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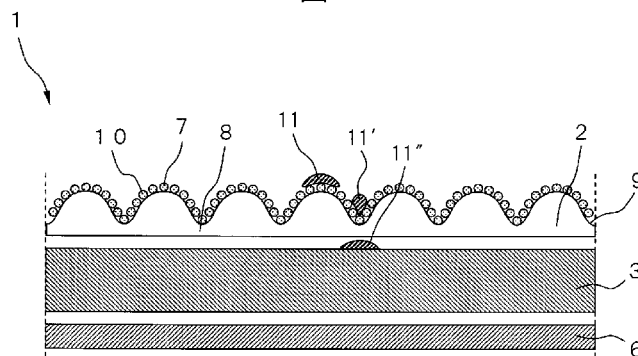
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(54) Title: ABSORBENT ARTICLE

(54) 発明の名称: 吸収性物品

[図2]

図2



(57) Abstract: The purpose of the present disclosure is to provide an absorbent article in which a liquid-permeable topsheet has an irregular structure comprising convexities and concavities on a skin contact surface, and by which even after highly viscous menstrual blood is absorbed the top sheet will not feel sticky and will feel smooth and dry. This absorbent article has the following configuration. An absorbent article including a liquid-permeable topsheet, a liquid-impermeable backsheet, and an absorber between the liquid-permeable topsheet and the liquid-impermeable backsheet; wherein the absorbent article is characterized in that the liquid-permeable topsheet includes in at least the convexities at the excretion orifice contact region a blood lubricity-imparting agent having a kinetic viscosity of 0.01-80 mm²/s at 40°C, a rate of water retention of 0.01-4.0 mass%, and a weight-average molecular weight less than 1,000.

(57) 要約:

[続葉有]



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NE, SN, TD, TG).

本開示は、液透過性のトップシートが、肌当接面に、凸部と、凹部とを含む凹凸構造を有し、そして粘度の高い経血を吸収した後であってもトップシートにべたつき感がなく、トップシートがサラサラしている吸収性物品を提供することを目的とする。本開示の吸収性物品は、以下の構成を有する。液透過性のトップシートと、液不透過性のバックシートと、上記液透過性のトップシート及び液不透過性のバックシートの間の吸収体とを有する吸収性物品であって、上記液透過性のトップシートが、排泄口当接域において、少なくとも凸部に、 40°C における $0.01 \sim 80 \text{ mm}^2/\text{s}$ の動粘度と、 $0.01 \sim 4.0$ 質量%の抱水率と、 $1,000$ 未満の重量平均分子量とを有する血液滑性付与剤を含むことを特徴とする吸収性物品。

DESCRIPTION

Title of the invention

ABSORBENT ARTICLE

Technical field

[0001]

The present disclosure relates to an absorbent article.

Background Art

[0001A]

A reference herein to a matter which is given as prior art is not to be taken as an admission or a suggestion the matter was known, or that the information it contains was part of the common general knowledge as at the priority date of any of the claims.

[0002]

As the basic performance of absorbent articles, such as sanitary napkins and panty liners has continued to improve with technological development over many years, leakage after absorption of excreta, such as menstrual blood has become a less frequent occurrence than in the past, and research is currently ongoing with the aim of achieving even higher performance, including a feel similar to underwear, and smoothness of the top sheet even after absorption of excreta, such as menstrual blood.

[0003]

Menstrual blood during menstruation, in particular, can also contain components of the endometrium which are highly viscous, and the top sheet preferably remains smooth and stick-free even after absorption of such highly viscous menstrual blood. Highly viscous menstrual blood usually remains on the top sheet in the form of masses, generally leaving the user with a visually unpleasant image, and therefore from this viewpoint as well it is preferred for no highly viscous menstrual blood to remain on the top sheet.

[0004]

In addition, the menstrual blood discharge during menstruation is not constant, but discharge of menstrual blood varies depending on the period after the start of

menstruation, at times involving a large amount of menstrual blood discharge at once, and at other times a small amount of menstrual blood discharge at once. Menstrual blood is not constantly discharged during menstruation, and there are even periods without discharge of menstrual blood.

Consequently, there are periods when a large amount of menstrual blood reaches the top sheet side at once, periods when small amounts of menstrual blood reach it at once, and periods when no menstrual blood reaches it at all, and preferably the menstrual blood migrates into the absorbent body without remaining on the top sheet, regardless of the amount of menstrual blood.

[0005]

Absorbent articles are known in the technical field which are coated with lotion compositions.

For example, PTL 1 discloses an absorbent article having a polypropylene glycol material-containing lotion composition situated on the inner surface of the top sheet (the clothing side surface), the inner surface of the back sheet (the body side surface), and on the base material between the inner surface of the top sheet and the inner surface of the back sheet.

Also, PTL 2 discloses an absorbent article wherein a polypropylene glycol material-containing lotion composition is applied on the outer surface of the top sheet (body side surface).

Citation List

Patent Literature

[0006]

PTL 1 Japanese Unexamined Patent Publication No. 2010-518918

PTL 2 Japanese Unexamined Patent Publication No. 2011-510801

Summary of Invention

Technical Problem

[0007]

However, the inventions described in PTL 1 and 2 are not designed so that menstrual blood migrates into the absorbent body without remaining on the top sheet regardless of the amount of menstrual blood, and nothing is mentioned regarding the relationship between the lotion composition and the top sheet, and especially the shape of the top sheet.

It is therefore desirable to provide an absorbent article without a sticky feel on the top sheet and with a smooth top sheet, not only when a large amount of menstrual blood has been absorbed, but even when a small amount of menstrual blood has been absorbed.

Solution to Problems

[0008]

As a result of diligent research directed toward solving the problems described above, the present inventors have discovered an absorbent article comprising a liquid-permeable top sheet, a liquid-impermeable back sheet and an absorbent body between the liquid-permeable top sheet and liquid-impermeable back sheet,

wherein the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on a skin contact surface thereof, and

the liquid-permeable top sheet comprises a blood slipping agent having a kinematic viscosity of 0.01 to 80 mm²/s at 40°C, a water holding percentage of 0.01 to 4.0 mass%, and a weight-average molecular weight of less than 1,000, on at least a projection in an excretory opening contact region, and a coating is configured to slip down from the liquid-permeable top sheet to the absorbent body along with menstrual blood.

[0008A]

In one form of the invention there is provided the use of an agent in a liquid permeable top sheet in an absorbent article for modifying blood on the liquid permeable top sheet, the absorbent article comprising a liquid-permeable top sheet, a liquid-impermeable back

sheet and an absorbent body between the liquid-permeable top sheet and liquid-impermeable back sheet,

wherein the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on a skin contact surface thereof, and

the liquid-permeable top sheet comprises a blood slipping agent having a kinematic viscosity of 0.01 to 80 mm²/s at 40°C, a water holding percentage of 0.01 to 4.0 mass%, and a weight-average molecular weight of less than 1,000, on at least a projection in an excretory opening contact region, and a coating is configured to slip down from the liquid-permeable top sheet to the absorbent body along with menstrual blood.

Advantageous Effects of Invention

[0009]

The absorbent article of the present disclosure has no sticky feel on the top sheet and has a smooth top

sheet, not only when a large amount of menstrual blood has been absorbed, but even when a small amount of menstrual blood has been absorbed.

5 Brief Description of Drawings

 [0010]

 Fig. 1 is a front view of a sanitary napkin, as an embodiment of an absorbent article of the invention.

 Fig. 2 is a cross-sectional view of section A of the
10 sanitary napkin 1 shown in Fig. 1, along X-X.

 Fig. 3 is an electron micrograph of the skin contact surface of a top sheet in a sanitary napkin wherein the top sheet comprises tri-C2L oil fatty acid glycerides.

 Fig. 4 is a pair of photomicrographs of menstrual
15 blood containing and not containing a blood slipping agent.

 Fig. 5 is a diagram illustrating a method of measuring surface tension.

20 Description of Embodiments

 [0011]

 The absorbent article of the present disclosure will now be explained in detail.

 [Liquid-permeable top sheet]

25 In the absorbent article of the present disclosure, the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on the skin contact surface. The mechanism of the present disclosure will be described below, but in order
30 for the menstrual blood that has reached the projection to migrate to the recess and subsequently into the absorbent body in the absorbent article of the present disclosure, the difference between the height of the projection and the height of the recess is preferably
35 constant.

 [0012]

 The height of the projection is preferably about 0.1

to about 15.0 mm higher, more preferably about 0.5 to about 5.0 mm higher and even more preferably about 0.5 to about 2.0 mm higher than the height of the recess. If the difference in heights is less than about 0.1 mm, it will be difficult for menstrual blood to migrate from the projection to the recess and subsequently into the absorbent body, while if the difference in heights is greater than about 15.0 mm, the projection will tend to be easily collapsable during wear.

The heights of the projection and the recess can be measured using a high precision laser displacement meter, such as an LJ-G Series two-dimensional laser displacement gauge (Model: LJ-G030) by Keyence Corp.

[0013]

Examples of top sheets having uneven structures include those in which the liquid-permeable top sheet has a ridge-furrow structure comprising a plurality of ridges and a plurality of furrows on the skin contact surface, such as the nonwoven fabrics described in Japanese Examined Patent Publication HEI No. 7-84697, Japanese Unexamined Patent Publication HEI No. 2-229255, Japanese Unexamined Patent Publication No. 2001-328191, Japanese Unexamined Patent Publication No. 2008-002034, No. 2008-023311, No. 2008-025078, No. 2008-025085, No. 2008-307179, No. 2009-030218, No. 2010-285735, No. 2011-038211, No. 2011-074515 and No. 2011-080178, and the porous films described in Japanese Unexamined Patent Publication SHO No. 64-34365 and Japanese Examined Patent Publication SHO No. 57-17081.

[0014]

For an embodiment in which the uneven structure is a ridge-furrow structure, the heights of the ridges are preferably about 0.1 to about 15.0 mm higher, more preferably about 0.5 to about 5.0 mm higher and even more preferably about 0.5 to about 2.0 mm higher than the heights of the furrows. The pitch of the ridges is preferably about 1.5 to about 17 mm, more preferably

about 2.0 to about 12 mm and even more preferably about 3 to about 8 mm. This is so that menstrual blood will slip down from the projection to the recess and then rapidly migrate into the absorbent body.

