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**Antimicrobial Efficacy of an Innovative Emulsion of Medium Chain Triglycerides against
Canine and Feline Periodontopathogens**

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Conflict of interest statement

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Structured Summary

Objectives

To test the *in vitro* antimicrobial efficacy of a non-toxic emulsion of free fatty acids against clinically relevant canine and feline periodontopathogens for the prevention of periodontitis and gingivitis in cats and dogs.

Methods

Antimicrobial kill kinetics were established utilising an alamarBlue® viability assay against ten strains of canine and feline periodontopathogens, in the biofilm mode of growth, at a concentration of 0.125% v/v medium chain triglyceride (ML:8) emulsion. The results were compared with 0.12% v/v chlorhexidine digluconate and a commercially available xylitol-containing dental formulation (Virbac Vet Aquadent®). Mammalian cellular cytotoxicity was also investigated for both the ML:8 emulsion and chlorhexidine digluconate (0.25 to 0.0625% v/v) using *in vitro* tissue culture techniques.

Results

No statistically significant difference was observed in the antimicrobial activity of 0.125% v/v ML:8 emulsion and 0.12% v/v chlorhexidine digluconate against all ten periodontopathogens tested; a high percentage kill rate (> 70%) was achieved within 5 minutes of exposure and at subsequent time points investigated. A statistically significant improvement in antibiofilm activity was seen with 0.125% v/v ML:8 emulsion when compared with a currently available xylitol containing drinking water additive (Virbac Vet Aquadent®). The ML:8 emulsion possessed a significantly lower ($P < 0.001$) toxicity profile when compared to 0.12% v/v chlorhexidine digluconate in cytotoxicity assays.

Clinical Significance

The ML:8 emulsion exhibited significant potential as a putative effective antimicrobial alternative to chlorhexidine- and xylitol- based products for the prevention of periodontal disease, which, when compared to chlorhexidine at equivalent concentration, exhibited significantly reduced cytotoxic characteristics.

Keywords: Periodontitis, antimicrobial, medium chain triglycerides, biofilm bacteria, fatty acids.

Introduction

Periodontitis is the most common described progressive inflammatory disease in companion animal practice, affecting more than 80-85% of dogs and cats above three years of age (Watson 2006). Within the oral cavity, the condition refers to inflammation of the tooth support structures leading to damage and loss of the periodontal membrane, alveolar bone and adjacent soft tissues; the resulting damage may potentially result in tooth loss. The severity and prognosis of dental disease is dependent on multiple factors including species, age, breed, genetics, nutritional status, the presence of irritants, chewing activity, co-morbidities, dental crowding, occlusion and oral microbial profile (Harvey and Emily 1993). The prospect for systemic and chronic diseases to develop subsequent to periodontal disease is high due to the dense vascular network of the gum tissue (DeBowes *et al.* 1996).

The tooth and its supporting structures provide an optimum environment for the growth and replication of transient microorganisms within the mouth (Wiggs and Lobprise 1997). Food particles collect between the teeth to provide a nutrient source for the development of a bacterial biofilm community (plaque) at the tooth's surface. Bacteria, growing as biofilms, are notoriously difficult to eradicate, often requiring bactericidal concentrations of 10-1000 times that of free-floating, planktonic bacteria in suspension. Exposure to sub-optimal or sub-therapeutic antimicrobial concentrations in the biofilm thereby increases the potential for antimicrobial resistance development (Stewart and Costerton 2001). Within days minerals in the saliva, such as calcium, combine with plaque to form calculus material (tartar) and an immune response is initiated by the host resulting in the inflammatory signs of gingivitis and periodontitis, indirect periodontal destruction, pain, halitosis and loss of appetite (Wiggs and Lobprise 1997).

The microbial ecology of the oral cavity of cats and dogs is vastly diverse with aerobic bacteria predominating in the early phase of gingivitis, followed by a predominantly anaerobic and Gram-negative bacterial profile when periodontitis becomes established (Hennet and Harvey 1991) (Harvey *et al.* 1995). The process of bacterial biofilm formation begins on the tooth surface immediately after successful scaling. Initially, Gram-positive cocci, including *Streptococci* (Leonhardt *et al.* 1992)

(Radice *et al.* 2006) become attached to the surface. Further growth and maturation of this aerobic or facultative flora leads to depletion of locally available oxygen and anaerobes such as *Porphyromonas gingivalis* and *Fusobacterium nucleatum* become more predominant (Cleland 2001). As gingival inflammation develops in response to the presence of bacteria, metabolic and inflammatory products such as endotoxins become constitute major components of the gingival fluid, contributing to local oral and systemic tissue destruction and dental bone loss (Holmstrom and others 2004). Invasive pathogens capable of active spread through the systemic blood supply may colonise other highly vascularised tissues including the kidneys, liver and heart. These micro-abscesses reduce overall long-term health for the animal, increasing risk factors associated with heart disease, hypertension and kidney disease (Glickman *et al.* 2011).

