

SPECIFICATION
COMPOSITION FOR SUPPOSITORIES

TECHNICAL FIELD

This invention relates to a composition for suppositories. More particularly, it relates to a composition for suppositories which comprises (A) a fatty base, (B) monodecanoyl-glycerol, (C) monolauroyl-glycerol, (D) a powder insoluble in fatty base and (E) a drug for suppositories.

BACKGROUND ART

Suppository bases are classified into fatty bases and water-soluble bases. Although they are both excellent bases, the fatty bases have been widely employed, since they are superior to the water-soluble ones in less irritation to the administration sites, etc. It has been a practice to design suppositories containing fatty bases to melt at the body temperature. When inserted into a body cavity, therefore, such a suppository would migrate upward from the administration site.

In the case of antihemorrhoidal suppository, drugs should be retained around the affected part. Accordingly, there have been reported a number of compositions for suppositories for preventing drugs from spreading over the rectum by, for example, JP-A-54-26325, JP-A-6-40889, JP-A-63-280016, JP-A-1-143825, JP-A-61-109710, JP-A-2-15024, JP-A-4-164023 and EP No. 103995 (the term "JP-A" as

used herein means an "unexamined published Japanese patent application").

However, there has been reported no suppository which sustains a melting point higher than the body temperature and thus never melts during storage but, when inserted into a body cavity, melts or gels at the body temperature owing to a decrease in its melting point.

DISCLOSURE OF THE INVENTION

Suppositories comprising fatty bases would melt at the body temperature after inserted into body cavity such as rectum or vagina and then release drugs contained therein, thus exerting the efficacy. That is to say, suppositories containing fatty bases are designed so that they melt at the human body temperature. Accordingly, it is frequently observed that such suppositories melt during transportation or storage at high temperatures in summer. After melting, suppositories are deformed or dented and thus become unusable. Moreover, the drug distribution becomes less uniform due to the sedimentation of the drug. It is therefore necessary to transport and store suppositories at low temperatures.

An object of the present invention is to provide compositions for suppositories which sustain a melting point higher than the body temperature under dry storage conditions and thus remain stable without melting during transportation or storage but, when inserted into body cavity, quickly melt or gel at the body temperature. Another object of the present invention is to provide compositions

for suppositories aiming at treating affected parts in body cavity such as hemorrhoids which are retained at the affected part without spreading out therefrom.

The present inventors conducted extensive studies on compositions for suppositories. As a result, they have successfully obtained compositions for suppositories which sustain a melting point higher than the body temperature under dry storage conditions but, when inserted into body cavity, become moist due to the moisture in the cavities and thus quickly melt or gel at the body temperature. The present invention has been completed based on this finding.

Accordingly, the present invention provides a composition for suppositories which comprises (A) a fatty base, (B) monodecanoyl-glycerol, (C) monolauroyl-glycerol, (D) a powder insoluble in fatty base and (E) a drug for suppositories.

The present invention further provides a composition for suppositories which aims at treating affected parts in body cavity such as hemorrhoids and is prepared by adding a retentive base for intracavity administration to the above-mentioned composition for suppositories.

By adding a vasoconstrictor as the drug for suppositories, furthermore, the obtained composition for suppositories can prevent the drug from spreading out from the affected tissue and elevate the drug concentration in the tissue, thus achieving a potentiated efficacy.

The fatty bases to be used in the present invention are fatty acid triglycerides exemplified by cacao fat, lanolin fat, medium-chain fatty acid triglycerides and hard fats. Examples of the hard fats include Witepsol (manufactured by Hüls Aktiengesellschaft), Saposyer (manufactured by Gattefoss), Isocacao (manufactured by Kao) and Pharmasol (manufactured by Nippon Oils and Fats).

The term "powder insoluble in fatty base" means a powder which is insoluble in fatty acid triglycerides. Examples thereof include anhydrous silicic acid, starches, crystalline cellulose, zinc oxide and alginic acid. It is preferable to use anhydrous silicic acid therefor.