5 [0015]

The liquid-permeable top sheet may have embossed sections formed by embossing at least the liquid-permeable top sheet. For example, the liquid-permeable top sheet may have embossed sections formed by embossing the liquid-permeable top sheet and the absorbent body, and for an embodiment in which the absorbent article of the present disclosure includes a second sheet, the liquid-permeable top sheet may have embossed sections formed by embossing the liquid-permeable top sheet, the second sheet and the absorbent body. If the top sheet has embossed sections, the blood slipping agent will slip down from the projection to the recess, together with menstrual blood, and menstrual blood will subsequently be able to rapidly migrate into the absorbent body.

20 [0016]

[Blood slipping agent]

For the absorbent article of the present disclosure, the liquid-permeable top sheet contains, at least in the projection, a blood slipping agent having kinematic viscosity of about 0.01 to about 80 mm²/s at 40°C, a water holding percentage of about 0.05 to about 4.0 mass%, and a weight-average molecular weight of less than about 1,000.

[0017]

30 The blood slipping agent has, at 40°C, a kinematic viscosity of about 0 to about 80 mm²/s, preferably a kinematic viscosity of about 1 to about 70 mm²/s, more preferably a kinematic viscosity of about 3 to about 60 mm²/s, even more preferably a kinematic viscosity of about 5 to about 50 mm²/s, and yet more preferably a kinematic viscosity of about 7 to about 45 mm²/s.

The kinematic viscosity tends to be higher with a) a

larger molecular weight of the blood slipping agent, b) a higher percentage of polar groups, such as carbonyl bonds (-CO-), ether bonds (-O-), carboxyl groups (-COOH) and hydroxyl groups (-OH), and c) a larger IOB.

5 [0018]

In order to have a kinematic viscosity of about 0 to about 80 mm²/s at 40°C, the melting point of the blood slipping agent is preferably 45°C or less. This is because the kinematic viscosity will tend to be higher if
10 the blood slipping agent contains crystals at 40°C.

As used herein, the "kinematic viscosity at 40°C" may be referred to simply as "kinematic viscosity".

[0019]

The significance of the kinematic viscosity of the
15 blood slipping agent will be explained below, but a kinematic viscosity exceeding about 80 mm²/s will tend to result in high viscosity of the blood slipping agent, such that it will not as easily slip down from the projection to the recess together with menstrual blood
20 that has reached the skin contact surface of the top sheet, and subsequently migrate into the absorbent body.

[0020]

The kinematic viscosity can be measured according to JIS K 2283:2000, "5. Kinematic Viscosity Test Method",
25 using a Cannon-Fenske reverse-flow viscometer, at a testing temperature of 40°C.

[0021]

The blood slipping agent has a water holding percentage of about 0.01 to about 4.0 mass%, preferably
30 it has a water holding percentage of about 0.02 to about 3.5 mass%, more preferably it has a water holding percentage of about 0.03 to about 3.0 mass%, even more preferably it has a water holding percentage of about 0.04 to about 2.5 mass%, and yet more preferably it has a
35 water holding percentage of about 0.05 to about 2.0 mass%.

[0022]

As used herein, "water holding percentage" means the percentage of water that can be held by a substance, and it may be measured in the following manner.

5 (1) A test tube, a rubber stopper, the substance to be measured and deionized water are allowed to stand for a day and a night in a thermostatic chamber at 40°C.

(2) Into the 20 mL test tube in the thermostatic chamber there are charged 5.0 g of the substance to be measured and 5.0 g of deionized water.

(3) The mouth of the test tube is sealed with the rubber stopper in the thermostatic chamber, and it is rotated once and allowed to stand for 5 minutes.

[0023]

15 (4) A 3.0 g portion of the layer of the substance to be measured (usually the upper layer) is sampled into a glass dish with a diameter of 90 mm (weight: W_0), in the thermostatic chamber.

(5) The dish is heated at 105°C for 3 hours in an oven to evaporate off the moisture, and the weight of each dish is measured (weight: W_1).

(6) The water holding percentage is calculated by the following formula.

$$\text{Water holding percentage (\%)} = 100 \times (W_0 - W_1) / 3.0$$

25 The measurement is conducted three times, and the average value is recorded.

[0024]

The significance of the water holding percentage of the blood slipping agent will be explained below, but a low water holding percentage will tend to lower the affinity between the blood slipping agent and menstrual blood, thus impeding its migration into the absorbent body together with menstrual blood that has reached the skin contact surface of the top sheet. If the water holding percentage is high, on the other hand, the affinity between menstrual blood and the blood slipping

agent will become very high, similar to a surfactant, and absorbed menstrual blood will tend to remain on the skin contact surface of the top sheet, resulting in more red coloration of the skin contact surface of the top sheet.

5 [0025]

The water holding percentage tends to be a larger value with a) a smaller molecular weight of the blood slipping agent, and b) a higher percentage of polar groups, such as carbonyl bonds (-CO-), ether bonds (-O-),
10 carboxyl groups (-COOH) and hydroxyl groups (-OH). This is because the blood slipping agent has greater hydrophilicity. The water holding percentage will tend to have a larger value with a greater IOB, i.e. with a higher inorganic value or with a lower organic value.
15 This is because the blood slipping agent will have greater hydrophilicity.

[0026]

The significance of the kinematic viscosity and water holding percentage of the blood slipping agent will
20 now be explained.

Fig. 1 is a front view of a sanitary napkin, as an embodiment of an absorbent article of the invention, as viewed from the skin contact side. The sanitary napkin 1 shown in Fig. 1 has its forward direction facing left in
25 the drawing. The sanitary napkin 1 shown in Fig. 1 has a liquid-permeable top sheet 2, an absorbent body 3, and a liquid-impermeable back sheet (not shown). The sanitary napkin 1 in Fig. 1 is also shown as having a side sheet 4 and embossed sections 5.

30 [0027]

In the sanitary napkin 1 shown in Fig. 1, the top sheet has a plurality of ridges and a plurality of furrows on the skin contact surface, extending in the
35 lengthwise direction of the absorbent article, and the ridges and furrows may be omitted as appropriate. In the sanitary napkin 1 shown in Fig. 1, the ridges and furrows are disposed in an alternating fashion in the widthwise

direction of the absorbent article. The side sheet 4 and embossed section 5 are shown for the sanitary napkin 1 illustrated in Fig. 1, but another embodiment of the absorbent article of the present disclosure is an absorbent article without a side sheet and embossed sections.

[0028]

Fig. 2 is a cross-sectional view of section A of the sanitary napkin 1 shown in Fig. 1, along X-X. The sanitary napkin 1 shown in Fig. 2 comprises a liquid-permeable top sheet 2, a liquid-impermeable back sheet 6, and an absorbent body 3 between the liquid-permeable top sheet 2 and liquid-impermeable back sheet 6. In Fig. 2, the top sheet 2 has projections 7 and recesses 8 on the skin contact surface 9, and a blood slipping agent 10 coated on the skin contact surface 9 of the top sheet 2. In Fig. 2, the blood slipping agent 10 is shown as droplets on the skin contact surface 9 of the top sheet 2 for convenience, but according to the absorbent article of the present disclosure, the form and distribution of the blood slipping agent is not limited to that shown in the drawing.

[0029]

As shown in Fig. 2, highly viscous menstrual blood 11 that has reached the projections 7 on the skin contact surface 9 of the top sheet 2 contacts with the blood slipping agent 10 that is present at least on the projections 7. The blood slipping agent 10 having the prescribed water holding percentage and kinematic viscosity slips down into the recesses 8 together with the menstrual blood 11, becoming menstrual blood 11', and then the menstrual blood 11' passes mainly through the recesses 8, reaching the absorbent body 3 to become menstrual blood 11'', and is rapidly absorbed into the absorbent body 3.

[0030]

More specifically, since the blood slipping agent 10

with a kinematic viscosity of about 0.01 to about 80 mm²/s at 40°C has very low viscosity near the body temperature of the wearer and has a constant affinity with the menstrual blood 11, it slips down from the projections 7 to the recesses 8 together with the menstrual blood 11, and utilizing the energy during slipping, the menstrual blood 11' is able to pass through the recesses 8 of the top sheet 2 to rapidly migrate into the absorbent body 3. Also, since the blood slipping agent 10 present in the projections 7 has a water holding percentage of about 0.01 to about 4.0 mass%, presumably it has no affinity with the hydrophilic component (blood plasma, etc.) in the menstrual blood 11, and therefore the menstrual blood 11 does not easily remain on the top sheet.

15 [0031]

When the menstrual blood 11 is a large amount of menstrual blood, the menstrual blood 11 easily migrates into the absorbent body 3, even when the kinetic energy of the menstrual blood 11 itself is high and the kinematic viscosity of the blood slipping agent 10 is relatively high so that it does not easily slip down together with the menstrual blood 11, or when the water holding percentage value is relatively high so that affinity with the hydrophilic components of the menstrual blood 11 is high, or when the weight-average molecular weight value is relatively high so that it does not easily slip down together with the menstrual blood 11, or when the skin contact surface of the top sheet does not have an uneven structure.

30 [0032]

When the menstrual blood 11 is a small amount of menstrual blood, on the other hand, the kinetic energy of the menstrual blood 11 is low, and menstrual blood that has reached the skin contact surface 9 of the top sheet 2 tends to easily pool in such cases. Consequently, the prescribed blood slipping agent slips down from the projections 7 into the recesses 8 together with the

menstrual blood 11, and the menstrual blood 11' is drawn into the top sheet 2 and then drawn into the absorbent body 3, so that the menstrual blood can rapidly migrate into the absorbent body.

5 [0033]

The blood slipping agent has a weight-average molecular weight of less than about 1,000, and preferably a weight-average molecular weight of less than about 900. This is because, if the weight-average molecular weight is about 1,000 or higher, tack may result in the blood slipping agent itself, tending to create a feeling of unpleasantness for the wearer. If the weight-average molecular weight increases, the viscosity of the blood slipping agent will tend to increase, and it will therefore be difficult to lower the viscosity of the blood slipping agent by heating to a viscosity suitable for coating, and as a result, the blood slipping agent may need to be diluted with a solvent.

[0034]

20 The blood slipping agent preferably has a weight-average molecular weight of about 100 or greater, and more preferably it has a weight-average molecular weight of about 200 or greater. This is because if the weight-average molecular weight is low, the vapor pressure of the blood slipping agent may be increased, gasification may occur during storage and the amount may be reduced, often leading to problems, such as odor during wear.

