INTRODUCTION

Several antimicrobial chemicals, incorporated into mouthwash, can be beneficial in the prevention of dental caries. Supplementation of mechanical brushing with effective antimicrobial mouthwash has proven beneficial in the control of plaque (Kornman, 1986a; Mandel, 1988). The greatest success has been with chlorhexidine, which is now considered the gold standard against which the other potential antiplaque agents are measured (Kornmann, 1986b; Mandel, 1988; Hearsman and Seymour, 1955). However, side effects may occur such as discoloration of the teeth or unpleasant aftertaste may occur when these chemicals are used for an extended period (Greenstein et al., 1986). At present, there are large numbers of mouthrinses containing antimicrobial agents available to the public but these rarely reach the rural areas. Many studies on the effect of these mouthrinses or toothpastes on salivary microbiology, plaque and gingivitis have been done (Collaert et al., 1992; Rosin et al., 2001a,b; Meurman et al., 2002; Rosin et al., 2002).

Several medicinal plants have been recently found to have antimicrobial properties. Extracts of these plants have been added to some toothpastes and mouthwashes to control dental caries and have been extensively surveyed (Fine et al., 2000; Khalesi et al., 2004; Matsumoto et al., 2004; Weiss et al., 2004). Albizia myriophylla Benth (commonly known as Cha-em Thai in Thailand) is used as a substitute for licorice due to its sweet taste. This plant is widely distributed in southeastern Asian countries, such as Thailand. Its water extract has antibacterial activity against Streptococcus mutans ATCC 25175 (Cholticha A, unpublished data). Cha-em Thai mouthwash was prepared to use by people in rural areas. The aim of this study was to determine the efficiency of mouthwash containing water extract of Cha-em Thai on mutans streptococci counts and total IgA in saliva.

MATERIALS AND METHODS

**Albizia myriophylla** (Cha-em Thai) extract

The stems of Cha-em Thai were locally col-
lected in Kanchanaburi Province, Thailand. The stems were washed, cut into small pieces and allowed to dry at room temperature for one week. It was then ground into powder at a ball mill. One hundred grams of powder was soaked in 300 ml of distilled water for 30 minutes and then boiled for 30 minutes. After cooling, the liquid extract was passed through filter paper (0.45 µm pore size) and then adjusted to a volume of 200 ml with deionized water to obtain a 50% w/v water extract. The 50% w/v extract was stored at 4°C before preparing the mouthwash.

Mouthwash preparation

From the 50% water extract of Cha-em Thai, 40% herbal mouthwash was prepared by adding menthol, glycerine, sodium benzoate and deionized water. A placebo mouthwash was also prepared similarly except there was no Cha-em Thai extract present.

Experimental design and subject selection

Sixty-seven schoolchildren, age 6-12 years, in Kanchanaburi Province with salivary MS counts ≥1x10^5 cfu/ml saliva were entered in this study. They were divided into 2 balanced groups according to their baseline levels of salivary MS counts (≥10^5 cfu/ml). Each group was randomly assigned to use either Cha-em Thai mouthwash or placebo mouthwash. The subjects were asked to rinse with 10 ml of their assigned mouthwash for 1 minute twice daily for 2 weeks. At baseline and after 2 weeks mouthrinsing, stimulated saliva were collected and analysed for MS counts and total IgA.

The determination of salivary levels of MS and total IgA

Paraffin wax stimulated saliva was collected from subjects in sterile cups. One milliliter of saliva was pipetted into a small sterile vial and kept in dry ice until sent to the microbiological laboratory to determine the total IgA levels. They were stored at -20°C prior to assay for total IgA.

The salivary levels of MS were obtained immediately after saliva sample collection using a modified micromethod of Westergren and Grasse (1978). MSB agar was used as selective media for MS counts. Twenty-five microliters of 10^-1 and 10^-2 dilutions of saliva were spotted onto MSB agar and incubating for 2 days at 37°C. MS colonies were counted and expressed as cfu/ml saliva.

