Microbial biodiversity of natural toothbrushes in Mali

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Abstract

Different oral hygiene practices are used to overcome endemic diseases such as dental caries and oral infections. In Mali (Africa), natural plant-based toothbrushes are used for eliminating bacterial biofilm. The repertoire of microorganisms associated with natural toothbrushes is unknown. The aim of our study is to study microbial flora in particular the methanogenic archaea associated with natural toothbrushes recently recognized as responsible for periodontitis and peri-implantitis. We investigated the methanogens and bacteria associated with 15 different natural plant toothbrushes collected in Bamako local market (Mali). Microbiological investigations consisted in culturing the bacteria on agar plates and searching archaea using molecular techniques. No archaea were demonstrated by molecular biology but 50 bacterial species, including 33 aero-anaerobic and 17 aerobic species, were isolated from natural toothbrushes. We isolated Pseudomonas sp., Staphylococcus sp. and Klebsiella pneumoniae, which are acknowledged as opportunistic human pathogens. This study has highlighted the likely impact of the use of natural toothbrushes in the spread of potentially pathogenic bacteria in the human oral cavity.

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Keywords: Bacteria, Mali, methanogen, miswak, natural toothbrushes, oral cavity

Original Submission: 3 October 2019; Revised Submission: 13 January 2021; Accepted: 20 January 2021

Article published online: 3 February 2021

The use of natural toothbrushes or chewing sticks became popular in the Muslim world, including several African countries including Mali. These natural toothbrushes, also called Miswak, are prepared from the root, stem, twigs, or bark of trees and are used to clean the oral cavity [3]. Miswak is an Arabic word meaning tooth cleaning stick, becoming a common designation for Salvadora persica [4]. Salvadora persica is the most used plant for oral hygiene [5]. Today, the use of natural toothbrushes is very common for reasons of low cost, ease of availability and tradition.

There are around 173 different types of trees that can be used as chewing sticks, belonging to the families Salvadoraceae, Acacia, Fabaceae, Terminalia, Combretaceae, Lasianthera, Icacinaceae, Gouania and Rhamnaceae [6–8]. Studies have shown that these plants confer an essential antibacterial role, especially against cariogenic bacteria and periodontal pathogens [6]. A recent study has shown that the aqueous extract of the roots of Salvadora persica contains thermostable components with significant anti-protozoal activity against Blastocystis sp. subtypes [9]. As the
correct decontamination of natural toothbrushes before use is not always carried out, this could constitute a pathway for the diffusion of environmental bacteria and archaea into the oral microbiota in humans. It is known that the main route of exposure of humans to resistant pathogens comes from other people, either in clinics or in community settings. Environmental dissemination routes for resistant bacteria have also been pointed out as potentially important for the spread of antibiotic resistance in humans [10]. Plants harbour a wide variety of microbiomes that are specific to each plant genotype, but also specific to each plant organ, for example roots, leaves and spheres [11]. Recent next-generation sequencing studies of the environmental microbiome revealed archaeal signatures in substantial amounts in diverse microbiomes associated with plants [12]. The microbial community associated with the use of natural toothbrushes has not been studied; hence the influence on the oral microbiota is unknown. In this preliminary study conducted in Mali, we examined for the first time the microbial flora that contaminates natural toothbrushes used for oral hygiene.

Materials and methods

Collection and identification of samples

Unused natural toothbrushes were collected in March 2018 from the local market in Bamako, Mali. Identification of the scientific names of plants was based on the literature by correspondence with local names [8,13,14]. The collection consisted of 15 natural toothbrushes from different plants, including Fagara xanthoxyloides, Balanites aegyptiaca, Salvadora persica, Guiera senegalensis, Prosopis africana, Burkea africana, Diospyros mespiliformis, Ziziphus mucronata, Ficus exasperata, Amona senegalensis, Cassia sieberiana, Ficus capsensis, Grewia bicolor, Detarium microcarpum and Loeuseriella africana (Table 1). The samples were analysed at the IHU Mediterranean Infection laboratory (Marseille, France) for microbiological investigations.

Culture of the samples

The different samples were rehydrated in Falcon tubes containing sterile phosphate-buffered saline for 24 hours at room temperature. The suspension was used to inoculate the agars at different dilutions. Culture of bacteria was performed at 37°C on 5% sheep-blood-enriched Columbia agar and PolyViteX agar (bioMérieux, Marcy l’Etoile, France) under aerobic and anaerobic atmospheres, for 48 hours.

