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(54) ORAL PREPARATION FOR THE PROPHYLACTIC AND THERAPEUTIC TREATMENT OF HELICOBACTER SP. INFECTION

ORALE ZUBEREITUNG ZUR PROPHYLAKTISCHEN/THERAPEUTISCHEN BEHANDLUNG VON HELICOBACTER SP. INFektionEN

PREPARATION ORALE POUR LA PROPHYLAXIE ET LE TRAITEMENT DES INFECTIONS PAR HELICOBACTER

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Description

[0001] The present invention relates to the use of at least one type of xanthophylls for the preparation of a medicament for the prophylactic and/or therapeutic treatment of inflammation in the mucous membrane of mammalian gastrointestinal tract caused by *Helicobacter sp.* infection. The medicament comprises as a xanthophyll, preferably naturally produced astaxanthin.

Background of the invention


[0003] *Helicobacter pylori* can cause drastic changes of the gastric mucous membrane barrier functions in an early infection (i.e. type B gastritis) with breakdown of the hydrophobic lining of the gastric epithelium. This can cause backflow of acid and pepsin from the lumen into the mucosa to cause peptic ulcers in the stomach and duodenum. It seems likely that this breakdown of the mucosa barrier also affects the uptake in the gastric mucosa of a number of substances in food such as certain food-associated carcinogens.

[0004] Xanthophylls, including astaxanthin, is a large group of carotenoids containing oxygen in the molecule in addition to carbon and hydrogen. The carotenoids are produced *de novo* by plants, fungi and some bacteria [Johnson E.A. and Schroeder W.A., 1995, Adv In Biochem Engin. Biotechn. 53: 119-178]. In biological tests astaxanthin has been shown to possess clearly the best antioxidative properties compared to other carotenoids [Miki W., 1991, Pure and Appl Chem 63(1): 141-146].

[0005] At present, the therapeutic treatment of inflammation in the mucous membranes of mammalian gastrointestinal tract caused by *Helicobacter sp.* infection, mainly involves the use of so-called proton pump inhibitors, such as Losec® (omeprazol), and in case of gastric ulcers different antibiotics (which may cause the development of resistant strains).

Description of the invention

[0006] The present invention provides the use of at least one type of xanthophylls for the preparation of a medicament for the prophylactic and/or therapeutic treatment of inflammation in the mucous membrane of mammalian gastrointestinal tract caused by *Helicobacter sp.* infection.

[0007] The medicament prepared according to the invention may comprise a mixture of different types of xanthophylls or different forms of the same xanthophyll, such as a mixture of synthetic astaxanthin and naturally produced astaxanthin.

[0008] In a particular embodiment of the invention the mammalian gastrointestinal tract is the human stomach, and the *Helicobacter sp.* is *H. pylori*.

[0009] The mechanism of the prophylactic and therapeutic effect of the xanthophylls in the treatment of inflammation in the mucous membrane of the mammalian gastrointestinal tract caused by *Helicobacter sp.* infection is not known, but it is believed that the antioxidative properties of the xanthophylls, which are soluble in fat/oil, play an important role in the protection of the hydrophobic lining of the mucous membrane so that *Helicobacter sp.* cannot colonize.

[0010] In a preferred embodiment of the invention, the xanthophyll is dissolved in an oil of food grade.

[0011] In another preferred embodiment the type of xanthophyll is astaxanthin, particularly astaxanthin in a form esterified with fatty acids.

[0012] In yet another preferred embodiment the astaxanthin derives from a natural source, particularly a culture of the alga *Haematococcus sp.* [Rensström B. et al, 1981, Phytochem 20(11) :2561-2564].

[0013] The medicament prepared according to the invention may further comprise carbohydrate structures, such as lipopolysaccharides, polysaccharides and glycoproteins.

[0014] At present, the most preferred embodiment of the invention comprises algal meal having astaxanthin in esterified form with fatty acids dissolved in small droplets of naturally occurring oil and naturally occurring carbohydrate structures in the partially disrupted cell walls.

[0015] The medicament prepared according to the invention is preferably an oral preparation that may comprise additional ingredients which are pharmaceutically acceptable inactive or active in prophylactic and/or therapeutic use, such as flavoring agents, and a prophylactically and/or therapeutically effective amount of a water soluble antioxidant,
especially ascorbic acid (vitamin C).

