within the tumor, physical size of samples, and sample handling, especially the need to avoid contamination with saliva, all need to be harmonized. Comparisons of healthy tissues within the same patient have value, but data from comparable anatomical sites from healthy, matched control patients are essential as well. Determination of adequate sample sizes in the statistical sense, standardized bioinformatic pipelines and analytic approaches are desirable to allow comparisons between studies. Ideally, metagenomic studies should also be designed to explore changes over time, in the total microbiome associated with oral potentially malignant disorders with high risk of malignant transformation. And finally an integrated approach with microbiomics, transcriptomics and proteomics/metabolomics will allow robust understanding of microbial role in carcinogenesis. These are demanding criteria, which will take many years to fulfill. We hope this review will encourage future collaborations towards meeting them.

Conflict of interest

None declared.

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