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Al-2 of Aggregatibacter actinomycetemcomitans inhibits Candida albicans biofilm formation

Endang W. Bachtiar1, Boy M. Bachtiar1, Lucia M. Jarosz2†, Lisa R. Amir1, Hari Sunarto3, Hadas Ganin4, Michael M. Meijler4 and Bastiaan P. Krom5*

1 Department of Oral Biology, Faculty of Dentistry, Universitas Indonesia, Jakarta, Indonesia
2 Department of Biomedical Engineering, The W.J. Koff Institute, University Medical Center Groningen and University of Groningen, Groningen, Netherlands
3 Department of Periodontology, Faculty of Dentistry, Universitas Indonesia, Jakarta, Indonesia
4 Department of Chemistry, Ben-Gurion University of the Negev, Be’erSheva, Israel
5 Department of Preventive Dentistry, Academic Centre for Dentistry Amsterdam (ACTA), University of Amsterdam and Free University Amsterdam, Amsterdam, Netherlands

*Correspondence: Bastiaan P Krom, Department of Preventive Dentistry, Academic Centre for Dentistry Amsterdam (ACTA), G. Mahierlaan 3004, 1081 LA Amsterdam, Netherlands. E-mail: b.krom@acta.nl

Present address: Lucia M. Jarosz, Department of Cell Biology, University Medical Center Groningen, Groningen, Netherlands

Aggregatibacter actinomycetemcomitans, a Gram-negative bacterium, and Candida albicans, a polymorphic fungus, are both commensals of the oral cavity but both are opportunistic pathogens that can cause oral diseases. A. actinomycetemcomitans produces a quorum-sensing molecule called autoinducer-2 (Al-2), synthesized by LuxS, that plays an important role in expression of virulence factors, in intra- but also in interspecies communication. The aim of this study was to investigate the role of Al-2 based signaling in the interactions between C. albicans and A. actinomycetemcomitans. A. actinomycetemcomitans adhered to C. albicans and inhibited biofilm formation by means of a molecule that was secreted during growth. C. albicans biofilm formation increased significantly when co-cultured with A. actinomycetemcomitans luxS strain, lacking Al-2 production. Addition of wild-type-derived spent medium or synthetic Al-2 to spent medium of the luxS strain, restored inhibition of C. albicans biofilm formation to wild-type levels. Addition of synthetic Al-2 significantly inhibited hypha formation of C. albicans possibly explaining the inhibition of biofilm formation. Al-2 of A. actinomycetemcomitans is synthesized by LuxS, accumulates during growth and inhibits C. albicans hypha- and biofilm formation. Identifying the molecular mechanisms underlying the interaction between bacteria and fungi may provide important insight into the balance within complex oral microbial communities.