[0016] The medicament or oral preparation is presented in a separate unit dose or in mixture with food. Examples of separate unit doses are tablets, gelatin capsules and predetermined amounts of solutions, e.g. oil solutions, or emulsions, e.g. water-in-oil or oil-in-water emulsions. Examples of foods in which the preparation of the invention may be incorporated is dairy products, such as yoghurt, chocolate and cereals.

[0017] The daily dosage of the active ingredient in the medicament prepared according to the invention will normally be in the range of 0.01 to 10 mg per kg body weight for a human calculated on the amount of astaxanthin, but the actual dosage will depend on the mammalian species and the individual species-specific biological effect.

Experiments

[0018] The xanthophyll in the oral preparation used in the experiments is astaxanthin which is commercially produced via culturing of the algae Haematococcus sp. by AstaCarotene AB, Gustavsberg, Sweden.

[0019] Astaxanthin from other sources, and other xanthophylls as well, are expected to be similarly useful for the purposes of the invention. An advantage of using astaxanthin from algae is, however, that the astaxanthin exists in a form esterified with fatty acids [Renström B. et al, ibid], which esterified astaxanthin thereby is more stable during handling and storage than free astaxanthin.

[0020] The naturally produced astaxanthin can be obtained also from fungi and crustaceans, in addition to from algae [Johnson E.A. and Schroeder W.A., ibid]. Fifty 6 - 8 weeks old Balb/cA mice weighing 28-30 g were infected with H. pylori by administration of 10^8 cfu in phosphate buffer through a gastric tube into the stomach. The treatment was repeated three times on one-day intervals [Aleljung P., et al., 1996, FEMS Immunol Med Microbiol. 13: 303-309].

[0021] After 14 days, 10 mice were sacrificed and cultures were made on stomach biopsies to isolate H. pylori. The culturing takes seven days.

[0022] Twenty-one days after the infection with H. pylori half of the remaining animals were given feed supplemented by algal meal corresponding to 0.3 mg astaxanthin per animal per day for a period of 10 days.

[0023] On day 30 half of the animals in each group were sacrificed and culturing was made in a similar way as disclosed above.

[0024] On day 40 the rest of the animals were sacrificed and culturing was made in a similar way as disclosed above.

[0025] The results are given in Table 1.

<table>
<thead>
<tr>
<th>Day</th>
<th>Treated animals</th>
<th>Control animals</th>
</tr>
</thead>
<tbody>
<tr>
<td>14</td>
<td>-</td>
<td>8/10</td>
</tr>
<tr>
<td>30</td>
<td>0/8</td>
<td>8/10</td>
</tr>
<tr>
<td>40</td>
<td>0/10</td>
<td>7/10</td>
</tr>
</tbody>
</table>

[0026] From the results in Table 1 it is evident that the algal meal containing astaxanthin has a therapeutic effect and can be used for prophylactic purposes.

Claims

1. Use of at least one type of xanthophylls for the preparation of a medicament for the prophylactic and/or therapeutic treatment of inflammation in the mucous membrane of mammalian gastrointestinal tract caused by a Helicobacter sp. infection.

2. Use according to claim 1, wherein the mammalian gastrointestinal tract is the human stomach and the Helicobacter sp. is H. pylori.

3. Use according to claim 1 or 2, wherein the xanthophyll is dissolved in an oil of food grade.

4. Use according to any one of claims 1-3, wherein the type of xanthophyll is astaxanthin.

5. Use according to claim 4, wherein the astaxanthin is in a form esterified with fatty acids.
6. Use according to claim 4 or 5, wherein the astaxanthin derives from a natural source.

7. Use according to claim 6, wherein the natural source is a culture of the alga *Haematococcus sp.*

8. Use according to any one of the claims 1-7, wherein the medicament further comprises carbohydrate structures.

9. Use according to claims 7 and 8, wherein the medicament comprises algal meal having astaxanthin in esterified form with fatty acids dissolved in small droplets of naturally occurring oil and naturally occurring carbohydrate structures in the partially disrupted algal cell walls.

10. Use according to any one of claims 1 - 9, wherein the medicament further comprises a prophylactically and/or therapeutically effective amount of a water soluble antioxidant.

11. Use according to claim 10, wherein the water soluble antioxidant is ascorbic acid (vitamin C).

12. Use according to any one of claims 1 - 11, wherein the medicament is presented in a separate unit dose or in mixture with food.

