



(86) Date de dépôt PCT/PCT Filing Date: 1998/09/05
(87) Date publication PCT/PCT Publication Date: 1999/03/25
(45) Date de délivrance/Issue Date: 2009/01/20
(85) Entrée phase nationale/National Entry: 2000/03/14
(86) N° demande PCT/PCT Application No.: EP 1998/005632
(87) N° publication PCT/PCT Publication No.: 1999/013851
(30) Priorité/Priority: 1997/09/15 (DE197 40 453.7)

(51) Cl.Int./Int.Cl. *A61K 8/26* (2006.01),
A61K 8/24 (2006.01), *A61K 8/25* (2006.01),
A61Q 11/00 (2006.01)
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(54) Titre : AGENT NETTOYANT A USAGE DENTAIRE COMPRENANT UNE COMBINAISON D'AGENTS POLISSANTS
A BASE D'ACIDE SILICIQUE ET D'OXYDE D'ALUMINIUM
(54) Title: CLEANING AGENT FOR DENTAL USE COMPRISING A COMBINATION OF POLISHING AGENTS BASED
ON A SILICIC ACID AND ALUMINIUM OXIDE

(57) **Abrégé/Abstract:**

The invention concerns a cleaning agent for dental use in the form of a pasty or liquid dispersion, containing between 10 and 30 wt. % of a combination of polishing agents comprising polishing agents based on silicic acid and aluminium oxide in a weight ratio of 10 : (0.2 - 2) and between 20 and 50 wt. % of a moistening agent from the group containing sorbitol, glycerin, propyleneglycol-1,2, polyethyleneglycol and their mixtures, comprising, for reinforcing the cleaning effect, between 2 and 12 wt. % of condensed phosphate from the group comprising tripolyphosphate, pyrophosphate or trimetaphosphate in the form of alkaline salt or ammonium salt. In spite of their low abrasive values, said cleaning agents have an excellent polishing effect and a high cleaning power.



**PCT**WELTORGANISATION FÜR GEISTIGES EIGENTUM
Internationales BüroINTERNATIONALE ANMELDUNG VERÖFFENTLICHT NACH DEM VERTRAG ÜBER DIE
INTERNATIONALE ZUSAMMENARBEIT AUF DEM GEBIET DES PATENTWESENS (PCT)

