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Anti-Microbial Breath Freshening Mouthrinse**Field of The Invention**

5 This invention relates to an anti-microbial breath freshening mouthrinse that contains cetylpyridinium chloride (abbreviated herein to "CPC") and zinc ion. More particularly the invention relates to a mouthrinse that contains CPC and zinc ion and an ionone terpene ketone.

10 **BACKGROUND OF THE INVENTION**

It has been known that a mouthrinse composition that contains zinc ions and alpha or beta ionone terpene is effective to reduce mouth odor in humans. This is discussed in U.S. Patent 4,689,214. The zinc ion is provided by a non-toxic organic zinc or inorganic
15 zinc compound. These compositions are effective in counteracting breath odors. However, it has been found that mouthrinse compositions that contain CPC and zinc ions, and in particular zinc acetate have an improved affect in providing fresher breath. This composition can also contain alpha or beta ionone terpenes for additional counteracting of breath odors. However, it is a synergistic anti-microbial effect of CPC
20 and zinc ions that is most effective in providing fresher breath.

Brief Summary of The Invention

A new mouthrinse containing cetylpridinium and zinc salt (CPC/Zn) significantly reduced mouthodor scores caused by volatile sulfur compounds in clinical studies versus
25 a control rinse. It also can contain an ionone terpene. This investigation examined the antimicrobial effects of these rinses in laboratory and clinical studies. Laboratory tests were performed with oral bacteria with alamar blue, a redox dye that rapidly detects bacterial viability. A Treatment of *A. viscosus*, *S. sanguis*, and *S. mutans* by CPC/Zn resulted in a 25, 35 and 16% decrease in bacterial viability respectively versus a control
30 In one study a randomized, double blind clinical study with volunteer subjects was conducted to confirm these laboratory results. After a washout phase, the subjects abstained from oral hygiene and provided a saliva sample for determining the total and



oral odorigenic producing bacteria on enriched and indicator plates. Mouthrinses were used once and the subjects sampled at 90 and 180 minutes post treatment for total and oral odorigenic bacteria. In a comparison to a control, the decrease of total bacteria following CPC/Zn use at 90 and 180 minutes were 59.9% and 32.1% respectively. The use of CPC/Zn resulted in 61.7% and 72.5% decrease of odorigenic bacteria at 90 and 180 minutes respectively versus a control rinse. The decrease in bacteria with a CPC/Zn rinse was significantly higher than the control rinse at all post-use time points.

Detailed Description of The Invention

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There are various strains of bacteria that are present in the mouth. Many of these aid in the production of volatile sulfur compounds that cause mouth odor. These include the following bacteria:

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Actinobacillus

Actinomycetemitans

Campylobacter recta

Capnocytophaga sp.

Eikenella corrodens

Fusobacterium nucleatum

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Porphyromonas gingivalis

Prevotella intermedia

Actinomyces naeslundii

Actinomyces viscosus

Actinomyces mutans

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Streptococcus mutans

Streptococcus sanguis

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For a mouthrinse the CPC will be in the mouthrinse composition in an amount of about 0.0005 percent to about 3 percent by weight, and preferably about 0.01 percent to about 1 percent by weight. The primary component of such a mouthwash usually will be water.

In accordance with a preferred embodiment, this invention relates to an oral composition consisting essentially of a non-toxic zinc compound, usually with salt, in an amount which provides at least about 0.01 mg of zinc ions in 1 ml of water. The ratio of zinc ions to the CPC being about 100:1 to 2:1 by weight, and preferably about 50:1 to about 4:1 by weight.

The zinc compounds that provide zinc ions for use in combination with the CPC may be any physiologically acceptable zinc compound including water soluble (inclusive of sparingly water soluble) organic and inorganic zinc compounds. The water-soluble zinc compounds (at least 1% soluble) are preferred. Examples of suitable zinc compounds that may be employed include:

zinc acetate	zinc fluoride
zinc ammonium sulfate	zinc formate
zinc bromide	zinc iodide
zinc chloride	zinc nitrate
zinc chromate	zinc phenol sulfonate
zinc citrate	zinc salicylate
zinc dithionate	zinc sulfate
zinc fluosilicate	zinc gluconate
zinc tartarate	zinc succinate
zinc glycerophosphate	

Other zinc compounds disclosed in U.S. Patent No. 4,138,477 having a solubility of at least about 0.01 mg of zinc ions per ml water are incorporated by reference.

The zinc compounds is present in amounts which provides about 0.01-5% by weight of zinc ions and preferably about 0.02-1% of zinc ions by weight in the oral composition.