**Patentansprüche**

1. Verwendung von Xanthophyllen mindestens eines Typs für die Herstellung eines Medikaments zur prophylaktischen und/oder therapeutischen Behandlung einer Entzündung in der Schleimhaut des Gastrointestinaltrakts von Säugetieren hervorgerufen durch eine Infektion mit *Helicobacter sp.*

2. Verwendung nach Anspruch 1, wobei es sich beim Gastrointestinaltrakt von Säugetieren um den menschlichen Magen und bei *Helicobacter sp.* um H. pylori handelt.

3. Verwendung nach Anspruch 1 oder 2, wobei das Xanthophyll in einem Öl von Lebensmittelqualität gelöst ist.

4. Verwendung nach einem der Ansprüche 1 bis 3, wobei der Typ des Xanthophylls Astaxanthin ist.

5. Verwendung nach Anspruch 4, wobei das Astaxanthin in mit Fettsäuren veresterter Form vorliegt.

6. Verwendung nach Anspruch 4 oder 5, wobei das Astaxanthin aus einer natürlichen Quelle stammt.

7. Verwendung nach Anspruch 6, wobei die natürliche Quelle eine Zucht der Alge *Haematococcus sp.* ist.

8. Verwendung nach einem der Ansprüche 1 bis 7, wobei das Medikament weiterhin Kohlenhydratstrukturen enthält.

9. Verwendung nach den Ansprüchen 7 und 8, wobei das Medikament Algenmehl mit Astaxanthin in mit Fettsäuren veresterter Form enthält, das in kleinen Tröpfchen natürlichen vorkommenden Öls gelöst ist, sowie mit natürlichen vorkommenden Kohlenhydratstrukturen in den teilweise aufgebrochenen Algenzellwänden.

10. Verwendung nach einem der Ansprüche 1 bis 9, wobei das Medikament weiterhin eine prophylaktisch und/oder therapeutisch wirksame Menge eines wasserlöslichen Antioxidans enthält.

11. Verwendung nach Anspruch 10, wobei das wasserlösliche Antioxidans Ascorbinsäure (Vitamin C) ist.

12. Verwendung nach einem der Ansprüche 1 bis 11, wobei das Medikament als separate Einzeldosis oder gemischt mit Nahrung vorliegt.

**Revendications**

1. Utilisation d'au moins un type de xanthophylles pour la préparation d'un médicament pour le traitement prophylactique et/ou thérapeutique d'une inflammation dans la membrane muqueuse du tractus gastro-intestinal de mammifères, provoquée par une infection avec *Helicobacter sp.*
2. Utilisation selon la revendication 1, où le tractus gastro-intestinal de mammifères est l'estomac humain et l'Heli-
obcobacter sp. est H. pylori.

3. Utilisation selon la revendication 1 ou 2, où la xanthophylle est dissoute dans une huile de qualité alimentaire.

4. Utilisation selon l'une quelconque des revendications 1 à 3, où le type de la xanthophylle est l'astaxanthine.

5. Utilisation selon la revendication 4, où l'astaxanthine est sous forme estérifiée avec des acides gras.

6. Utilisation selon la revendication 4 ou 5, où l'astaxanthine provient d'une source naturelle.

7. Utilisation selon la revendication 6, où ladite source naturelle est une culture de l'algue Haematococcus sp.

8. Utilisation selon l'une quelconque des revendications 1 à 7, où le médicament contient en outre des structures d'hydrates de carbon.

9. Utilisation selon les revendications 7 et 8, où le médicament contient de la farine d'algues avec de l'astaxanthine sous forme estérifiée avec des acides gras qui est dissoute dans des petites gouttelettes d'huile d'occurrence naturelle, ainsi qu'avec des structures d'hydrates de carbon d'occurrence naturelle dans les parois cellulaires partiellement brisées des algues.

10. Utilisation selon l'une quelconque des revendications 1 à 9, où le médicament contient en outre une quantité prophylactiquement et/ou thérapeutiquement active d'un antioxydant soluble dans l'eau.

11. Utilisation selon la revendication 10, où l'antioxydant soluble dans l'eau est l'acide ascorbique (vitamine C).

12. Utilisation selon l'une quelconque des revendications 1 à 11, où le médicament est présenté sous forme d'une dose unitaire ou sous forme mélangée à un aliment.