(51) Internationale Patentklassifikation ⁶ : A61K 7/16	A1	(11) Internationale Veröffentlichungsnummer: WO 99/13851 (43) Internationales Veröffentlichungsdatum: 25. März 1999 (25.03.99)
(21) Internationales Aktenzeichen: PCT/EP98/05632 (22) Internationales Anmeldedatum: 5. September 1998 (05.09.98) (30) Prioritätsdaten: 197 40 453.7 15. September 1997 (15.09.97) DE (71) Anmelder (für alle Bestimmungsstaaten ausser US): HENKEL KOMMANDITGESELLSCHAFT AUF AKTIEN [DE/DE]; Henkelstrasse 67, D-40589 Düsseldorf (DE). (72) Erfinder; und (75) Erfinder/Anmelder (nur für US): LEINEN, Hans-Theo [DE/DE]; Gertrudisstrasse 2, D-40229 Düsseldorf (DE). WÜLK NITZ, Peter [DE/DE]; Im Erlengrund 9, D-42799 Leichlingen (DE).		(81) Bestimmungsstaaten: AU, CA, CN, CZ, HU, JP, KR, NO, PL, SK, US, europäisches Patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Veröffentlicht <i>Mit internationalem Recherchenbericht. Vor Ablauf der für Änderungen der Ansprüche zugelassenen Frist; Veröffentlichung wird wiederholt falls Änderungen eintreffen.</i>
(54) Title: CLEANING AGENT FOR DENTAL USE COMPRISING A COMBINATION OF POLISHING AGENTS BASED ON A SILICIC ACID AND ALUMINIUM OXIDE (54) Bezeichnung: ZAHNREINIGUNGSMITTEL MIT EINER POLIERMITTELKOMBINATION AUS KIESELSÄURE-POLIERMITTELN UND ALUMINIUMOXID (57) Abstract The invention concerns a cleaning agent for dental use in the form of a pasty or liquid dispersion, containing between 10 and 30 wt. % of a combination of polishing agents comprising polishing agents based on silicic acid and aluminium oxide in a weight ratio of 10 : (0.2 - 2) and between 20 and 50 wt. % of a moistening agent from the group containing sorbitol, glycerin, propyleneglycol-1,2, polyethyleneglycol and their mixtures, comprising, for reinforcing the cleaning effect, between 2 and 12 wt. % of condensed phosphate from the group comprising tripolyphosphate, pyrophosphate or trimetaphosphate in the form of alkaline salt or ammonium salt. In spite of their low abrasive values, said cleaning agents have an excellent polishing effect and a high cleaning power. (57) Zusammenfassung Zahnreinigungsmittel in Form einer pastösen oder flüssigen Dispersion mit einem Gehalt von 10-30 Gew.-% einer Poliermittelkombination aus Kieselsäure-Poliermitteln und Aluminiumoxid im Gewichtsverhältnis 10 : (0,2-2) und 20-50 Gew.-% eines Feuchthaltemittels aus der Gruppe Sorbit, Glycerin, Propylenglycol-1,2, Polyethylenglycol und Gemischen davon, die zur Erhöhung der Reinigungswirkung 2-12 Gew.-% eines kondensierten Phosphats aus der Gruppe Tripolyphosphat, Pyrophosphat oder Trimetaphosphat in Form eines Alkali- oder Ammoniumsalzes enthalten, zeigen trotz niedriger Abriebwerte eine sehr gute Polierwirkung sowie ein hohes Reinigungsvermögen.		

**Cleaning Agent for Dental Use Comprising a Combination of
Polishing Agents Based on a Silicic Acid and Aluminium Oxide**

This invention relates to a dentifrice in the form of a water-containing, paste-form or liquid cream containing 10 to 30% by weight of a combination of polishing agents and 20 to 50% by weight of a humectant, of which the cleaning performance has been further improved by an addition
5 of condensed phosphates

Toothpastes are used in the daily cleaning of teeth by brushing with a toothbrush. Toothpastes are intended above all to support the removal from the tooth surfaces of food remains, discoloration, for example by tobacco or tea, and firmly adhering bacterial films, so-called plaque. This is
10 mainly done by the polishing agents present in the toothpaste and, to a lesser extent, also by the surfactants present. In order to develop their cleaning and polishing effect, the polishing agents have to have a certain abrasiveness towards the tooth surface. However, it is extremely important that the abrasiveness towards dental enamel and dentin is kept at low
15 levels to avoid damage to the tooth surface by the daily use of the toothpastes. Above all, the polishing agents used should not cause any deep scratches on the tooth surface. On the contrary, they should have a smoothing effect on any roughness present in the tooth surface.

Condensed phosphates are known from numerous publications as
20 scale inhibitors and as demineralization inhibitors in dentifrices.

The cleaning effect of water-soluble pyrophosphate salts on teeth discolored by tea and coffee is also known from **US 4,822,599 A**.

Dentifrices containing an abrasive mixture of a silica polishing agent and a calcined aluminium oxide are known as particularly effective tooth
25 cleaning preparations from **DE 27 58 548 C2**. This document recommends the addition of certain inorganic electrolyte salts to prevent abrasion of the

dental enamel. In spite of this, extremely high dentin abrasion values (RDA) and enamel abrasion values (REA) of 300 to 500 were obtained.

Although the abrasion values can be reduced by reducing or omitting the aluminium oxide polishing agent, cleaning performance is significantly impaired at the same time. Accordingly, there was a need to develop a dentifrice which would combine very low abrasiveness (RDA and REA values below 100) with a very good cleaning effect (CRS values of around 100).