Identification by MALDI-TOF MS

All microbial colonies that grew on agar plates were identified by matrix-assisted laser desorption–ionization time-of-flight mass spectrometry (MALDI-TOF MS) using a Microflex spectrometer (Bruker Daltonics, Bremen, Germany) [15,16].

| Vernacular name (Bambarra) | Scientific name | Plant parts used | Bacteria |
|----------------------------|-----------------|------------------|----------|
| Wo                         | Fagara xanthoxyloides | Root and twig | Enterococcus faecium; Pedloococcus aegyptiacol; Enterococcus casseliflavus; Pantoea septica; Cronobacter sakazakii; Pantoea massiliaens; Staphylococcus epidermidis |
| Ziguène ou Séréné         | Balanites aegyptiaca | Rod             | Enterobacter cloacae; Enterococcus faecium; Bacillus mariotavus; Bacillus megaterium; Bacillus pumilus |
| Sourala gisè             | Salvatore persico | Branch          | Pantoea internitis; Klebsiella pneumoniae; Pantoea sepitca; Pantoea massiliaens; Pseudomonas balearica; Cronobacter sakazakii; Enterococcus faecium; Pantoea dispersa; Entrobacter cloacae |
| Kounjè                    | Guiera senegalensis | Root            | Entrobacter tabaco; Entrobacter cloacae; Entrobacter asburiae; Bacillus pumilus; Pantoea massiliaens; Pantoea septica; Cronobacter sakazakii; Escherichia coli; Kocuria rosea; Bacillus megaterium |
| Gwele                     | Prosopis africana   | Rod             | Pantoea sepitca; Klebsiella pneumoniae; Pantoea massiliaens; Bacillus firmus; Bacillus oiz; Pseudobacilla yumanenensis; Pantoea dispersa; Pantoea eucnica; Enterococcus faecium; Entrobacter cloacae; Escherichia coli; Cronobacter sakazakii; Pantoea massiliaens; Cronobacter heiloticus; Raoultella arthromyloides |
| Siri                      | Burkea africana     | Branch          | Klebsiella pneumoniae; Cassia sieberiana; Staphylococcus pasteuri; Staphylococcus epidermidis; Entrobacter cloacae; Pantoea septica; Cronobacter sakazakii; Pantoasia intestinalis |
| Sundun gisè              | Diospyros mespiliformis | Branch        | Klebsiella pneumoniae; Entrobacter cloacae; Pantoea calida; Pantoea massiliaens; Cronobacter sakazakii; Pantoasia intestinalis |
| Surukou latonono          | Ziziphus mucronata  | Branch          | Bacillus megaterium; Bacillus pumilus; Pantoea massiliaens; Bacillus megaterium; Escherichia coli; Kocuria rosea; Pantoea intestinalis |
| Surukou niénié           | Ficus exasperata    | Branch          | Pantoea septica; Staphylococcus epidermidis; Pseudomonas aeruginosa; Klebsiella pneumoniae; Entrobacter cloacae; Cronobacter sakazakii; Stenotrophomonas maltophilia; Enterococcus casseliflavus; Enterococcus faecium; Enterococcus gallinarum; Acinetobacter baumannii |
| Mandé sundun             | Amona senegalensis  | Branch          | Escherichia coli; Kocuria rosea; Pantoea septica; Pantoasia intestinalis; Bacillus oiz; Pseuobacilla yumanenensis; Pantoea dispersa; Pantoea eucnica; Enterococcus faecium; Entrobacter cloacae; Entrobacter asburiae; Bacillus pumilus; Pantoea massiliaens; Pantoea septica; Cronobacter sakazakii; Pantoasia intestinalis |
| Sinjan                   | Cassia sieberiana   | Roots           | Entrobacter casseliflavus; Entrobacter cloacae; Klebsiella pneumoniae; Entrobacter asburiae; Acinetobacter baumannii |
| Seretoro                 | Ficus capensis      | Branch          | Enterococcus casseliflavus; Entrobacter cloacae; Klebsiella pneumoniae; Acinetobacter calcoaceticus; Bacillus circulare; Bacillus flexus |
| Nogo nogo                | Grewia bicolor      | Branch          | Pantoea septica; Pantoea massiliaens; Escherichia coli; Kocuria rosea; Pantoea septica; Pantoasia intestinalis |
| Tanba kuriba             | Detarium microcarpum | Branch        | Klebsiella pneumoniae; Escherichia coli; Kocuria rosea; Pantoea septica; Pantoea massiliaens; Bacillus pumilus; Cronobacter sakazakii; Raoultella arthromyloides |
| Mankana                  | Loeuseriella africana | Branch       | Klebsiella pneumoniae; Pantoea septica; Pantoea massiliaens; Bacillus pumilus; Cronobacter sakazakii; Raoultella arthromyloides |
PCR sequencing

Methanogens were searched for by PCR sequencing. A 0.3-g quantity of acid-washed beads (B106 mm; Sigma, Saint-Quentin Fallavier, France) was added to each tube containing 250 μL of toothbrush sample, the suspension was shaken to achieve a mechanical lysis in a FastPrep BIO 101 apparatus (Qbiogene, Strasbourg, France) at level 6.5 for 2 min. Then 200 μL of buffer TL and OB Protease Solution (1.5 mL) from the E.Z.N.A. Tissue DNA Kit (OMEGA, bio-tek, Norcross, GA, USA) were added. The mixture was incubated overnight at 56°C. After a second cycle of mechanical lysis, the mixture was incubated for 60 min at 70°C. Extracted DNA was eluted with 100 μL of elution buffer and the DNA was stored at –20°C.