[0035]

30 In addition, as used herein, "weight-average molecular weight" includes the concept of a polydisperse compound (for example, a compound produced by stepwise polymerization, an ester formed from a plurality of fatty acids and a plurality of aliphatic monohydric alcohols), and a simple compound (for example, an ester formed from one fatty acid and one aliphatic monohydric alcohol), and in a system comprising N_i molecules with molecular weight M_i ($i = 1, \text{ or } i = 1, 2, \dots$), it refers to M_w determined

by the following formula.

$$M_w = \Sigma N_i M_i^2 / \Sigma N_i M_i$$

[0036]

As used herein, the weight-average molecular weights
5 are the values measured by gel permeation chromatography
(GPC), based on polystyrene.

The GPC measuring conditions may be the following,
for example.

Device: Lachrom Elite high-speed liquid chromatogram
10 by Hitachi High-Technologies Corp.

Columns: SHODEX KF-801, KF-803 and KF-804, by Showa
Denko K.K.

Eluent: THF

Flow rate: 1.0 mL/min

15 Driving volume: 100 μ L

Detection: RI (differential refractometer)

The weight-average molecular weights listed in the
examples of the present specification were measured under
the conditions described below.

20 [0037]

The blood slipping agent can have an IOB of about
0.00 to about 0.60.

The IOB (Inorganic Organic Balance) is an indicator
of the hydrophilic-lipophilic balance, and as used
25 herein, it is the value calculated by the following
formula by Oda et al.:

$$IOB = \text{inorganic value} / \text{organic value}.$$

[0038]

The inorganic value and the organic value are based
30 on the organic paradigm described in "Organic compound
predictions and organic paradigms" by Fujita A., Kagaku
no Ryoiki (Journal of Japanese Chemistry), Vol.11, No.10
(1957) p.719-725.

The organic values and inorganic values of major
35 groups, according to Fujita, are summarized in Table 1
below.

[0039]

Table 1

Group	Inorganic value	Organic value
-COOH	150	0
-OH	100	0
-O-CO-O-	80	0
-CO-	65	0
-COOR	60	0
-O-	20	0
Triple bond	3	0
Double bond	2	0
CH ₂	0	20
<i>iso</i> -branch	0	-10
<i>tert</i> -branch	0	-20
Light metal (salt)	≥500	0
Heavy metal (salt), amine, NH ₃ salt	≥400	0

[0040]

For example, in the case of an ester of
5 tetradecanoic acid which has 14 carbon atoms and dodecyl
alcohol which has 12 carbon atoms, the organic value is
520 (CH₂, 20 × 26) and the inorganic value is 60 (-COOR,
60 × 1), and therefore IOB = 0.12.

[0041]

10 The IOB of the blood slipping agent is preferably
between about 0.00 and 0.60, more preferably between
about 0.00 and 0.50, even more preferably between about
0.00 and 0.40 and most preferably between about 0.00 and
0.30. If the IOB is within this range, it will be easier
15 to meet the aforementioned conditions for the water-
holding capacity and kinematic viscosity.

[0042]

The blood slipping agent preferably has a melting
point of 45°C or less. If the blood slipping agent has a
20 melting point of 45°C or less, the blood slipping agent
will more easily exhibit a kinematic viscosity in the
aforementioned range.

[0043]

As used herein, the term "melting point" refers to the peak top temperature for the endothermic peak during conversion from solid to liquid, upon measurement with a differential scanning calorimetry analyzer at a temperature-elevating rate of 10°C/min. The melting point may be measured using a Model DSC-60 DSC measuring apparatus by Shimadzu Corp., for example.

[0044]

If the blood slipping agent has a melting point of about 45°C or less, it may be either liquid or solid at room temperature (25°C), or in other words, the melting point may be either about 25°C or higher or below about 25°C, and for example, it may have a melting point of about -5°C or about -20°C. The reason for a melting point of about 45°C or less for the blood slipping agent will be explained below.

[0045]

The blood slipping agent does not have a lower limit for the melting point, but the vapor pressure is preferably low. The vapor pressure of the blood slipping agent is preferably about 0-200 Pa, more preferably about 0-100 Pa, even more preferably about 0-10 Pa, even more preferably about 0-1 Pa, and even more preferably about 0.0-0.1 Pa at 25°C (1 atmosphere).

[0046]

Considering that the absorbent article of this disclosure is to be used in contact with the human body, the vapor pressure is preferably about 0-700 Pa, more preferably about 0-100 Pa, even more preferably about 0-10 Pa, even more preferably about 0-1 Pa, and even more preferably 0.0-0.1 Pa, at 40°C (1 atmosphere). If the vapor pressure is high, gasification may occur during storage and the amount of blood slipping agent may be reduced, and as a consequence problems, such as odor during wear, may be created.

[0047]

The melting point of the blood slipping agent may be selected depending on the weather or duration of wear. For example, in regions with a mean atmospheric temperature of about 10°C or less, using a blood slipping agent with a melting point of about 10°C or less may help the blood slipping agent function after excretion of menstrual blood, even if it has been cooled by the ambient temperature.

[0048]

Also, when the absorbent article is to be used for a prolonged period of time, the melting point of the blood slipping agent is preferably at the high end of the range of about 45°C or less. This is so that the blood slipping agent will not be easily affected by sweat or friction during wearing, and will not easily become biased even during prolonged wearing.

[0049]

In the technical field, the skin contact surfaces of top sheets are coated with surfactants in order to alter the surface tension of menstrual blood and promote rapid absorption of menstrual blood. However, the top sheet coated with the surfactant has very high affinity for the hydrophilic components (blood plasma, etc.) in menstrual blood, and acts to attract them, tending to cause menstrual blood instead to remain on the top sheet. The blood slipping agent, unlike conventionally known surfactants, has low affinity with menstrual blood and therefore does not cause residue of menstrual blood on the top sheet and allows rapid migration into the absorbent body.

[0050]

Preferably, the blood slipping agent is selected from the group consisting of following items (i)-(iii), and any combination thereof:

(i) a hydrocarbon;

(ii) a compound having (ii-1) a hydrocarbon moiety,

and (ii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

5 (iii) a compound having (iii-1) a hydrocarbon moiety, (iii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii-3) one or more,
10 same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen of the hydrocarbon moiety.

[0051]

15 As used herein, "hydrocarbon" refers to a compound composed of carbon and hydrogen, and it may be a chain hydrocarbon, such as a paraffinic hydrocarbon (containing no double bond or triple bond, also referred to as alkane), an olefin-based hydrocarbon (containing one
20 double bond, also referred to as alkene), an acetylene-based hydrocarbon (containing one triple bond, also referred to as alkyne), or a hydrocarbon comprising two or more bonds selected from the group consisting of double bonds and triple bonds, and cyclic hydrocarbon,
25 such as aromatic hydrocarbons and alicyclic hydrocarbons.

[0052]

 Preferred as such hydrocarbons are chain hydrocarbons and alicyclic hydrocarbons, with chain hydrocarbons being more preferred, paraffinic
30 hydrocarbons, olefin-based hydrocarbons and hydrocarbons with two or more double bonds (containing no triple bond) being more preferred, and paraffinic hydrocarbons being even more preferred.

 Chain hydrocarbons include linear hydrocarbons and
35 branched hydrocarbons.

[0053]

 When two or more oxy groups (-O-) are inserted in

the compounds of (ii) and (iii) above, the oxy groups (-O-) are not adjacent each other. Thus, compounds (ii) and (iii) do not include compounds with continuous oxy groups (i.e., peroxides).

5 [0054]

In the compounds of (iii), compounds in which at least one hydrogen on the hydrocarbon moiety is substituted with a hydroxyl group (-OH) are preferred over compounds in which at least one hydrogen on the hydrocarbon moiety is substituted with a carboxyl group (-COOH). This is because the carboxyl groups bond with metals and the like in menstrual blood, increasing the water holding percentage of the blood slipping agent, which may sometimes exceed the prescribed range. The same is true from the viewpoint of the IOB as well. As shown in Table 1, the carboxyl groups bond with metals and the like in menstrual blood, drastically increasing the inorganic value from 150 to 400 or greater, and therefore a blood slipping agent with carboxyl groups can increase the IOB value to more than about 0.60 during use.

[0055]

More preferably, the blood slipping agent is selected from the group consisting of following items (i')-(iii'), and any combination thereof:

(i') a hydrocarbon;

(ii') a compound having (ii'-1) a hydrocarbon moiety, and (ii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii') a compound having (iii'-1) a hydrocarbon moiety, (iii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the

hydrocarbon moiety, and (iii'-3) one or more, same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen on the hydrocarbon moiety.

5 [0056]

When 2 or more same or different bonds are inserted in the compound of (ii') or (iii'), i.e., when 2 or more same or different bonds selected from the group consisting carbonyl bonds (-CO-), ester bonds (-COO-),
10 carbonate bonds (-OCOO-) and ether bonds (-O-) are inserted, the bonds are not adjacent to each other, and at least one carbon atom lies between each of the bonds.
[0057]

The blood slipping agent has more preferably about
15 1.8 or less carbonyl bonds (-CO-), about 2 or less ester bonds (-COO-), about 1.5 or less carbonate bonds (-OCOO-), about 6 or less ether bonds (-O-), about 0.8 or less carboxyl groups (-COOH) and/or about 1.2 or less hydroxyl groups (-OH), per 10 carbon atoms in the hydrocarbon
20 moiety.

[0058]

Even more preferably, the blood slipping agent is selected from the group consisting of following items (A)-(F), and any combination thereof:

25 (A) an ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain
30 hydrocarbon moiety;

(B) an ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl
35 group substituting for a hydrogen on the chain hydrocarbon moiety;

(C) an ester of (C1) a carboxylic acid, hydroxy

acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (C2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(D) a compound having a chain hydrocarbon moiety and one bond selected from the group consisting of ether bonds (-O-), carbonyl bonds (-CO-), ester bonds (-COO-) and carbonate bonds (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety;

(E) a polyoxy C₃-C₆ alkylene glycol, or ester or ether thereof; and

(F) a chain hydrocarbon.

The blood slipping agent in accordance with (A) to (F) will now be described in detail.

[0059]

[(A) Ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety]

In the (A) ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety (hereunder also referred to as "compound (A)"), it is not necessary for all of the hydroxyl groups to be esterified so long as the kinematic viscosity, water holding percentage and weight-average molecular weight are within the aforementioned ranges.