DE 34 25 152 describes a combination of polishing agents consisting of a silica polishing agent and a lightly calcined aluminium oxide with which reduced abrasiveness can be combined with a good cleaning effect.

It has now been found that condensed phosphates are ideally suitable for increasing the cleaning performance of toothpastes containing a combination of silica polishing agents and aluminium oxide polishing agents to such an extent that the need stated above is satisfied without any significant increase in the abrasion values.

Accordingly, the present invention relates to dentifrices in the form of water-containing, paste-form or liquid dispersions containing 10 to 30% by weight of a combination of silica polishing agents and aluminium oxide in a ratio by weight of 10:0.2 to 10:2 and 20 to 50% by weight of a humectant selected from the group consisting of sorbitol, glycerol, 1,2-propylene glycol and mixtures thereof, characterized in that they contain a condensed phosphate selected from the group consisting of tripolyphosphate, pyrophosphate and trimetaphosphate in the form of the alkali metal or ammonium salts in a quantity of 2 to 12% by weight in order to increase the cleaning effect.

In one embodiment, the condensed phosphate is selected from the group consisting of tripolyphosphate, pyrophosphate, trimetaphosphate and mixtures thereof.

Suitable silica polishing agents are, for example, silica gels which are obtained by reacting sodium silicate solutions with strong aqueous mineral acids to form a hydrosol, ageing the hydrosol to form the hydrogel, washing and drying. If drying is carried out under moderate conditions to

water contents of 15 to 35% by weight, so-called hydrogel silicas, which are known for example from **US 4,153,680**, are obtained.

Drying to water contents below 15% by weight results in irreversible shrinkage of the previously loose structure of the hydrogel to the dense structure of the so-called xerogel. Xerogel silicas are described, for example, in **US 3,538,230**.

A second preferred group of silica polishing agents are the precipitated silicas. Precipitated silicas are obtained by precipitation of silica from dilute alkali metal silicate solutions by addition of strong acids under conditions where aggregation to the sol and gel cannot occur. Suitable processes for the production of precipitated silicas are described, for example, in **DE-OS 25 22 486** and in **DE-OS 31 14 493**. Preferred precipitated silicas are, for example, those which have a mean particle size of 5 to 20 μm , a sieve residue (45 μm) of less than 1% by weight and a specific surface (BET) of 100 to 300 m^2/g .

By virtue of the special combination of polishing agents, the dentifrices according to the invention have an excellent cleaning effect, even against discoloration of the teeth by tea and nicotine. At the same time, a good polishing effect (smoothing of roughness) is obtained despite only moderate dentin and enamel abrasion. In spite of the presence of a relatively hard polishing component, namely aluminium oxide, the dentifrices according to the invention have hardly any scratching effect.

A preferred aluminium oxide polish is a lightly calcined alumina with a content of at least 10% by weight of α -aluminium oxide of various so-called γ -aluminium oxide modifications.

Suitable lightly calcined aluminas are prepared by calcination from aluminium hydroxide. Aluminium hydroxide is converted by calcination into $\alpha\text{-Al}_2\text{O}_3$ which is thermodynamically stable at temperatures above 1200°C. The thermodynamically unstable Al_2O_3 modifications occurring at temperatures of 400 to 1000°C are known as γ -forms (cf. **Ullmann, Enzyklopädie**

der technischen Chemie, 4th Edition (1974), Vol. 7, page 298). The degree of calcination, i.e. the conversion into the thermodynamically stable α - Al_2O_3 , can be adjusted as required through the choice of the temperature and the duration of the calcination process. Light calcination gives an alumina with a γ - Al_2O_3 content which is lower, the higher the calcination temperature and the longer the calcination time. Lightly calcined aluminas differ from pure α - Al_2O_3 in the lower hardness of the agglomerates, in a larger specific surface and larger pore volumes.