The mouthrinse composition preferably also will contain an alpha or beta ionone terpene. Ionone is a ketone terpene derivative containing one ketonic carbonyl group. The basic ionone formula is $(\text{CH}_3)_3 (\text{C}_6\text{H}_5)_2 (\text{CH CO CH}_3)_3$. It is available as alpha-ionone (b.p. 120 degree. C.) and beta-ionone (b.p. 135.degree.), both of which are colorless liquids and slightly soluble in water. It is employed in oral compositions in amounts such that the ratio of zinc ions to ionone is about 1,000: 1 to 10 :1 by weight. Other variants of ionone such as gamma-ionone dihydroionone and alphas-methyl ionone may also be employed. These include isomeric forms of ionones, e.g. irone. It is convenient to employ it in oral compositions in amounts of about 0.0005-1% by weight, preferably about 0.001-1%.

10 Alpha-ionone is the preferred mouthrinse ionone variant.

Any suitable flavoring or sweetening materials may be employed in the mouthrinse or other composition. The presence of flavoring oil improves the taste of the zinc-containing product. Examples of suitable supplemental flavoring oils include oils of spearmint, peppermint, wintergreen, sassafras, clove, sage eucalyptus, marjoram, cinnamon, lemon and orange, as well as methylsalicylate. Mint oils such as oil of peppermint is most preferred. Suitable sweetening agents include sucrose, lactose, maltose, sorbitol, sodium cyclamate, saccharine, acetosulfam, N-1 alpha.-aspartyl-1-phenylalanine-methyl ester ("aspartame"), xylitol, chalcone materials. Suitably, flavor and sweetening agents may together comprise from about 0.01 to 5% by weight or more of the compositions of the instant invention, each typically being about 0.005-2.5%.

A fluorine-containing compound having a beneficial effect on the care and hygiene of the oral cavity, e.g., diminution of enamel solubility in acid and protection of the teeth against decay may also be incorporated in the composition. Examples thereof include sodium fluoride, potassium fluoride, potassium stannous fluoride ($\text{SnF}_2\text{-KF}$), sodium hexafluorostannous, stannous chlorofluoride, sodium fluorozirconate, and sodium monofluorophosphate. These materials, which disassociate or release fluorine-containing ions in water, suitably may be present in an effective but non-toxic amount, usually within the range of about 0.01 to 1% by weight of the water-soluble fluorine content thereof.

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When the oral preparation is a liquid such as mouth rinse or mouth spray which typically contains 20-99% by weight of an aqueous vehicle comprising non-toxic lower aliphatic, preferably having about 1-30% by weight alcohol such as ethanol, n-propyl, or isopropyl alcohol with the remainder water. Flavor and/or sweetener and surface active agent are
5 also generally present. Mouth sprays also contain a suitable amount of an orally acceptable propellant materials such as a fluorocarbon, e.g. Freon or isobutane to propel the spray from a pressurized container. The CPC/Zn/ionone can also be delivered in dentifrices, gums, lozenges, breathe strips or any other technique suitable to deliver the material into the mouth.

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The oral presentations should have a PH practicable for use. The pH range of about 4-9, preferably about 5-7.5, is considered the most practicable for use.

The following examples further illustrate preferred embodiments of the invention.

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Example 1

Two mouthrinse formulations were formulated. Mouthrinse A contains CPC and mouthrinse B no CPC. Each contains zinc acetate and alpha ionone. The formulations are as set out in Table 1.

Table 1
Mouthrinse Compositions (%)

Ingredients	<u>Mouthrinse A</u>	<u>Mouthrinse B</u>
	<u>CPC/Zinc Acetate/Alpha ionone</u>	<u>Zinc Acetate/Alpha ionone</u>
5 Ethyl Alcohol	10.00	10.00
Cetylpyridinium Chloride	0.05	0.00
Zinc Acetate Dihydrate	0.40	0.40
Alpha Ionone	0.022	0.022
10 Flavor	0.198	0.198
Glycerine	7.50	7.50
Sorbitol	7.50	7.50
Sodium Saccharin	0.030	0.030
Benzoic Acid	0.05	0.05
15 PEG 40 Sorbitan Diisostearate	1.00	2.00
Hydrochloric Acid	0.08	0.08
Water (Deionized/Purified)	73.1695	72.2195
FD&C Blue No. 1	0.0005	0.0005

- 20 The antibacterial effect of mouthrinses A and B was evaluated against oral bacteria by measuring maximum inhibitory dilution (MID) of the mouthrinse that inhibit bacterial growth. The results, summarized in Table 2, showed that mouthrinse A has higher MID values than mouthrinse B against representative oral bacteria which are implicated in the production of mouth malodor, plaque and gingivitis. These results clearly showed that
- 25 addition of CPC to Zn acetate enhanced the antibacterial efficacy when compared to Zn acetate and alpha ionone.