[0060]

Examples of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety (hereunder

also referred to as "compound (A1)") include chain hydrocarbon tetraols, such as alkanetetraols, including pentaerythritol, chain hydrocarbon triols, such as alkanetriols, including glycerins, and chain hydrocarbon diols, such as alkanediols, including glycols.

[0061]

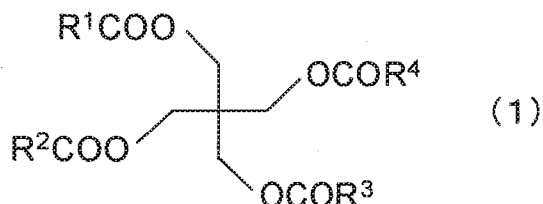
Examples of (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety include compounds in which one hydrogen on the hydrocarbon is substituted with one carboxyl group (-COOH), such as fatty acids.

Examples for compound (A) include (a₁) an ester of a chain hydrocarbon tetraol and at least one fatty acid, (a₂) an ester of a chain hydrocarbon triol and at least one fatty acid, and (a₃) an ester of a chain hydrocarbon diol and at least one fatty acids.

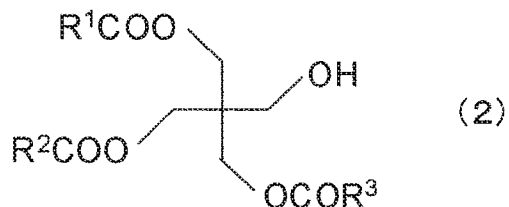
[0062]

[(a₁) Esters of a chain hydrocarbon tetraol and at least one fatty acid]

Examples of an ester of a chain hydrocarbon tetraol and at least one fatty acid include tetraesters of pentaerythritol and fatty acids, represented by the following formula (1):

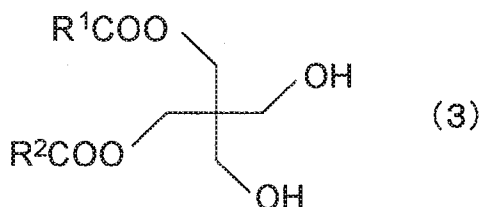


triesters of pentaerythritol and fatty acids, represented by the following formula (2):

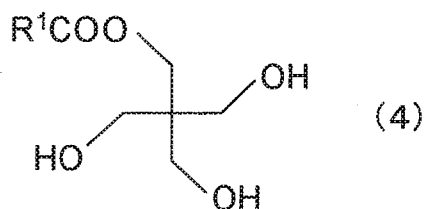


diesters of pentaerythritol and fatty acids,

represented by the following formula (3):



and monoesters of pentaerythritol and fatty acids, represented by the following formula (4).



5

In the formulas, $\text{R}^1\text{-R}^4$ each represent a chain hydrocarbon.

[0063]

The fatty acids consisting of the esters of
 10 pentaerythritol and fatty acids (R^1COOH , R^2COOH , R^3COOH ,
 and R^4COOH) are not particularly restricted so long as the
 pentaerythritol and fatty acid esters satisfy the
 conditions for the kinematic viscosity, water holding
 percentage and weight-average molecular weight, and for
 15 example, there may be mentioned saturated fatty acids,
 such as a $\text{C}_2\text{-C}_{30}$ saturated fatty acids, including acetic
 acid (C_2) (C_2 representing the number of carbons,
 corresponding to the number of carbons of each of R^1C ,
 R^2C , R^3C or R^4C , same hereunder), propanoic acid (C_3),
 20 butanoic acid (C_4) and isomers thereof, such as 2-
 methylpropanoic acid (C_4), pentanoic acid (C_5) and isomers
 thereof, such as 2-methylbutanoic acid (C_5) and 2,2-
 dimethylpropanoic acid (C_5), hexanoic acid (C_6), heptanoic
 acid (C_7), octanoic acid (C_8) and isomers thereof, such as
 25 2-ethylhexanoic acid (C_8), nonanoic acid (C_9), decanoic
 acid (C_{10}), dodecanoic acid (C_{12}), tetradecanoic acid
 (C_{14}), hexadecanoic acid (C_{16}), heptadecanoic acid (C_{17}),
 octadecanoic acid (C_{18}), eicosanoic acid (C_{20}), docosanoic
 acid (C_{22}), tetracosanoic acid (C_{24}), hexacosanoic acid

(C₂₆), octacosanoic acid (C₂₈), triacontanoic acid (C₃₀), as well as isomers thereof which are not described above.

[0064]

5 The fatty acid may also be an unsaturated fatty acid. Examples of unsaturated fatty acids include C₃-C₂₀ unsaturated fatty acids, such as monounsaturated fatty acids including crotonic acid (C₄), myristoleic acid (C₁₄), palmitoleic acid (C₁₆), oleic acid (C₁₈), elaidic acid (C₁₈), vaccenic acid (C₁₈), gadoleic acid (C₂₀) and
10 eicosenoic acid (C₂₀), di-unsaturated fatty acids including linolic acid (C₁₈) and eicosadienoic acid (C₂₀), tri-unsaturated fatty acids including linolenic acids, such as α -linolenic acid (C₁₈) and γ -linolenic acid (C₁₈), pinolenic acid (C₁₈), eleostearic acids, such as α -
15 eleostearic acid (C₁₈) and β -eleostearic acid (C₁₈), Mead acid (C₂₀), dihomog γ -linolenic acid (C₂₀) and eicosatrienoic acid (C₂₀), tetra-unsaturated fatty acids including stearidonic acid (C₂₀), arachidonic acid (C₂₀) and eicosatetraenoic acid (C₂₀), penta-unsaturated fatty
20 acids including bosseopentaenoic acid (C₁₈) and eicosapentaenoic acid (C₂₀), and partial hydrogen adducts of the foregoing.

[0065]

25 Considering the potential for degradation by oxidation and the like, the ester of pentaerythritol and a fatty acid is preferably an ester of pentaerythritol and a fatty acid, which is derived from a saturated fatty acid, i.e., an ester of pentaerythritol and a saturated fatty acid.

30 Also, from the viewpoint of lowering the water holding percentage, the ester of pentaerythritol and a fatty acid is preferably a diester, triester or tetraester, more preferably a triester or tetraester, and even more preferably a tetraester.

35 [0066]

From the viewpoint of the IOB being from about 0.00

to about 0.60, in a tetraester of pentaerythritol and a fatty acid, the total number of carbons of the fatty acid composing the tetraester of the pentaerythritol and fatty acid, i.e. the total number of carbons of the R^1C , R^2C , R^3C and R^4C portions in formula (1), is preferably about 15 (the IOB is 0.60 when the total number of carbon atoms is 15).

[0067]

Examples of tetraesters of pentaerythritol and fatty acids include tetraesters of pentaerythritol with hexanoic acid (C_6), heptanoic acid (C_7), octanoic acid (C_8), such as 2-ethylhexanoic acid (C_8), nonanoic acid (C_9), decanoic acid (C_{10}) and/or dodecanoic acid (C_{12}).

[0068]

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a triester of pentaerythritol and a fatty acid, the total number of carbons of the fatty acid composing the triester of the pentaerythritol and fatty acid, i.e. the total number of carbons of the R^1C , R^2C and R^3C portions in formula (2), is preferably about 19 or greater (the IOB is 0.58 when the number of carbon atoms is 19).

[0069]

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a diester of pentaerythritol and a fatty acid, the total number of carbons of the fatty acid composing the diester of the pentaerythritol and fatty acid, i.e. the total number of carbons of the R^1C and R^2C portion in formula (3), is preferably about 22 or greater (the IOB is 0.59 when the number of carbon atoms is 22).

[0070]

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a monoester of pentaerythritol and a fatty acid, the total number of carbons of the fatty acid composing the monoester of the pentaerythritol and fatty acid, i.e. the number of carbons of the R^1C portion in formula (4), is preferably about 25 or greater (the IOB

is 0.60 when the number of carbon atoms is 25).

The effects of double bonds, triple bonds, iso-branches and tert-branches are not considered in this calculation of the IOB (same hereunder).

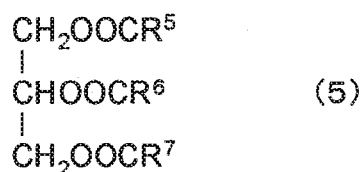
5 [0071]

Commercial products which are esters of pentaerythritol and fatty acids include UNISTAR H-408BRS and H-2408BRS-22 (mixed product) (both products of NOF Corp.).

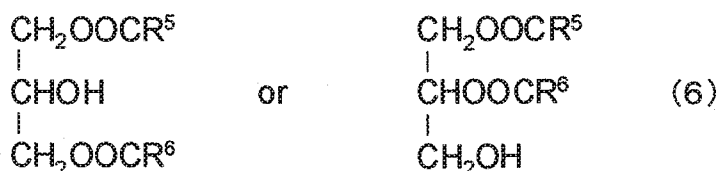
10 [0072]

[(a₂) Ester of a chain hydrocarbon triol and at least one fatty acid]

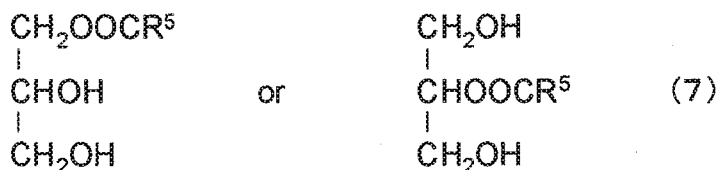
15 Examples of esters of a chain hydrocarbon triol and at least one fatty acid include triesters of glycerin and fatty acids, represented by formula (5):



diesters of glycerin and fatty acids, represented by the following formula (6):



20 and monoesters of glycerin and fatty acids, represented by the following formula (7):



wherein R⁵-R⁷ each represent a chain hydrocarbon.

[0073]

25 The fatty acid consisting of the ester of glycerin and a fatty acid (R⁵COOH, R⁶COOH and R⁷COOH) is not particularly restricted so long as the ester of glycerin and a fatty acid satisfies the conditions for the

kinematic viscosity, water holding percentage and weight-average molecular weight, and for example, there may be mentioned the fatty acids mentioned for the "(a₁) Ester of a chain hydrocarbon tetraol and at least one fatty acid",
5 namely saturated fatty acids and unsaturated fatty acids, and in consideration of the potential for degradation by oxidation and the like, the ester is preferably a glycerin and fatty acid ester, which is derived from a saturated fatty acid, i.e., an ester of glycerin and a
10 saturated fatty acid.