Amplification of the archaea 16S rRNA gene (primers used: SDArch0333aS15, 5'-TCCAGGCCCTACGGG-3' and SDArch0958aA19, 5'-YCCGGCGTTGAMTCCAATT-3') and the methyl-coenzyme M (mcrA) gene (primers used: mcrAFor, 5'-GCTCTACGACCAGATMTGGCTTGG-3' and mcrARev, 5'-CCGTAGTACGTGAAGTCATCCAGCA-3') was performed as previously described [17]. Sequencing reactions were carried-out using the Big-Dye Terminator, version 1.1, cycle sequencing kit DNA according to the manufacturer’s instructions (Applied Biosystems, Foster City, CA, USA). Nucleotide sequences were assembled using CHROMAS PRO software, version 1.7 (Technelysium Pty Ltd., Tewantin, QLD, Australia) and compared with the GenBank database by similarity search using the BLASTN program (http://www.ncbi.nlm.nih.gov/blast/).

Results and discussion

In this study, we were interested in the different types of natural toothbrushes used in Mali. We obtained a collection of 15 natural toothbrushes from different plants that are frequently used in Mali (Table 1). The natural toothbrushes that presented the most bacteria were: Prosopis africana and Guiera senegalensis with respectively 15 and 13 bacterial species (Table 1).

We isolated 50 bacterial species from natural toothbrushes, including 33 aero-anaerobic and 17 aerobic species. Among them, a variety of potential pathogenic bacteria [16–18] of the oral cavity were found. Among all bacteria isolated by culture, 70% can cause pathologies in humans, including seven bacterial genera that could potentially be responsible for respiratory infection (Fig. 1). Enterobacteria were the most represented in our study. Among these bacteria, 15 Klebsiella pneumoniae strains were isolated followed by 12 Cronobacter sakazakii, 12 Enterobacter cloacae, four Escherichia coli and three Raoultella ornithinolytica (Fig. 1). Klebsiella pneumoniae is an opportunistic bacterial pathogen known for its high frequency and diversity of antimicrobial resistance genes [19]. Natural toothbrushes can contribute to the spread of multidrug-resistant strains in humans from the environment [19].

Raoultella ornithinolytica is a bacterium often found in aquatic environments, soil, insects, fish, ticks, and termites [20]. This bacterium is involved in several pathologies in humans, including in the oral cavity [20,21].

![FIG. 1. Potential opportunistic pathogens isolated on natural toothbrushes collected in Bamako, Mali. The x-axis indicates the different bacterial species potentially pathogenic for humans, the y-axis indicates the number of bacteria isolated for each species.](image-url)
In this study, we did not detect the presence of methanogens and archaea in general on the toothbrushes using molecular biology, suggesting that natural toothbrushes are not a source for methanogens commonly found in the oral cavity, chiefly *Methanobrevibacter oralis* [22–24], *Methanobrevibacter smithii* and *Methanobrevibacter massiliensis* [18].

We isolated nine different bacterial species from the natural toothbrush *Salvadora persica* despite its previously reported antimicrobial activity against both aerobic and anaerobic bacteria [4,6,25]. However, previous studies had shown some disadvantages of using natural toothbrushes. A study by Eid et al. [26] showed that natural toothbrush users had significantly more sites with gingival recession than industrial toothbrush users, and this may influence the periodontal health. The gingival recession may have different origins, such as the absence or low thickness of keratinized tissue, the traction of a frenum or even a thin external table of bone, or traumatic or excessive brushing [27]. Another study showed that the antimicrobial substances contained in natural toothbrushes or chewing sticks provided no additional benefits to those produced by the antimicrobial activity of commercially available toothpastes [28]. The mechanical effect exerted by natural toothbrushes during dental hygiene could lead to micro-wounds through which methanogens and bacteria that are present in the oral flora could enter the body and cause systemic diseases [29]. These practices may pose a risk to the elderly, immunosuppressed individuals and those following chemotherapy or radiotherapy, who are more likely to develop both oral cavity and systemic diseases.

The mechanical effect made by natural toothbrushes during dental hygiene, as mentioned above, could cause micro-wounds and promote the haematogenous spread of microorganisms in the body.

Natural toothbrushes contaminated with *Pseudomonas* sp., *Staphylococcus* sp. and *Klebsiella pneumoniae* may be a source for these opportunistic pathogens, posing a threat to oral as well as general health in natural toothbrush users.

This pilot study made it possible to show the microbial diversity contaminating natural toothbrushes. In this work, we have also shown that natural toothbrushes could lead to the spread of multidrug-resistant bacteria in humans.

Further studies are needed to compare the oral flora of individuals using natural toothbrushes and industrial toothbrushes. In order to test the hypotheses formulated in this study, it would also be interesting to test the sensitivity of environmental bacteria to antibiotics and to determine the resistance genes circulating for a better oral health perspective.

## Conflict of interest

There is no conflict of interest.

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