- Table 2 shows that addition of CPC to zinc acetate and alpha ionone provided improved antibacterial efficacy of the combination compared to zinc acetate and alpha ionone. This
- 30 is an enhanced destruction of mouth bacteria in the combination of CPC and zinc. Mouthrinse A is effective in killing odorgenic bacteria at much lower concentrations is seen in the MID values.

Table 2

Maximum Inhibitory Dilution

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	Bacterial Strain	<u>Mouthrinse A</u>	<u>Mouthrinse B</u>
		<u>CPC/Zinc Acetate/Alpha ionone</u>	<u>Zinc Acetate/Alpha ionone</u>
	Actinobacillus-		
	Actinomycetemcomitans	107	64
10	Campylobacter recta	128	24
	Capnocytophaga sp.	43	13
	Eikenella corrodens	128	16
	Fusobacterium nucleatum	128	96
	Porphyromonas gingivalis	256	128
15	Prevotella intermedia	427	235
	Actinomyces naeslundii	96	8
	Actinomyces viscosus	91	8
	Streptococcus mutans	192	6
	Streptococcus sanguis	64	24

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Example 2

The mouthrinses of the Table 1 also were tested clinically on volunteer subjects. Mouth odor reduction efficacy of the rinses was evaluated in two clinical studies conducted in humans. Mouth odor of the subjects was scored by human judges. The subjects rinsed in the morning after a baseline reading and then were evaluated again at 4 hours and 8 hours. The subjects rinsed again before bed and evaluated again in the next morning for the overnight scores.

subjects according to malodor scores 0 through 5. 0 is equal to no odor, 1 is equal to questionable odor, 2 is equal to faint odor, 3 is equal to moderate odor, 4 is equal to strong odor and 5 is equal to very strong odor. Results of the clinical studies are summarized in Table 1. Mouthrinse B containing Zn acetate and (X-ionone significantly

reduced mouth odor at 4 hours, 8 hours and overnight compared to baseline value.

However, mouth odor level at each time point was above moderate odor and close to strong odor. Mouthrinse A containing CPC, Zn acetate and (alpha ionone also

significantly reduced mouth odor at 4 hours, 8 hours, and overnight compared to

5 baseline. However, in the case if mouthrinse A, mouth odor level at each time point was

lower than mouthrinse B and it was in the range of the faint odor to moderate odor. The

effect of mouthrinse A and B on mouth odor was also expressed in terms of percent

reduction from baseline and shown as (%) next to each time point in Table 1. Results

clearly showed that mouthrinse B provided higher percent reduction than mouthrinse A at

10 each time point.

Table 3

Study#	Treatment	No. of	<u>Mouth Odor Scores</u>				<u>Significance</u>
	<u>Rinse</u>	<u>Subjects</u>	<u>Baseline</u>	<u>4 hours</u>	<u>8 hours</u>	<u>overnight</u>	
A	Zn Acetate/ionone	44	4.19	3.11 (26%)	3.27% (22%)	3.64 (13%)	p<0.01
B	CPC/Zn acetate/ionone	47	3.96	2.51 (37%)	2.89% (27%)	3.06 (22%)	p<0.01

Example 3

A randomized 2 cell double blind clinical study was conducted on 17 volunteer subjects using a placebo mouthrinse and a mouthrinse containing active ingredients 0.05% CPC, 0.4% zinc acetate and 0.022% alpha ionone. The study was directed to a reduction in volatile sulfur compounds (VCS).

The subjects had a 5 day washout period before the study and between the use of each mouthrinse. An initial baseline evaluation then was conducted. Each volunteer then rinsed their mouths with 15 mls of the assigned mouthrinse for 30 seconds. They returned in three hours for a post-treatment evaluation. Then that night prior to bed time each rinsed again with the same amount of the assigned mouthrinse for the same period of time. Each of the volunteer subjects was evaluated the following morning. The evaluations consisted of each subject keeping their mouths closed and breathing only through their nostrils for 10 minutes. A Teflon tube then was inserted through the subjects lips and two samples of breaths taken from the mouths by means of a pump. The breath samples were sent to a gas chromatograph and analyzed for the volatile sulfur compounds hydrogen sulfide, methyl mercaptan and dimethyl sulfide. These are components of breath malodor.

The three hour breath freshening results with regards to the presence of volatile sulfur compounds is given in Table 3 with overnight results given in Table 4.