For both cats and dogs, treatment of established infection centres on the use of antibiotics and anti-inflammatory agents, along with dental scaling and polishing (physical removal of calculus) performed by a veterinary practitioner. Scaling and removal of subgingival plaque is particularly problematic, often requiring administration of a general anaesthetic with the procedure repeated regularly throughout the animal's lifetime (Harvey 2005). Preventative therapy is typically centred on mechanical removal of adhered bacteria with a routine of regular toothbrushing aided by veterinary toothpastes containing antimicrobials ranging from chlorhexidine digluconate and cetylpyridium chloride to enzyme-based formulations. However, in addition to pet compliance issues, the effective removal of plaque requires the pet owner to be manually dexterous and patient (Iacono *et al.* 1998). In real-life practice, these factors often limit successful compliance.

This study describes the formulation of an antimicrobial emulsion, intended for buccal application in companion animals, comprising of a medium chain triglyceride (ML:8) oil phase dispersed in water that displays *in vitro* efficacy at a low concentration (0.125% v/v) against resistant biofilm forms of ten periodontopathogens clinically implicated in canine and feline dental disease (Elliot *et al.* 2005) (Kolenbrander *et al.* 2002). This antimicrobial emulsion has the potential to be utilised as a drinking water additive to increase ease of use for the pet owner and subsequent aid compliance, with the

overall aims of reducing long-term oral bacterial bioburden and the incidence of periodontal disease in both cats and dogs.

Materials and Methods

Formulation of ML:8 Emulsion

The antimicrobial composition of ML:8 consists of an oil in water emulsion. A mixture of free fatty acids solubilised in water is promoted by the addition of membrane lipids, in this case lecithin. Lower melting point fatty acids such as caprylic and oleic acid were utilised to a final concentration of 6.375% w/w. The ratio of free fatty acids to membrane lipid was 1.275:1. The final formulation was freshly diluted 1 in 51 (0.125% v/v free fatty acids) in sterile water before analysis.

Comparator Substances

Chlorhexidine digluconate was obtained from Sigma-Aldrich (Dorset, UK). A commercially available drinking water additive (Virbac Vet Aquadent®) containing xylitol 0.5% concentration (with chlorhexidine <0.01%) was obtained from Virbac (Bury St Edmonds, UK).

Microbial Isolates Investigated

Haemophilus actinomycetemcomitans (NCTC 10979), *Streptococcus sanguinis* (NCTC 10904) were obtained from HPA Culture Collections (Salisbury, UK). *Porphyromonas cangingivalis* (VPB 4874), *Porphyromonas salivosa* (VPB 3313), *Porphyromonas gingivalis* (VPB 5089), *Fusobacterium nucleatum* (VPB 4888), *Eikenella corrodens* (VPB 3935), *Bacteroides fragilis* (VPB 3371), *Prevotella intermedia* (VPB 3321) and *Tannerella forsythesis* (VPB 4947) were obtained from Dr Denise Wigney, Faculty of Veterinary Science, University of Sydney, Australia. All isolates of microorganisms were stored at -80°C in 10% glycerol.

Rate of Reduction in Viability of Periodontal Biofilms using an alamarBlue® Assay

Microorganisms under investigation were grown over 48-72 hours at 37°C in Tryptone Soya broth in a Biotat Class II Microbiological anaerobic Safety Cabinet (Don Whitely Scientific Ltd., Shipley, UK).