[0074]

Also, from the viewpoint of lowering the water holding percentage and result in greater hydrophobicity, the ester of glycerin and a fatty acid is preferably a
15 diester or triester, and more preferably a triester.

[0075]

A triester of glycerin and a fatty acid is also known as a triglyceride, and examples include triesters of glycerin and octanoic acid (C₈), triesters of glycerin
20 and decanoic acid (C₁₀), triesters of glycerin and dodecanoic acid (C₁₂), triesters of glycerin and 2 or more different fatty acids, and mixtures of the foregoing.

[0076]

Examples of triesters of glycerin and 2 or more
25 fatty acids include triesters of glycerin with octanoic acid (C₈) and decanoic acid (C₁₀), triesters of glycerin with octanoic acid (C₈), decanoic acid (C₁₀) and dodecanoic acid (C₁₂), and triesters of glycerin with octanoic acid (C₈), decanoic acid (C₁₀), dodecanoic acid
30 (C₁₂), tetradecanoic acid (C₁₄), hexadecanoic acid (C₁₆) and octadecanoic acid (C₁₈).

[0077]

In order to obtain a melting point of about 45°C or less, preferred triesters of glycerin and fatty acids are
35 those with about 40 or less as the total number of carbons of the fatty acid consisting of the triester of glycerin and the fatty acid, i.e., the total number of

carbons of the R^5C , R^6C and R^7C sections in formula (5).

[0078]

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a triester of glycerin and a fatty acid, the total number of carbons of the fatty acid composing the triester of the glycerin and fatty acid, i.e. the total number of carbons of the R^5C , R^6C and R^7C portions in formula (5), is preferably about 12 or greater (the IOB is 0.60 when the total number of carbon atoms is 12).

Triesters of glycerin and fatty acids, being aliphatic and therefore potential constituent components of the human body, are preferred from the viewpoint of safety.

[0079]

Commercial products of triesters of glycerin and fatty acids include tri-coconut fatty acid glycerides, NA36, PANACET 800, PANACET 800B and PANACET 810S, and tri-C2L oil fatty acid glycerides and tri-CL oil fatty acid glycerides (all products of NOF Corp.).

[0080]

A diester of glycerin and a fatty acid is also known as a diglyceride, and examples include diesters of glycerin and decanoic acid (C_{10}), diesters of glycerin and dodecanoic acid (C_{12}), diesters of glycerin and hexadecanoic acid (C_{16}), diesters of glycerin and 2 or more different fatty acids, and mixtures of the foregoing.

[0081]

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a diester of glycerin and a fatty acid, the total number of carbons of the fatty acid composing the diester of the glycerin and fatty acid, i.e. the total number of carbons of the R^5C and R^6C portions in formula (6), is preferably about 16 or greater (the IOB is 0.58 when the total number of carbon atoms is 16).

[0082]

Monoesters of glycerin and fatty acids are also known as monoglycerides, and examples include glycerin and octadecanoic acid (C₁₈) monoester, and glycerin and docosanoic acid (C₂₂) monoester.

5 In a monoester of glycerin and a fatty acid, the IOB is 0.59 if the total number of carbons of the fatty acid consisting the monoester of the glycerin and fatty acid, i.e. the number of carbons of the R⁵C portion in formula (7), is 19. Thus, when the number of carbons of the
10 fatty acid consisting the monoester of the glycerin and fatty acid is approximately 19 or greater, the IOB satisfies the condition of being about 0.00 to 0.60.

[0083]

15 From the viewpoint of the IOB being from about 0.00 to about 0.60, in a monoester of glycerin and a fatty acid, the total number of carbons of the fatty acid composing the monoester of the glycerin and fatty acid, i.e. the number of carbons of the R⁵C portion in formula (7), is preferably about 19 or greater (the IOB is 0.59
20 when the number of carbon atoms is 19).

[0084]

[(a₃) Ester of a chain hydrocarbon diol and at least one fatty acid]

25 Examples of an ester of a chain hydrocarbon diol and at least one fatty acid include monoesters and diesters of fatty acids with C₂-C₆ chain hydrocarbon diols, such as C₂-C₆ glycols, including ethylene glycol, propylene glycol, butylene glycol, pentylene glycol and hexylene glycol.

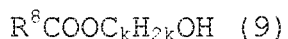
30 [0085]

Specifically, examples of an ester of a chain hydrocarbon diol and at least one fatty acid include diesters of C₂-C₆ glycols and fatty acids, represented by the following formula (8):

35
$$R^8COOC_kH_{2k}OCOR^9 \quad (8)$$

wherein k represents an integer of 2-6, and R⁸ and R⁹ each represent a chain hydrocarbon,

and monoesters of C₂-C₆ glycols and fatty acids,
represented by the following formula (9):



wherein k represents an integer of 2-6, and R⁸ is a
chain hydrocarbon.

[0086]

The fatty acid to be esterified in an ester of a C₂-
C₆ glycol and a fatty acid (corresponding to R⁸COOH and
R⁹COOH in formula (8) and formula (9)) is not particularly
restricted so long as the ester of the C₂-C₆ glycol and
fatty acid satisfies the conditions for the kinematic
viscosity, water holding percentage and weight-average
molecular weight, and for example, there may be mentioned
the fatty acids mentioned above for the "(a₁) Ester of a
chain hydrocarbon tetraol and at least one fatty acid",
namely saturated fatty acids and unsaturated fatty acids,
and in consideration of the potential for degradation by
oxidation and the like, it is preferably a saturated
fatty acid.

[0087]

From the viewpoint of the IOB being from about 0.00
to about 0.60, in a diester of butylene glycol
represented by formula (8) (k = 4) and a fatty acid, the
total number of carbons of the R⁸C and R⁹C portions is
preferably about 6 or greater (the IOB is 0.60 when the
total number of carbon atoms is 6).

[0088]

From the viewpoint of the IOB being from about 0.00
to about 0.60, in a monoester of ethylene glycol
represented by formula (9) (k = 2) and a fatty acid, the
number of carbons of the R⁸C portion is preferably about
12 or greater (the IOB is 0.57 when the number of carbon
atoms is 12).

[0089]

Considering the potential for degradation by
oxidation and the like, the ester of the C₂-C₆ glycol and
fatty acid is preferably a C₂-C₆ glycol and fatty acid

ester derived from a saturated fatty acid, or in other words, an ester of a C₂-C₆ glycol and a saturated fatty acid.

[0090]

5 Also, from the viewpoint of lowering the water holding percentage, the ester of the C₂-C₆ glycol and fatty acid is preferably a glycol and fatty acid ester derived from a glycol with a greater number of carbons, such as an ester of a glycol and a fatty acid derived
10 from butylene glycol, pentylene glycol or hexylene glycol.

 Also, from the viewpoint of lowering the water holding percentage, the ester of a C₂-C₆ glycol and fatty acid is preferably a diester.

15 Examples of commercial products of esters of C₂-C₆ glycols and fatty acids include COMPOL BL and COMPOL BS (both products of NOF Corp.).

[0091]

20 [(B) Ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety]

25 In the (B) ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain
30 hydrocarbon moiety (hereunder also referred to as "compound (B)"), it is not necessary for all of the hydroxyl groups to be etherified so long as the kinematic viscosity, water holding percentage and weight-average molecular weight are within the aforementioned ranges.

35 [0092]

 Examples of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting

for hydrogens on the chain hydrocarbon moiety (hereunder also referred to as "compound (B1)") include those mentioned for "compound (A)" as compound (A1), such as pentaerythritol, glycerin and glycol.

5 [0093]

Examples of (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety (hereunder also referred to as "compound (B2)") include compounds
10 wherein 1 hydrogen on the hydrocarbon is substituted with 1 hydroxyl group (-OH), such as aliphatic monohydric alcohols, including saturated aliphatic monohydric alcohols and unsaturated aliphatic monohydric alcohols.

[0094]

15 Examples of saturated aliphatic monohydric alcohols include C₁-C₂₀ saturated aliphatic monohydric alcohols, such as methyl alcohol (C₁) (C₁ representing the number of carbon atoms, same hereunder), ethyl alcohol (C₂), propyl alcohol (C₃) and isomers thereof, including isopropyl
20 alcohol (C₃), butyl alcohol (C₄) and isomers thereof, including sec-butyl alcohol (C₄) and tert-butyl alcohol (C₄), pentyl alcohol (C₅), hexyl alcohol (C₆), heptyl alcohol (C₇), octyl alcohol (C₈) and isomers thereof, including 2-ethylhexyl alcohol (C₈), nonyl alcohol (C₉),
25 decyl alcohol (C₁₀), dodecyl alcohol (C₁₂), tetradecyl alcohol (C₁₄), hexadecyl alcohol (C₁₆), heptadecyl alcohol (C₁₇), octadecyl alcohol (C₁₈) and eicosyl alcohol (C₂₀), as well as their isomers other than those mentioned.

[0095]

30 Unsaturated aliphatic monohydric alcohols include those wherein 1 C-C single bond of a saturated aliphatic monohydric alcohol mentioned above is replaced with a C=C double bond, such as oleyl alcohol, and for example, such alcohols are commercially available by New Japan Chemical
35 Co., Ltd. as the RIKACOL Series and UNJECOL Series.

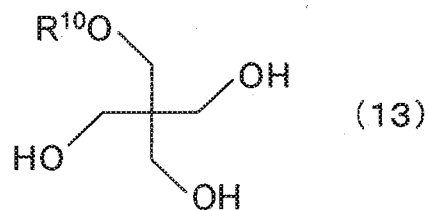
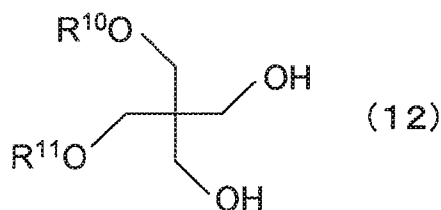
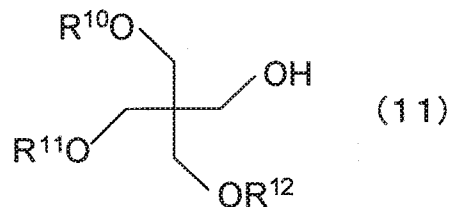
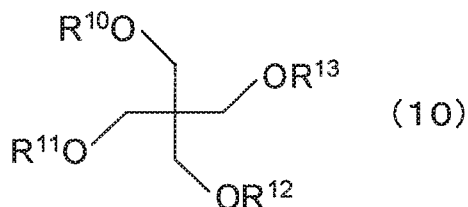
[0096]

Examples for compound (B) include (b₁) an ether of a

chain hydrocarbon tetraol and at least one aliphatic monohydric alcohol, such as monoethers, diethers, triethers and tetraethers, preferably diethers, triethers and tetraethers, more preferably triethers and tetraethers and even more preferably tetraethers, (b₂) an ether of a chain hydrocarbon triol and at least one aliphatic monohydric alcohol, such as monoethers, diethers and triethers, preferably diethers and triethers and more preferably triethers, and (b₃) an ether of a chain hydrocarbon diol and at least one aliphatic monohydric alcohol, such as monoethers and diethers, and preferably diethers.