The dentin abrasion (RDA) of the relatively lightly calcined aluminas to be used in accordance with the invention containing 10 to 50% by weight of γ - Al_2O_3 is only 30 to 60% of the dentin abrasion of a heavily calcined, pure α - Al_2O_3 (as measured in a standard toothpaste containing 20% by weight alumina as sole polishing agent).

In contrast to α - Al_2O_3 , γ - Al_2O_3 can be dyed red with an aqueous-ammoniacal solution of Alizarin S (1,2-dihydroxy-9,10-anthraquinone-4-sulfonic acid). The degree of dyeability can be used as a measure of the degree of calcination or rather the percentage content of δ - Al_2O_3 in a calcined alumina:

Ca. 1 g Al_2O_3 , 10 ml of a solution of 2 g/l Alizarin S in water and 3 drops of an aqueous 10% by weight solution of NH_3 are introduced into a test tube and briefly boiled. The Al_2O_3 is then filtered off, washed, dried and examined under a microscope or evaluated by colorimetry.

Suitable lightly calcined aluminas containing 10 to 50% by weight γ - Al_2O_3 can be colored pale to deep pink by this method.

Aluminium oxide polishing agents with various degrees of calcination, fineness and bulk densities are commercially obtainable, for example the "Poliertonerden (polishing aluminas)" of Giulini-Chemie and ALCOA.

A particularly suitable quality "Poliertonerde P10 feinst" has an agglomerate size below 20 μm , a mean primary crystal size of 0.5 to 1.5 μm and a bulk density of 500 to 600 g/l.

5 Sorbitol, xylitol, glycerol, propylene glycol or mixtures of these polyols may be present as humectants. Polyethylene glycols with molecular weights of 400 to 2000 may also be partly present as humectant components. Sorbitol in a quantity of 20 to 40% by weight is preferably present as the humectant.

10 The condensed phosphates are present in the form of their alkali metal salts, preferably their sodium or potassium salts. The aqueous solutions of these phosphates show an alkaline reaction on account of hydrolytic effects. The dentifrices according to the invention are adjusted to a pH of 7.5 to 9 by addition of an acid. Suitable acids are, for example, citric acid, phosphoric acid or acidic salts, for example NaH_2PO_4 . However, 15 acidic salts of the condensed phosphates, i.e. for example $\text{K}_2\text{H}_2\text{P}_2\text{O}_7$, may also be partly used to adjust the dentifrice to the required pH value.

Mixtures of various condensed phosphates or hydrated salts of the condensed phosphates may also be used. However, the specified quantities of 2 to 12% by weight are based on the water-free salts. A 20 sodium or potassium tripolyphosphate in a quantity of 5 to 10% by weight of the composition is preferably present as the condensed phosphate.

The cleaning effect of the dentifrices according to the invention may be further improved by addition of a suitable surfactant. The addition of a surfactant may also be desirable for producing a foam during brushing of 25 the teeth, for stabilizing the dispersion of polishing agents and for emulsifying or solubilizing the flavoring oils. Suitable surfactants which develop a certain foaming effect are the nonionic surfactants, for example sodium alkyl sulfates containing 12 to 18 carbon atoms in the alkyl group. These surfactants also have a certain enzyme-inhibiting effect on the 30 bacterial metabolism of plaque. Other suitable surfactants are alkali metal

salts, preferably sodium salts, of alkyl polyglycol ether sulfate containing 12 to 16 carbon atoms in the linear alkyl group and 2 to 6 glycol ether groups in the molecule, of linear alkane (C₁₂₋₁₈) sulfonate, of sulfosuccinic acid monoalkyl (C₁₂₋₁₈) esters, of sulfated fatty acid monoglycerides, sulfated
5 fatty acid alkanolamides, sulfoacetic acid alkyl (C₁₂₋₁₆) esters, acyl sarcosines, acyl taurides and acyl isethionates containing 8 to 18 carbon atoms in the acyl group.

