

COMPOSITIONS

The present invention is concerned with a sweetness enhancing composition and with an orally consumable composition containing same. The present invention is also concerned with methods of enhancing the sweetness of orally consumable
5 compositions.

Steviol glycosides are responsible for the sweet taste of the stevia plant (*Stevia Rebaudiana*, (Bertoni) Bertoni, family Compositae). The leaves of Stevia plant contain at least 10 steviol glycosides. The major ingredient is stevioside with a natural abundance around 4-14% by weight, followed by rebaudioside A (typically
10 2-4% by weight), and rebaudioside C (or called dulcoside B, ca 1%). Other minor steviol glycosides include dulcoside A, rebaudiosides B, D, E, and F, rubusoside, and steviobioside. The total contents of steviol glycosides and the individual components in the leaves vary depending upon its species and conditions at the location of cultivation, such as temperature, latitude as well as the harvesting time.
15 The abundance of individual steviol glycosides can be specifically enriched through natural hybrid or genetic modification.

Rebaudioside A is the most potent sweetener and is regarded as having the best taste amongst all the steviol glycosides. Indeed, it is the first natural sweetener to be used in beverage and food applications in the United States. However, despite it
20 being generally regarded as having the most desirable taste profile, this material still has associated with its use an unwanted or unpleasant after taste that has been described as a lingering, liquorice-type bitterness. This remains an issue regarding its use in food and beverage applications. Further, just like other high potency sweeteners, rebaudioside A lacks the upfront sweetness, body and
25 possesses the lingering after-taste even when used at low levels as a sweetness enhancer.

Much work has been undertaken to reduce or eliminate the unpleasant after taste of associated with steviol glycosides, and with regard to rebaudioside A, current practice is to separate rebaudioside A from other steviol glycosides and employ it
30 in a highly pure form. Unfortunately, obtaining rebaudioside A in highly pure form from stevia plant extracts is complex, time-consuming and expensive

There remains a need to provide sweetness enhancing compositions that perform substantially similarly to or better than rebaudioside A, but which are less costly.

Applicant arrived at a solution to this problem in a surprising manner.

Accordingly, the invention provides in a first aspect a composition comprising a
5 mixture of stevioside and rebaudioside A in a stevioside to rebaudioside A ratio of about 1:1 to about 4:1, more particularly about 2:1 to 3:1.

The invention resides in the surprising discovery that mixtures relatively rich in stevioside and relative poor in rebaudioside A nevertheless can enhance
10 sweetness of sweetened compositions in a way that is substantially similar to or even better than pure rebaudioside A. Applicant arrived at this surprising discovery whilst working on fractionation of a commercial stevia extract. Applicant generated a fraction that was prepared for tasting and discovered to its surprise that it was at least as good as, or even better than pure rebaudioside A in terms of sweetness enhancement on top of a sugar solution. Additionally, the flavour profile was
15 assessed as less lingering and lacked the liquorice bitterness of rebaudioside A, and it had a profile substantially similar to sugar itself.

Upon analysis, applicant found this fraction to contain stevioside and rebaudioside A. Surprisingly, the fraction was relatively rich in stevioside and relatively poor in rebaudioside A. This result was indeed surprising given that it is known in the
20 literature that rebaudioside A is far the most potent sweetener of the steviol glycosides, with what is generally considered to be the best taste profile, whereas stevioside is of much lower potency and of poorer taste quality. Additional experimentation confirmed a range of mixtures of stevioside and rebaudioside A that provide desirable sugar-like profiles that are at least as good as or superior
25 even to rebaudioside A in its pure form. More particularly, the mixtures of the present invention were found to have about the same sweetness as rebaudioside A, but they displayed more upfront impact, less licorice lingering off-taste and a much more sugar-like profile than rebaudioside A alone.

Both stevioside and rebaudioside A are known sweeteners. In addition, the sweetness enhancing properties of both rebaudioside A and stevioside are known in the art (see for example US patent US20100092638).

However, applicant is unaware of any prior art teaching related to the effect of
5 mixing stevioside and rebaudioside A on the taste profiles of these mixtures, and in particular the sweetness enhancement effects of the mixtures. Indeed, applicant believes that it is entirely counter-intuitive to consider mixing stevioside with rebaudioside A in the expectation of achieving a desirable result, when the current focus of the flavour industry is concerned with obtaining and using pure forms of
10 rebaudioside A. Why, once having gone to the trouble and expense of isolating pure rebaudioside A, would the skilled person then combine it with what is generally considered to be the poorer performing stevioside?

The mixtures of the present invention can be used in amounts up to 250 ppm or greater if desired. However, at levels higher than 250 ppm bitter up-front notes,
15 believed to be associated with the high levels of stevioside start to become apparent. The acceptability or otherwise of these bitter off-notes will depend somewhat on the nature of the consumable compositions in which the mixtures are to be employed. However, it is preferred to employ the mixture in consumable compositions in amounts lower than 250 ppm.

20 Indeed, at levels above 250 ppm, our observations suggest that it would be necessary to alter the proportions of stevioside and rebaudioside A in the mixtures such that a much higher amount of the latter and a much lower amount of the former would be necessary if the mixtures were to find any use as sweeteners. In other words, the ratio of ingredients in the mixtures of the present invention would
25 not be appropriate when the mixture is employed at levels of 250 ppm or greater.

It is preferred to employ mixtures of the present invention in amounts of 150 ppm or less, more particularly 90 ppm or less, more particularly 60 ppm or less, more particularly 45 ppm or less, more particularly 40 ppm or less, still more particularly 35 ppm or less.

It is particularly preferred to use the mixtures in amounts wherein they are not sweet-tasting in and of themselves, that is, it is preferred to use the mixtures at or near the sweetness detection threshold.

The sweetness detection threshold of rebaudioside A and stevioside is discussed in US20100092638, which is incorporated by reference. The term sweetness detection threshold is known in the art, and a discussion of this can be found in US20100092638. In relation to the present invention, it refers to a concentration at which the mixture is not perceived as sweet. However, the mixtures at their sweetness detection threshold nevertheless can exert on a composition containing a sweetener.