Upon visual confirmation of growth, the inoculum was adjusted to an optical density of 0.3 at 550nm in QSR solution, which was equivalent to 1×10^8 colony forming units per milliliter (CFU/mL). This suspension was further diluted in Tryptone Soya broth (TSB) (1 in 50) and dispensed aseptically in 100µL aliquots to each well of the microtitre plate. The inoculated plates were placed in Anaerogen sachets (Oxoid, Hampshire, UK) and the air tight sachets containing the inoculated plates were removed from the cabinet and biofilm was formed on the surface of the well under shear stress provided by a Gallenkamp gyratory incubator at 37°C. After an inoculation period of 48 hours, the Anaerogen sachets containing the biofilm growth plates were transferred to the anaerobic cabinet, cultures were decanted and the plates irrigated twice with 200µL of sterile autoclaved 0.9% w/v sodium chloride (NaCl) in each well. Washed plates were tapped gently upside down on a sterile paper towel to remove residual wash. The prepared biofilm was treated for evaluation of viability using alamarBlue® in a method similar to that used by Pettit *et al.* (2005). The viability of established biofilms was assessed by re-charging wells from above immediately after washing and without drying, with 0.1mL fresh TSB broth containing 20% v/v alamarBlue® (AbD Serotec, Oxford, UK), incubating the plate at 37°C for one hour followed by spectrophotometric measurement of absorbance at a wavelength of 570nm. AlamarBlue® is resazurin, a redox indicator which is reduced by metabolic activity of viable microbial cells to pink fluorescent resorufin. The reduction of viability (killing) of established biofilm with the test formulations was evaluated by loading wells containing washed biofilm from above immediately after washing with 0.1mL of 0.125% v/v ML:8 emulsion, 0.12% v/v chlorhexidine digluconate or xylitol followed by incubation at 37°C. Time points selected for analysis of ML:8 emulsion/chlorhexidine digluconate/xylitol activity on established biofilms were 0, 5, 10, 20, 30, 40, 50 and 60 minutes, followed by 2, 4 and 24 hours. Each control well had 8 replicates at each time point. Blank determinations (100% viability: positive control) were conducted using sterile distilled water. Immediately following incubation, the exposed biofilms were washed twice by irrigation with sterile 0.9% w/v NaCl and recharged with fresh TSB containing 20% v/v alamarBlue® (sterilized by passage twice through a syringe fitted with a 0.22µm membrane filter) and 3% w/v Tween 80 (polysorbate), incubated at 37°C for 1 hour and the development of UV absorbance was then measured spectrophotometrically at 570nm. A positive control (100% microbicidal effect) was also

included in the assay using 2% chlorhexidine digluconate. Reduction in viability of biofilm following exposure to the test formulations was expressed as a percentage based on the percentage reduction between untreated (blank) wells and treated wells using the following equation:

$$\% \text{reduction in viability} = \frac{Abs_{570nm} \text{ ML:8 Emulsion / Chlorhexidine digluconate / Virbac Vet Aquadent®} - Abs_{570nm} \text{ Blank}}{Abs_{570nm} \text{ Chlorhexidine digluconate 2\%} - Abs_{570nm} \text{ Blank}} \times 100$$

Cell Survival Analysis (cytotoxicity assays)

Two individual mammalian cell lines were selected for cytotoxicity analysis following exposure to the test formulations. These were Het-1A (ATCC CRL-2692) human oesophageal cells and NCTC Clone 929 (ATCC CCL 1) murine fibroblast subcutaneous connective tissue cells; areolar and adipose International Standard cell lines. Both cell lines were obtained from LGC Standards (London, UK). NCTC Clone 929 (ATCC CCL 1) cell line was cultured in Minimum Essential Medium (MEM) containing phenol red with Earle's Salts, L-Glutamine, supplemented with 10% Horse Serum, 100iu/mL penicillin and 100µg/mL streptomycin supplied by Invitrogen (Paisley, UK). The Het-1A (ATCC CRL-2692) cell line was cultured in Bronchial Epithelial cell Basal Medium (BEBM) supplemented by Bronchial Epithelial Cell Growth Medium (BEGM) SingleQuot Kit and Growth Factors (Lonza, Basle, Switzerland). Cells were grown at 37°C and 5% CO₂ and subcultured at 80 – 90 % confluency. Subculturing consisted of removal of spent medium, rinsing of the adherent cell surface with sterile autoclaved phosphate buffered saline (PBS) before treatment with a 0.05% Trypsin/0.53mM EDTA·4Na solution (Invitrogen, Paisley, UK) for cell monolayer detachment. For the Het-1A (ATCC CRL-2692) cell line Trypsin/EDTA required inclusion of 0.5% polyvinylpyrrolidone (Sigma-Aldrich, Dorset, UK) and preparation flasks were required to be precoated with 0.01mg/mL fibronectin, 0.03mg/mL bovine collagen type I and 0.01mg/mL bovine serum albumin (all supplied by Invitrogen, Paisley, UK) to facilitate attachment.

Cell viability was assessed by means of a quantitative alamarBlue® assay, using a modification of the method of O'Brien *et al.* (2000). Cells were cultured (until at least third passage) and inoculated into 96-well tissue culture treated microtitre plates at a concentration of 1 x 10⁴ cells/well and incubated at

37°C and 5% CO₂ for 24 ± 1 hour, until approximately 90% confluency as described above. After this time, the medium was removed and replaced with required fresh growth medium, containing doubling dilutions of ML:8 emulsion at final concentrations of 0.25-0.0625% v/v with eight replicates at each concentration. Chlorhexidine digluconate was tested over the same concentration range and acted as a comparative control. Time points selected for analysis of ML:8 emulsion/chlorhexidine digluconate activity on established cell lines were 5, 30 and 60 minutes. Absorption was measured at 570nm in a Tecan Sunrise® plate reader after a development time of 10 hours for NCTC Clone 929 (ATCC CCL 1) cell line and 4 hours for Het-1A (ATCC CRL-2692) cell line. A positive control (100% reduction in viability) was also included in the assay using 90% ethanol (Sigma-Aldrich, Dorset, UK); the negative control consisted of untreated cell line wells percentage cell viability was calculated relative to untreated control wells after subtraction of the blank value corresponding to untreated cells in the absence of alamarBlue® reagent.

$$\% \text{ reduction in viability} = \frac{Abs_{570nm} \text{ ML:8 Emulsion / Chlorhexidine digluconate} - Abs_{570nm} \text{ Blank}}{Abs_{570nm} \text{ ethanol 90\%} - Abs_{570nm} \text{ Blank}} \times 100$$

Statistical Analysis

Statistical analyses were performed using GraphPad InStat 3. Standard deviations were obtained at each concentration/timepoint of antimicrobials tested based on eight replicates for both quantitative biofilm and cell cytotoxicity viability assays and mean values obtained. Further statistical analysis was employed using a one way Analysis of Variance (ANOVA), with a Tukey-Kramer multiple comparisons test used to identify individual differences between the antibiofilm activity of 0.125% v/v ML:8 emulsion and 0.12% v/v chlorhexidine digluconate, and also the antibiofilm activity of 0.125% v/v ML:8 emulsion and 2.4% v/v Virbac Vet Aquadent® (as directed by the manufacturer), at relative timepoints. ANOVA with a Tukey-Kramer multiple comparisons test was also utilised for statistical analysis of cytotoxicity data of ML:8 emulsion and chlorhexidine digluconate at the same concentrations (0.25-0.06125% v/v) and relative timepoints. ANOVA assumes that the data is sampled from populations that follow Gaussian distributions. Data was shown to be normally

distributed using the Kolmogorov and Smirnov method. In all cases a probability of $P \leq 0.05$ denoted significance.

Results

The 0.125% v/v ML:8 emulsion displayed significant activity against biofilm forms of the 10 periodontopathogens investigated within 5 to 10 minutes exposure. Antibiofilm efficacy was significantly greater than the 2.4% v/v Virbac Vet Aquadent® (xylitol containing formulation) and statistically similar to 0.12% v/v chlorhexidine digluconate. Antibiofilm activity for 0.125% v/v ML:8 emulsion, 0.12% v/v chlorhexidine digluconate and xylitol containing formulation are displayed in Figures 1-10. ML:8 emulsion was significantly less cytotoxic than chlorhexidine digluconate at similar concentrations ($P < 0.001$ for all assays). Cytotoxicity of ML:8 emulsion and chlorhexidine digluconate at the same concentrations (0.25-0.06125% v/v) against CCL 1 (NCTC Clone 929) murine fibroblast subcutaneous connective tissue monolayer cells and human oesophageal tissue monolayer cells (ATCC CRL-2692) are shown in Figures 11 and 12, respectively.

Discussion

The results of the current study demonstrate that ML:8 emulsion displayed a high degree of potency against 48 hour biofilm forms of the 10 periodontopathogens investigated. High percentage kill rates ($> 70\%$) were achieved against the majority of test organisms within 5 minutes of exposure, and at all subsequent time points. The selection of biofilm forms of bacteria was purposeful in order to test the ability of the formulation to eradicate this more resistant bacterial phenotype present within the oral cavity (Hojo *et al.* 2009). The majority of previously reported dental-related studies have centred on human plaque and less resistant liquid planktonic forms of bacteria (Stanley *et al.* 1989) (McBain *et al.* 2004), and whilst contributing valuable information, their clinical relevance may be limited in comparison to biofilm-based data. The bacteria selected for assessment in the current study were derived from an extensive literature search for relevant canine and feline periodontopathogens, and as such, have direct relevance to the clinical microbiota encountered in canine/feline periodontal disease

(Syed and Svanberg 1981)(Svanberg *et al.* 1982)(Hennet and Harvey 1991)(Leonhardt *et al.* 1992)(Okuda and Harvey 1992)(Harvey *et al.* 1995)(Harvey 1998)(Harvey 2005).