Keywords: oral microbiology, interspecies interaction, quorum sensing

INTRODUCTION

Aggregatibacter actinomycetemcomitans is a non-motile, Gram-negative coccobacillus, which can be found as a commensal in the oral cavity. In addition, it is also the principal cause of aggressive periodontal disease (Saarela et al., 1999). A. actinomycetemcomitans uses chemical signals to sense cell density and alter expression of virulence factors (Novak et al., 2010). This microbial communication is known as quorum sensing (QS) and the only identified cell-cell signaling molecule in A. actinomycetemcomitans this far is autoinducer 2 (AI-2). AI-2, which has been proposed to be a general interspecies, concentration dependent signal, is synthesized by LuxS as a precursor, 4,5-dihydroxy-2,3-pentanedione (DPD), followed by secretion into the medium where it spontaneously undergoes cyclization into AI-2 which accumulates. Interestingly, DPD can thus be converted into several structures that can be recognized by different species (Miller et al., 2004). For instance, Vibrio harveyi produces an unusual furanosyl borate diester (3aS,6S,6AR)-2,2,6,6a-tetrahydroxy-3a-methyltetrahydrofururo[3,2-d][1,3,2]dioxaborol-2-uide), while Salmonella Typhimurium recognizes (2R,4S)-2-methyl-2,3,3,4-tetrahydroxytetrahydrofuran. In addition, the polymorphic fungus Candida albicans is one of the most commonly isolated fungi from the oral cavity (Cannon and Chaffin, 1999). In healthy individuals, C. albicans grows as a commensal, mostly in the unicellular yeast morphology, but in immune-compromised individuals this species is capable of producing multicellular filamentous forms, a pathogenic morphology (Odds, 1988). The morphological transition from the yeast-to-hyphal mode of growth is influenced by many factors, including pH, nutrient availability, temperature and the presence of QS molecules, that stimulate or repress filamentation as a function of cell density (Hornby et al., 2001; Hazan et al., 2002). Recently, the interaction of bacteria with C. albicans through QS molecules has received increasing attention (Shirliff et al., 2009). In the oral cavity two Gram-positive bacteria have been shown to affect C. albicans biofilm formation through QS molecules. Streptococcus mutans was shown to inhibit hyphal formation through the QS molecule competence-stimulating peptide (CSP) (Jarosz et al., 2009) and the fatty acid signaling molecule trans-2-decenolic acid (Vilchez et al., 2010). In addition, S. gordonii was shown to secrete AI-2, which was shown to repress C. albicans QS by inhibiting the action of farnesol (Bamford et al., 2009) upon
contact. In the healthy oral cavity a multitude of bacterial species co-exist with *C. albicans*, both Gram-negative and Gram-positive (Zaura et al., 2009). In contrast to the limited information on the effect of Gram-positive oral bacteria on *C. albicans*, no information is available on the effect of QS molecules of Gram-negative oral bacteria. Therefore, in the present study, it was our aim to investigate the effect of AI-2 produced by the Gram-negative oral bacterium *A. actinomycetemcomitans* on *C. albicans*.

**MATERIALS AND METHODS**

**MICROBIAL STRAINS AND GROWTH CONDITIONS**

*A. actinomycetemcomitans* spp. were routinely cultured for 18 h at 37°C under microaerobic conditions (10% CO₂) on trypticase soy agar or broth containing 0.6% yeast extract (TSB-YE/Difco). When appropriate, standardized cell suspensions were prepared and placed on agar plates and determined at the optical density measured at 655 nm (OD₆₅₅). *Escherichia coli* was routinely grown in Luria-Bertani broth at 37°C with constant aeration. For solid medium, 15 g agar per liter was added to the liquid medium. When required, ampicillin (100 μg mL⁻¹) or kanamycin (30 μg mL⁻¹) was added to the medium. *C. albicans* strain ATCC 10231 or SC5314 were taken from stock cultures frozen in 15% glycerol and incubated at 37°C and sub-cultured twice onto yeast peptone agar plates with 2% glucose (YPD) or when indicated in yeast nitrogen base medium pH 7, supplemented with 50 mM glucose (YNB).

**ADHESION OF A. ACTINOMYCETEMCOMITANS TO C. ALBICANS**

Adhesion of *A. actinomycetemcomitans* Y4 to *C. albicans* strain SC5314 was studied using a Bioflux 1000Z setup. Briefly, *C. albicans* was seeded into a 48-well microfluidics plate (Fluxion Biosciences) at an initial optical density measured at 600 nm of 0.2 (OD₆₀₀ = 0.2). After 30 min adhesion to the bottom plate at 37°C, flow with prewarmed YNB was started at 0.5 dyne cm⁻² for 4 h. A bacterial suspension at an initial OD₆₀₀ = 0.2 in phosphate buffered saline (PBS; 10 mM potassium phosphate, 0.15 M sodium chloride, pH 7) containing 0.2 μL mL⁻¹ Syto9 and 0.2 μL mL⁻¹ propidium iodide (Baclight, Invitrogen) was flowed through the microfluidics channels at 0.5 dyne cm⁻² and images were captured every 30 s using the appropriate filter settings. Images were translated to AVI-movies using ImageJ (1.46r).

**CONSTRUCTION OF AN A. ACTINOMYCETEMCOMITANS luxS MUTANT**

*A. actinomycetemcomitans* (serotype b) was isolated in our periodontal clinic (Universitas Indonesia, Jakarta) from periodontitis patients, with their consent. The isolated bacteria were identified by means of Gram staining, characteristic star-positive colonies on agar plates and tight adherence to surfaces when grown in broth (Slots et al., 1982). Positive clones were confirmed and serotyped using PCR (Suzuki et al., 2001). A single isolate, *A. actinomycetemcomitans* UI-09, was selected as wild-type strain for all other experiments. In addition, the commonly used *A. actinomycetemcomitans* Y4 was used as a reference in certain experiments and showed similar results compared to *A. actinomycetemcomitans* UI-09.

All primers for plasmid construction were designed using *A. actinomycetemcomitans* database (http://www.oralgen.org, ID for luxS in *A. actinomycetemcomitans* HK 1651 is AA00516). In order to create a luxS defective mutant, a suicide vector was constructed using the neighbor-joining technique as previously described for *Campylobacter jejuni* (Bachtir et al., 2007). Firstly, a 229-bp DNA fragment containing the upstream sequence adjacent to luxS was PCR-amplified using the primer EcoRI-L1 (ACGAATTCATCCACCGCATCTT, forward) and primer BamHI-L2 (TCCGATCCAGTTTTCTTGATG, reverse). The PCR product was cloned into pBluescript in the forward direction via the EcoRI and BamHI sites, and confirmed by restriction analysis. The construct (pBl-L1) was subsequently introduced into *E. coli* JM 107. Positive clones were selected on LB agar supplemented with ampicillin, X-Gal and IPTG. Secondly, a 409-bp DNA fragment containing the downstream flanking region of luxS was amplified using primers BamHI-L3 (CATGGATCCAGTTTTCTTGATG, forward) and Xbal-L4 (ATCTAGAGCAATTTCTTGATG, reverse). The amplified fragment was inserted into pBl-L1 between BamHI and Xbal sites. The resulted intermediate plasmid (pBl-L2) was cut at the unique BamHI site and a 1.4-kbp fragment containing a kanamycin cassette (obtained from vector pMW2 digested with BamHI) was ligated into the plasmid to obtain the suicide plasmid, pBLkm².

**NATURAL TRANSFORMATION**

A biphasic system for *A. actinomycetemcomitans* transformation was performed as described previously (Wang et al., 2002). Transformation was done by incubating a suspension of 10⁶ CFU mL⁻¹ of *A. actinomycetemcomitans* at 37°C under microaerobic conditions for 3 h. Subsequently, 10 μg of the suicide vector was added and cells were incubated for 3 h at 37°C. Cells were then harvested and plated on medium supplemented with kanamycin and incubated at 37°C, under microaerobic conditions for 2 days to select for transformants. Homologous recombination and disruption of luxS was confirmed using PCR analysis with primers flanking the target site (EcoRI-L1 and Xbal-L4).

**SPENT MEDIUM PREPARATION**

Spent medium of *A. actinomycetemcomitans* cultures was prepared as described previously (Jarosz et al., 2009). Protein concentration in the spent medium was measured using the Bradford method. Spent medium was diluted in PBS to yield 10 and 100 μg mL⁻¹ concentrations and used immediately or stored for short periods of time at −20°C. The pH of the spent medium was adjusted to pH 7.

**SYNTHESIS OF (S)-4,5-DIHYDROXY-2,3-PENTANEDIONE (DPD)**

DPD was synthesized following a procedure published by Ganin et al. (2009). Lyophilized DPD was dissolved in DMSO at 33 mM stock concentrations and stored at −20°C until required. Synthetic DPD concentrations used in the described experiments are assumed to be physiologically relevant as they are in line with concentrations of AI-2 reported to be produced in saliva-fed natural oral biofilms (Rickard et al., 2008).

**BIOFILM FORMATION OF C. ALBICANS AND A. ACTINOMYCETEMCOMITANS**

Quantification of *A. actinomycetemcomitans* biofilms was achieved by staining with Crystal Violet (CV).
A. actinomyctemcomitans strains (5 µL) were used to inoculate wells of 96-well (flat-bottom) cell culture plates (Sumitomo Bakelite Co., Ltd, Tokyo, Japan) containing 95 µL TSB-YE in each well. After 18 h of incubation under microaerobic condition, the culture medium containing planktonic cells was removed and the wells were carefully washed with 200 µL of distilled water. Adherent bacteria were stained with 50 µL of 0.1% CV for 15 min at room temperature. After rinsing twice with 200 µL of distilled water, CV bound to the biofilm was extracted with 200 µL of 99% ethanol for 20 min and quantified by measuring the absorbance at 655 nm with a microplate reader (Model 3550, Bio-Rad Laboratories, Hercules, CA, USA).

Biofilm formation of C. albicans was induced in YNB as previously reported (Krom et al., 2007). Mixed species biofilms of C. albicans (2 × 10^6 CFU mL^-1) and A. actinomyctemcomitans (2.1 × 10^7 CFU mL^-1) were grown in medium containing 70% YNB and 30% TSB-YE (vol/vol). Where indicated, the 30% fresh TSB-YE fraction was replaced by spent medium as described previously (Jarosz et al., 2009). In addition, when indicated, sterile spent medium from the A. actinomyctemcomitans was added to YNB at 1, 10, and 100 µg mL^-1 protein concentrations. After 24 and 48 h of growth, biofilm formation on the well of microtiter plates were washed once with PBS and metabolic activity of the biofilms was quantified using [3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide] (MTT) as described previously (Krom et al., 2007). Since C. albicans rapidly metabolized MTT, in contrast to A. actinomyctemcomitans, this assay could be used to determine C. albicans biofilm formation even in co-cultures with A. actinomyctemcomitans. All assays were carried out on at least two separate occasions with at least duplicate determinations on each occasion. Microtiter wells containing only YNB broth but no cells were used as negative controls. To mimic the conditions in the oral cavity initial experiments were also performed in the presence of pooled sterilized human saliva, however no differences were observed relative to experiments without saliva. Therefore, all other experiments were performed without saliva. 

INHIBITION OF HYPHA FORMATION

Hypha formation was assayed as described previously (Jarosz et al., 2009) using C. albicans strain SC5314. The morphology of cells was analyzed using a 20x objective on an inverted microscope (Olympus, Tokyo, Japan).

STATISTICAL ANALYSES

Differences between means were analyzed for statistical significance using two-tailed Student’s t-tests. Differences were considered significant when p ≤ 0.05 level. For multiple comparisons, a One-Way ANOVA followed by a TUKEY HSD test was performed (http://vassarstats.net/anovalu.html).

RESULTS

ADHESION OF A. ACTINOMYCTEMCOMITANS TO C. ALBICANS

Adhesion to surfaces is a critical initial step in microbial biofilm formation. Using a microfluidics setup, adhesion under flow conditions were studied. Single A. actinomyctemcomitans Y4 cells adhere to hyphae and yeast cells of C. albicans SC5314, as illustrated by the increased green fluorescent spots associated with C. albicans (Figure 1). In addition, adhesion of A. actinomyctemcomitans to the glass surface could be observed.

EFFECTS OF luxS DELETION ON BIOFILM FORMATION OF A. ACTINOMYCTEMCOMITANS

A. actinomyctemcomitans luxS formed significantly less biofilm compared to the wild type strain (Figure 2) but growth rate in planktonic cultures was not affected, in line with previous reports (Novak et al., 2010). Spent medium of a 4 h-old culture of the wild type strain was able to rescue biofilm formation of the luxS strain at 100 µg protein mL^-1 (Figure 2, left panel). Medium of 6 h-old cultures also rescued the phenotype to a similar, but not medium of 8 and 24 h-old cultures. Addition of synthetic DPD rescued this phenotype specifically at 100 nM DPD, but not at lower or higher concentrations (Figure 2, right panel).

FIGURE 1 | Real-time microscopic analysis of adhesion of A. actinomyctemcomitans to C. albicans SC5314. Hyphae of C. albicans SC5314 were allowed to form on the bottom of a microfluidics plate. Bacteria were stained with Syto9 and PI and allowed to adhere to C. albicans SC5314 while flowing at 0.5 dyne cm^-2. Images were captured every 30 s for a total of 10 min (montage shows images every 210 s). Images were edited for brightness and contrast.
**ROLE OF AI-2 ON MIXED SPECIES BIOFILMS OF A. ACTINOMYCETEMCOMITANS AND C. ALBICANS**

Compared to mono-species biofilm formation of *C. albicans*, mixed species biofilms of *C. albicans* with *A. actinomycetemcomitans* resulted in a reduction of more than 50% in metabolic activity after 24 h of culturing (Figure 3A). A similar decrease in biofilm formation was observed after 48 h of culturing (not shown). Mixed species biofilms of *C. albicans* with *A. actinomycetemcomitans luxS* had no significant effect compared to mono-species biofilms.

**EFFECT OF SECRETED FACTORS ON C. ALBICANS BIOFILM FORMATION**

To further determine whether the inhibition effect was modulated by secreted compounds of the bacteria, the spent medium of the wild type *A. actinomycetemcomitans* was used as a source of secreted molecules to complement *A. actinomycetemcomitans luxS*. When spent medium from 4 and 6 h-old wild-type cultures was added, the *A. actinomycetemcomitans luxS* mutant was able to inhibit *C. albicans* biofilm formation, but this inhibition was not observed for spent medium derived from 8 and 24-h old cultures (Figure 3B). Conversely, when spent medium of *A. actinomycetemcomitans luxS* was added to mixed species biofilms of *C. albicans* and *A. actinomycetemcomitans luxS*, no significant inhibitions of biofilm formation was observed (Supplementary Figure S2).

**EFFECT OF SYNTHETIC DPD ON C. ALBICANS BIOFILM GROWTH AND HYPHA FORMATION**

When synthetic DPD was added to spent medium of *A. actinomycetemcomitans luxS* during *C. albicans* biofilm growth, a concentration dependent inhibition of *C. albicans* biofilm formation was observed with the maximum inhibition reached at 100 nM DPD (Figure 4, right panel). Hypha formation is a key process in biofilm formation. Synthetic DPD was added to *C. albicans* under hypha inducing conditions. DPD inhibited hypha formation by 30 and 70% at 100 nM and 1 µM, respectively (Figure 4, left panel).

**A. ACTINOMYCETEMCOMITANS SPENT MEDIUM DISRUPTS ESTABLISHED C. ALBICANS BIOFILM**

To test the ability of *A. actinomycetemcomitans* to disrupt *C. albicans* biofilms, we challenged established biofilms (24 h old) of *C. albicans* with spent medium from *A. actinomycetemcomitans* WT and luxS cultures of increasing age. A culture age dependent decrease in viability was observed when *C. albicans* biofilms were exposed to spent medium of cultures of the WT strain, but not to spent medium derived from the same aged cultures of the luxS strain (Figure 5).