The sweetness detection threshold may not be absolute for the mixture, and may depend on the presence of other ingredients that may be present in a composition containing said mixture, or on other stimuli that a consumer experiences during consumption of the mixture. In addition, it is known that sweetness detection thresholds can vary somewhat from individual to individual. For example, some individuals are able to detect the sweetness of sucrose in a very low concentration, e.g. at 0.4%, whereas others can only detect at 0.7% or even higher. Accordingly, it is typical to perform such assessments objectively using sweet sensitive panelists able to detect at least 0.5% of sucrose or less. As a matter of generality, such panelists would rate a 1.00% sucrose solution as being practically lacking sweetness. Accordingly, the sweetness detection threshold of a sample can be determined as the concentration of that sample that is iso-intense as a 1.25 % or lower sucrose solution, more particularly a 1.00% lower sucrose solution. The concentration of a sweetness enhancer at or near its detection threshold can be easily and objectively tested by organoleptic titration techniques that are well known in the field of sensory analysis as more fully described in US20100092638.

In a particular embodiment of the present invention the mixture can be employed in an amount that is iso-sweet with a 1 % sucrose solution.

The mixture according to the present invention, when used in amounts referred to hereinabove, and even at its sweetness detection threshold, can exert an

enhancing effect on a sweetened composition that adds approximately 1% sucrose equivalence.

- If desired, one can add additional sweetness enhancers to mixtures of the present invention. Additional sweetness enhancers may be selected from those disclosed
 5 in US20100092638, i.e. trilobatin, hesperetin dihydrochalcone glucoside, naringin dihydrochalcone, mogroside V, Luo Han Guo extract, rubusoside, rubus extract or mixtures thereof. US20100092638, which is hereby incorporated by reference, in addition to disclosing these materials, also proposes usage levels at or below their sweetness detection thresholds.
- 10 Other additional sweetness enhancers may include, glycyphyllin, isomogroside V, mogroside IV, siamenoside I, neomogroside, mukurozioside IIb, (+)-hernandulcin, 4 β -hydroxyhernandulcin, baiyunoside, phlomisioside I, bryodulcoside, bryoside bryonoside, abrusosides A-E, cyclocarioside A, cyclocaryoside I, albiziasaponins A-E, glycyrrhizin, araboglycyrrhizin, periandrins I-V, pterocaryosides A and B,
 15 osladin, polypodosides A and B, telosmoside A8-18, phyllodulcin, huangqioside E neoastilbin, monatin, 3-acetoxy-5,7-dihydroxy-4'-methoxyflavanone, 2*R*,3*R*-(+)-3-Acetoxy-5,7,4'-trihydroxyflavanone, (2*R*,3*R*)-dihydroquercetin 3-*O*-acetate, dihydroquercetin 3-*O*-acetate 4'-methyl ether, brazzein, curculin, mabinlin, monellin, neoculin, pentadin, thaumatin or combinations thereof.
- 20 Each of the aforementioned additional sweetness enhancers may be employed in a concentration around its sweetness detection threshold. In the case of trilobatin, hesperitin dihydrochalcone-4-beta-glucoside (HDG), naringin dihydrochalcone, mogroside V, iso-mogroside V, siamenoside I, mogroside IV, neomogroside, Luo Han Guo extract, rubusoside, rubus extract, and NHDC, the sweetness detection
 25 thresholds are respectively for trilobatin from 3 to 100 ppm, HDG from 1 to 75 ppm, naringin dihydrochalcone from 2 to 60 ppm, for rubusoside from 1.4 ppm to 56 ppm, for rubus extract from 2 ppm to 80 ppm, for mogroside V from 0.4 ppm to 12.5 ppm, iso-mogroside V from 0.3 to 12 ppm, siamenoside I from 0.2 to 10 ppm, for Luo Han Guo extract from 2 to 60 ppm, and for NHDC from 1 to 5 ppm.
- 30 The mixture according to the present invention may be added to a consumable composition containing a sweetener.

The mixture may be added to the consumable composition in order to exert a sweetness enhancing effect. The mixture may be added to a consumable composition alone, or it may form part of a flavour composition, which can be added to the consumable composition.

- 5 The flavour composition may comprise, in addition to the mixture of the present invention, one or more sweeteners, one or more additional sweetness enhancers as described above, as well as any flavourants and other standard ingredients commonly used in such compositions as more fully described herein below.

The invention provides in another of its aspects, a consumable composition
10 comprising a sweetener and a sweetness-enhancing composition comprising a mixture of stevioside and rebaudioside A in a ratio of about 1:1 to about 4:1, more particularly about 2:1 to 3:1, and wherein said mixture is employed in said consumable composition at a concentration of 250 ppm or less, more particularly 150 ppm or less, 90 ppm or less, still more particularly 60 ppm or less, more
15 particularly 45 ppm or less, more particularly 40 ppm or less, still more particularly 35 ppm or less.

In an embodiment of the invention said consumable composition contains the mixture in a concentration at or near the sweetness detection threshold, in particular at a level such that the mixture is iso-sweet with 1.25 % sucrose or less,
20 still more particularly iso-sweet with 1 % sucrose.

The sweetener employed in said consumable composition may be selected from the group consisting of sucrose, fructose, glucose, high fructose corn syrup, corn syrup, xylose, arabinose, rhamnose, erythritol, xylitol, mannitol, sorbitol, inositol, acesulfame potassium, aspartame, neotame, sucralose, and saccharine.

- 25 The sweetener may be present in amounts of 0.0001 to 15% by weight of a consumable composition.

A consumable composition of the present invention is any solid or liquid composition that is consumed for at least one of nourishment and pleasure. It may be placed in the mouth before being consumed, or it may be held in the mouth for a
30 period of time before being discarded. Broadly, the consumable composition

includes, but is not limited to foodstuffs of all kinds, confectionery, baked goods, sweet goods, dairy products and beverages.

Confectionery includes, but is not limited to chewing gum (which includes sugarized gum, sugar-free gum, functional gum and bubble gum), centre-fill
 5 confections, chocolate and other chocolate confectionery, medicated confectionery, lozenges, tablets, pastilles, mints, standard mints, power mints, chewy candies, hard candies, boiled candies, breath and other oral care films or strips, candy canes, lollipops, gummies, jellies, fudge, caramel, hard and soft panned goods, toffee, taffy, liquorice, gelatin candies, gum drops, jelly beans, nougats, fondants,
 10 or combinations of one or more of these, or edible compositions incorporating one or more of these.

Baked goods include, but are not limited to alfajores, bread, packaged/industrial bread, unpackaged/artisanal bread, pastries, cakes, packaged/industrial cakes, unpackaged/artisanal cakes, cookies, chocolate coated biscuits, sandwich biscuits,
 15 filled biscuits, savory biscuits and crackers, bread substitutes.

Sweet goods include, but are not limited to breakfast cereals, ready-to-eat ("rte") cereals, family breakfast cereals, flakes, muesli, other rte cereals, children's breakfast cereals and hot cereals.

Dairy products include, but are not limited to ice cream, impulse ice cream, single
 20 portion dairy ice cream, single portion water ice cream, multi-pack dairy ice cream, multi-pack water ice cream, take-home ice cream, take-home dairy ice cream, ice cream desserts, bulk ice cream, take-home water ice cream, frozen yoghurt, artisanal ice cream, dairy products, milk, fresh/pasteurized milk, full fat fresh/pasteurized milk, semi skimmed fresh/pasteurized milk, long-life/uht milk, full
 25 fat long life/uht milk, semi skimmed long life/uht milk, fat-free long life/uht milk, goat milk, condensed/evaporated milk, plain condensed/evaporated milk, flavored, functional and other condensed milk, flavored milk drinks, dairy only flavored milk drinks, flavored milk drinks with fruit juice, soy milk, sour milk drinks, fermented dairy drinks, coffee whiteners, powder milk, flavored powder milk drinks, cream,
 30 yoghurt, plain/natural yoghurt, flavored yoghurt, fruited yoghurt, probiotic yoghurt,

drinking yoghurt, regular drinking yoghurt, probiotic drinking yoghurt, chilled and shelf-stable desserts, dairy-based desserts and soy-based desserts.

Other foodstuff include, but are not limited to chilled snacks, sweet and savory snacks, fruit snacks, chips/crisps, extruded snacks, tortilla/corn chips, popcorn,
 5 pretzels, nuts, other sweet and savory snacks, snack bars, granola bars, breakfast bars, energy bars, fruit bars, other snack bars, meal replacement products, slimming products, convalescence drinks, ready meals, canned ready meals, frozen ready meals, dried ready meals, chilled ready meals, dinner mixes, frozen pizza, chilled pizza, soup, canned soup, dehydrated soup, instant soup, chilled
 10 soup, uht soup, frozen soup, pasta, canned pasta, dried pasta, chilled/fresh pasta, noodles, plain noodles, instant noodles, cups/bowl instant noodles, pouch instant noodles, chilled noodles, snack noodles, dried food, dessert mixes, sauces, dressings and condiments, herbs and spices, spreads, jams and preserves, honey, chocolate spreads, nut-based spreads, and yeast-based spreads.

15 Beverages include any drinkable liquid or semi-liquid, including for example flavored water, soft drinks, fruit drinks, coffee-based drinks, tea-based drinks including tea, black tea, green tea, oolong tea, herbal tea, cocoa-based drinks, juice-based drinks (includes fruit and vegetable), milk-based drinks, gel drinks, carbonated or non-carbonated drinks, powdered drinks and alcoholic or non-
 20 alcoholic drinks.

In the preparation of the abovementioned flavour compositions or consumable compositions, there may be used all or any of the standard ingredients found in such compositions, used in art-recognised quantities. Examples of such ingredients include (but are by no means limited to) solvents and co-solvents; surfactants and
 25 emulsifiers; viscosity and rheology modifiers; thickening and gelling agents; preservative materials; pigments, dyestuffs and colouring matters; extenders, fillers and reinforcing agents; stabilisers against the detrimental effects of heat and light, bulking agents, flavoring and flavor-enhancing agents, warming agents, breath fresheners, mouth moisteners, coloring agents, acidulants, buffering agents and
 30 antioxidants.

Suitable bulking agents may include mineral adjuvants, which may serve as fillers and textural agents. Suitable mineral adjuvants include calcium carbonate, magnesium carbonate, alumina, aluminum hydroxide, aluminum silicate, talc, tricalcium phosphate, tricalcium phosphate and the like, which can serve as fillers
 5 and textural agents.

Additional bulking agents (carriers, extenders) suitable for use include monosaccharides, disaccharides, polysaccharides, sugar alcohols, polydextrose, and maltodextrins; minerals, such as calcium carbonate, talc, titanium dioxide, dicalcium phosphate; and combinations thereof.

- 10 Fillers modify the texture and aid processing. Examples of such fillers include magnesium and aluminum silicates, clay, alumina, talc, titanium oxide, cellulose polymers, and the like.

Specific examples of flavouring agents include, but are not limited to natural flavors, artificial flavors, spices, seasonings, and the like. Exemplary flavoring
 15 agents include synthetic flavor oils and flavoring aromatics and/or oils, oleoresins, essences, distillates, and extracts derived from plants, leaves, flowers, fruits, and so forth, and a combination comprising at least one of the foregoing.

Flavour oils include spearmint oil, cinnamon oil, oil of wintergreen (methyl salicylate), peppermint oil, Japanese mint oil, clove oil, bay oil, anise oil, eucalyptus
 20 oil, thyme oil, cedar leaf oil, oil of nutmeg, allspice, oil of sage, mace, oil of bitter almonds, and cassia oil; useful flavoring agents include artificial, natural and synthetic fruit flavors such as vanilla, and citrus oils including lemon, orange, lime, grapefruit, yuzu, sudachi, and fruit essences including apple, pear, peach, grape, blueberry, strawberry, raspberry, cherry, plum, prune, raisin, cola, guarana, neroli,
 25 pineapple, apricot, banana, melon, apricot, ume, cherry, raspberry, blackberry, tropical fruit, mango, mangosteen, pomegranate, papaya and so forth. Additional exemplary flavors imparted by a flavoring agent include a milk flavor, a butter flavor, a cheese flavor, a cream flavor, and a yogurt flavor; a vanilla flavor; tea or coffee flavors, such as a green tea flavor, an oolong tea flavor, a tea flavor, a
 30 cocoa flavor, a chocolate flavor, and a coffee flavor; mint flavors, such as a peppermint flavor, a spearmint flavor, and a Japanese mint flavor; spicy flavors,

such as an asafetida flavor, an ajowan flavor, an anise flavor, an angelica flavor, a fennel flavor, an allspice flavor, a cinnamon flavor, a chamomile flavor, a mustard flavor, a cardamom flavor, a caraway flavor, a cumin flavor, a clove flavor, a pepper flavor, a coriander flavor, a sassafras flavor, a savory flavor, a Zanthoxyli Fructus flavor, a perilla flavor, a juniper berry flavor, a ginger flavor, a star anise flavor, a horseradish flavor, a thyme flavor, a tarragon flavor, a dill flavor, a capsicum flavor, a nutmeg flavor, a basil flavor, a marjoram flavor, a rosemary flavor, a bayleaf flavor, and a wasabi (Japanese horseradish) flavor; a nut flavor such as an almond flavor, a hazelnut flavor, a macadamia nut flavor, a peanut flavor, a pecan flavor, a pistachio flavor, and a walnut flavor; alcoholic flavors, such as a wine flavor, a whisky flavor, a brandy flavor, a rum flavor, a gin flavor, and a liqueur flavor; floral flavors; and vegetable flavors, such as an onion flavor, a garlic flavor, a cabbage flavor, a carrot flavor, a celery flavor, mushroom flavor, and a tomato flavor.

Generally any flavoring or food additive such as those described in Chemicals Used in Food Processing, publication 1274, pages 63-258, by the National Academy of Sciences, can be used. This publication is incorporated herein by reference.

There may additionally be present in nutraceuticals and medicaments.

Suitable nutraceuticals may include herbs and botanicals such as aloe, bilberry, bloodroot, calendula, capsicum, chamomile, cat's claw, echinacea, garlic, ginger, ginkgo, goldenseal, various ginseng, green tea, guarana, kava kava, lutein, nettle, passionflower, rosemary, saw palmetto, St. John's wort, thyme, and valerian. Also included are mineral supplements such as calcium, copper, iodine, iron, magnesium, manganese, molybdenum, phosphorous, zinc, and selenium. Other nutraceuticals may include fructooligosaccharides, glucosamine, grapeseed extract, cola extract, guarana, ephedra, inulin, phytosterols, phytochemicals, catechins, epicatechin, epicatechin gallate, epigallocatechin, epigallocatechin gallate, isoflavones, lecithin, lycopene, oligofructose, polyphenols, flavanoids, flavanols, flavonols, and psyllium as well as weight loss agents such as chromium picolinate and phenylpropanolamine. Exemplary vitamins and co-enzymes include water or fat soluble vitamins such as thiamin, riboflavin, nicotinic acid, pyridoxine,

pantothenic acid, biotin, folic acid, flavin, choline, inositol and para-aminobenzoic acid, carnitine, vitamin C, vitamin D and its analogs, vitamin A and the carotenoids, retinoic acid, vitamin E, vitamin K, vitamin B6, and vitamin B12. Combinations comprising at least one of the foregoing nutraceuticals may be used.

- 5 Suitable medicaments may include oral care agents, throat care agents, allergy relief agents, and general medical care agents. General medical care agents may include antihistamines, decongestants (sympathomimetics), antitussives (cough suppressants), antiinflammatories, homeopathic agents, expectorants, anesthetics, demulcents, analgesics, anticholinergics, throat- soothing agents, antibacterial
- 10 agents, antiviral agents, antifungal agents, antacids, antinauseants, chemotherapeutics, diuretics, psychotherapeutic agents, cardiovascular agents, various alkaloids, laxatives, appetite suppressants, ACE-inhibitors, anti-asthmatics, anti-cholesterolemics, antidepressants, anti-diarrhea preparations, anti-hypertensives, anti-lipid agents, acne drugs, amino acid preparations, anti-uricemic
- 15 drugs, anabolic preparations, appetite stimulants, bone metabolism regulators, contraceptives, endometriosis management agents, enzymes, erectile dysfunction therapies such as sildenafil citrate, fertility agents, gastrointestinal agents, homeopathic remedies, hormones, motion sickness treatments, muscle relaxants, osteoporosis preparations, oxytocics, parasympatholytics, parasympathomimetics,
- 20 prostaglandins, respiratory agents, sedatives, smoking cessation aids such as bromocriptine or nicotine, tremor preparations, urinary tract agents, anti-ulcer agents, anti-emetics, hyper- and hypoglycemic agents, thyroid and anti-thyroid preparations, terine relaxants, erythropoietic drugs, mucolytics, DNA and genetic modifying drugs, and nutritional supplements, including nutraceuticals,
- 25 micronutrients, vitamins and co-enzymes. The pharmaceutically acceptable salts and prodrugs of the medicaments are also included unless specified otherwise. Some of these medicaments may serve more than one purpose. Combinations of the foregoing types of optional medicaments can be used. Two or more medicaments that have activity against the same or different symptoms can be
- 30 used together in a combination.

Other suitable and desirable additives are described in standard texts, such as "Handbook of Industrial Chemical Additives", ed. M. and I. Ash, 2nd Ed., (Synapse 2000).

One can prepare the mixtures according to the present invention by means of
5 extraction from the stevia plant according to techniques well known in the art.

Generally, steviol glycosides can be extracted from the leaves of stevia plant with water or alcohols (ethanol or methanol). The leaves may be first crushed into small pieces and then extracted with hot water. Generally this procedure is repeated at least twice. The extraction can be carried out at room temperature but preferably
10 at an elevated temperature, up to 95 degrees centigrade. The extraction time can be from 2 to 10 hours depending on the extraction conditions. When under hot water extraction, 2-4 hours would suffice. After filtration to remove the plant residue, the solution can be passed through an adsorption resin column. After washing with deionized water, the steviol glycosides can be eluted from the resin
15 by ethanol or methanol, or an alcohol-water solution. The extract may then be filtered, desalinated, and concentrated before it being de-colored with activated carbon or ion exchange resin. This refined solution can be either vacuum dried or spray dried to furnish an off-white powder with steviol glycoside content of 95% or higher.

20 A typical composition of steviol glycosides generated through the above process consists of stevioside as the major ingredient followed by rebaudioside A, rebaudioside C, dulcoside A along with other minor steviol glycosides.

The mixture of stevioside/rebaudioside A according to the present invention can be generated from a mixture of steviol glycosides purification methods generally
25 known in the art. An example of a suitable method is reversed-phased chromatography. Reversed-phased chromatography refers to any chromatographic methods that use a non-polar stationary phase, in which the binding of mobile phase solute to an immobilised n-alkyl hydrocarbon occurs via hydrophobic interaction. The most common reversed-phase material is octadecyl
30 carbon chain (C18) bonded silica, followed by C-8 bonded silica. Other stationary phases that can also be employed are polystyrene divinylbenzene polymer based

resin, such as Dow AMBERCHROM™ chromatographic grade resin or Mitsubishi Chemical synthetic absorbents such as DIAION HP-20. The mobile phase usually consists of water and water-miscible organic solvents. The most commonly used organic solvents are methanol, acetonitrile, ethanol and isopropanol. Elution can be isocratic, gradient or step-wise increasing the percentage of organic solvents. The pH of the mobile phase is usually kept at neutral.

The temperature can impact on the elution time of steviol glycosides. In a particular embodiment of the invention, the temperature is held constant within a range of 30 to 60 degrees centigrade. Chromatographic separation can be monitored by UV at 210 nm or evaporative light scattering detector (ELSD) and the fractions containing a desired mixture of stevioside/rebaudioside A can be combined accordingly.

Through such chromatographic procedures, the desired mixture of stevioside/rebaudioside A can be produced and the minor steviol glycosides, such as rebaudioside C and dulcoside A, which can contribute bitterness off-taste at above certain levels, can be removed.

Alternatively, the stevioside/rebaudioside A mixture according to the present invention can also be produced from a steviol glycosides mixture using crystallization processes.

In a particular embodiment, a steviol glycoside extract is dissolved in 50-90% methanol or ethanol-water at a temperature in the range of 60-90 degrees centigrade under stirring for 1-3 hours. After the complete solubilization of the extract, the solution is kept at room temperature overnight. The precipitate formed can be filtered and dried. The precipitate can contain upwards of 80% stevioside, with lower amounts of rebaudioside A and a trace of rebaudioside C (below 1%).

Addition of appropriate amounts of rebaudioside A, or a mixture rich in rebaudioside A, to this precipitate can be made in order to arrive at mixtures of the present invention.

A specific example to generate a mixture of stevioside/rebaudioside A is given in Example 1, below.

However, the mixtures according to the present invention can also be prepared simply by admixing in the required ratios the single ingredients stevioside and rebaudioside A. Whereas it is counter-intuitive, to mix the potent rebaudioside A in a highly pure form with stevioside when the latter is regarded as being of lower
 5 potency, the advantage of using pure stevioside and rebaudioside A to produce a mixture consisting essentially of these two individual ingredients within the aforementioned ratio will reside in constant and reproducible sweetness enhancing effects that can be assured by mixing single ingredients.

Rebaudioside A may be employed in a mixture according to the invention in a
 10 purity of greater than 80 %, more particularly greater than 90 %, still more particularly greater than 97 %, and more particular still greater than 99 %. Commercially available rebaudioside A products are well known to a person skilled in the art and not further discussion is merited here.

Similarly stevioside may be employed in a mixture according to the invention in a
 15 purity of greater than 80 %, more particularly greater than 90 %, still more particularly greater than 95 %. Once again, commercially available stevioside products are well known to a person skilled in the art and not further discussion is merited here.

There now follows a series of examples that serve to illustrate the invention.

20 Example 1:

Preparation of a mixture of stevioside and rebaudioside A

The commercial steviol glycoside extract marketed as KAL brand by Nutraceutical Corp. was purified using a Flash chromatography system made by Biotage. The cartridge employed was C-18 40+ M and gradient steps with increasing methanol
 25 content in water as shown below. Flow rate 30 mL/min. UV detection at 210 nm and 254 nm. The eluent was collected into test tubes (27 mL in each).

	MeOH in water (%)	Flow Rate (mL/min)	Volume (mL)

Step 1	40	30	264
Step 2	40-80	30	1320
Step 3	80	30	264
Step 4	100	30	264

The chromatographic fractions were analyzed by analytical HPLC and combined accordingly. The fractions were pooled and lyophilized after removing the methanol. ^1H NMR and HPLC analyses indicated that the combined fraction
 5 consisted of a mixture of stevioside and rebaudioside A in a ratio of 3:1.

Initial tasting indicated that the above mixture had more upfront sweetness, had more body and had a less lingering profile.

Example 2:

Comparison of the upfront sweetness of stevioside/rebaudioside A (3:1) and (2:1)
 10 with stevioside and rebaudioside A.

Two mixtures, stevioside/rebaudioside A (3:1) and (2:1) were evaluated for their upfront sweetness in comparison with pure stevioside and rebaudioside A. Sensory evaluation was carried out with 5 experienced tasters. 30 ppm of stevioside/rebaudioside A (3:1), (2:1), and Rebaudioside A on top of 5% sucrose
 15 solution were prepared along with 20, 35, 45, 75 ppm stevioside in 5% sucrose solutions. Samples were prepared fresh and at room temperature and were paired and tasted with water rinsing in-between.

Stevioside/rebaudioside A (3:1) was ranked with the most upfront sweetness followed by stevioside/rebaudioside A (2:1), rebaudioside A, steviosides. The
 20 sample containing 75 ppm stevioside was ranked as the sweetest but not the most upfront one. The pure stevioside samples were described as delayed onset and lack of the upfront sweetness. Therefore, stevioside and rebaudioside A individually can not provide the desire upfront sweetness, body, less lingering as the mixtures do.

Example 3:

Sweetness threshold determination of stevioside/rebaudioside A (3:1) and (2:1)

Stevioside/rebaudioside A (3:1) and (2:1) were prepared at 30, 35, 40, and 45 ppm levels and evaluated by 4 experienced tasters in water for their iso-intensities to
 5 sucrose solutions (0.5, 1.0, and 1.5% sucrose). Samples and references were paired and tasted with rinsing of the mouth with water in-between.

It was found that 40-45 ppm level of stevioside/Rebaudioside A (3:1) and 35-40 ppm of stevioside/Rebaudioside A (2:1) was close to 1% sucrose, the sweetness threshold level. The exact sucrose equivalence levels around the sweetness
 10 threshold were determined as shown below in Examples 4 and 5.

Example 4:

Ranking Test of 30 and 45 ppm stevioside/rebaudioside A (3:1) in water, determining its sucrose iso-intensity using a formal sensory panel.

The sensory evaluation was conducted using a ranking method. Samples at
 15 ambient temperature were randomly presented in 20 ml blind aliquots (unidentifiable by panelists). Panels consisted of 20 sweet sensitive subjects and samples were presented in 2 replications over 1 session. After tasting each sample, the mouth was rinsed thoroughly with water at ambient temperature prior to tasting the next sample. Panelists were presented with 0.5%, 1.0%, and 1.5%
 20 sucrose solutions in water and two samples of stevioside/rebaudioside A at 30 and 45 ppm in water, respectively. Subjects were asked to rank the samples from low to high with respect to perceived sweet taste. R-indices were calculated for 30 and 45 ppm stevioside/rebaudioside A samples versus either 0.5%, 1.0% or 1.5% sucrose.

25 Stevioside/rebaudioside A (3:1) at 30 ppm in water

Sample sweetness	Sucrose solutions [% wt/wt]	R-index [%]	Critical Value Range [%]	p-value
sweeter	0.5%	19.50	37.76-62.24	P<0.05

less sweet	1.0%	66.62	37.76-62.24	P<0.05
less sweet	1.5%	96.12	37.76-62.24	P<0.05

An R-index 19.50%, which is below the critical value (37.76), means that the 30 ppm stevioside/rebaudioside A (3:1) sample is more sweet than 0.5% sucrose. An R-index of 66.62% or 96.12%, which is above the critical value (62.24%), means
 5 the sample is significantly less sweet than 1.0% or 1.5% sucrose.

By interpolation, the sweetness of 30 ppm stevioside/rebaudioside A (3:1) is equivalent to about 0.75% sucrose.

Stevioside/rebaudioside A (3:1) at 45 ppm

Sample sweetness	sucrose solutions [% wt/wt]	R-index [%]	Critical Value Range [%]	p-value
sweeter	0.5%	1.94	37.76-62.24	P<0.05
sweeter	1.0%	19.94	37.76-62.24	P<0.05
less sweet	1.5%	70.88	37.76-62.24	P<0.05

10 An R-index 1.94 and 19.94%, which is below the critical value (37.76), means that the 45 ppm stevioside/rebaudioside A (3:1) sample is more sweet than 0.5% and 1.0% sucrose solutions. An R-index of 70.88%, which is above the critical value (62.24%), means the 45 ppm stevioside/rebaudioside A (3:1) sample is significantly less sweet than 1.5% sucrose.

15 By interpolation, the sweetness of 45 ppm stevioside/rebaudioside A (3:1) is equivalent to about 1.25% sucrose.

Example 5:

Ranking Test of 35 and 40 ppm stevioside/rebaudioside A (3:1) in water, determining its sucrose iso-intensity using a formal sensory panel.

The sensory evaluation was conducted using the same method as described in Example 4. Panels consisted of 20 sweet sensitive subjects and samples were presented in 2 replications over 1 session. Panelists were presented with 20 mL of 0.5%, 1.0%, and 1.5% sucrose solutions in water and two samples of

5 stevioside/rebaudioside A (3:1) at 35 and 40 ppm in water, respectively. Subjects were asked to rank the samples from low to high with respect to perceived sweet taste. R-indices were calculated for 35 and 40 ppm stevioside/rebaudioside A samples versus either 0.5%, 1.0% or 1.5% sucrose.

Stevioside/rebaudioside A (3:1) at 35 ppm in water

Sample sweetness	Sucrose solutions [% wt/wt]	R-index [%]	Critical Value Range [%]	p-value
sweeter	0.5%	16.50	37.76-62.24	P<0.05
Iso-sweet	1.0%	54.74	37.76-62.24	P<0.05
less sweet	1.5%	92.75	37.76-62.24	P<0.05

10

An R-index 16.50%, which is below the critical value (37.76), means that the 35 ppm stevioside/rebaudioside A (3:1) sample is more sweet than 0.5% sucrose. An R-index of 54.74%, which is between the critical values (37.76-62.24%), means the 35 ppm stevioside/rebaudioside A (3:1) sample is iso-sweet to 1.0% sucrose. An

15 R-index of 92.75%, which is above the critical value (62.24), means the sample is significantly less sweet than 1.5% sucrose in water.

Stevioside/rebaudioside A (3:1) at 40 ppm

Sample sweetness	sucrose solutions [% wt/wt]	R-index [%]	Critical Value Range [%]	p-value
sweeter	0.5%	8.06	37.76-62.24	P<0.05
sweeter	1.0%	36.78	37.76-62.24	P<0.05
less sweet	1.5%	86.13	37.76-62.24	P<0.05

An R-index 8.06 and 36.78%, which is below the critical value (37.76), means that the 40 ppm stevioside/rebaudioside A (3:1) sample is more sweet than 0.5% and 1.0% sucrose solutions. An R-index of 86.13%, which is above the critical value
 5 (62.24%), means the 40 ppm stevioside/rebaudioside A (3:1) sample is significantly less sweet than 1.5% sucrose.

By interpolation, the sweetness of 40 ppm stevioside/Rebaudioside A (3:1) is equivalent to about 1.25% sucrose.

Example 6:

- 10 Iso-intensity of 35 ppm stevioside/rebaudioside A (3:1) in 5.0% sucrose solution using a formal sensory panel

The sensory evaluation was conducted using the same method as described in Example 4. Panels consisted of 20 sweet sensitive subjects and samples were presented in 2 replications over 1 session. Panelists were presented with 20 mL of
 15 5.5%, 6.0%, 6.5%, 7.0%, 8.0% sucrose solutions and one sample of 35 ppm stevioside/rebaudioside A in 5.0% sucrose, respectively. Subjects were asked to rank the samples from low to high with respect to perceived sweet taste. R-indices were calculated for 35 ppm stevioside/rebaudioside A in 5.0% sucrose sample versus either 6.0%, 6.5%, 7.0% or 8.0% sucrose.

- 20 Sweetness enhancing effect for 35 ppm stevioside/rebaudioside A (3:1) in 5% sucrose

Sample sweetness	sucrose solutions [% wt/wt]	R-index [%]	Critical Value Range [%]	p-value
sweeter	6.0%	23.47	37.76-62.24	P<0.05
sweeter	6.5%	31.91	37.76-62.24	P<0.05
Iso-sweet	7.0%	60.56	37.76-62.24	P<0.05
less sweet	8.0%	91.56	37.76-62.24	P<0.05

An R-index 23.47% and 31.91%, which is below the critical value (37.76), means that the 35 ppm stevioside/rebaudioside A (3:1) in 5.0% sucrose sample is more sweet than 6.0% and 6.5% sucrose. An R-index of 60.56%, which is between the critical values (37.76-62.24%), means the sample is iso-sweet to 7.0%. An R-index of 91.56%, which is above the critical value (62.24), means the sample is significantly less sweet than 8.0% sucrose in water. 35 ppm stevioside/rebaudioside A (3:1) in water was determined to be iso-sweet to 1.0% sucrose (Example 5). 35 ppm stevioside/rebaudioside A (3:1) in 5.0% sucrose tasted significantly sweeter than 6% sucrose, which is the sucrose concentration that the 35 ppm stevioside/rebaudioside A (3:1) in 5% sucrose sample would be expected to taste iso-sweet to, assuming a merely additive effect. However, 35 ppm stevioside/rebaudioside A (3:1) on top of 5% sucrose was determined to be iso-sweet to a 7% sucrose solution.

15

Accordingly, the mixture provided an additional 1.0% sucrose equivalence, clearly showing a sweetness enhancing effect.

Example 7:

20 Stevioside iso-intensity

Two ounces of each sample was presented in random order to 20 sweet sensitive panelists. In two replications (over 1 session), Panelists were asked to rank the samples from least sweet to most sweet. The data were subjected to an R-index analysis.

Solutions compared	R-index	Critical Value Range (Non-significant range)	Significantly different ($p < 0.05$)	More Sweet Sample
30 ppm Stevioside vs. 0.5% sucrose	25.625	37.76% to 62.24%	Yes	30 ppm Stevioside
30 ppm Stevioside vs. 1.0% sucrose	66.938	37.76% to 62.24%	Yes	1.0% sucrose

30 ppm Stevioside vs. 1.5% sucrose	94.750	37.76% to 62.24%	Yes	1.5% sucrose
40 ppm Stevioside vs. 0.5% sucrose	11.375	37.76% to 62.24%	Yes	40 ppm Stevioside
40 ppm Stevioside vs. 1.0% sucrose	29.094	37.76% to 62.24%	Yes	40 ppm Stevioside
40 ppm Stevioside vs. 1.5% sucrose	82.219	37.76% to 62.24%	Yes	1.5% sucrose
30 ppm Stevioside vs. 40 ppm Stevioside	80.188	37.76% to 62.24%	Yes	40 ppm Stevioside

30 ppm Stevioside was perceived to be more sweet than 0.5% sucrose in water and significantly less sweet than 1.0% and 1.5% sucrose in water (the calculated R-index values exceeding the critical values at $p < 0.05$). From these results it was
 5 determined that 30 ppm Stevioside was perceived to be isosweet to 0.75% sucrose by interpolation.

40 ppm Stevioside was perceived to be more sweet than 0.5% and 1.0% sucrose in water and significantly less sweet than 1.5% sucrose in water (the calculated R-index values exceeding the critical values at $p < 0.05$). From these results it can be
 10 assumed that 40 ppm Stevioside was perceived to be isosweet to 1.25% sucrose by interpolation. 30 ppm Stevioside was perceived to be significantly less sweet than 40 ppm Stevioside in water (the calculated R-index values exceeding the critical values at $p < 0.05$).

From these data it was determined that 35 ppm Stevioside is isosweet to 1.0%
 15 sucrose.

Example 8:

35 ppm stevioside iso-intensity in 5 % sucrose solution

The objective of this test was to determine the extent to which stevioside enhances sweetness perception of a 5 % sucrose solution.

compared	R-index	Critical Value Range (Non-significant range)	Significantly different ($p < 0.05$)	More Sweet Sample
5% sucrose + 35 ppm Stevioside vs. 5.0% sucrose	8.281	37.76% to 62.24%	Yes	5% sucrose + 35 ppm Stevioside
5% sucrose + 35 ppm Stevioside vs. 6.0% sucrose	21.094	37.76% to 62.24%	Yes	5% sucrose + 35 ppm Stevioside
5% sucrose + 35 ppm Stevioside vs. 7.0% sucrose	78.125	37.76% to 62.24%	Yes	7.0% sucrose

5

5% sucrose + Stevioside was perceived to be significantly more sweet than 5.0% and 6.0% sucrose and significantly less sweet than 7.0% sucrose (the calculated R-index values exceeding the critical values at $p < 0.05$). From these data it can be derived that 5% sucrose + Stevioside is perceived to be iso-sweet to 6.5% sucrose in water by interpolation. As we see in in Example 7, 35 ppm Stevioside was shown to be isosweet to 1.0% sucrose in water. Accordingly, 35 ppm Stevioside on top of 5% sucrose is showing an additional 0.5% sucrose equivalence, which is less than the enhancing effect observed for the stevioside/rebaudioside A mixtures of the present invention.

15 Example 9:

Rebaudioside A iso-intensity

In an iso-intensity study, 30 ppm rebaudioside A was found to be iso-intense with 1 % sucrose.

This example examined the effect of 30 ppm rebaudioside A on top of 5 % sucrose.

Solutions compared	R-Index	Critical value range	significant	More sweet sample
5% sucrose + 30 ppm RebA vs. 5% sucrose	1.313	37.76% to 62.24%	Yes	5% sucrose + 30 ppm RebA
5% sucrose + 30 ppm RebA vs. 6% sucrose	22.625	37.76% to 62.24%	Yes	5% sucrose + 30 ppm RebA
5% sucrose + 30 ppm RebA vs. 7% sucrose	89.000	37.76% to 62.24%	Yes	7% sucrose

- 5 The data show that 5% Sucrose + 30 ppm rebaudioside A was perceived to be significantly more sweet than 5% Sucrose and significantly less sweet than 7% Sucrose (the calculated R-index values exceeding the critical values at $p < 0.05$). By interpolation, 5% Sucrose + 30 ppm rebaudioside A was isosweet to 6.5% sucrose. Given that 30 ppm rebaudioside A was found to be isosweet to 1.0%
- 10 sucrose, it follows that when placed on top of 5 % sucrose there is a 0.5% sucrose equivalence in enhancement, which is less than the enhancing effect observed for the stevioside/rebaudioside A mixtures of the present invention.

Claims:

1. A mixture consisting essentially of the individual ingredients stevioside and rebaudioside A in a stevioside to rebaudioside A ratio of about 1:1 to about 4:1.
2. A mixture according to claim 1 wherein the ratio is about 2:1 to 3:1.
- 5 3. A mixture according to claim 1 or claim 2 in the form of a stevia extract.
4. A mixture according to claim 1 or claim 2 formed by mixing together stevioside and rebaudioside A in a ratio of 1:1 to 4:1, more particularly 2:1 to 3:1.
5. A consumable composition comprising a mixture according to any of the preceding claims, wherein said mixture is added at a concentration of less than 250
10 ppm, more particularly 150 ppm, or less more particularly 90 ppm or less, more particularly 60 ppm or less, more particularly 45 ppm or less, more particularly 40 or less, more particularly 35 ppm or less, more particularly 30 ppm or less .
6. A consumable composition according to claim 5 wherein the mixture is present in an amount of between 45 ppm and 30 ppm.
- 15 7. A consumable composition according to claim 5 or 6 wherein the mixture is added at a concentration at which it is at or near its sweetness detection threshold.
8. A consumable composition according to claim 7 wherein the mixture is added at a concentration at which it is iso-sweet with less than 1.25 % sucrose solution, more particularly iso-sweet with 1 % sucrose solution.
- 20 9. A consumable composition according to any of the claims 5 to 8 additionally comprising a sweetener.
10. A consumable composition according to claim 9 wherein the sweetener is selected from the group consisting of sweetener is selected from the group consisting of sucrose, fructose, glucose, high fructose corn syrup, corn syrup,
25 xylose, arabinose, rhamnose, erythritol, xylitol, mannitol, sorbitol, inositol, acesulfame potassium, aspartame, neotame, sucralose, and saccharine.

11. A consumable composition according to claim 9 or claim 10 wherein the sweetener is present in an amount of 0.0001 % to 15 % by weight of said composition.
12. A consumable composition according to any of the claims 5 through 11
 5 comprising one or more compounds selected from the group consisting of trilobatin, hesperetin dihydrochalcone glucoside, naringin dihydrochalcone, mogroside V, Luo Han Guo extract, rubusoside, rubus extract, glycyphyllin, isomogroside V, mogroside IV, siamenoside I, neomogroside, mukurozioside IIb, (+)-hernandulcin, 4 β -hydroxyhernandulcin, baiyunoside, phlomisioside I,
 10 bryodulcoside, bryoside bryonoside, abrusosides A-E, cyclocarioside A, cyclocaryoside I, albiziasaponins A-E, glycyrrhizin, araboglycyrrhizin, periandrins I-V, pterocaryosides A and B, osladin, polypodosides A and B, telosmoside A8-18, phyllodulcin, huangqioside E neoastilbin, monatin, 3-acetoxy-5,7-dihydroxy-4'-methoxyflavanone, 2*R*,3*R*-(+)-3-Acetoxy-5,7,4'-trihydroxyflavanone, (2*R*,3*R*)-
 15 dihydroquercetin 3-*O*-acetate, dihydroquercetin 3-*O*-acetate 4'-methyl ether, brazzein, curculin, mabinlin, monellin, neoculin, pentadin, thaumatin or combinations thereof.
13. A method of forming a mixture defined in any of the claims 1 through 4
 comprising the step of mixing stevioside and rebaudioside A in the ratio of 1:1 to 4:1,
 20 more particularly 2:1 to 3:1.
14. A method of sweetening a consumable composition according to any of the claims 5 through 12, comprising the step of adding thereto a mixture as defined in any of the claims 1 through 4 at a concentration of less than 250 ppm, more particularly 150 ppm, or less more particularly 90 ppm or less, more particularly 60
 25 ppm or less, more particularly 45 ppm or less, more particularly 40 or less, more particularly 35 ppm or less, more particularly 30 ppm or less .
15. A method according to claim 14 wherein the mixture is present in an amount of between 45 ppm and 30 ppm.
16. A method according to claim 14 or 15 wherein the mixture is added at a
 30 concentration at which it is at or near its sweetness detection threshold.

17. A method according to claim 14 through 16 wherein the mixture is added at a concentration at which it is iso-sweet with less than 1.25 % sucrose solution, more particularly iso-sweet with 1 % sucrose solution.