Table 3

Treatment Rinse	<u>N</u>	Initial <u>Mean</u> (ng/ml)	Final <u>Mean</u> (ng/ml)	<u>% Reduction</u>
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CPC/Zn/ionone	17	15.31 (± 3.92)	5.69 (± 2.62)	62.8% (p<0.05)
Placebo (T) —	17	14.42 (± 3.15)	10.32 (± 4.18)	28.4 (p<0.05)

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Table 4

	Treatment	N	Initial	Final	% Reduction
10	<u>Rinse</u>		<u>Mean</u> (ng/ml)	<u>Mean</u> (ng/ml)	

CPC/Zn/ionone	17	15.31 (± 3.92)	7.38 (± 2.71)	51.8% (p<0.05)
Placebo (T)	17	14.42 (± 3.15)	13.17 (± 4.02)	8.67% (p>0.05)

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The 3 hour and overnight results for the CPC/Zn/alpha ionone mouthrinse shows a significant improvement in the redirection of volatile sulfur compounds which are a primary component in mouth odor when zinc ions are present. The zinc ions have a synergistic affect in the reduction of mouth odor.

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Example 4

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A clinical study was conducted on volunteer subjects for the decrease in oral bacteria with mouthwash that contained zinc ions CPC and alpha ionone. The concentrations were the same as in Example 3. The volunteer subjects after a 7 day washout period using a Colgate toothpaste with fluoride. The volunteer subjects then gargled 15 ml of mouthwash for 30 seconds. Each then was tested at 90 minutes and 180 minutes for the

presence of *A. viscosus*, *S. sanguis*, and *S. mutans*. Each volunteer subject refrained from eating and drinking for the three hour period of the test. Saliva samples were taken from each volunteer subject and plated on ? media (to enumerate total salivary bacteria) and on indicator plates to enumerate odorigenic hydrogen sulfide producing bacteria.

- 5 The analysis for the above bacteria was conducted using the Alamar blue fluorescence Method. The results as a percent decrease in bacteria is given in Table 5.

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Table 5

Parameter	90 Minutes	180 Minutes
15 Salivary Bacteria	59.9	32.1
Odorigenic Bacteria	61.7	72.5

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The decrease is more evident at 90 minutes with some bacterial regenerations at 180 minutes.

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CLAIMS

We Claim:

1. An oral synergistic volatile sulfur compound breath-odor controlling composition comprising a non-toxic zinc compound and cetylpyridinium chloride, the ratio of zinc ions to cetylpyridinium chloride being about 1000:1 to 2:1 by weight, and an ionone ketone terpene, said zinc compound being present in the amount of about 0.01 to 5% by weight, said cetylpyridinium chloride being present in amount of 0.0005 to 3% by weight and said ionone ketone being present in an amount of 0.0005% to 2% by weight.
2. An oral composition as in claim 1 wherein said zinc compound is present in amount of 0.02 to 2% by weight of zinc compound and cetylpyridinium chloride is present in amounts of about 0.001 to 1% by weight.
3. An oral composition as in claim 2 wherein said zinc compound is selected from the group consisting of zinc chloride and zinc acetate.
4. An oral composition as in claim 1 wherein said zinc compound is present in an amount of 0.02 to 2% by weight, cetylpyridinium chloride is present in an amount of 0.001 to 1 % by weight and said ionone is present in an amount of 0.001 to 1% by weight.
5. An oral composition as in claim 4 wherein said ionone terpene ketone is selected from the group consisting of alpha-ionone, beta-ionone, gamma-ionone, dihydroionone, alpha-methylionone and ionone.
6. An oral composition as claimed in claim 5 wherein said ionone ketone terpene is alpha-ionone.
7. An oral composition as claimed in claim 4 wherein said zinc compound is selected from the group consisting of zinc chloride and zinc acetate.
8. An oral composition as claimed in claim 7 wherein said compound salt is zinc chloride.

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9. An oral composition as claimed in claim 1 wherein said composition comprises a liquid portion comprising water and humectant and a solid portion comprising a gelling agent and said oral composition is a dentifrice.
10. An oral composition as in claim 1 wherein said ionone ketone terpene derivative is present in a component of a flavoring oil in amounts of about 0.5 to 1% by weight of said flavoring oil.
11. An oral composition as in claim 10 wherein said flavoring oil is oil of peppermint which oil of peppermint is present in said oral composition in amounts of about 0.005 to 2% by weight.
12. An oral composition as in claim 1 wherein said composition comprises an aqueous alcohol composition wherein said alcohol is a non-toxic lower aliphatic alcohol and said oral preparation is a mouthrinse or mouthspray.
13. A method of synergistically countering volatile sulfur compound breath odor which comprises dispersing in the oral cavity an oral preparation comprising a zinc compound which provides 0.01 to 5% by weight of a zinc ion and 0.0005 to 1% by weight of cetylpyridinium chloride the ratio of zinc ions to ionone ketone terpene being 1000:1 to 2:1 by weight.
14. A method in claim 11 wherein said ionone ketone derivative is alpha-ionone and said zinc compound is zinc chloride.