[0097]

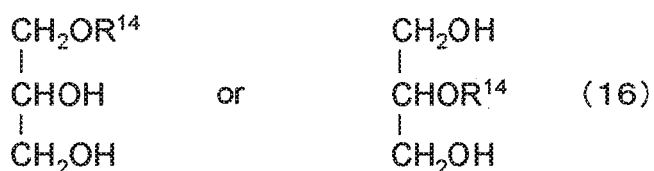
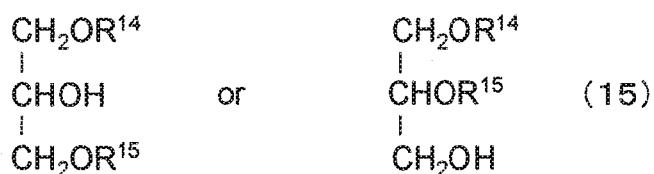
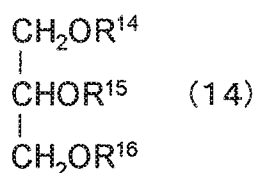
Examples of an ether of a chain hydrocarbon tetraol and at least one aliphatic monohydric alcohol include tetraethers, triethers, diethers and monoethers of pentaerythritol and aliphatic monohydric alcohols, represented by the following formulas (10)-(13):



wherein R¹⁰-R¹³ each represent a chain hydrocarbon.

[0098]

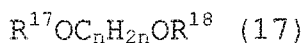
Examples of an ether of a chain hydrocarbon triol and at least one aliphatic monohydric alcohol include triethers, diethers and monoethers of glycerin and aliphatic monohydric alcohols, represented by the following formulas (14)-(16):



wherein R^{14} - R^{16} each represent a chain hydrocarbon.

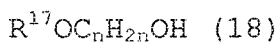
[0099]

Examples of an ether of a chain hydrocarbon diol and at least one aliphatic monohydric alcohol include diethers of C_2 - C_6 glycols and aliphatic monohydric alcohols, represented by the following formula (17):



wherein n is an integer of 2-6, and R^{17} and R^{18} are each a chain hydrocarbon,

and monoethers of C_2 - C_6 glycols and aliphatic monohydric alcohols, represented by the following formula (18):



wherein n is an integer of 2-6, and R^{17} is a chain hydrocarbon.

[0100]

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a tetraether of pentaerythritol and an aliphatic monohydric alcohol, the total number of carbon atoms of the aliphatic monohydric alcohol composing the tetraether of pentaerythritol and the aliphatic monohydric alcohol, i.e. the total number of carbon atoms of the R^{10} , R^{11} , R^{12} and R^{13} portions in formula (10), is preferably about 4 or greater (the IOB

is 0.44 when the total number of carbon atoms is 4).

[0101]

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a triether of pentaerythritol and an aliphatic monohydric alcohol, the total number of carbon atoms of the aliphatic monohydric alcohol composing the triether of pentaerythritol and the aliphatic monohydric alcohol, i.e. the total number of carbon atoms of the R^{10} , R^{11} and R^{12} portions in formula (11), is preferably about 9 or greater (the IOB is 0.57 when the total number of carbon atoms is 9).

[0102]

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a diether of pentaerythritol and an aliphatic monohydric alcohol, the total number of carbon atoms of the aliphatic monohydric alcohol composing the diether of pentaerythritol and the aliphatic monohydric alcohol, i.e. the total number of carbon atoms of the R^{10} and R^{11} portions in formula (12), is preferably about 15 or greater (the IOB is 0.60 when the total number of carbon atoms is 15).

[0103]

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a monoether of pentaerythritol and an aliphatic monohydric alcohol, the number of carbon atoms of the aliphatic monohydric alcohol composing the monoether of pentaerythritol and the aliphatic monohydric alcohol, i.e. the number of carbon atoms of the R^{10} portion in formula (13), is preferably about 22 or greater (the IOB is 0.59 when the number of carbon atoms is 22).

[0104]

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a triether of glycerin and an aliphatic monohydric alcohol, the total number of carbon atoms of the aliphatic monohydric alcohol composing the triether of glycerin and the aliphatic monohydric

alcohol, i.e. the total number of carbon atoms of the R^{14} , R^{15} and R^{16} portions in formula (14), is preferably about 3 or greater (the IOB is 0.50 when the total number of carbon atoms is 3).

5 **[0105]**

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a diether of glycerin and an aliphatic monohydric alcohol, the total number of carbon atoms of the aliphatic monohydric alcohol composing the
10 diether of glycerin and the aliphatic monohydric alcohol, i.e. the total number of carbon atoms of the R^{14} and R^{15} portions in formula (15), is preferably about 9 or greater (the IOB is 0.58 when the total number of carbon atoms is 9).

15 **[0106]**

From the viewpoint of the IOB being between about 0.00 and about 0.60, in a monoether of glycerin and an aliphatic monohydric alcohol, the number of carbon atoms of the aliphatic monohydric alcohol composing the
20 monoether of glycerin and the aliphatic monohydric alcohol, i.e. the number of carbon atoms of the R^{14} portion in formula (16), is preferably 16 or greater (the IOB is 0.58 when the number of carbon atoms is 16).

[0107]

25 From the viewpoint of the IOB being from about 0.00 to about 0.60, in a diether of butylene glycol represented by formula (17) ($n = 4$) and an aliphatic monohydric alcohol, the total number of carbon atoms of the R^{17} and R^{18} portions is preferably about 2 or greater
30 (the IOB is 0.33 when the total number of carbon atoms is 2).

From the viewpoint of the IOB being from about 0.00 to about 0.60, in a monoether of ethylene glycol represented by formula (18) ($n = 2$) and an aliphatic
35 monohydric alcohol, the number of carbon atoms of the R^{17} portion is preferably about 8 or greater (the IOB is 0.60 when the number of carbon atoms is 8).

[0108]

Compound (B) may be produced by dehydrating condensation of compound (B1) and compound (B2) in the presence of an acid catalyst.

5 [0109]

[(C) Ester of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety and (C2) a compound
10 having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety]

In the (C) ester of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain
15 hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety and (C2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety (hereunder also referred to as
20 "compound (C)"), it is not necessary for all of the carboxyl groups to be esterified so long as the kinematic viscosity, water holding percentage and weight-average molecular weight are within the aforementioned ranges.

[0110]

25 Examples of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety (hereunder also referred to as "compound (C1)") include chain hydrocarbon
30 carboxylic acids with 2-4 carboxyl groups, such as chain hydrocarbon dicarboxylic acids including alkanedicarboxylic acids, such as ethanedioic acid, propanedioic acid, butanedioic acid, pentanedioic acid, hexanedioic acid, heptanedioic acid, octanedioic acid,
35 nonanedioic acid and decanedioic acid, chain hydrocarbon tricarboxylic acids, including alkanetricarboxylic acids, such as propanetricarboxylic acid, butanetricarboxylic acid,

pentanetricioic acid, hexanetricioic acid, heptanetricioic acid, octanetricioic acid, nonanetricioic acid and decanetricioic acid, and chain hydrocarbon tetracarboxylic acids, including alkanetetracarboxylic acids, such as butanetetraoic acid, pentanetetraoic acid, hexanetetraoic acid, heptanetetraoic acid, octanetetraoic acid, nonanetetraoic acid and decanetetraoic acid.

[0111]

Compound (C1) includes chain hydrocarbon hydroxy acids with 2-4 carboxyl groups, such as malic acid, tartaric acid, citric acid and isocitric acid, chain hydrocarbon alkoxy acids with 2-4 carboxyl groups, such as O-acetylcitric acid, and chain hydrocarbon oxoacids with 2-4 carboxyl groups.

(C2) Compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety includes those mentioned for "compound (B)", such as aliphatic monohydric alcohols.

[0112]

Compound (C) may be (C₁) an ester, for example a monoester, diester, triester or tetraester, preferably a diester, triester or tetraester, more preferably a triester or tetraester and even more preferably a tetraester, of a chain hydrocarbon tetracarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 4 carboxyl groups, and at least one aliphatic monohydric alcohol, (C₂) an ester, for example, a monoester, diester or triester, preferably a diester or triester and more preferably a triester, of a chain hydrocarbon tricarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 3 carboxyl groups, and at least one aliphatic monohydric alcohol, or (C₃) an ester, for example, a monoester or diester, and preferably a diester, of a chain hydrocarbon dicarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 2 carboxyl groups, and at least one aliphatic monohydric alcohol.

Examples for compound (C) include dioctyl adipate,

diisostearyl malate, tributyl citrate and tributyl O-acetylcitrate, of which commercially available products exist.

[0113]

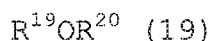
5 [(D) Compound having a chain hydrocarbon moiety and one bond selected from the group consisting of an ether bond (-O-), carbonyl bond (-CO-), ester bond (-COO-) and carbonate bond (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety]

10 The (D) compound having a chain hydrocarbon moiety and one bond selected from the group consisting of an ether bond (-O-), carbonyl bond (-CO-), ester bond (-COO-) and carbonate bond (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety (hereunder
15 also referred to as "compound (D)") may be (d₁) an ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol, (d₂) a dialkyl ketone, (d₃) an ester of a fatty acid and an aliphatic monohydric alcohol, or (d₄) a dialkyl carbonate.

20 [0114]

[(d₁) Ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol]

Ethers of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol include compounds having the
25 following formula (19):



wherein R¹⁹ and R²⁰ each represent a chain hydrocarbon.

