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(54) Titre : COMPOSITIONS PHARMACEUTIQUES CONTENANT DES INHIBITEURS DE LIPASES ET DU CHITOSANE
(54) Title: PHARMACEUTICAL COMPOSITIONS CONTAINING LIPASE INHIBITORS AND CHITOSAN

(57) Abrégé/Abstract:

Orally administrable pharmaceutical composition containing an inhibitor of gastrointestinal lipases, one (or more) additional compound(s) of the group consisting of chitosan, its derivatives and salts thereof, and auxiliary excipients.



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PHARMACEUTICAL COMPOSITIONS CONTAINING LIPASE INHIBITORS AND CHITOSAN

The present invention relates to pharmaceutical compositions containing an inhibitor of gastrointestinal lipases, one (or more) additional compound(s) of the group consisting of chitosan, its derivatives and salts thereof, and auxiliary excipients.

An example of an inhibitor of gastrointestinal lipases is orlistat, previously known as tetrahydrolipstatin or THL. It reduces the absorption of dietary fat. Its use for the control or prevention of obesity and hyperlipaemia, is described in US patent 4 598 089. Orlistat is the N-formyl-L-leucine ester with (3S,4S)-3-hexyl-4-[(2S)-2-hydroxytridecyl]-2-oxetanone.

Anal leakage of oil (oily spotting) is an adverse effect which is occasionally observed in patients treated with lipase inhibitors. It results from physical separation of some liquid unabsorbed dietary fat from the bulk of the fecal mass in the lower large intestine. This effect can be prevented with the pharmaceutical compositions of the present invention.

In the US patent 5 447 953 it has been demonstrated that by combining a lipase inhibitor with substantial amounts of water insoluble crude fibers, the inhibiting effect on fat absorption can be increased.

Surprisingly, it has now been found that by combining the lipase inhibitor with low amounts of chitosan or a derivative or a salt thereof, the phenomenon of anal leakage of oil can be strongly reduced.

The invention is also concerned with the use of chitosan or a derivative or a salt thereof for the combined simultaneous, separate or chronologically spaced use with an inhibitor of gastrointestinal lipases, such as orlistat, in the treatment of obesity and hyperlipaemia and their comorbidities, such as type II diabetes mellitus.

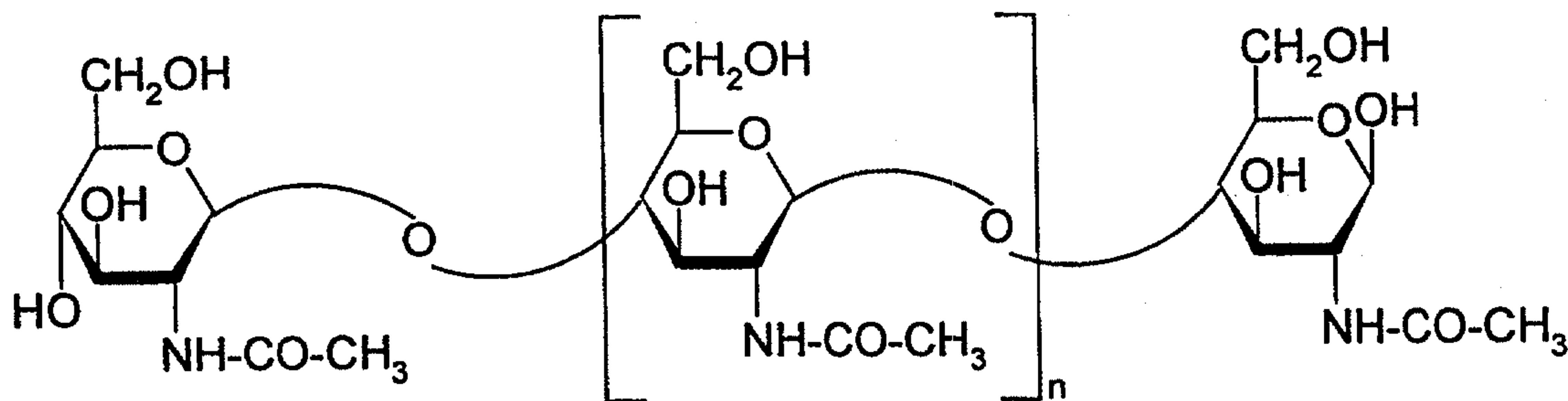
Artificial non-absorbed fats, mostly sucrose polyester, are used in the food industry for the production of low fat foods, such as low fat potatoe chips, low fat

cookies, low fat salad dressings and low fat ice cream. The ingestion of higher amounts of such foodstuffs containing non-absorbable fats can induce oily leakage.

The invention is further concerned with the use of chitosan or a derivative or a salt thereof for treating or preventing the syndrome of anal leakage of oil occurring after the administration of an inhibitor of gastrointestinal lipases, such as orlistat, or after ingestion of food containing poorly absorbable or non-absorbable fats or oils or of undigestible oily fat substitutes.

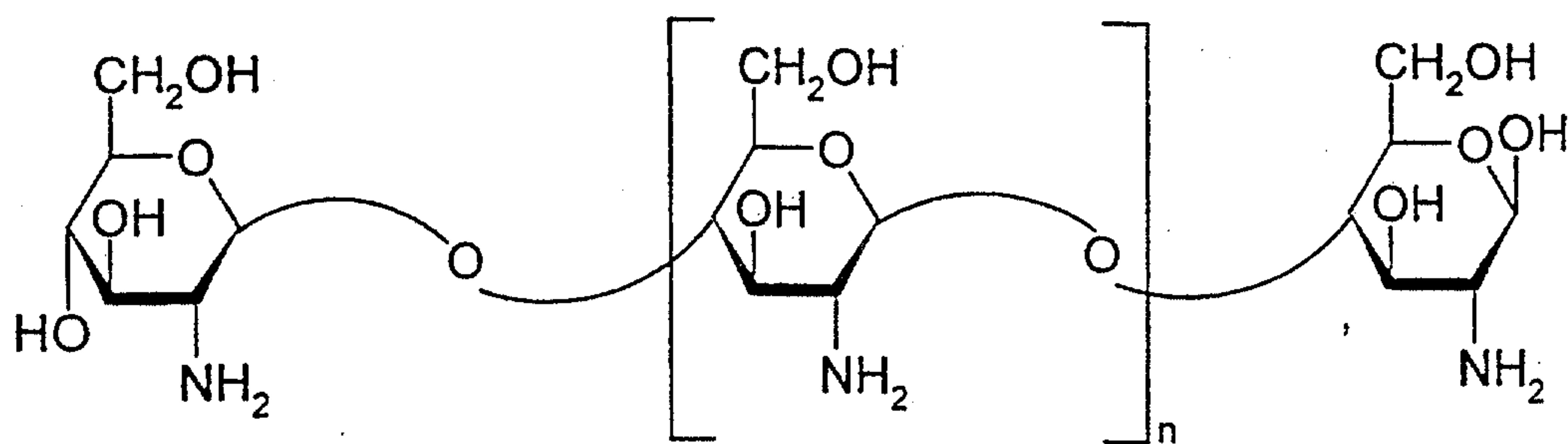
Chitosan is derived from chitin, a polysaccharide composed of (1→4)-linked 2-acetamido-2-deoxy-β-D-glucopyranosyl residues isolated from natural sources, by complete or partial deacetylation and partial depolymerization. Chitosan has a molecular weight of the order of 10^4 to 10^5 . Chitosan is soluble at gastric pH and insoluble or gel-like at intestinal pH. Examples of chitosan derivatives are medium or long chain N-alkyl- or N-alkanoyl-chitosan. The term "medium chain N-alkyl- or N-alkanoyl" refers to C_{8-13} -N-alkyl- or -N-alkanoyl chains, the term "long chain N-alkyl- or N-alkanoyl" refers to C_{14-18} -N-alkyl- or -N-alkanoyl chains. Examples of salts of chitosan are those with acetic, citric, formic and tartaric acid, as well as with diluted mineral acids.

Synonyms of chitosan (the Merck Index, 11th ed., #2052, 1989) are poly-D-glucosamine; poly-[1→4]-β-D-glucosamine and deacetylated chitin. The deacetylation of chitin which has the formula



to chitosan of the formula

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can be performed in hot concentrated NaOH solution (40-50%). Chitosan is commercially available from Pronova Biopolymer, Inc., 135 Commerce Way, Suite 201, Portsmouth, NH03801, e.g. as SEACURE^{*} 142, 242 or 342 with a viscosity interior to 20 cps, from 20 to 200 cps and from 200 to 800 cps, respectively.

Examples of auxiliary excipients which can be used in the pharmaceutical compositions of the invention are binders, diluents and lubricants, such as AVICEL^{*}, polyvinyl pyrrolidone (povidone), talc and sodium stearyl fumarate; sweeteners, such as sorbitol, glucose, saccharose, saccharine-sodium salt and sodium cyclamate; flavour agents, such as passion fruit, citron and limette; flavour enhancers, such as citric acid, monosodium citrate, sodium chloride and chinine sulfate; effervescing agents, such as sodium bicarbonate and tartaric acid, disintegrants, antimicrobial agents, such as p-hydroxybenzoic acid methyl or propyl ester; detergents and colouring agents, such as β -carotene.

AVICEL^{*} essentially comprises microcrystalline cellulose. It is available from FMC Corporation, Pharmaceutical Division, 1735 Market Street, Philadelphia, PA 19103, e.g. as AVICEL^{*} RC-591 or CL-611, which are mixtures of microcrystalline cellulose (about 92%) and carboxymethylcellulose sodium (about 8%), AVICEL^{*} PH 101 or PH 105, which is microcrystalline cellulose with an average particle size of 50 or 20 μ , respectively; AVICEL^{*} CE-15 a mixture of microcrystalline cellulose and guar gum.

The superiority of chitosan over microcrystalline cellulose, e.g. AVICEL^{*}, in reducing anal leakage of oil is shown in the following experiment:

The experiment for loss of free fecal oil is based on the observation that mice, due to steadily grooming their furs, distribute any excreted free fecal oil all over their bodies. This results in an easily visible brownish coloring of the fur (oily fur greasing). In mice weighing 20-25 g, excretion of free oil was provoked by administering an excessive dose of orlistat (300 μ mol/kg/day) together with a diet

Trademark^{*}

containing 7% fat, resulting in a daily fat intake of 1 g/day. The diet consisted of mashed Hamburger, butter, French fries and string beans.

The following time schedule of treatment was followed:

- 5
- days 1-5: feeding groups of 3 mice on the above diet and food additives (increasing doses of chitosan or AVICEL), and a control group without food additives
- days 3-4: adding orlistat (300 μ mol/kg/day)
- day 5: registration by photography of oily fur greasing

Oily fur greasing was evaluated by daily scoring units 1 to 4:

- 10
- 1 representing 0-25% skin involvement, 2: 26-50% skin involvement, 3: 50-75%, and 4: 75-100% skin involvement.

15

The skin involvement of the individual mice was added up for days 3-5. (At e.g. 100% skin involvement for days 3-5 the scoring would be maximal, i.e. 4+4+4=12 units). The oily fur greasing score in the controls was 6.3 ± 1.6 units (mean $\pm$ SE).

In 7 groups of mice receiving chitosan or AVICEL against oily fur greasing, its extent was first evaluated according to the scoring above, then expressed in % of the control groups and finally averaged within the animals of each group. The results are given in the table below:

Oily fur greasing (% of controls) in mice ingesting orlistat (300 μ mol/kg/day)

food additive (g/100 g feed)	chitosan	AVICEL
15	n.d.	17 $\pm$ 4
10	n.d.	29 $\pm$ 7
5	n.d.	53 $\pm$ 24
3	6 $\pm$ 3	n.d.
1	37 $\pm$ 16	n.d.
0.3	53 $\pm$ 5	n.d.
0.1	63 $\pm$ 9	n.d.

n.d. = not determined

When the diet contained chitosan, the extent of oily fur greasing was reduced. For instance, as compared to controls, the same 53% inhibition of oily fur greasing is obtained with a feed containing as much as 5 weight % of AVICEL and only 0.3 weight % of chitosan.

The compositions of the invention conveniently contain from 10 to 50 , preferably from 20 to 40 parts by weight of chitosan or a derivative or a salt thereof and from 10 to 200, preferably from 20 to 80 parts by weight of auxiliary excipients for 1 part by weight of an inhibitor of gastrointestinal lipase, such as orlistat.

The composition of the invention can also be in form of a commercial pack containing an inhibitor of gastrointestinal lipase and chitosan or a derivative or a salt thereof, with instructions for its use for the simultaneous, separate or chronologically spaced use in the treatment of obesity or hyperlipaemia.

For the treatment or prevention of obesity or hyperlipaemia, a composition of the invention containing from 10 mg to 500 mg of an inhibitor of gastrointestinal lipase, such as orlistat, and from 500 mg to 20 g, preferably from 2 g to 10 g, of an additional compound, such as chitosan, can be administered orally once, twice or three times per day.

The compositions of the invention can be in form of drinkable formulations, such as solutions or suspensions prepared from powder, granules, pellets, tablets to be reconstituted or effervescent tablets; or in form of chewable formulations, such

as tablets, capsules or lozenges. They can also be incorporated into food preparations, such as wafers, crackers or bread, or can be in form of swallowable formulations, such as tablets or capsules.

5 A preferred composition of the invention is a chewable tablet for the treatment of obesity, consisting essentially of orlistat as the active ingredient and of chitosan as the additional compound, wherein the dosage is from 10 to 120 mg of orlistat and from 0.5 to 5 g of chitosan. Most preferably, the chewable tablet consists essentially of about 60 mg of orlistat and about 2.5 g of chitosan.

10 A further preferred composition of the invention is a wafer for the treatment of obesity, consisting essentially of orlistat as the active ingredient and of chitosan as the additional compound, wherein the dosage is from 10 to 200 mg of orlistat and from 1 to 10 g of chitosan. Most preferably, the wafer consists essentially of about 120 mg of orlistat and about 5 g of chitosan.

15 A preferred method of preventing the syndrome of anal leakage of oil occasionally occurring after the oral administration of orlistat, comprises orally administering orlistat and chitosan in a dosage amount from 10 to 200 mg of orlistat and from 0.5 to 10 g of chitosan per fat containing main meal. Most conveniently, this method comprises orally administering orlistat and chitosan, the dosage amount being from 10 to 120 mg of orlistat and from 2 to 6 g of chitosan, particularly about 60 mg of orlistat and about 2.5 g of chitosan per fat containing
20 main meal.

The following non-limiting examples illustrate pharmaceutical preparations that can be produced by conventional procedures:

Example 1

25 Chitosan granules or pellets for the simultaneous, separate or chronologically spaced administration of orlistat are prepared as follows:

50 g of chitosan (SEACURE 342) and 50 g of AVICEL RC-591 are mixed and kneaded with demineralized water to a suitable consistency. The wet mass is sieved and then dried in a fluidized bed to give granules. Alternatively the wet
30 mass is extruded and spheronized and then dried in a fluidized bed to give pellets. A quantity of 5 g or 10 g of granules or pellets is filled into sachets as unit dose.

Alternatively, this material is filled into appropriate containers. The dosing may be performed with appropriate spoons.

Example 2

Powder for reconstitution:

5	Orlistat	0.12 g
	Low viscosity chitosan (SEACURE 142)	5 g
	Sorbitol	7.11 g
	AVICEL CL 611	1.20 g
	β -carotene	0.06 g
10	Citric acid	0.10 g
	p-Hydroxybenzoic acid methyl ester	0.15 g
	p-Hydroxybenzoic acid propyl ester	0.03 g
	Flavouring agent (passion fruit)	0.13 g
	AVICEL PH 105	8.00 g
15	Monosodium citrate	1.00 g
	Saccharine-sodium salt	<u>0.10 g</u>
	Total	23 g

An oral suspension is obtained by adding tap-water to the above powder to a volume of about 100 ml.

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Example 3

Granulates or pellets:

Orlistat	0.120 g
Chitosan (SEACURE 242)	5.0 g
AVICEL PH 101	4.88 g

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The above ingredients are mixed and kneaded with demineralized water to obtain a suitable consistency. The wet mass is sieved and dried in a fluidized bed at a temperature below 35°C to give granules. Alternatively, the wet mass is extruded and spheronized and then dried in a fluidized bed to give pellets. A quantity of 10 g the granules or pellets is filled into sachets as unit dose. Alternatively, the material

30 is filled into appropriate containers. The dosing can be performed with appropriate spoons.

Example 4

Effervescent tablets:

	Orlistat	0.120 g
	Saccharose powder	1.669 g
5	Low viscosity chitosan (SEACURE 142)	2.5 g
	Sodium cyclamate	0.115 g
	Saccharine sodium salt	0.004 g
	Sodium bicarbonate	0.7 g
	Tartaric acid (crystallized)	1.12 g
10	Sodium chloride (milled)	0.04 g
	Chinine sulfate	0.007 g
	Flavoring agent	<u>0.025 g</u>
	Total	6.3 g

Orlistat, saccharose, chitosan, sodium cyclamate and saccharin sodium are mixed and sieved. The mixture is kneaded with a mixture of ethanol and demineralized water, granulated and dried at a temperature below 35°C in a fluidized bed to give a mixture A. Sodium bicarbonate, tartaric acid, sodium chloride, chinine sulfate and the flavoring agent are mixed and sieved to give a mixture B. A and B are mixed and compressed to effervescent tablets of 6.3 g and a diameter of 30 mm.

Example 5

Chewable tablets:

	Orlistat	0.060 g
	Chitosan (SEACURE 242)	2.5 g
25	Sorbitol	1.84 g
	AVICEL CE-15	1.0 g
	Talc	0.480 g
	Sodium stearyl fumarate	<u>0.120 g</u>
	Total	6.0 g

Orlistat, chitosan, sorbitol and AVICEL CE-15 are mixed and sieved. Talc and sodium stearyl fumarate are sieved and mixed with the first obtained mixture and then compressed to chewable tablets of 6.0 g and a diameter of 2 cm.

Example 6

Chitosan wafers for the simultaneous, separate or chronologically spaced administration of orlistat are prepared as follows:

5 Corn flour (5 g) and 5 g of chitosan are mixed. Soybean oil (2 g) is added and the mixture is mixed for 15 minutes. Water is added to form a wet mass which is then extruded. The wet wafers are dried in an oven at 35°C and then packaged.

Example 7

Wafers:

10	Chitosan	5 g
	Soybean oil	2 g
	Corn flour	5 g
	Orlistat	120 mg

15 The process is the same as in Example 6 but orlistat is first dissolved in soybean oil and added to the blend. After the wet massing with water and extrusion, the wafers are dried at 35°C.

Example 8

Wafers: the proportions of the ingredients and the procedure are the same as in Example 7, but molten tripalmitin is substituted for soybean oil.

Example 9

20 Chitosan wafers for the simultaneous, separate or chronologically spaced administration of orlistat are prepared as follows:

Chitosan (5 g) and 5 g of maltodextrin are mixed. Molten tripalmitin (2 g) is added to the mixture. The mass is then wetted with water and the wet mass is extruded. The wafers are dried at 35°C.

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Example 10

Wafers:

Chitosan	5 g
Maltodextrin	5 g

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Triplamitin

2 g

Orlistat

120 mg

The process is the same as in Example 9 but orlistat is dissolved in molten tripalmitin and then added to the blend. The wafers are dried at 35°C.

The Embodiments Of The Invention In Which An Exclusive Property Or Privilege Is Claimed Are Defined As Follows:

1. Orally administrable pharmaceutical composition containing orlistat as inhibitor of gastrointestinal lipases, one additional compound of the group consisting of chitosan, medium and long chain N-alkyl- and N-alkanoyl-chitosan and salts thereof, and auxiliary excipients, wherein said composition contains from 500 mg to 20 g of the additional compound and from 10 mg to 500 mg of orlistat.
2. A composition as in claim 1 comprising from 2 g to 10 g of the additional compound.
3. A composition as in claim 1 wherein the additional compound is chitosan.
4. The use of chitosan for treating or preventing the syndrome of anal leakage of oil occasionally occurring after the oral administration of orlistat as inhibitor of gastrointestinal lipases, or after ingestion of food containing poorly absorbable or non-absorbable fats or oils or of undigestible oily fat substitutes.
5. A composition as in claim 3 which is a chewable tablet for the treatment of obesity, consisting essentially of orlistat as the active ingredient and of chitosan as the additional compound, wherein the dosage is from 10 to 120 mg of orlistat and from 0.5 to 5 g of chitosan.
6. The composition as in claim 5, wherein the dosage in the chewable tablet is about 60 mg of orlistat and about 2.5 g of chitosan.
7. A composition as in claim 3 which is a wafer for the treatment of obesity, consisting essentially of orlistat as the active ingredient and of chitosan as the additional compound, wherein the dosage is from 10 to 200 mg of orlistat and from 1 to 10 g of chitosan.

8. The composition as in claim 7, wherein the dosage in the wafer is about 120 mg of orlistat and about 5 g of chitosan.

9. Use of the composition as defined by any one of claims 1 to 3 and 7 for preventing the syndrome of anal leakage of oil occurring after the oral administration of orlistat.

10. Use of the composition as defined by any one of claims 1 to 3 and 7 in the preparation of a medicament for preventing the syndrome of anal leakage of oil occurring after the oral administration of orlistat.

11. The use of claim 9 or 10, wherein the dosage amount is from 10 to 120 mg of orlistat and from 2 to 6 g of chitosan per fat containing main meal.

12. The use of claim 11, wherein the dosage amount is about 60 mg of orlistat and about 2.5 g of chitosan per fat containing main meal.

13. The use of chitosan for the production of a medicament for treating or preventing the syndrome of anal leakage of oil occasionally occurring after the oral administration of orlistat as inhibitor of gastrointestinal lipases, or after ingestion of food containing poorly absorbable or non-absorbable fats or oils or of undigestible oily fat substitutes.

14. A use of orlistat with a compound selected from the group consisting of chitosan, medium chain N-alkyl-chitosan, long chain N-alkyl-chitosan, N-alkanoyl-chitosan and salts thereof, for simultaneous, separate or chronologically spaced administration in the treatment of obesity, hyperlipaemia, or a comorbidity of obesity or hyperlipaemia.

15. A use of orlistat and a compound selected from the group consisting of chitosan, medium chain N-alkyl-chitosan, long chain N-alkyl-chitosan, N-alkanoyl-chitosan and salts thereof, separately or in combination, for preparation of one or more medicaments for administration simultaneously, separately or in a chronologically spaced manner with said compound in the treatment of obesity,

hyperlipaemia, or a comorbidity of obesity or hyperlipaemia.

16. A use of orlistat with a compound selected from the group consisting of chitosan, medium chain N-alkyl-chitosan, long chain N-alkyl-chitosan, N-alkanoyl-chitosan and salts thereof, for simultaneous, separate or chronologically spaced administration in the treatment of type II diabetes mellitus.

17. A use of orlistat and a compound selected from the group consisting of chitosan, medium chain N-alkyl-chitosan, long chain N-alkyl-chitosan, N-alkanoyl-chitosan and salts thereof, separately or in combination, for preparation of one or more medicaments for administration simultaneously, separately or in a chronologically spaced manner with said compound in the treatment of type II diabetes mellitus.