Zwitterionic and ampholytic surfactants may also be used, preferably in combination with anionic surfactants. However, it is particularly preferred
10 to use nonionic surfactants to promote the cleaning effect. Suitable nonionic surfactants are, for example, products of the addition of ethylene oxide with fatty alcohols, with fatty acids, with fatty acid monoglycerides, with sorbitan fatty acid monoesters or with methyl glucoside fatty acid monoesters. The quantity of ethylene oxide added on should be so large
15 that the surfactants are soluble in water, i.e. at least 1 g/l should be soluble in water at 20°C. Another group of suitable surfactants are the alkyl (oligo)-glycosides containing 8 to 16 carbon atoms in the alkyl group and having a degree of oligomerization of the glycoside unit of 1 to 4. Alkyl (oligo)glycosides, their production and use as surfactants are known, for example, from
20 **US-A-3,839,318, DE-A-20 36 472, EP-A-77 167 or WO-A-93/10132.**

So far as the glycoside unit is concerned, monoglycosides ($x = 1$) where a monosaccharide unit is attached to a C₁₀₋₁₆ fatty alcohol by a glycoside linkage and oligomeric glycosides with a degree of oligomerization x of up to 10 are suitable. The degree of oligomerization is a
25 statistical mean value on which the homolog distribution typical of such technical products is based.

A particularly suitable alkyl (oligo)glycoside is an alkyl (oligo)glucoside with the formula $RO(C_6H_{10}O)_x-H$, where R is an alkyl group containing 12 to 14 carbon atoms and x has a mean value of 1 to 4.

A nonionic solubilizer from the group of surface-active compounds may be necessary, particularly for solubilizing the generally water-insoluble flavoring oils. Particularly suitable nonionic solubilizers are, for example, ethoxylated fatty acid glycerides, ethoxylated fatty acid sorbitan partial esters or fatty acid partial esters of glycerol or sorbitan ethoxylates. Solubilizers from the group of ethoxylated fatty acid glycerides include above all products of the addition of 20 to 60 moles of ethylene oxide onto mono- and diglycerides of linear fatty acids containing 12 to 18 carbon atoms or onto glycerides of hydroxy fatty acids, such as hydroxystearic acid or ricinoleic acid. Other suitable solubilizers are ethoxylated fatty acid sorbitan partial esters, i.e. preferably products of the addition of 20 to 60 moles ethylene oxide onto sorbitan monoesters and sorbitan diesters of fatty acids containing 12 to 18 carbon atoms. Other suitable solubilizers are fatty acid partial esters of glycerol or sorbitan ethoxylates, i.e. preferably monoesters and diesters of C₁₂₋₁₈ fatty acids and products of the addition of 20 to 60 moles ethylene oxide onto 1 mole glycerol or onto 1 mole sorbitol.

The dentifrices according to the invention preferably contain products of the addition of 20 to 60 moles ethylene oxide onto hydrogenated or non-hydrogenated castor oil (i.e. onto hydroxystearic acid or ricinoleic acid triglyceride), onto glycerol monostearate and/or distearate or onto sorbitan monostearate and/or distearate as solubilizers for any flavoring oils present.

Suitable flavoring components are, for example, sweeteners and/or flavoring oils. Suitable flavoring oils are any of the natural and synthetic flavors typically used in oral and dental care preparations. Natural flavors may be used both in the form of the essential oils isolated from the drugs and in the form of the individual components isolated therefrom. The dentifrice should preferably contain at least one flavoring oil from the group consisting of peppermint oil, spearmint oil, anise oil, Japanese anise oil,

caraway oil, eucalyptus oil, fennel oil, cinnamon oil, clove oil, geranium oil, sage oil, pimento oil, thyme oil, marjoram oil, basil oil, citrus oil, gaultheria oil or one or more components of these oils isolated from them or synthetically produced. The most important components of the oils mentioned are, for example, menthol, carvone, anethol, cineol, eugenol, cinnamaldehyde, caryophyllene, geraniol, citronellol, linalool, salvia, thymol, terpinene, terpineol, methyl chavicol and methyl salicylate. Other suitable flavors are, for example, menthyl acetate, vanillin, ionone, linalyl acetate, rhodinol and piperitone.

10 In addition, the dentifrices according to the invention may contain a therapeutic agent for controlling caries, scale, parodontitis or other diseases of the mouth and teeth. An active principle preferably present is a caries-inhibiting fluorine compound, preferably from the group of fluorides or monofluorophosphates in a quantity of 0.1 to 0.5% by weight fluorine.

15 Suitable fluorine compounds are, for example, sodium fluoride, potassium fluoride, tin fluoride, sodium monofluorophosphate ($\text{Na}_2\text{PO}_3\text{F}$), potassium monofluorophosphate or the fluoride or an organic amino compound.

Other suitable therapeutic agents are, for example scale inhibitors, for example organophosphates, such as 1-azacycloheptane-2,2-diphosphonic acid (Na salt) or 1-hydroxyethane-1,1-diphosphonic acid (Na salt), and antimicrobial plaque inhibitors such as, for example, hexachlorophene, chlorhexidine, hexetidine, triclosan, bromochlorophene, phenyl salicylate. Substances effective in promoting remineralization and the closure of dental lesions, for example dicalcium phosphate dihydrate, preferably in combination with magnesium ions, may also be present in the dentifrices according to the invention.

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In one preferred embodiment, the toothpastes according to the invention additionally contain, for example, 1 to 10% by weight of dicalcium phosphate dihydrate (brushite) and 0.1 to 0.5% by weight of magnesium ions, preferably in the form of a water-soluble magnesium salt, for example

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magnesium sulfate, magnesium fluoride or magnesium monofluorophosphate.

Finally, the toothpastes according to the invention may contain other components which are normally encountered in dentifrices and which do not reduce the effects according to the invention. Such typical toothpaste additives are, for example,

- other polishing agents in relatively small quantities of, for example, 1 to 10% by weight, for example calcium carbonate (chalk), insoluble sodium metaphosphate, calcium pyrophosphate, hydroxyl apatite, aluminium hydroxide, sodium aluminium silicates (zeolite A) or particulate organic polymers, for example polymethacrylate,
- pigments, for example titanium dioxide or zinc oxide,
- dyes,
- pH regulators and buffers, for example sodium citrate or sodium bicarbonate, sodium benzoate,
- wound-healing and anti-inflammatory agents such as, for example, allantoin, urea, panthenol, azulene or camomile extract,
- preservatives such as, for example, sorbic acid salts, p-hydroxybenzoic acid ester.

The following Examples are intended to illustrate the invention.

Examples

The following toothpastes were prepared:

Composition	1	2	3	4	5
Sident TM 8 (1)	14.0	-	-	-	-
Sorbosil TM AC 39 (2)	-	14.0	-	-	14.0
Zeodent TM 113 (3)	-	-	14.0	-	-
Zeodent TM 623 (4)	-	-	-	14.0	-
Poliertonerde TM P10 feinst (5)	1.0	1.0	1.0	1.0	0.5
Na ₅ P ₃ O ₁₀ (Na tripolyphosphate)	5.0	5.0	5.0	5.0	5.0
Na ₂ PO ₃ F (Na monofluorophosphate)	-	0.8	-	-	-
NaF	0.24	-	0.24	0.24	0.24
Na saccharinate	0.1	0.1	0.1	0.1	0.1
Titanium dioxide	0.5	0.5	0.5	0.5	0.5
PHB methyl ester	0.1	0.1	0.1	0.1	0.1
Carboxymethyl cellulose	1.25	1.25	1.25	1.25	1.25
Sorbitol TM (70%)	32.0	32.0	32.0	32.0	32.0
1,2-Propylene glycol	5.0	5.0	5.0	5.0	5.0
Cremophor TM RH 60 (7)	1.0	-	1.0	1.0	1.0
Arlatone TM 289 (8)	-	1.0	-	-	-
Flavoring oil	0.8	0.8	0.8	0.8	0.8
Water	to 100	to 100	to 100	to 100	to 100

Composition	6	C1	C2	C3
Sorbosil TM AC 39 (2)	14	14	14	14
Poliertonerde TM P 10 feinst (5)	1	-	-	-
Precarb TM 100 (6)	-	2	2	2
Na ₅ P ₃ O ₁₀	10	10	5	0
Na ₂ PO ₃ F	0.8	0.8	0.8	0.8
NaF	-	-	-	-
Na saccharinate	0.1	0.1	0.1	0.1
Titanium dioxide	0.5	0.5	0.5	0.5
PHB methyl ester	0.1	0.1	0.1	0.1
Carboxymethyl cellulose	1.25	1.25	1.25	1.25
Sorbitol (70%)	32.0	32.0	32.0	32.0
1,2-Propylene glycol	5.0	5.0	5.0	5.0
Arlatone TM 289 (8)	1.0	1.0	1.0	1.0
Flavoring oil	0.8	0.8	0.8	0.8
Water	to 100	to 100	to 100	to 100

The following commercial products were used:

- (1) Sident[®] 8 : silica polish
 (DEGUSSA) mean particle size : 10.0 µm
 sieve residue (45 µm) : ≤ 0.3%
 compacted bulk density : 300 g/l
- (2) Sorbosil AC 39 : silica polish
 (Crosfield Ltd.) particle size: 9-13 µm
- (3) Zeodent 113 : silica polish
 (Huber Chemicals) mean particle size : 12 µm

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sieve residue (45 μm): 1.0% max.spec. surface (BET) : 150 m^2/g

(4) Zeodent 623
5 (Huber Chemicals)

: silica polish

mean particle size : 12 μm sieve residue (45 μm): 0.5% max.spec. surface (BET) : 250 m^2/g

(5) Poliertonerde P10 feinst
10 (Giulini Chemie)

: aluminium oxide polish

mean agglomerate size: < 20 μm (min. 99%)primary particle size: ca. 1 μm

degree of calcination: low

15 (6) Precarb 100

: calcium carbonate (chalk)

compacted bulk density 0.5 g/ml

spec. surface: 9.0 m^2/g

20 (7) Cremophor RH 60

: hydr. castor oil + 60 moles EO

(8) Arlatone 289

: hydr. castor oil + 54 moles EO

melting point: 39°C

HLB value: 14.4

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Determination of the cleaning effect (CRS)**Method:**

The surface of bovine teeth was conditioned, stained with tea under defined conditions and cleaned with the toothpaste to be tested under defined conditions. The lightening in color obtained was measured by
30 colorimetry and was compared with the lightening effect of a standard

toothpaste.

Sample preparation

5 Bovine incisors were cut into 7 x 7 mm blocks which were then mounted with wax on PlexiglasTM squares (1 x 2.5 x 2.5 cm) in such a way that only the enamel surface was exposed. The enamel surface was polished until it appeared uniformly smooth.

Sample conditioning

10 The mounted tooth blocks were successively immersed in 0.12 N hydrochloric acid (60 seconds), saturated Na₂CO₃ solution (120 seconds) and in 1% phytic acid solution (60 seconds). After each treatment, the samples were rinsed with deionized water and dried by blotting with absorbent paper. The tooth samples were then fixed to the staining
15 apparatus and moved for 5 days through a solution of black tea at 20°C. The tea solution had been prepared by extracting a 1.5 g teabag with 300 g of boiling water (10 minutes) and was renewed twice a day.

Polishing tests

20 The mounted tooth blocks were then placed in a Grabenstetter V8 brushing machine and were brushed in a toothpaste suspension of 20 g paste and 40 g deionized water using a soft Oral-BTM toothbrush (pressure applied 150 g). A suspension of 10 g of calcium pyrophosphate in 50 g of swollen carboxymethyl cellulose (0.5% CMC, 10% glycerol, 89.5% water)
25 was used as the polishing standard. The cleaning effect of this standard is defined as the 100% value.

Measurement of the cleaning effect

The lightening effect was measured to DIN 5033 using a Dr. Lange
30 color difference measuring instrument (type Micro Color (DC 8334)). A

xenon lamp producing D 65 standard light corresponding to daylight was used. Barium sulfate was used as the color standard.

Double measurements of a circular area 7 mm in diameter on the sample surface were carried out. Eight stained tooth samples were used
5 for each toothpaste and average values were formed. The test parameter used was the standard color value Y determined to DIN 5033 which is a measure of the lightness of a color.

Lightness measurement was carried out after 1000 brush strokes. The cleaning effect CRS (cleaning ratio soft) is calculated as follows:

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$$\text{CRS [\%]} = \frac{\text{average standard color value Y - increment test paste}}{\text{average standard color value Y - increment polishing standard}} \cdot 100 [\%]$$

15 **Determination of abrasiveness (RDA)**

Abrasiveness was determined by the RDA method (radioactive dentin abrasion) using the Grabenstetter process of Missouri Analytical Laboratories, St. Louis, MO. (cf. J. Dent. Res. 17, 1060 - 1068 (1958).

20 **Results:**

Example	1	2	3	4	5	6	C1	C2	C3
CRS %	90	88	103	98	73	97	80	64	31
RDA %	n.d.*	66	55	95	69	n.d.	n.d.	57	n.d.

(* n.d. not determined)

The results show that, without aluminium oxide (C1 - C3), a weaker
25 cleaning effect is obtained than with the toothpastes according to the invention (1-4), despite high Na tripolyphosphate contents (C1), and that dentin abrasion shows hardly any increase in relation to the comparison pastes despite the high cleaning performance.

CLAIMS:

1. Dentifrices in the form of water-containing, paste-form or liquid dispersions containing 10 to 30% by weight of a combination of silica polishing agents and aluminium oxide in a ratio by weight of 10:0.2 to 10:2 and 20 to 50% by weight of a humectant selected from the group consisting of sorbitol, glycerol, 1,2-propylene glycol and mixtures thereof, characterized in that they contain a condensed phosphate selected from the group consisting of tripolyphosphate, pyrophosphate and trimetaphosphate in the form of an alkali metal or ammonium salt in a quantity of 2 to 12% by weight in order to increase the cleaning effect.
2. A dentifrice as claimed in claim 1, characterized in that a lightly calcined alumina containing α -aluminium oxide, of various Al_2O_3 modifications, is present as the aluminium oxide in a quantity of at least 10% by weight of the total weight of the dentifrice.
3. A dentifrice as claimed in claim 1 or 2, characterized in that sorbitol is present as humectant in a quantity of 25 to 40% by weight of the total weight of the dentifrice.
4. A dentifrice as claimed in any one of claims 1 to 3, characterized in that sodium or potassium tripolyphosphate is present as the condensed phosphate in a quantity of 5 to 10% by weight of the total weight of the dentifrice.
5. A dentifrice as claimed in any one of claims 1 to 4, characterized in that a precipitated silica with a particle size of 5 to 20 μm and a specific surface (BET) of 100 to 300 g/l is present as the silica polishing agent.
6. A dentifrice as claimed in any one of claims 1 to 5, characterized in that a fluorine compound is present in a quantity of 0.1 to 0.5% by weight fluorine of the total weight of the dentifrice.
7. A dentifrice as claimed in claim 6, characterized in that the fluorine compound is selected from the group consisting of fluorides and monofluoro-phosphates.