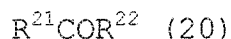
[0115]

30 The aliphatic monohydric alcohol consisting of the ether (corresponding to R¹⁹OH and R²⁰OH in formula (19)) is not particularly restricted so long as the ether satisfies the conditions for the kinematic viscosity, water holding percentage and weight-average molecular
35 weight, and for example, it may be one of the aliphatic monohydric alcohols mentioned for "compound (B)".

[0116]

[(d₂) Dialkyl ketone]

The dialkyl ketone may be a compound of the following formula (20):



5 wherein R²¹ and R²² are each an alkyl group.

The dialkyl ketone may be a commercially available product, or it may be obtained by a known method, such as by oxidation of a secondary alcohol with chromic acid or the like.

10 [0117]

[(d₃) Ester of a fatty acid and an aliphatic monohydric alcohol]

15 Examples of esters of a fatty acid and an aliphatic monohydric alcohol include compounds having the following formula (21):



wherein R²³ and R²⁴ each represent a chain hydrocarbon.

[0118]

20 Examples of fatty acids consisting of these esters (corresponding to R²³COOH in formula (21)) include the fatty acids mentioned for the "(a₁) an ester of a chain hydrocarbon tetraol and at least one fatty acids", and specifically these include saturated fatty acids and
25 unsaturated fatty acids, with saturated fatty acids being preferred in consideration of the potential for degradation by oxidation and the like. The aliphatic monohydric alcohol consisting of the ester (corresponding to R²⁴OH in formula (21)) may be one of the aliphatic
30 monohydric alcohols mentioned for "compound (B)".

[0119]

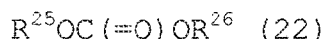
35 Examples of esters of such fatty acids and aliphatic monohydric alcohols include esters of dodecanoic acid (C₁₂) and dodecyl alcohol (C₁₂) and esters of tetradecanoic acid (C₁₄) and dodecyl alcohol (C₁₂), and examples of commercial products of esters of such fatty acids and aliphatic monohydric alcohols include ELECTOL WE20 and

ELECTOL WE40 (both products of NOF Corp.).

[0120]

[(d₄) Dialkyl carbonate]

5 The dialkyl carbonate may be a compound of the following formula (22):



wherein R²⁵ and R²⁶ are each an alkyl group.

10 The dialkyl carbonate may be a commercially available product, or it may be synthesized by reaction between phosgene and an alcohol, reaction between formic chloride and an alcohol or alcoholate, or reaction between silver carbonate and an alkyl iodide.

[0121]

15 From the viewpoint of the water holding percentage and vapor pressure, the weight-average molecular weight is preferably about 100 or greater and more preferably about 200 or greater, for (d₁) an ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol, (d₂) a dialkyl ketone, (d₃) an ester of a fatty acid and
20 an aliphatic monohydric alcohol, and (d₄) a dialkyl carbonate.

If the total number of carbon atoms is about 8 in a (d₂) dialkyl ketone, the melting point will be approximately -50°C and the vapor pressure will be about
25 230 Pa at 20°C, in the case of 5-nonanone, for example.

[0122]

[(E) Polyoxy C₃-C₆ alkylene glycol, or alkyl ester or alkyl ether thereof]

30 The (E) polyoxy C₃-C₆ alkylene glycol, or alkyl ester or alkyl ether thereof (hereunder also referred to as "compound (E)") may be (e₁) a polyoxy C₃-C₆ alkylene glycol, (e₂) an ester of a polyoxy C₃-C₆ alkylene glycol and at least one fatty acid, or (e₃) an ether of a polyoxy C₃-C₆ alkylene glycol and at least one aliphatic
35 monohydric alcohol. These will now be explained.

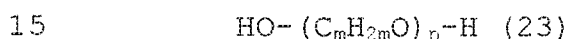
[0123]

[(e₁) Polyoxy C₃-C₆ alkylene glycol]

Polyoxy C₃-C₆ alkylene glycols refer to i) one or more homopolymers having a unit selected from the group consisting of oxy C₃-C₆ alkylene units, such as oxypropylene unit, oxybutylene unit, oxypentylene unit and oxyhexylene unit and having hydroxyl groups at both ends, ii) one or more block copolymers having 2 or more units selected from oxy C₃-C₆ alkylene units described above and oxyhexylene unit and having hydroxyl groups at both ends, or iii) random copolymers having 2 or more units selected from oxy C₃-C₆ alkylene units described above and having hydroxyl groups at both ends.

[0124]

The polyoxy C₃-C₆ alkylene glycol can be represented by the following formula (23):



wherein m represents an integer of 3-6.

[0125]

The present inventors have found that with polypropylene glycol (corresponding to a homopolymer of formula (23) where m = 3), the condition for the water holding percentage is not satisfied when the weight-average molecular weight is less than about 1,000. Therefore, polypropylene glycol homopolymer is not included in the scope of the blood slipping agent described above, and propylene glycol should be included in the (e₁) polyoxy C₃-C₆ alkylene glycol only as a copolymer or random polymer with another glycol.

[0126]

Incidentally, investigation by the present inventors suggests that with polyethylene glycol (corresponding to a homopolymer of formula (23) where m = 2), the condition for the kinematic viscosity and water holding percentage cannot be satisfied when the weight-average molecular weight is less than about 1,000.

[0127]

From the viewpoint of the IOB being about 0.00 to about 0.60, when formula (23) is polybutylene glycol (a

homopolymer where $m = 4$), for example, preferably $n \geq$ about 7 (when $n = 7$, the IOB is 0.57).

[0128]

5 Examples of commercial products of poly C_3-C_6 alkylene glycols include UNIOL™ PB-500 and PB-700 (all products of NOF Corp.).

[0129]

[(e₂) Ester of a polyoxy C_3-C_6 alkylene glycol and at least one fatty acid]

10 Examples of an ester of a polyoxy C_3-C_6 alkylene glycol and at least one fatty acids include the polyoxy C_3-C_6 alkylene glycols mentioned for "(e₁) Polyoxy C_3-C_6 alkylene glycol" in which one or both OH ends have been esterified with fatty acids, i.e. monoesters and
15 diesters.

[0130]

Examples of fatty acids to be esterified in the ester of a polyoxy C_3-C_6 alkylene glycol and at least one fatty acid include the fatty acids mentioned for the "(a₁)
20 Ester of a chain hydrocarbon tetraol and at least one fatty acid", and specifically these include saturated fatty acids and unsaturated fatty acids, with saturated fatty acids being preferred in consideration of the potential for degradation by oxidation and the like.

25 [0131]

[(e₃) Ether of a polyoxy C_3-C_6 alkylene glycol and at least one aliphatic monohydric alcohol]

Examples of an ether of a polyoxy C_3-C_6 alkylene glycols and at least one aliphatic monohydric alcohol
30 include the polyoxy C_3-C_6 alkylene glycols mentioned for "(e₁) Polyoxy C_3-C_6 alkylene glycol" wherein one or both OH ends have been etherified by an aliphatic monohydric alcohol, i.e. monoethers and diethers.

In an ether of a polyoxy C_3-C_6 alkylene glycol and at
35 least one aliphatic monohydric alcohol, the aliphatic monohydric alcohol to be etherified may be an aliphatic monohydric alcohol among those mentioned for "compound

(B)".

[0132]

[(F) Chain hydrocarbon]

5 Examples of chain hydrocarbons include (f₁) chain
alkanes, such as straight-chain alkanes and branched
chain alkanes. Straight-chain alkanes with melting
points of about 45°C or less have up to about 22 carbon
atoms, and at a vapor pressure of 1 atmosphere and no
greater than about 0.01 Pa at 25°C, the number of carbon
10 atoms is 13 or greater. Branched chain alkanes tend to
have lower melting points than chain alkanes, given the
same number of carbon atoms. Branched chain alkanes may
therefore include those with 22 and more carbon atoms,
even with melting points of below about 45°C.

15 Examples of commercially available hydrocarbon
products include PARLEAM 6 (NOF Corp.).

[0133]

The liquid-permeable top sheet may be any one that
is commonly used in the art without any particular
20 restrictions, so long as it has an uneven structure
comprising at least one projection and at least one
recess on the skin contact surface, and for example, it
may be a sheet-like material having a structure that
allows permeation of liquids, such as a film, woven
25 fabric, nonwoven fabric or the like. The fibers
composing such a woven fabric or nonwoven fabric may be
natural fibers or chemical fibers, with examples of
natural fibers including cellulose, such as ground pulp
and cotton, and examples of chemical fibers including
30 regenerated cellulose, such as rayon and fibril rayon,
semi-synthetic cellulose, such as acetate and triacetate,
thermoplastic hydrophobic chemical fibers, and
hydrophilicized thermoplastic hydrophobic chemical
fibers.

35 [0134]

Examples of thermoplastic hydrophobic chemical

fibers include polyethylene (PE), polypropylene (PP) and polyethylene terephthalate (PET) monofilaments, and fibers including PE and PP graft polymers.

Examples of nonwoven fabrics include air-through nonwoven fabrics, spunbond nonwoven fabrics, point bond nonwoven fabrics, spunlace nonwoven fabrics, needle punching nonwoven fabrics and meltblown nonwoven fabrics, as well as combinations thereof (such as SMS and the like).

[0135]

Liquid-impermeable back sheets include films comprising PE and PP, air-permeable resin films, air-permeable resin films bonded to spunbond or spunlace nonwoven fabrics, and multilayer nonwoven fabrics, such as SMS. In consideration of flexibility of the absorbent article, a low-density polyethylene (LDPE) film with a basis weight of about 15-30 g/m², for example, is preferred.

[0136]

According to one embodiment of the absorbent article of the present disclosure, the absorbent article may comprise a second sheet between the liquid-permeable top sheet and the absorbent body. The second sheet may be any of the same examples as for the liquid-permeable top sheet.

[0137]

The first example of the absorbent body is one having an absorbent core covered with a core wrap.

Examples of components for the absorbent core include hydrophilic fibers, including cellulose, such as ground pulp or cotton, regenerated cellulose, such as rayon or fibril rayon, semi-synthetic cellulose, such as acetate or triacetate, particulate polymers, filamentous polymers, thermoplastic hydrophobic chemical fibers, and hydrophilicized thermoplastic hydrophobic chemical fibers, as well as combinations of the foregoing. The component of the absorbent core may also be a super

absorbent polymer, such as granules of a sodium acrylate copolymer or the like.