The term "drug for suppositories" means a drug which is usually administered in the dosage form of suppositories. Examples thereof include anti-inflammatory, antipyretic and analgesic agents such as acetylsalicylic acid, acetaminophen, buprenorphine hydrochloride, indomethacin, ibuprofen, ketoprofen, piroxicam, diclofenac sodium, morphine hydrochloride, lysozyme hydrochloride and glycyrrhetic acid; antibiotics such as penicillin, cephalosporin, tetracycline and macrolides; antitumor agents such as 5-fluorouracil and futraful; antifungal agents such as econazole, econazole nitrate, miconazole, miconazole nitrate, clotrimazole, bifonazole, terbinafine hydrochloride and butenafine hydrochloride; steroids such as hydrocortisone, hydrocortisone acetate, prednisolone, dexamethasone and dexamethasone acetate;

local anesthetics such as ethyl aminobenzoate, lidocaine, lidocaine hydrochloride, dibucaine, dibucaine hydrochloride, procaine, procaine hydrochloride, meprylcaine, meprylcaine hydrochloride and mepivacaine; astringents such as zinc oxide, tannic acid, albumin tannate and aluminum potassium sulfate; antihistaminic agents such as diphenhydramine, diphenhydramine hydrochloride and chlorpheniramine maleate; wound healing promoters such as allantoin and aluminum chlorhydroxy allantoin; bactericides such as chlorhexidine hydrochloride, cetrimide, decalinium chloride or benzalkonium chloride; sulfa drugs such as sulfisomidine, sulfisomidine sodium, homosulfamine or sulfadiazine; vitamins such as liver oil, ergocalciferol, riboflavin, pyridoxine hydrochloride and tocopherol acetate; refrigerants such as d-camphor, dl-camphor, l-menthol, dl-menthol, peppermint oil and eucalyptus oil; antiemetic agents such as domperidone; defecation promoters such as bisacodyl; bronchodilating agents such as theophylline; peptides such as insulin; and vasoconstrictors such as tetrahydrozoline hydrochloride, naphazoline hydrochloride, phenylephrine hydrochloride, ephedrine hydrochloride and oximetazoline hydrochloride.

When vasoconstrictors such as tetrahydrozoline hydrochloride, naphazoline hydrochloride, phenylephrine hydrochloride, ephedrine hydrochloride and oxymetazoline hydrochloride are added to compositions for suppositories containing retentive bases for intracavity administration, the retention of the drugs can be

enhanced in the affected sites located in the lower parts of body cavity.

The term "retentive base for intracavity administration" means a base ingredient allowing the retention of drugs around affected sites in the lower parts of body cavity. Examples thereof include acrylic acid polymers, Polygam alkali metal salts, layered silicate minerals, starch acrylate, polyvinyl alcohol, pectin, cellulose derivatives (methylcellulose, carboxymethylcellulose, etc.), polyvinyl pyrrolidone, pullulan and tragacanth gum. It is preferable to use acrylic acid polymers therefor. Among all, carboxyvinyl polymer may be cited as the most desirable one.

Based on the total weight of the composition for suppositories, the content of a fatty base (A) is from 25 to 85 % by weight, preferably from 40 to 70 % by weight; the content of monodecanoyl-glycerol (B) is from 0.1 to 30 % by weight, preferably from 3 to 10 % by weight; the content of monolauroyl-glycerol (C) is from 10 to 70 % by weight, preferably from 15 to 50 % by weight; the content of a powder insoluble in fatty base (D) is from 0.1 to 20 % by weight, the content of anhydrous silicic acid, if employed, is from 0.5 to 10 %; and the content of a drug for suppositories (E) is from 0.1 to 20 % by weight.

When a retentive base for intracavity administration is further added, the content of a fatty base (A) is from 30 to 85 % by weight, preferably from 40 to 70 % by weight; the content of monodecanoyl-glycerol (B) is from 0.1 to 30 % by weight, preferably

from 1 to 10 % by weight; the content of monolauroyl-glycerol (C) is from 5 to 65 % by weight, preferably from 10 to 45 % by weight; the content of a powder insoluble in fatty base (D) is from 0.1 to 20 % by weight, the content of anhydrous silicic acid, if employed, is from 0.5 to 10 %; the content of a drug for suppositories (E) is from 0.1 to 20 % by weight and the content of a retentive base for intracavity administration (F) is from 0.1 to 20 % by weight, each based on the total weight of the composition for suppositories.

The content of carboxyvinyl polymer, which is the most desirable example of the retentive base for intracavity administration, is from 0.2 to 15 % by weight, still preferably from 1 to 10 % by weight, based on the total weight of the composition for suppositories.

The content of a vasoconstrictor is from 0.005 to 2.0 % by weight based on the total weight of the composition for suppositories. More particularly speaking, it is preferable to use from 0.005 to 0.1 % by weight of tetrahydrozoline hydrochloride, naphazoline hydrochloride or oximetazoline hydrochloride.

During storage, the composition for suppositories of the present invention has a melting point preferably falling within a range of from about 40 to 50 °C. When inserted into a body cavity, it preferably has a melting point of from about 36 to 37 °C.

By using the composition for suppositories of the present invention, suppositories may be produced in the following manner. First, a fatty base, monodecanoyl-glycerol and monolauroyl-

glycerol are mixed together in a melted state optionally with a retentive base for intracavity administration. Next, a drug and a powder insoluble in fatty base are added thereto and uniformly mixed by stirring. Then the resultant mixture is filled in containers, molds, etc. where it is then solidified by cooling. The mixing may be performed by an arbitrary method without restriction.

The suppositories produced by using the compositions for suppositories of the present invention can be administered intrarectally, intravaginally or urethrally.

INDUSTRIAL APPLICABILITY

The suppositories produced by using the compositions for suppositories of the present invention have a melting point higher than the body temperature under the dry storage conditions and thus remain stable without melting upon changes in temperature during transportation or storage. When inserted into body cavity, the suppositories become moist due to the moisture in the cavities and thus the melting point thereof is lowered. As a result, the suppositories melt or gel at the body temperature. Namely, the suppositories with the use of the compositions for suppositories of the present invention are highly useful ones which can be easily handled during transportation and storage and ensure good melting and efficacious drug-release in body cavity.

To further illustrate the present invention in greater detail, the following Examples will be given.

Example 1

(Formulation)

ketoprofen	50 mg
monodecanoyl-glycerol	75 mg
monolauroyl-glycerol	225 mg
light anhydrous silicic acid	10 mg
Pharmasol B115	450 mg.

(Production method)

To the fatty base (Phamasol B115) melted by heating (50 - 80 °C) were successively added other ingredients under stirring and dispersed therein. After cooling to about 50 °C, the mixture was filled into suppository containers and molded by further cooling to give suppositories.

Example 2

(Formulation)

ketoprofen	50 mg
monodecanoyl-glycerol	37.5 mg
monolauroyl-glycerol	225 mg
light anhydrous silicic acid	10 mg
Pharmasol B115	487.5 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 3

(Formulation)

lidocaine	60 mg
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hydrocortisone acetate	5 mg
allantoin	20 mg
tocopherol acetate	50 mg
monodecanoyl-glycerol	80 mg
monolauroyl-glycerol	560 mg
light anhydrous silicic acid	20 mg
Pharmasol B115	905 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 4

(Formulation)

acetaminophen	100 mg
monodecanoyl-glycerol	35 mg
monolauroyl-glycerol	140 mg
light anhydrous silicic acid	10 mg
Witepsol H15	525 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 5

(Formulation)

acetaminophen	100 mg
monodecanoyl-glycerol	35 mg
monolauroyl-glycerol	140 mg
light anhydrous silicic acid	10 mg
macrogol 4000	35 mg

Witepsol H15 490 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 6

(Formulation)

diclofenac sodium	25 mg
monodecanoyl-glycerol	35 mg
monolauroyl-glycerol	220 mg
light anhydrous silicic acid	10 mg
macrogol 4000	35 mg
Witepsol H15	475 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 7

(Formulation)

miconazole nitrate	100 mg
monodecanoyl-glycerol	45 mg
monolauroyl-glycerol	225 mg
light anhydrous silicic acid	10 mg
macrogol 4000	45 mg
Witepsol H15	575 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 8

(Formulation)

tetrahydrozoline hydrochloride	1 mg
lidocaine	60 mg
hydrocortisone acetate	5 mg
allantoin	20 mg
tocopherol acetate	60 mg
light anhydrous silicic acid	20 mg
monodecanoyl-glycerol	80 mg
monolauroyl-glycerol	560 mg
Pharmasol B115	960 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 9

(Formulation)

phenylephrine hydrochloride	1.3 mg
dibucaine	1.8 mg
hydrocortisone acetate	1.3 mg
zinc oxide	40 mg
light anhydrous silicic acid	20 mg
carboxyvinyl polymer	20 mg
monodecanoyl-glycerol	165 mg
monolauroyl-glycerol	500 mg
Pharmasol B115	905 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 10

(Formulation)

naphazoline hydrochloride	1 mg
lidocaine	60 mg
prednisolone acetate	1 mg
allantoin	20 mg
tocopherol acetate	60 mg
light anhydrous silicic acid	20 mg
carboxyvinyl polymer	20 mg
monodecanoyl-glycerol	80 mg
monolauroyl-glycerol	420 mg
Witepsol H15	930 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 11

(Formulation)

naphazoline hydrochloride	1 mg
lidocaine	60 mg
hydrocortisone acetate	5 mg
diphenhydramine hydrochloride	10 mg
allantoin	20 mg
tocopherol acetate	50 mg
zinc oxide	100 mg
Laponite	70 mg
monodecanoyl-glycerol	70 mg
monolauroyl-glycerol	260 mg

Witepsol H15

1000 mg.

(Production method)

The procedure of Example 1 was repeated.

Example 12

(Formulation)

tetrahydrozoline hydrochloride	1 mg
lidocaine	60 mg
prednisolone acetate	1 mg
crotamiton	50 mg
chlorhexidine hydrochloride	5 mg
aluminum chlorohydroxy allantoinate	5 mg
tocopherol acetate	50 mg
starch acrylate	65 mg
monodecanoyl-glycerol	150 mg
monolauroyl-glycerol	450 mg
Witepsol W35	820 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 1

(Formulation)

ketoprofen	50 mg
light anhydrous silicic acid	10 mg
Pharmasol B115	750 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 2

(Formulation)

ketoprofen	50 mg
monolauroyl-glycerol	300 mg
light anhydrous silicic acid	10 mg
Pharmasol B115	450 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 3

(Formulation)

ketoprofen	50 mg
monodecanoyl-glycerol	300 mg
light anhydrous silicic acid	10 mg
Pharmasol B115	450 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 4

The formulation and procedure of Example 8 were repeated but the carboxyvinyl polymer was omitted and the amount of the suppository base (Witepsol W35) was increased to compensate therefor.

Comparative Example 5

The formulation and procedure of Example 8 were repeated but the tetrahydrozoline hydrochloride was omitted and the amount of the suppository base (Witepsol W35) was increased to compensate

therefor.

Comparative Example 6

The formulation and procedure of Example 8 were repeated but the tetrahydrozoline hydrochloride and carboxyvinyl polymer were omitted and the amount of the suppository base (Witepsol W35) was increased to compensate therefor.

Comparative Example 7

(Formulation)

phenylephrine hydrochloride	1.3 mg
dibucaine	1.8 mg
hydrocortisone acetate	1.3 mg
zinc oxide	40 mg
light anhydrous silicic acid	20 mg
carboxyvinyl polymer	20 mg
Pharmasol B115	1570 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 8

(Formulation)

phenylephrine hydrochloride	1.3 mg
dibucaine	1.8 mg
hydrocortisone acetate	1.3 mg
zinc oxide	40 mg
light anhydrous silicic acid	20 mg
carboxyvinyl polymer	20 mg

monolauroyl-glycerol	630 mg
Pharmasol B115	940 mg.

(Production method)

The procedure of Example 1 was repeated.

Comparative Example 9

(Formulation)

phenylephrine hydrochloride	1.3 mg
dibucaine	1.8 mg
hydrocortisone acetate	1.3 mg
zinc oxide	40 mg
light anhydrous silicic acid	20 mg
carboxyvinyl polymer	20 mg
monodecanoyl-glycerol	630 mg
Pharmasol B115	940 mg.

(Production method)

The procedure of Example 1 was repeated.

Test Example 1: Measurement of melting point of suppository

(Sample)

As the samples, the suppositories of Examples 1, 2 and 9 were used. Also, the suppositories of Comparative Examples 1 to 3 and 7 to 9 were used as the comparative samples.

(Test method)

The melting point of each suppository was measured in accordance with the method of Muranishi et al. ["Zazai: Seizai kara Rinsho Oyo made "Suppositories: From Production to Clinical

Application)", page 64, 1985, Nanzan-do] with the use of a suppository penetrometer (Model PM3, manufactured by ERWEKA, Germany). A test tube containing a suppository was immersed in water bath at 30 °C for 10 minutes. Next, the temperature was elevated at a rate of 0.2 °C/min. The water temperature at which a test needle penetrated through the suppository was referred to as the melting point thereof. The melting point determined by adding 5 ml of water to the test tube was referred to as the melting point under the moist conditions in body cavity, while the one determined without adding water was referred to as the melting point under dry storage conditions. Also, the conditions of the suppositories were monitored at 37 °C corresponding to the body temperature in human body cavity.

(Results)

Table 1 summarizes the data determined under dry and moist conditions and at 37 °C.

	<u>TABLE 1</u>			
	<u>Melting point (°C)</u>		<u>Conditions at 37 °C</u>	
	<u>Dry</u>	<u>Moist</u>	<u>Dry</u>	<u>Moist</u>
Ex. 1	40.0	33.6	solid	liquid
Ex. 2	45.1	36.8	solid	liquid
Ex. 9	43.5	37.0	solid	gel
Comp. Ex. 1	34.4	34.5	liquid	liquid
Comp. Ex. 2	48.9	39.5	solid	solid
Comp. Ex. 3	34.1	32.0	liquid	liquid

Comp. Ex. 7	37.0	36.8	gel	gel
Comp. Ex. 8	50.3	47.2	solid	solid
Comp. Ex. 9	36.5	34.0	gel	gel

At 37 °C, the suppositories of Examples 1, 2 and 9 were in a solid state under dry conditions but liquefied or gelled under moist conditions. On the other hand, the suppositories of Comparative Examples 1, 3, 7 and 9 were in a state of liquid or gel both under dry and moist conditions. These comparative samples showed little difference in melting point between dry and moist conditions. Although the suppositories of Comparative Examples 2 and 8 showed differences in melting point between dry and moist conditions, these samples were in a solid state under both conditions.

Test Example 2: Measurement of drug-releasing rate

(Sample)

The suppository of Example 1 was used as the sample.

(Test method)

A suppository was placed on a dialysis membrane and the bottom of the membrane was closed with a closer provided with a weight (manufactured by SPECTRUM, USA). Next, it was immersed in a test solution (pH 7.0, 1,000 ml) at 37 °C and the drug thus released into the solution was measured by high performance liquid chromatography.

(Results)

Table 2 shows the ketoprofen-releasing rates after 4 hours.

The suppositories of Example 1 showed a high ketoprofen-releasing rate.

TABLE 2

	<u>Ex. 1</u>	<u>C.Ex. 1</u>	<u>C.Ex. 2</u>	<u>C.Ex. 3</u>
Releasing rate(%)	90.8	95.2	52.8	89.8

Test Example 3: Retention test *in vitro*

(Sample)

The suppository of Example 8 was used as the sample, while that of Comparative Example 4 was used as the comparative sample.

(Test method)

To evaluate the retention of the suppository in an affected part conveniently *in vitro*, a test was performed by applying the method of Sentikar-Fantelli.

A cellulose membrane (dialysis membrane, size 36: Viskase Sales Corporation) having been well washed with deionized water was tied with a string at the bottom and then fixed in a glass tube (2 cm in diameter x 20 cm). Then the sample and 5 ml of water were poured thereinto from the top. Immediately thereafter, warm water (37 °C) was circulated under a hydraulic pressure of 15 ± 2 cm H₂O. The location of the suppository was examined 10, 30 and 60 minutes thereafter. To judge the migration distance of the suppository, the circulating water was rapidly cooled simultaneously with the termination of the test so as to solidify the suppository. Then the sample was taken out together with the cellulose membrane and

thoroughly dried. Subsequently, the suppository was weighed in the section 0 to 8 cm from the tied site.

(Results)

Table 3 shows the weight ratios (%) of the suppositories. The sample of Example 8 showed a retention rate exceeding 90 % at every point (10, 30 and 60 minutes), while the sample of Comparative Example 4 showed a retention rate lower than 50 % at every point.

TABLE 3

<u>Time (min)</u>	<u>Ex. 8</u>	<u>C.Ex. 4</u>
10	99.3	47.2
30	98.0	40.9
60	93.8	49.4

Test Example 4: Edema inhibition test

(Sample)

The suppository of Example 8 was used as the sample, while those of Comparative Examples 5 and 6 were used as the comparative samples.

(Test method)

After fasting for 24 hours, male Wistar rats (weight: 150 - 170 g) were arbitrarily classified into groups each having 12 animals. A swab immersed in 0.16 ml of an inflammation-inducing solution (6 % croton oil in ether : distilled water : pyridine : ether = 10 : 1 : 4 : 5) was inserted into the anus of each animal for 10 seconds to

thereby induce inflammation. Immediately thereafter, the sample or the comparative samples (each 3 mm in diameter, 10 mm in length per 100 g body weight) were administered to the rats. After the administration of the suppositories, the anus of each animal was clipped to prevent the suppositories from leakage. After 6 hours, the rectum was taken out and the tissues 5 to 20 mm apart from the anus were collected. These tissues were weighed in the wet state and the recto-anus coefficient (RAC) was calculated as an indication of edema. Moreover, the edema inhibitory rate was calculated from the RAC thus determined.

(Results)

Compared with the comparative samples, the suppository of Example 8 showed an extremely high edema inhibitory rate, as Table 4 shows.

$$\text{RAC} = \frac{\text{wet weight of rectum/anus}}{\text{body weight (g)}} \times 1000.$$

$$\begin{array}{l} \text{Edema} \\ \text{inhibitory} \\ \text{ratio (\%)} \end{array} = \frac{1 - (\text{RAC of test group} - \text{RAC of untreated group})}{\text{RAC of comparative group} - \text{RAC of untreated group}} \times 100.$$

TABLE 4

<u>Group</u>	<u>RAC</u>	<u>Edema inhibitory ratio (%)</u>
untreated	0.79 ± 0.03	-
comparative	2.15 ± 0.07	-
Ex. 8	1.19 ± 0.04	70.6
C.Ex. 5	1.58 ± 0.03	41.9
C.Ex. 6	1.99 ± 0.04	11.8

CLAIM

1. A composition for suppositories which comprises (A) a fatty base, (B) monodecanoyl-glycerol, (C) monolauroyl-glycerol, (D) a powder insoluble in the fatty base and (E) a drug for suppositories.

2. The composition for suppositories as claimed in Claim 1, wherein the content of a fatty base (A) is from 25 to 85 % by weight, the content of monodecanoyl-glycerol (B) is from 0.1 to 30 % by weight, the content of monolauroyl-glycerol (C) is from 10 to 70 % by weight, the content of a powder insoluble in the fatty base (D) is from 0.1 to 20 % by weight and the content of a drug for suppositories (E) is from 0.1 to 20 % by weight, each based on the total weight of the composition for suppositories.

3. A composition for suppositories which comprises (A) a fatty base, (B) monodecanoyl-glycerol, (C) monolauroyl-glycerol, (D) a powder insoluble in the fatty base, (E) a drug for suppositories and (F) a retentive base for intracavity administration.

4. The composition for suppositories as claimed in Claim 3, wherein the content of a fatty base (A) is from 30 to 85 % by weight, the content of monodecanoyl-glycerol (B) is from 0.1 to 30 % by weight, the content of monolauroyl-glycerol (C) is from 5 to 65 % by weight, the content of a powder insoluble in the fatty base (D) is from 0.1 to 20 % by weight, the content of a drug for suppositories (E) is from 0.1 to 20 % by weight and the content of a retentive base for intracavity administration (F) is from 0.1 to 20 % by weight, each



based on the total weight of the composition for suppositories.

5. The composition for suppositories as claimed in Claim 3 or 4, wherein said retentive base for intracavity administration is one or more members selected from among acrylic acid polymers, Polygam alkali metal salts, layered silicate minerals and starch acrylate.

6. The composition for suppositories as claimed in Claim 5, wherein said acrylic acid polymer is carboxyvinyl polymer.

7. A composition for suppositories prepared by adding a vasoconstrictor to the composition for suppositories as claimed in Claim 3 or 4.