The antimicrobial activity of free fatty acids has been widely reported previously in the literature (Kabara *et al.* 1972). Research conducted by Sun *et al.* (2002) concluded that caprylic (C₈), capric (C₁₀) and lauric acid (C₁₂) displayed antimicrobial activity with lauric and caprylic acid shown to be most efficacious against Gram-positive and Gram-negative bacteria, respectively. The ML:8 emulsion formulation described here displayed rapid antimicrobial efficacy, showing high potential to be an effective drinking water additive for periodontal disease prevention at low concentrations (0.125% v/v), despite the limited exposure times that can be achieved within the oral cavity. To test this hypothesis further, we compared the anti-biofilm activity of the ML:8 emulsion with the gold standard in human/veterinary dental hygiene (0.12% chlorhexidine digluconate) and another commercially available veterinary dental formulation containing 0.5% xylitol (prediluted). With the exception of the 0 minute timepoints for *Porphyromonas gingivalis* (VPB 5089), *Eikenella corrodens* (VPB 3935) and *Tannerella forsythensis* (VPB 4947), there was no significant difference in the antibiofilm activity of 0.125% v/v ML:8 emulsion and 0.12% v/v chlorhexidine digluconate (P>0.05). However, although chlorhexidine has been a mainstay in the control and treatment of dental pathogens in human health (Roberts *et al.* 2002), similar effective concentrations cannot be employed in canine drinking water formulations as the ingestion/swallowing of solutions containing 0.12% v/v chlorhexidine on a daily basis is likely to be associated with significant cytotoxicity, as demonstrated by the results generated in this study (*see later*). At concentrations significantly below the traditional 0.12% v/v threshold employed in human products, the range of chlorhexidine-based drinking water additives available on the veterinary market have debatable *in vitro* and *in vivo* efficacy (Roudebush *et al.* 2005).

Virbac Vet Aquadent® contains xylitol and <0.05% chlorhexidine digluconate (the chlorhexidine component is not included as an active ingredient). Our results show Aquadent® to have limited efficacy against biofilm forms of periodontal bacteria tested up to 24 hours under the conditions of this assay. At the 24 hour timepoint, the reduction in viable biofilm reached a maximum of 53% against

Eikenella corrodens (VPB 3935). The majority of bacteria/timepoints studied showed a mean reduction of biofilm of less than 10%. Overall, the 0.125% v/v ML:8 emulsion showed a statistically significant increase in biofilm reduction when compared with Aquadent® in 93.6% of the comparative time points/bacteria tested (103 out of 110 sample points; $P < 0.001$). Although issues have been raised in relation to the potential toxic effects of xylitol ingestion in dogs (Murphy *et al.* 2012), the dose levels employed in Aquadent® have not been associated with any reported toxic effects in the published literature.

Figures 11 and 12 show that at therapeutic concentrations (0.12% v/v) chlorhexidine digluconate demonstrated toxicity against mammalian cell lines; therefore, its long-term use or suitability as a drinking water additive may be limited by potential gastrointestinal and oral mucosal damage. At the same concentrations and correlating to the same exposure times and cell lines, the toxicity demonstrated by ML:8 emulsion was significantly lower ($P < 0.001$) than for chlorhexidine digluconate. After up to 60 minutes exposure to varying concentrations of ML:8 emulsion (0.25 to 0.0625% v/v), the relative percentage kill of CCL 1 (NCTC Clone 929) murine fibroblast cells was absent (0%). The results obtained for chlorhexidine digluconate against both human oesophageal tissue (ATCC CRL-2692) and the International cytotoxicity standard CCL 1 (NCTC Clone 929) murine fibroblast cells showed a statistically significant increase ($P < 0.001$) in the cytotoxic activity of chlorhexidine digluconate relative to ML:8 at all time points studied. Quantitative evaluations such as the alamarBlue® assay determine that a reduction of cell viability by more than 30% is indicative of cytotoxicity (International Standard ISO10993-5). Selection of the International cytotoxicity standard CCL 1 (NCTC Clone 929) murine fibroblast cell line allowed this novel formulation to be assessed for cytotoxicity in general, whilst selection of a mammalian oesophageal cell line also allowed the toxicity of ML:8 to be compared to a clinically relevant cell line.

Conclusions

The formulation and 1 in 51 dilution of this novel 6.375% v/v ML:8 emulsion to drinking water allows the active free fatty acids to be present at an antimicrobially active and non-cytotoxic 0.125 %

v/v final concentration. Such a product can increase compliance and ease of use allowing daily administration to help prevent periodontal disease, with superior *in vitro* results compared with a currently available xylitol-based drinking water additive. The findings of this study validate the use of the ML:8 emulsion as part of an ever increasing evidence-based approach to biofilm control in veterinary dental applications (Hamp and others 1973). Although *in vivo* clinical trials are now indicated to corroborate these findings, our initial laboratory results show large scope and promise for continuing research in this area.

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Figure/Legends

Fig. 1. Mean percentage reduction in 48 hour biofilm of *Haemophilus actinomycetemcomitans* (NCTC 10979) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v

Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®,
 ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, ***: P<0.001, significant difference
 between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or
 2.4%v/v Virbac Vet Aquadent® at same timepoint.

Fig. 2. Mean percentage reduction in 48 hour biofilm of *Streptococcus sanguinis* (NCTC 10904) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 3. Mean percentage reduction in 48 hour biofilm of *Porphyromonas cangingivalis* (VPB 4874) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 4. Mean percentage reduction in 48 hour biofilm of *Porphyromonas salivosa* (VPB 3313) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 5. Mean percentage reduction in 48 hour biofilm of *Porphyromonas gingivalis* (VPB 5089) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 6. Mean percentage reduction in 48 hour biofilm of *Fusobacterium nucleatum* (VPB 4888) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 7. Mean percentage reduction in 48 hour biofilm of *Eikenella corrodens* (VPB 3935) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 8. Mean percentage reduction in 48 hour biofilm of *Bacteroides fragilis* (VPB 3371) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 9. Mean percentage reduction in 48 hour biofilm of *Prevotella intermedia* (VPB 3321) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 10. Mean percentage reduction in 48 hour biofilm of *Tanerrella forsythesis* (VPB 4947) over a period of 24 hour exposure to food grade 0.125%v/v ML:8 Emulsion, 0.12%v/v Chlorhexidine digluconate and 2.4%v/v Virbac Vet Aquadent® utilising an alamarBlue® assay. Results are displayed as the mean of 8 replicates. Key: ■: 0.12%v/v Chlorhexidine digluconate, ▲: 0.125%v/v ML:8 Emulsion, ●: 2.4%v/v Virbac Vet Aquadent®, ns: no significant difference (P>0.05) *: P<0.05, **: P<0.01, *: P<0.001, significant difference between efficacy of 0.125%v/v ML:8 Emulsion and 0.12%v/v Chlorhexidine digluconate or 2.4%v/v Virbac Vet Aquadent® at same timepoint.**

Fig. 11. The percentage kill of CCL 1 [NCTC clone 929]- murine fibroblasts subcutaneous connective tissue monolayer cells after 5, 30 and 60 minutes exposure to varying concentrations of ML:8 emulsion and Chlorhexidine digluconate (CHX). Results are obtained via the use of an alamarBlue® assay (10 hour development time). Key:

◆ 0.25% v/v ML:8	■ 0.125% v/v ML:8	▲ 0.0625% v/v ML:8
✕ 0.25% v/v CHX	— 0.125% v/v CHX	● 0.0625% v/v CHX

ns: no significant difference ($P > 0.05$), *: $P < 0.05$, **: $P < 0.01$, ***: $P < 0.001$, significant difference between cytotoxicity of ML:8 Emulsion and Chlorhexidine digluconate at same time point and concentration.

Fig. 12. The percentage kill of ATCC CRL-2692- human oesophageal tissue monolayer cells after 5, 30 and 60 minutes exposure to varying concentrations of ML:8 emulsion and Chlorhexidine digluconate (CHX). Results are obtained via the use of an alamarBlue® assay (10 hour development time). Key:

◆ 0.25% v/v ML:8 ■ 0.125% v/v ML:8 ▲ 0.0625% v/v ML:8
 ✕ 0.25% v/v CHX — 0.125% v/v CHX ● 0.0625% v/v CHX

ns: no significant difference ($P > 0.05$), *: $P < 0.05$, **: $P < 0.01$, ***: $P < 0.001$, significant difference between cytotoxicity of ML:8 Emulsion and Chlorhexidine digluconate at same time point and concentration.