[0138]

5 The core wrap is not particularly restricted so long
as it is a substance that is liquid-permeable and with a
barrier property that does not allow permeation of the
polymer absorber, and it may be a woven fabric or
nonwoven fabric, for example. The woven fabric or
nonwoven fabric may be made of a natural fiber, chemical
10 fiber, tissue, or the like.

[0139]

 A second example of the absorbent body is one formed
from an absorbing sheet or polymer sheet, with a
thickness of preferably about 0.3-5.0 mm. The absorbing
15 sheet or polymer sheet may usually be used without any
particular restrictions so long as it is one that can be
used in an absorbent article, such as a sanitary napkin.

 The blood slipping agent may be present at any
location in the planar direction of the top sheet, such
20 as across the entire top sheet, or at the center region
near the vaginal opening.

[0140]

 As regards the region in the planar direction
wherein the liquid-permeable top sheet includes a blood
25 slipping agent, according to one embodiment of the
absorbent article of the present disclosure, the liquid-
permeable top sheet comprises a blood slipping agent in
the excretory opening contact region. According to
another embodiment of the absorbent article of the
30 present disclosure, the liquid-permeable top sheet also
comprises a blood slipping agent in regions other than
the excretory opening contact region, in addition to the
excretory opening contact region, and for example, it may
comprise the blood slipping agent across the entire
35 surface of the top sheet.

[0141]

 Similarly, as regards the region in the planar

direction wherein the liquid-permeable top sheet includes a blood slipping agent, according to one embodiment of the absorbent article of the present disclosure, the liquid-permeable top sheet comprises a blood slipping agent on at least the projection of the uneven structure on the skin contact surface. If the blood slipping agent is present on the projection, the blood slipping agent present on the projection will slip down into the recess together with the menstrual blood that has reached the projection, and menstrual blood can then migrate into the absorbent body.

[0142]

According to another embodiment of the absorbent article of the present disclosure, the liquid-permeable top sheet comprises a blood slipping agent on both the projection and the recess of the uneven structure. If the blood slipping agent is present on both the projection and the recess, the blood slipping agent present on the projection will slip down into the recess together with the menstrual blood that has reached the projection, and then the blood slipping agent present in the recess can cause menstrual blood that has slipped down into the recess to migrate into the absorbent body.

[0143]

As regards the region in the thickness direction in which the liquid-permeable top sheet includes a blood slipping agent, according to an embodiment of the absorbent article of the present disclosure, the liquid-permeable top sheet comprises a blood slipping agent on the surface of the skin side, i.e. on the skin contact surface. If the blood slipping agent is present on the skin contact surface of the top sheet, menstrual blood that has reached the projection will slip down into the recess and be able to migrate into the absorbent body.

According to another embodiment of the absorbent article of the present disclosure, particularly an embodiment in which the liquid-permeable top sheet is a woven fabric or

nonwoven fabric, the liquid-permeable top sheet includes a blood slipping agent on the skin contact surface and in the interior between the skin contact surface and the clothing side surface. According to yet another embodiment of the absorbent article of the present disclosure, the liquid-permeable top sheet includes a blood slipping agent over the entire thickness direction, i.e. on the skin contact surface, in the interior between the skin contact surface and the clothing side surface, and on the clothing side surface. If the blood slipping agent is present in the top sheet interior and/or on the clothing side surface, menstrual blood present on the skin contact surface will be able to rapidly migrate into the absorbent body.

[0144]

When the liquid-permeable top sheet is formed from a nonwoven fabric or woven fabric, the blood slipping agent preferably does not obstruct the voids between the fibers of the nonwoven fabric or woven fabric, and for example, the blood slipping agent may be attached as droplets or particulates on the surface of the nonwoven fabric or woven fabric fibers, or covering the surfaces of the fibers.

[0145]

On the other hand, when the liquid-permeable top sheet is formed from a porous film, the blood slipping agent preferably does not obstruct the holes in the porous film, and for example, the blood slipping agent may be attached as droplets or particulates on the surface of the porous film. This is because if the blood slipping agent obstructs the holes in the porous film, migration of the absorbed liquid into the absorbent body may be inhibited.

In order for the blood slipping agent to slip down together with the absorbed menstrual blood, it preferably has a large surface area, and a blood slipping agent present as droplets or particulates preferably has a

small droplet/particle size.

[0146]

According to another embodiment of the absorbent article of the present disclosure, the absorbent article
5 has a second sheet comprising a blood slipping agent. According to yet another embodiment of the absorbent article of the present disclosure, the absorbent article has an absorbent body comprising a blood slipping agent.

[0147]

10 In this absorbent article, the top sheet comprises the blood slipping agent at a basis weight in the range of preferably between about 1 and about 30 g/m², more preferably between about 2 and about 20 g/m² and more preferably between about 3 and about 10 g/m². If the
15 basis weight of the blood slipping agent is lower than about 1 g/m², the absorbed menstrual blood will tend to remain in the top sheet, while if the basis weight of the blood slipping agent is greater than 30 g/m², there will tend to be an increase in sticky feel during wear.

20 [0148]

When the material to be coated with the blood slipping agent, such as the top sheet, is a nonwoven fabric, woven fabric or porous film made of a synthetic resin, it is preferably subjected to hydrophilicizing
25 treatment by coating the surface with a hydrophilic agent, or by combining it with a synthetic resin or a film. This is because, if the original material is hydrophilic, there will be lipophilic regions due to the blood slipping agent and hydrophilic regions due to the
30 hydrophilic agent sparsely dispersed on the top sheet, which will facilitate slipping down of menstrual blood onto the projection and recess of the top sheet, and its subsequent migration into the absorbent body.

[0149]

35 In an embodiment in which the domed section comprises a blood slipping agent, there are no particular restrictions on the method of coating the blood slipping

agent, and coating may be accomplished with heating as necessary, using a non-contact coater, such as for example, a spiral coater, curtain coater, spray coater or dip coater, or a contact coater or the like. A non-
5 contact coater is preferred from the viewpoint of uniformly dispersing the droplet or particulate of the blood slipping agent throughout, and from the viewpoint of not causing damage in the material. The blood slipping agent may be coated directly, if it is a liquid
10 at room temperature, or it may be heated to lower the viscosity, and when it is a solid at room temperature, it may be heated to liquefaction and coated through a control seam hot melt adhesive (HMA) gun. By increasing the air pressure of the control seam HMA gun, it is
15 possible to coat the blood slipping agent as fine particulates.

[0150]

In an embodiment in which the domed section comprises a blood slipping agent, the blood slipping
20 agent may be coated during production of the material for the top sheet and/or second sheet, such as the nonwoven fabric, or it may be coated in the manufacturing line for production of the absorbent article. In an embodiment in which the domed section comprises a blood slipping agent,
25 from the viewpoint of minimizing equipment investment, the blood slipping agent is preferably coated in the manufacturing line for the absorbent article, and in order to prevent shedding of the blood slipping agent which may contaminate the line, the blood slipping agent
30 is preferably coated during a step downstream from the manufacturing line, and specifically, immediately before encapsulation of the product in an individual package.

[0151]

The blood slipping agent also has an effect as a
35 lubricant. When the top sheet is a nonwoven fabric, therefore, the blood slipping agent can reduce friction between fibers, thereby improving the flexibility of the

nonwoven fabric as a whole. When the top sheet is a resin film, the blood slipping agent can reduce friction between the top sheet and the skin.

[0152]

5 According to a preferred embodiment of the absorbent article of the present disclosure, the absorbent article is one that is intended for absorption of blood, such as a sanitary napkin or panty liner.

10 An absorbent article of the present disclosure does not require components, such as emollients and immobilizing agents, unlike in an absorbent article containing a known skin care composition, lotion composition or the like, and the blood slipping agent alone may be applied to the top sheet.

15 [Examples]

[0153]

 The blood slipping agents used for testing are listed below.

20 [(a₁) Ester of a chain hydrocarbon tetraols and at least one fatty acid]

· UNISTAR H-408BRS, product of NOF Corp.

 Pentaerythritol tetra(2-ethylhexanoate), weight-average molecular weight: approximately 640

· UNISTAR H-2408BRS-22, product of NOF Corp.

25 Mixture of pentaerythritol tetra(2-ethylhexanoate) and neopentylglycol di(2-ethylhexanoate) (58:42 as weight ratio), weight-average molecular weight: approximately 520

[0154]

30 [(a₂) Ester of a chain hydrocarbon triols and at least one fatty acid]

· Tri-C2L oil fatty acid glyceride, product of NOF Corp.

35 Glycerin and fatty acid triester with C₈ fatty acid:C₁₀ fatty acid:C₁₂ fatty acid at a mass ratio of about 37:7:56, weight-average molecular weight: approximately 570

· Tri-CL oil fatty acid glyceride, product of NOF Corp.

Glycerin and fatty acid triester with C₈ fatty acid:C₁₂ fatty acid at a mass ratio of about 44:56, weight-average molecular weight: approximately 570
[0155]

- 5 · PANACET 810s, product of NOF Corp.

Glycerin and fatty acid triester with C₈ fatty acid:C₁₀ fatty acid at a mass ratio of about 85:15, weight-average molecular weight: approximately 480
· PANACET 800, product of NOF Corp.

- 10 Glycerin and fatty acid triester with octanoic acid (C₈) as the entire fatty acid portion, weight-average molecular weight: approximately 470
[0156]

· PANACET 800B, product of NOF Corp.

- 15 Glycerin and fatty acid triester with 2-ethylhexanoic acid (C₈) as the entire fatty acid portion, weight-average molecular weight: approximately 470
· NA36, product of NOF Corp.

- 20 Glycerin and fatty acid triester with C₁₆ fatty acid:C₁₈ fatty acid:C₂₀ fatty acid (including both saturated fatty acids and unsaturated fatty acids) at a mass ratio of about 5:92:3, weight-average molecular weight: approximately 880
[0157]

- 25 · Tri-coconut fatty acid glyceride, product of NOF Corp.

- Glycerin and fatty acid triester with C₈ fatty acid:C₁₀ fatty acid:C₁₂ fatty acid:C₁₄ fatty acid:C₁₆ fatty acid (including both saturated fatty acids and unsaturated fatty acids) at a mass ratio of about
30 4:8:60:25:3, weight-average molecular weight: 670
· Caprylic acid diglyceride, product of NOF Corp.

- Glycerin and fatty acid diester with octanoic acid as the fatty acid, weight-average molecular weight: approximately 340

- 35 [0158]

[(a₃) