

University of Groningen

## Methods and compositions to support the growth or maintenance of oxygen-sensitive bacteria in the gastrointestinal tract

Gabarrini, Giorgio; Harmsen, Hermanus Jozef Martinus; Sadaghian Sadabad, Mehdi; Dijn, van, Jan Maarten

**IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.**

*Document Version*

Publisher's PDF, also known as Version of record

*Publication date:*  
2019

[Link to publication in University of Groningen/UMCG research database](#)

*Citation for published version (APA):*

Gabarrini, G., Harmsen, H. J. M., Sadaghian Sadabad, M., & Dijn, van, J. M. (2019). Methods and compositions to support the growth or maintenance of oxygen-sensitive bacteria in the gastrointestinal tract. (Patent No. WO2019234150).

### Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

### Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.



(51) International Patent Classification:

A61K 31/375 (2006.01) A61K 9/20 (2006.01)  
A61K 31/525 (2006.01) A61K 31/661 (2006.01)

(21) International Application Number:

PCT/EP2019/064781

(22) International Filing Date:

06 June 2019 (06.06.2019)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

18176697.3 08 June 2018 (08.06.2018) EP

(71) Applicants: **DSM IP ASSETS B.V.** [NL/NL]; Het Overloon 1, 6411 TE HEERLEN (NL). **ACADEMISCH ZIEKENHUIS GRONINGEN** [NL/NL]; Hanzeplein 1, 9713 GZ Groningen (NL). **RIJKSUNIVERSITEIT GRONINGEN** [NL/NL]; Broerstraat 5, 9712 CP Groningen (NL).

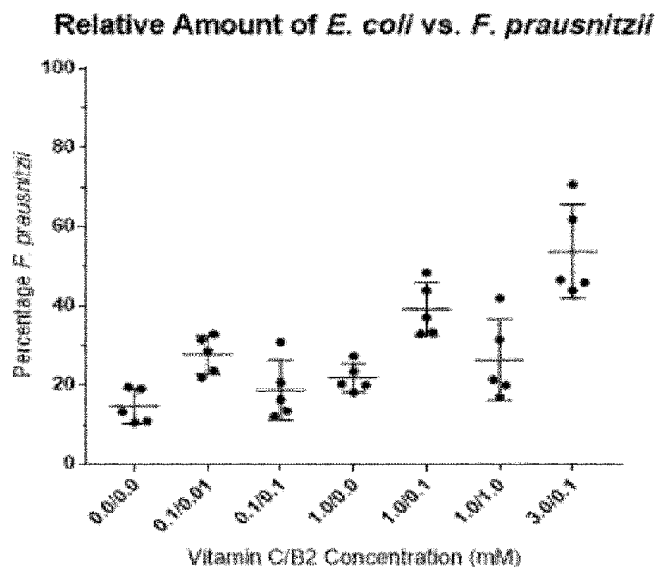
(72) Inventors: **GABARRINI, Giorgio**; c/o University Medical Center Groningen, PO Box 30001, 9700 RB Groningen (NL). **HARMSEN, Hermanus Jozef Martinus**; c/o University Medical Center Groningen, PO Box 30001, 9700 RB Groningen (NL). **SADAGHIAN SADABAD, Mehdi**; c/o University Medical Center Groningen, PO Box 30001, 9700 RB Groningen (NL). **VAN DIJL, Jan Maarten**; c/o University Medical Center Groningen, PO Box 30001, 9700 RB Groningen (NL).

(74) Agent: **KURT, Manfred**; DSM Nutritional Products Ltd, Patent Department, Wurmisweg 576, 4303 Kaiseraugst (CH).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA,

(54) Title: METHODS AND COMPOSITIONS TO SUPPORT THE GROWTH OR MAINTENANCE OF OXYGEN-SENSITIVE BACTERIA IN THE GASTROINTESTINAL TRACT

Figure 2



(57) Abstract: The present invention relates to methods and compositions to support the growth or maintenance of oxygen-sensitive bacteria in the gastrointestinal tract of an animal, preferably a mammal. Particularly, the invention relates to means and methods for selectively enhancing the growth of beneficial anaerobic butyrate-producing bacteria, such as Faecalibacterium prausnitzii (F. prausnitzii).



SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,  
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

**(84) Designated States** (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

**Declarations under Rule 4.17:**

- *as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))*
- *of inventorship (Rule 4.17(iv))*

**Published:**

- *with international search report (Art. 21(3))*

**Methods and Compositions to Support the Growth or Maintenance of Oxygen-sensitive Bacteria in the Gastrointestinal Tract**

5 The present invention relates to methods and compositions to support the growth or maintenance of oxygen-sensitive bacteria in the gastrointestinal tract of an animal, preferably a mammal. Particularly, the invention relates to means and methods for selectively enhancing the growth of beneficial anaerobic butyrate-producing bacteria, such as *Faecaliumbacterium prausnitzii* (*F. prausnitzii*).