The total IgA levels in saliva were measure by solid-phase ELISA, performed in 96-well flat-bottomed plates (maxisorp micro-titer plates Nune, Roskilde, Denmark). Saliva samples were centrifuged at 10,000g at 4°C for 10 minutes. The clear supernatants were collected and frozen at -20°C until used. The Maxisorp micro-titer plate was coated with monoclonal antibody against human secretory component (IgA). One hundred microliters of saliva samples were applied to individual wells and incubated for 1.5 hours at 37°C. After washing, 100 µl of biotin-conjugated anti-human IgA antibody was added and incubated for 1 hour at 37°C, then the plate was rewashed thoroughly. One hundred microliters per well of avidin-peroxidase conjugate was then added and incubated at 37°C for another hour. The unbound enzyme conjugate was removed by washing with washing buffer. The antigen-antibody complex was visualized by adding 100 µl/well of substrate (3, 3’, 5, 5’-tetramethylbenzidine – TMB) and incubated in the dark at room temperature for 30-45 minutes to allow the color to develop vividly. The enzymatic reaction was finally terminated by the addition of 50 µl of 0.5M H₂SO₄. The color was measured at a wavelength of 450 nm using a micro-titer plate spectrophotometer (BIO-TEX MQX 200, BIO-TEX Instruments, Inc, Highland Park, Vermont, USA). The quantity of total IgA was expressed in milligrams/milliliter of saliva (mg/ml).

Statistical analysis

A log transformation of MS counts (cfu/ml saliva) was carried out. Within group comparison of MS counts and IgA in saliva were calculated by Wilcoxon signed rank test and paired t-test respectively.

RESULTS

Sixty-seven schoolchildren with a mean MS count of ≥1x10^5 cfu/ml saliva or ≥5.0 log cfu/ml saliva, were divided into 2 groups: Cha-em Thai mouthwash and placebo groups. The Cha-em Thai mouthwash group had a mean MS count of 5.636 log cfu/ml and a mean salivary IgA of
0.372 mg/ml at baseline, while the placebo group had a mean MS count of 5.627 log cfu/ml and a mean salivary IgA of 0.351 mg/ml at baseline. Comparison among the groups for MS counts and IgA in saliva before and after Cha-em Thai mouthwash is presented in Table 1. A significant reduction in MS count \( (p<0.05) \) was found, but no significant difference in IgA levels was seen \( (p>0.05) \).

The comparison among the placebo group of salivary MS counts and IgA levels is shown in Table 2. After 2 weeks of placebo mouthwash, no significant differences were found for either the MS counts or the IgA levels in the saliva \( (p>0.05) \).

**DISCUSSION**

This study was designed to evaluate the short-term effect of 40% Cha-em Thai mouthwash compared to placebo control. The 40% aqueous Cha-em Thai solution showed a significant reduction in mean salivary MS counts after 2 weeks use but did not affect the total salivary IgA. In Thai folk medicine, Cha-em Thai is used as licorice due to its sweet taste. Yoshikawa et al (2002) isolated five new triterpene saponins from the stem of Cha-em Thai collected in Kanchanaburi Province, Thailand. This study also used the stem of Cha-em Thai collected from the same province to prepare 40% aqueous mouthwash and was studied in the same Province to investigate the efficiency of this mouthwash in schoolchildren. Aqueous Cha-em Thai mouthwash significantly reduced the amount of mutans streptococci from 5.6 log cfu/ml saliva to 3.7 after 2 weeks \( (p<0.05) \), which was more effective than placebo. This observation confirms that Cha-em Thai mouthwash can be used to lower salivary MS in schoolchildren. The levels of total IgA in saliva for both groups were not affected after 2 weeks of use. Koga-Ito et al (2004) reported a correlation among mutan streptococci counts, dental caries and IgA in saliva, but no correlation between mutans streptococci counts and anti-streptococcus mutants IgA levels were found. Camling et al (1987) showed that IgA to Streptococcus mutans did not reflect the quantity of mutans streptococci at the moment of saliva collection. These study results are similar to those studies. Total IgA levels in saliva were
nearly the same, even if the mutans streptococci levels were significantly different (p<0.05). Our results show that total IgA levels in saliva do not reflect the salivary mutans streptococci concentration and Cha-em Thai mouthwash does not alter salivary levels of total IgA.

Due to the sweet taste of Cha-em Thai, Yoshikawa et al (2002) tested the sweetness relative to sucrose by a human sensory panel. They found that Albiziasaponin A and B have a potent sweetness intensity relative to sucrose (600 times). Our aqueous Cha-em Thai mouthwash preparation has a good taste compared to chlorhexidine. The effectiveness in lowering mutans streptococci suggests that Cha-em Thai mouthwash can be used short-term to reduce salivary levels of mutans streptococci in high caries-risk schoolchildren.

REFERENCES