**DISCUSSION**

*A. actinomycetemcomitans* is related to severe periodontitis. In addition, several studies have reported on the isolation of a large number *Candida* spp., from periodontal pockets of patients with periodontitis (Jarvensivu et al., 2004; Urzua et al., 2008). Recently, co-isolation of *Candida* spp. and *A. actinomycetemcomitans* was correlated with the occurrence of severe periodontitis (Brusca et al., 2010). However, due to limited information it is currently unclear if *C. albicans* participates in the etiology of any kind of periodontitis. Biofilms are the most common mode of growth of *Candida* spp., as observed in vivo. Therefore, interspecies interactions occur preferentially in mixed species biofilms such as those found in the periodontal pocket. A better fundamental understanding of interspecies interaction in the oral cavity is therefore of great relevance. Here we show for the first time that a commonly isolated periodontal pathogen *A. actinomycetemcomitans* adheres to *C. albicans* under flow condition (Figure 1). Physical interactions, such as adhesion are initial and probably critical stages in microbial biofilm formation. Following adhesion growth...
**FIGURE 3** | The effect of co-culture with *A. actinomycetemcomitans* on *C. albicans* ATCC 10231 biofilm formation. (A) *C. albicans* biofilms were grown without any bacteria or with *A. actinomycetemcomitans* wild-type or *luxS* mutant. Biofilm formation, quantified using the MTT assay, was measured after 24 h of growth. Data represent the mean and standard deviations of six biofilms grown on two separate occasions. (B) The effect of spent medium on co-cultures between *C. albicans* and *A. actinomycetemcomitans* *luxS*. Spent medium of *A. actinomycetemcomitans* wild-type cultures grown for different times was added to the mixed species culture of *C. albicans* and *A. actinomycetemcomitans* *luxS*. Biofilm formation was quantified after 24 h of growth using the MTT assay. The relative biofilm formation compared to *C. albicans* + *A. actinomycetemcomitans* *luxS* without any spent medium was calculated. Data represent the mean and standard deviations of six biofilms grown on two separate occasions. *Indicates significantly different from control.

**FIGURE 4** | Effect of synthetic DPD on *C. albicans* SC5134. Hypha-formation was induced by switching fresh cultures to 37°C for 3–4 h. Representative microscopic images show a decreased hypha formation (A = control, B = 0.1 µM DPD, C = 1.0 µM DPD; bar represent 40 µm for all images). Hyphae and yeast morphologies were counted and plotted as % of all cells (D). The results represent the mean of two independent experiments, each consisting of at least 100 cells per sample. Synthetic DPD added to spent medium of *A. actinomycetemcomitans* *luxS* inhibits *C. albicans* SC5314 biofilm formation (E). *Indicates significantly different from control.
and accumulation results in interspecies chemical interactions, amongst others mediated by quorum sensing (Jarosz and Krom, 2011).

AI-2 activity has been discovered in spent culture supernatants of many oral bacteria (Fong et al., 2001; Blehert et al., 2003; Wen and Burne, 2004). However, the function of AI-2 as a general bacterial signaling molecule is an issue that is yet to be resolved. Several studies have suggested that AI-2 is involved in biofilm formation (Chung et al., 2001; Yoshida et al., 2005). On the other hand, it has also been suggested that the main function of the LuxS enzyme is in the regulation of metabolic processes (Winzer et al., 2002). Inactivation of luxS in A. actinomyctecomitans does not affect growth rates but does result in phenotypic alterations, including biofilm formation and reduced colonization in various experimental infection models (Hardie and Heurlier, 2008). Our data demonstrated that biofilm formation of the A. actinomycetecomitans luxS strain was decreased as compared with its parent strain. Complementation of the luxS deletion by both spent medium of wild-type A. actinomycetecomitans as well as by addition of synthetic DPD would indicate that the luxS mutant phenotype is due to absence of the quorum sensing molecule and not related to any metabolic defect. It should however be noted that it is not straightforward to uncouple signaling and metabolic functions of LuxS (Redanz et al., 2012). The remarkable concentration dependent complementation of the luxS mutant by DPD—reaching a maximum at 100 nM, followed by a sharp decrease at higher concentrations - is interesting and currently we are unable to explain this specific behavior. It is however important to notice that such a behavior has been observed in another study. Rickard and coworkers described a similar sharp concentration optimum for synthetic AI-2 in a mixed-species biofilm model consisting of Actinomyces naeslundii and Streptococcus oralis be it at approximately 100-fold lower concentration (Rickard et al., 2006). Spent medium of 8 h cultures and older no longer inhibited C. albicans biofilm formation (see Figure 5). This could be related to decreased AI-2 presence at later stages of growth which is in line with the growth-phase specific production of AI-2, higher in the first 6 h of growth compared to later stages, observed in several oral species (Fong et al., 2001; Blehert et al., 2003; Wen and Burne, 2004).

Bamford et al. showed that S. gordonii increases C. albicans biofilm formation in a LuxS dependent fashion, however, no effect of synthetic DPD on C. albicans hypha formation was observed (Bamford et al., 2009). In contrast, we observed that the exogenous addition of AI-2/DPD to the growth medium restored the luxS phenotype in a dose-dependent manner and that addition of synthetic DPD inhibited hypha formation of C. albicans. This effect was only seen in medium consisting of 70% YNB/30% TSB, and not in YNB alone. In contrast to Bamford we did not use saliva in the germ-tube assay. Both minimal medium and saliva are strong inducers of hypha formation and this might explain the observed different response of C. albicans to synthetic DPD. Additionally, use of different C. albicans strains in the present study compared to Bamford and coworkers could account for the observed differences in response. Alternatively, the different response under different hypha inducing conditions could indicate a difference in the chemical structure of AI-2 produced by S. gordonii or A. actinomycetecomitans, and intriguingly that C. albicans is able to recognize this chemical difference. Detailed chemical characterization of the AI-2 for both species should be performed to further support this statement.

Communication within oral microorganisms involves several classes of signal molecules; autoinducing peptides synthesized by Gram-positive bacteria, such as competence-stimulating peptide of S. mutans (Jarosz et al., 2009), fatty acid signaling molecules such as farnesol and trans-2-decenoic acid (SDSF) (Vilchez et al., 2010) and AI-2 as a proposed universal signal produced by both Gram-positive and Gram-negative bacteria (Federle and Bassler, 2003). The different responses of C. albicans to AI-2, depending on its origin, is intriguing and could illustrate that the already complicated interspecies chemical communication might be even more complicated, but a more extensive study is needed to provide solid evidence for such a tempting, but speculative conclusion.

Oral biofilms are very complex multi-species communities in which we assume that interspecies interaction plays a role in establishing and maintaining a balance. A recent study on bacterial and yeast colonization in a group of mucositis patients showed a complete absence of A. actinomycetecomitans and a more than average presence of C. albicans (Laheij et al., 2012). As hypha formation is a pivotal step in C. albicans biofilm formation (Krueger et al., 2004) and adhesion, the LuxS mediated interaction with A. actinomyctecomitans would decrease fungal biofilm formation. Additionally, the function of LuxS is related to the regulation of bacteria virulence factors, including cytolethal distending toxin (CDT), as reported in Campylobacter jejuni (Jeon et al., 2005). For A. actinomyctecomitans it has been shown that CDT (CDTB) is toxic to the yeast Saccharomyces cerevisiae (Matangkasombut et al., 2010). There are therefore potentially multiple systems, one based on
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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: http://www.frontiersin.org/journal/10.3389/fcimb.2014.00094/abstract

Supplementary Figure S1 | Comparison of MTT and XTT reduction by C. albicans SC5314 and A. actinomycetemcomitans. Metabolic activity of C. albicans yields a purple or orange color upon reduction of MTT and XTT respectively. Incubation of A. actinomycetemcomitans does not result in a significant color change.

Supplementary Figure S2 | Complementation of the luxS-effect using spent medium of wild-type A. actinomycetemcomitans. Addition of 4 and 6 h-old wt-spend medium to mixed species biofilms of C. albicans ATCC 10231 and A. actinomycetemcomitans results in inhibition of biofilm formation. Addition of spent medium of A. actinomycetemcomitans luxS did not affect biofilm formation.

Supplementary Figure S3 | Effect of spent medium of wild-type A. actinomycetemcomitans on growth kinetics of C. albicans SC5134. An overnight culture of C. albicans was diluted 1:20 or 1:50 in medium consisting of 70% YNB + 30% filter sterilized 4 h-old spent medium of wild-type A. actinomycetemcomitans. As a control, the same inoculum was grown in 70% YNB + 30% fresh TSB + YE. No significant differences were observed illustrating the absence of any toxicity related agent in the spent medium of wild-type A. actinomycetemcomitans.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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