10 A normal gastrointestinal microbiota is important to prevent inflammatory disorders such as inflammatory bowel disease and other diseases related to microbiota imbalance.

15 Probiotics are microorganisms that are claimed to provide health benefits when consumed by maintaining a normal host gastrointestinal microbiota. Probiotic formulations have been used as a dietary supplement for many years. So far, many different probiotic strains and combinations thereof exist, but nearly all of them employ relatively-oxygen tolerant strains for instance *Bifidobacterium* sp., *Lactobacillus* sp. and *Saccharomyces* sp.

20

Prebiotics are typically compounds that pass undigested through the upper part of the gastrointestinal tract and stimulate the growth or activity of advantageous bacteria that colonize the large bowel by acting as substrate for them. Typical examples of known prebiotics are oligosaccharides, such as fructooligosaccharides, galactooligosaccharides and inulin. Synbiotics refer to nutritional supplements combining  
25 probiotics and prebiotics in a form of synergism.

- 2 -

Numerous probiotics, prebiotics and symbiotic formulations are known in the art to maintain or stimulate the level of beneficial oxygen-tolerant bacteria in the gut. In contrast, very few studies have dealt with stimulating oxygen-sensitive bacteria.

5 Recent research in gut microbiology explores new horizons for pre- and probiotic applications, such as anti-inflammatory treatments including treatment of Crohn's disease. Beneficial bacteria including *F. prausnitzii* typically utilize a variety of carbohydrates and produce butyrate as a major fermentative end product. Butyrate is well known for its role in promoting and maintaining gut health.

10

In fact, *F. prausnitzii* was found to exhibit anti-inflammatory effects on cellular and colitis models, partly due to secreted metabolites able to block NF- $\kappa$ B activation and IL-8 production. *F. prausnitzii* is one of the most abundant human colon bacteria with numbers ranging from 5-20% of the total microbiota in stools of healthy individuals. It was found that a reduction of *F. prausnitzii* is associated with a higher risk of postoperative recurrence of Crohn's Disease. The current idea is that counterbalancing dysbiosis including low levels of *F. prausnitzii* is a promising strategy for treatment of inflammatory bowel diseases such as Crohn's Disease or any other diseases related to microbiota dysbiosis by preventing reoccurrence of exacerbations.

20

Inflammatory Bowel Disease (IBD) and Irritable Bowel Syndrome (IBS) are a chronic and debilitating illnesses. It is characterized by chronic intestinal inflammation that often shows an intermittent course with acute attacks followed by periods of remission. Clinical symptoms during acute attacks include diarrhea, bleeding, abdominal pain, fever, joint pain, and weight loss. These symptoms can range from mild to severe, and may gradually and subtly develop from an initial minor discomfort, or may present themselves suddenly in full-blown form. IBD can manifest itself in a variety of forms, the most common of which are Crohn's disease and ulcerative colitis. Both of these diseases are similar in terms of clinical symptoms, even though their inflammation patterns are distributed differently in the GI tract. Crohn's disease

30

- 3 -

is a chronic transmural inflammation of the bowel, which can affect the whole gastrointestinal tract, usually in a discontinuous pattern. The initial location of CD is most commonly in the lower ileum. From here the inflammation typically spreads towards proximal parts of the small intestine. However, the colon is also of-ten involved.

Ulcerative colitis is a chronic inflammatory bowel disease affecting only the colon and shows a continuous distribution in the gastrointestinal mucosa. In most patients the focal point of the inflammation is in the distal part of the colon and the rectum. From this origin, the inflammation often spreads proximally. In the most severe cases, the whole colon is affected which is called as "pancolitis". About 30% of patients suffer from this severe form of UC.

*F. prausnitzii* and some other strict anaerobe bacteria needed for the aforementioned anti-inflammatory and other beneficial effects are extremely sensitive to oxygen and cannot survive an exposure to ambient air for more than a few minutes. As a consequence, probiotic compositions containing *F. prausnitzii* have not been described thus far despite their promising therapeutic application.

Recognizing the therapeutic potential of beneficial anaerobic bacteria, the present inventors set out to identify new means and methods to enhance the population of butyrate-producing anaerobic bacteria including *F. prausnitzii* in the gastrointestinal tract. In particular, they aimed at providing novel nutritional ingredient formulations for selectively stimulating butyrate-producing anaerobic bacteria, preferably *F. prausnitzii*.

It is known i.e. that the intake of riboflavin (also known as vitamin B2) has a positive effect on the gut microbiota (maintaining a "normal" gut microbiota or treating gut microbiota, which is "in bad shape").

30

- 4 -

Now, it was surprisingly found that a combination of riboflavin (also known as vitamin B2) and vitamin C (also known as ascorbic acid) in a very specific ratio, is capable of increasing the concentration of *F. prausnitzii*.

This results in a surprisingly positive effect on the gut microbiota. This is very helpful  
5 in treating and/or preventing IBD (such as Crohn's disease and ulcerative colitis)

Therefore the present invention relates to a method for the selective growth of *F. prausnitzii* in the gastrointestinal tract in an animal, comprising administering to the  
10 animal a combination of

- (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and

wherein the ratio of (i) : (ii) is between 1:1 to 50:1

15

Furthermore, the present invention relates to a composition comprising a combination of

- (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt  
20 thereof, and

wherein the ratio of (i) : (ii) is between 1:1 to 50:1.

Furthermore, the present invention relates to a pharmaceutical composition comprising a combination of

- 25 (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and

wherein the ratio of (i) : (ii) is between 1:1 to 50:1 and at least one therapeutically inert carrier.

30

- 5 -

Furthermore, the present invention relates to a pharmaceutical composition comprising a combination of

- (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and

wherein the ratio of (i) : (ii) is between 1:1 to 50:1 for the use as therapeutically active substance for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis..

10 Furthermore, the present invention relates to the use of a pharmaceutical composition comprising a combination of

- (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and

15 wherein the ratio of (i) : (ii) is between 1:1 to 50:1 for the use as therapeutically active substance for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis.

Furthermore, the present invention relates to the use of a pharmaceutical composition comprising a combination of

- (i) vitamin C,
- (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and

20 wherein the ratio of (i) : (ii) is between 1:1 to 50:1 and at least one and at least one  
25 therapeutically inert carrier for the preparation of a medicament for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis.

Furthermore the present invention relates to a method for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis., which



- 6 -

method comprises administering a therapeutically effective amount of a pharmaceutical combination a combination of

- (i) vitamin C,
  - (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt
- 5 thereof, and

wherein the ratio of (i) : (ii) is between 1:1 to 50:1.

All the ratios are related to the amount (weight).

10 The overall amount consumption of vitamin C is around 1 g vitamin C/day. Due to the fact, that there is no recommended daily intake for Vitamin C and Vitamin B2, the amount could also be higher.

It is obvious that the amount could also be lower.

15 The amount in a formulation, which is consumed by a patient, depends on the galenical formulation (it can be any formulation, preferred are solid formulation, like a tablet, a granule, powder, etc). This means a tablet should not be too big to be swallowed and it should also be possible to formulate the amount in a stable application form.

20

Riboflavin, also known as vitamin B2, is a micronutrient with a key role in maintaining health in humans and other mammals. It is the central component of the cofactors FAD and FMN, and is therefore required by all flavoproteins. As such, riboflavin is required for a wide variety of cellular processes. It plays a key role in energy

25 metabolism, and for the metabolism of fats, ketone bodies, carbohydrates, and proteins. Riboflavin is found naturally in asparagus, popcorn, bananas, persimmons, okra, chard, cottage cheese, milk, yogurt, meat, eggs, fish, and green beans. Other sources specify cheese, leafy green vegetables, liver, kidneys, legumes, tomatoes, yeast, mushrooms, and almonds.

30

Vitamin C (also known as ascorbic acid or salts thereof) is commonly used as anti-oxidant food additives.

The vitamin C can be natural (extracted from a plant/fruit) or it can be synthetic one (chemically or biochemically produced). It is clear also mixture of both can be used.

5

In the context of the present invention, the intake of riboflavin and the vitamin C can take place in such ways that

(a) a mixture of both compounds can be consumed or

(b) both compounds can be consumed individually (the sequence is not crucial)

10

As said before component (i) and component (ii) can also be a mixture of different forms (i.e. different salts of the components).

In case the riboflavin and the vitamin C are eaten individually, this should take place in a short period of time.

15 The easiest (and most convenient) way is that a mixture of the (i) and (ii) are consumed.

This can happen any form. This could be a powder of both or the two ingredients (or mixture of the ingredients) can be formulated. They can be incorporated in any desirable formulation suitable for oral intake. The formulation can be liquid or solid.

20 Preferably, it is a food product, pharmaceutical composition, food or dietary supplement.

The invention provides a method for the selective growth of *F. prausnitzii* in the gastrointestinal tract in an animal, e.g. a mammal, in need thereof, comprising administering to the animal a mixture of vitamin C and riboflavin, riboflavin phosphate

25 or a physiologically acceptable salt thereof, in an amount effective to selectively stimulate the growth of *F. prausnitzii* in the gastrointestinal tract.

The animal can be a human, pet or livestock. Thus, veterinary use of the present invention is also encompassed.

Preferably, the animal is a human subject. In one embodiment, the human subject is suffering from an inflammatory gastrointestinal disease, in particular Crohn's disease or a related colitis, or any other diseases related to microbiota dysbiosis. Accordingly, also provided is a method for preventing, treating or reducing the symptoms associated with an inflammatory gastrointestinal disorder, comprising administering to a subject in need thereof an amount of riboflavin effective to maintain, support or stimulate the growth of *F. prausnitzii* in the gastrointestinal tract. Individuals with exemplary inflammatory gastrointestinal disorders, who may benefit from increasing *F. prausnitzii* numbers in the GI tract by riboflavin, include patients with Crohn's disease, inflammatory bowel disease and ulcerative colitis. Also encompassed is the treatment of other diseases, conditions or disorders where patients benefit from restoring or increasing *F. prausnitzii* numbers in the GI tract.

As stated above the gastrointestinal disorders are Crohn's disease, inflammatory bowel disease and ulcerative colitis or any other diseases related to microbiota dysbiosis.

As stated above the formulations, which are described in this patent application can also be in a galenical form.

The galenical formulation can comprise any pharmaceutically acceptable auxiliary agents, which are necessary, needed or desired to form such a galenical formulation.

The galenical formulation can be in any form, which is suitable for patients. Most commonly it is in a solid form (such as a tablet, powder or similar).

By "pharmaceutically acceptable" is meant a material that is not biologically or otherwise undesirable.

Pharmaceutically acceptable excipients include but are not limited to binders, diluents, lubricants, glidants and surface-active agents.

Such pharmaceutically acceptable excipients are used when Mesalamine and riboflavin are integrated into a suitable form for administration.

The amount of additive employed will depend upon how much active agent is to be used. One excipient can perform more than one function.

Binders include, but are not limited to, starches such as potato starch, wheat starch, corn starch; microcrystalline cellulose; celluloses such as hydroxypropyl cellulose, hydroxyethyl cellulose, hydroxypropylmethyl cellulose (HPMC), ethyl cellulose, sodium carboxy methyl cellulose; natural gums like acacia, alginic acid, guar gum; liquid glucose, dextrin, povidone, syrup, polyethylene oxide, polyvinyl pyrrolidone and the like and mixtures thereof.

Fillers or diluents, which include, but are not limited to confectioner's sugar, compressible sugar, dextrans, dextrin, dextrose, fructose, lactitol, mannitol, sucrose, starch, lactose, xylitol, sorbitol, talc, microcrystalline cellulose, calcium carbonate, calcium phosphate dibasic or tribasic, calcium sulphate, and the like can be used.

Lubricants may be selected from, but are not limited to, those conventionally known in the art such as Mg, Al or Ca or Zn stearate, polyethylene glycol, glyceryl behenate, mineral oil, sodium stearyl fumarate, stearic acid, hydrogenated vegetable oil and talc.

Glidants include, but are not limited to, silicon dioxide; magnesium trisilicate, powdered cellulose, starch, talc and tribasic calcium phosphate, calcium silicate, magnesium silicate, colloidal silicon dioxide, silicon hydrogel and other materials known to one of ordinary skill in the art.

The pharmaceutical formulation according to the present invention include but is not limited to tablets (single layered tablets, multilayered tablets, MUPS, mini tablets,

bioadhesive tablets, caplets, matrix tablets, tablet within a tablet, mucoadhesive tablets, modified release tablets, pulsatile release tablets, timed release tablets), pellets, beads, granules, sustained release formulations, capsules, microcapsules, tablets in capsules and microspheres, matrix formulations, microencapsulation and  
5 powder/pellets/granules for suspension.

The galenical formulation of the invention can optionally have one or more coatings such as film coating, sugar coating, enteric coating, bioadhesive coating and other coatings known in the art. These coatings help pharmaceutical formulations to re-  
10 lease the drug at the required site of action. In one example, the additional coating prevents the dosage from contacting the mouth or esophagus. In another example, the additional coating remains intact until reaching the small intestine (e.g., an enteric coating). Premature exposure of a bioadhesive layer or dissolution of a pharmaceutical dosage form in the mouth can be prevented with a layer or coating of  
15 hydrophilic polymers such as HPMC or gelatin. Optionally, Eudragit FS 3OD or other suitable polymer may be incorporated in coating composition to retard the release of the drug to ensure drug release in the colon.

These coating layers comprises one or more excipients selected from the group  
20 comprising coating agents, opacifiers, taste-masking agents, fillers, polishing agents, coloring agents, antitacking agents and the like.

The galenical formulations of the invention can be coated by a wide variety of methods. Suitable methods include compression coating, coating in a fluidized bed or a  
25 pan and hot melt (extrusion) coating. Such methods are well known to those skilled in the art.

Non-permeable coatings of insoluble polymers, e.g., cellulose acetate, ethylcellulose, can be used as enteric coatings for delayed/modified release (DR/MR) by inclusion of soluble pore formers in the coating, e.g., PEG, PVA, sugars, salts, detergents, triethyl citrate, triacetin, etc.

5

Also, coatings of polymers that are susceptible to enzymatic cleavage by colonic bacteria are another means of ensuring release to distal ileum and ascending colon. Materials such as calcium pectinate can be applied as coatings to dosage form and multiparticulates and disintegrate in the lower gastrointestinal tract, due to bacterial  
10 action. Calcium pectinate capsules for encapsulation of bioadhesive multiparticulates are also available.

In embodiments of the present invention, a pharmaceutical combinations of Mesalamine or a pharmaceutically acceptable salt or prodrugs thereof and riboflavin and  
15 at least one swellable polymer. Swellable polymers include, but are not limited to, a crosslinked poly(acrylic acid), a poly(alkylene oxide), a polyvinyl alcohol), a polyvinyl pyrrolidone); a polyurethane hydrogel, a maleic anhydride polymer, such as a maleic anhydride copolymer, a cellulose polymer, a polysaccharide, starch, and starch based polymers.

20

The pharmaceutical compositions of the present invention can optionally include one or more solubilizers, i.e., additives to increase the solubility of the pharmaceutical active ingredient or other composition components in the solid carrier. Suitable solubilizers for, use in the compositions of the present invention include: alcohols  
25 and polyols, such as ethanol, isopropanol, butanol, benzyl alcohol, ethylene glycol, propylene glycol, butanediols and isomers thereof, glycerol, pentaerythritol, sorbitol, mannitol, transcitol, dimethyl isosorbide, polyethylene glycol, polypropylene glycol, polyvinylalcohol, hydroxypropyl methylcellulose and other cellulose derivatives, cyclodextrins and cyclodextrin derivatives; ethers of polyethylene glycols having an  
30 average molecular weight of about 200 to about 6000, such as tetrahydrofurfuryl

- 12 -

alcohol PEG ether (glycofurol, available commercially from BASF under the trade name Tetraglycol) or methoxy PEG (Union Carbide); amides, such as 2-pyrrolidone, 2-piperidone, .epsilon.-caprolactam, N-alkylpyrrolidone, N-hydroxyalkylpyrrolidone, N-alkylpiperidone, N-alkylcaprolactam, dimethylacetamide, and polyvinylpyrrolidone; esters, such as ethyl propionate, tributylcitrate, acetyl triethylcitrate, acetyl tributyl citrate, triethylcitrate, ethyl oleate, ethyl caprylate, ethyl butyrate, triacetin, propylene glycol monoacetate, propylene glycol diacetate, .epsilon.-caprolactone and isomers thereof, .delta.-valerolactone and isomers thereof, .beta.-butyrolactone and isomers thereof; and other solubilizers known in the art, such as dimethyl acetamide, dimethyl isosorbide (Arlasolve DMI (ICI)), N-methyl pyrrolidones (Pharmasolve (ISP)), monooctanoin, diethylene glycol monoethyl ether (available from Gattefosse under the trade name Transcutol), and water.

Preferred solubilizers include triacetin, triethylcitrate, ethyl oleate, ethyl caprylate, dimethylacetamide, N-methylpyrrolidone, N-hydroxyethylpyrrolidone, polyvinylpyrrolidone, hydroxypropyl methylcellulose, hydroxypropyl cyclodextrins, ethanol, polyethylene glycol 200-600, glycofurol, transcutol, propylene glycol, and dimethyl isosorbide. Particularly preferred solubilizers include sorbitol, glycerol, triacetin, ethyl alcohol, SLS, polyethylene glycols glycofurol and propylene glycol. Cyclodextrins polyoxomers, surfactants and like

All formulations as well as the Galenical formulation described and disclosed above can be produced by using well-known methods and processes.

25

### Example

In in vitro experiments, the effect of both vitamin C and vitamin B2 on *Faecalibacterium prausnitzii* separately (Figure 1) as well as on a co-culture of *E. coli* and *F. prausnitzii* (Figure 2); was studied. Inoculation of *F. prausnitzii* was in the anaerobic chamber and *E. coli* was aerobically inoculated. All the growth were aerobic (outside of anaerobic chamber). Optical densities (ODs) were measured before (t=0h) and after (t=24h) incubation. Medium broths with a maximum OD of 2 at t=0h were excluded from analysis. Differences in OD at t=0h and t=24h were graphed to see changes in growth after vitamin intervention. To quantify the growth of each strain in the aerobic co-cultures, a FISH analysis was performed.

For *F. prausnitzii*, the increase in aerobic growth was larger after addition of a combination of vitamin C with low vitamin B2 (0.1 and 1.0 mM) concentrations compared to only vitamin C or vitamin B2, or combination of vitamin C and higher vitamin B2 concentrations (3.0 mM) (Figure 1).

All *F. prausnitzii* and *E. coli* co-culture samples showed a significantly increased percentage of *F. prausnitzii* compared to controls after addition of different vitamin concentration. The highest ratio was with 3.0/0.1 mM (30:1) of vitamin C/Vitamin B2 (Figure 2).



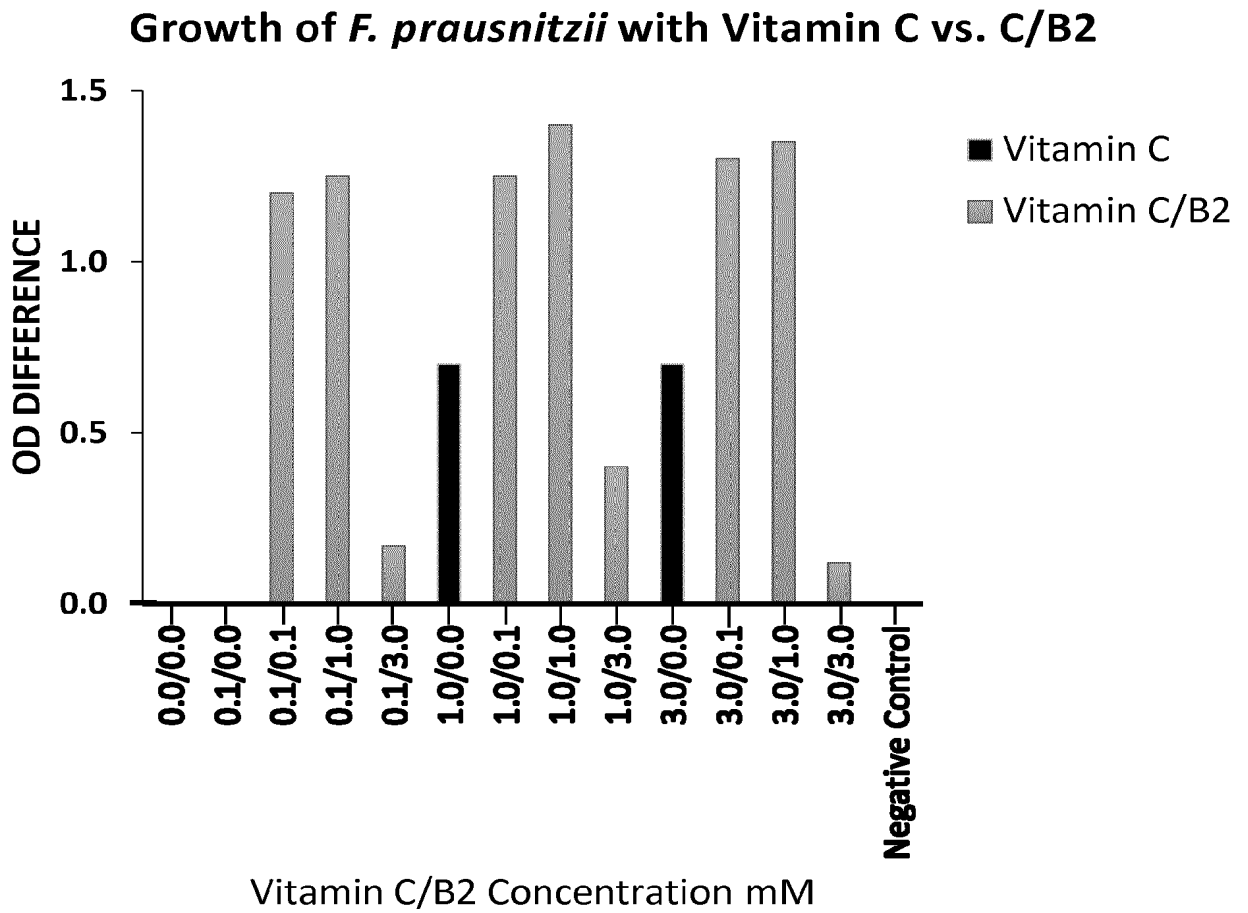
**Claims**

1. A composition comprising a combination of
  - (i) vitamin C,
  - 5 (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, andwherein the ratio of (i) : (ii) is between 1:1 to 50:1.
  
2. Composition according to claim 1, which is a pharmaceutical composition  
10 comprising at least one therapeutically inert carrier.
  
3. Pharmaceutical composition according to claim 2 for the use as therapeutically active substance for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis.  
15
  
4. Composition according to claim 1 or pharmaceutical composition according to claim 2 or claim 3, which is solid.
  
5. Composition according to claim 4, which is a tablet.  
20
  
6. The use of composition according to any of the preceding claims for the use as therapeutically active substance for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis.
  
- 25 7. Use of a composition according to claim 1 or a pharmaceutical composition according to claim 2, 3 or claim 4 for the preparation of a medicament for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis.

- 15 -

8. A method for the treatment or prophylaxis of IBD and/or IBS or any other diseases related to microbiota dysbiosis., which method comprises administering a therapeutically effective amount of a pharmaceutical combination a combination of
- (i) vitamin C,
  - 5 (ii) riboflavin, riboflavin phosphate and/or a physiologically acceptable salt thereof, and
- wherein the ratio of (i) : (ii) is between 1:1 to 50:1.

Figure 1



### Growth of *F. prausnitzii* with Vitamin B2 vs. C/B2

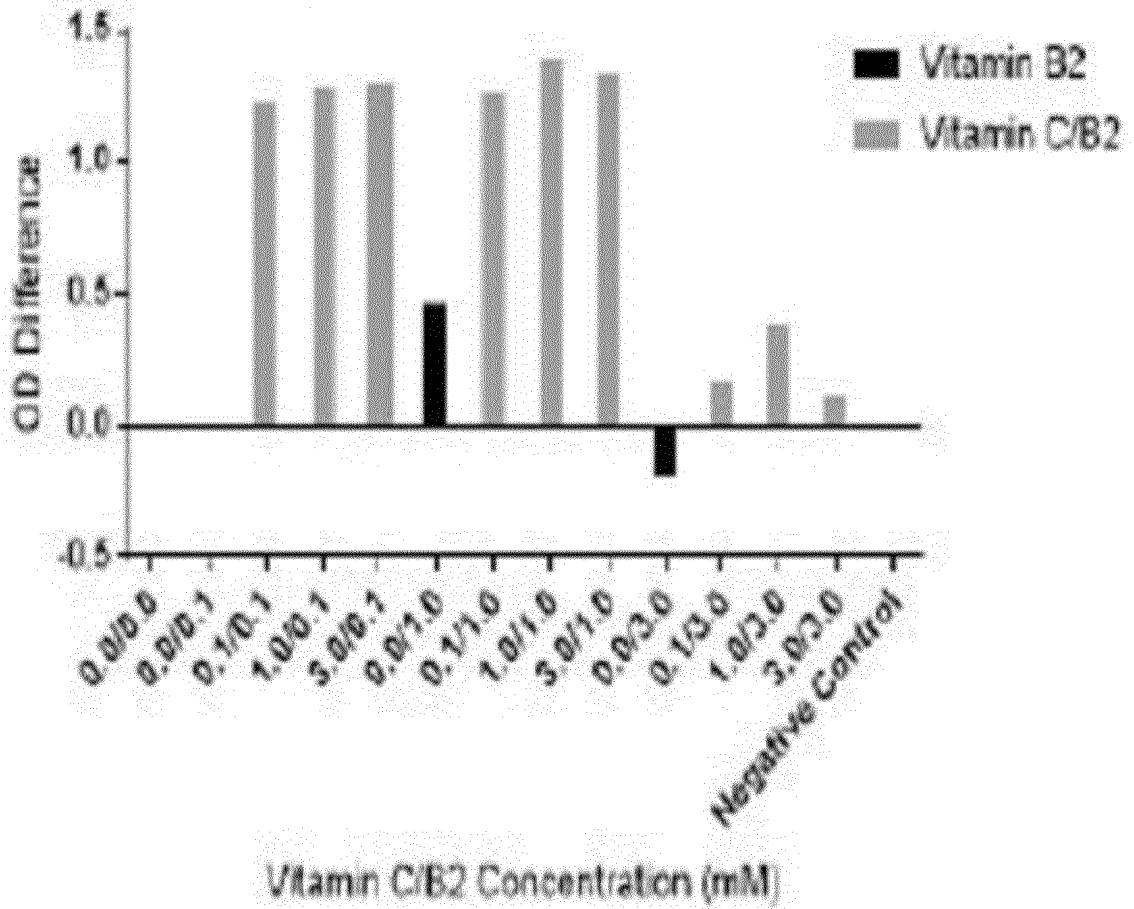
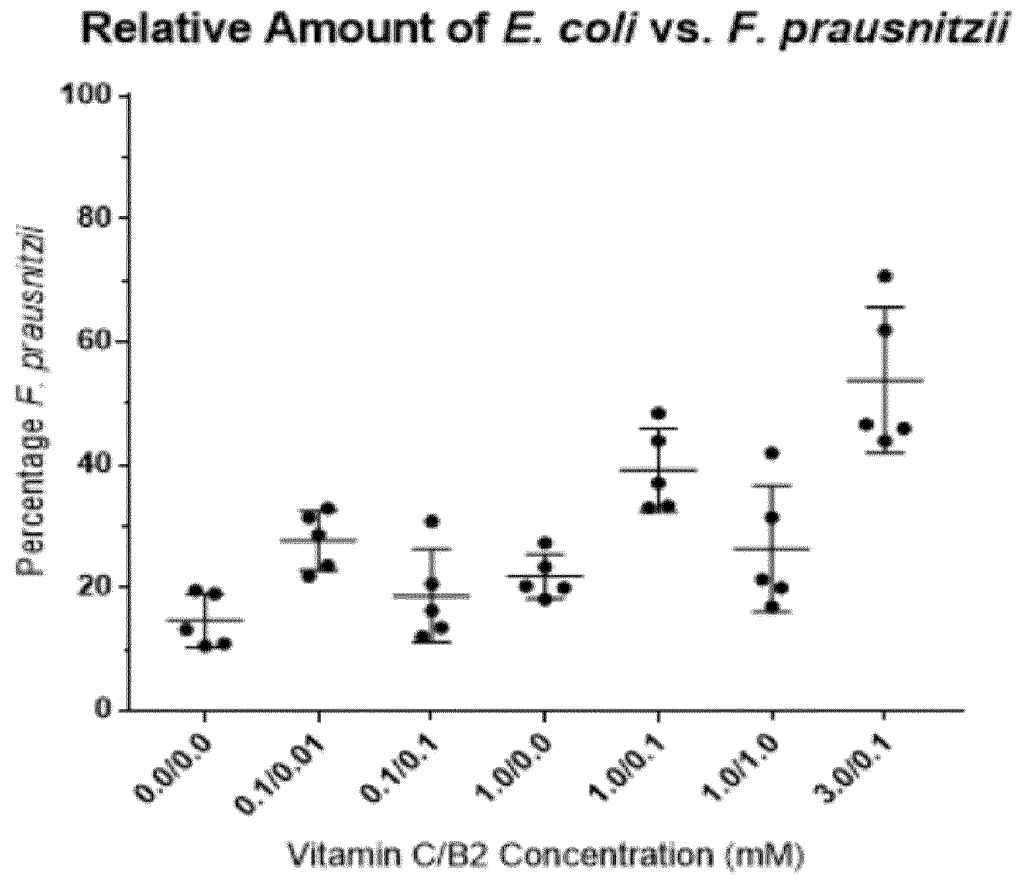


Figure 2



INTERNATIONAL SEARCH REPORT

International application No  
PCT/EP2019/064781

A. CLASSIFICATION OF SUBJECT MATTER  
INV. A61K31/375 A61K31/525 A61K9/20 A61K31/661  
ADD.  
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED  
Minimum documentation searched (classification system followed by classification symbols)  
A61K  
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 3 308 787 A1 (ZENSUN SHANGHAI SCIENCE & TECH CO LTD [CN]) 18 April 2018 (2018-04-18) page 15, line 41 - line 44; claim 8 page 6, line 50, paragraph [0031] - line 58	1-8
X	----- WO 01/24642 A1 (SNOWDEN ROBERT B [US]) 12 April 2001 (2001-04-12) page 12, line 6 - line 10; table 2 page 17; claim 9 page 18; claim 10 page 20; claim 17 ----- -/--	1-8

Further documents are listed in the continuation of Box C.  See patent family annex.

\* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search <b>17 September 2019</b>	Date of mailing of the international search report <b>25/09/2019</b>
---	---

Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer <b>Giró, Annalisa</b>
--	---

## INTERNATIONAL SEARCH REPORT

International application No  
PCT/EP2019/064781

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2017/001703 A1 (NOVOZYMES AS [DK]; DSM IP ASSETS BV [NL]) 5 January 2017 (2017-01-05) page 53, line 3 - page 53, line 3; table 4 page 60 - page 60; table 8 page 57, line 14 - page 57; table 6 page 10, line 11 - line 15 page 10, line 35 - page 11, line 14 -----	1-8
X	HASSN AHMED ET AL: "Medium-Term Results of Combined Laparoscopic Sleeve Gastrectomy and Modified Jejunio-Ileal Bypass in Bariatric Surgery", OBESITY SURGERY, RAPID COMMUNICATIONS OF OXFORD, OXFORD, GB, vol. 26, no. 10, 27 February 2016 (2016-02-27), pages 2316-2323, XP036053192, ISSN: 0960-8923, DOI: 10.1007/S11695-016-2098-Z [retrieved on 2016-02-27] page 2321, column 1, line 27 -----	1-8

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2019/064781

Patent document cited in search report	Publication date	Patent family member(s)	Publication date	
EP 3308787	A1	18-04-2018	AU 2016276482 A1	18-01-2018
			BR 112017026834 A2	14-08-2018
			CA 2989214 A1	15-12-2016
			CN 106265696 A	04-01-2017
			EP 3308787 A1	18-04-2018
			JP 2018521031 A	02-08-2018
			US 2018353469 A1	13-12-2018
			WO 2016197889 A1	15-12-2016
-----				
WO 0124642	A1	12-04-2001	CA 2387787 A1	12-04-2001
			US 6214373 B1	10-04-2001
			WO 0124642 A1	12-04-2001
-----				
WO 2017001703	A1	05-01-2017	AU 2016288515 A1	21-12-2017
			EP 3316703 A1	09-05-2018
			US 2018184688 A1	05-07-2018
			WO 2017001703 A1	05-01-2017
-----				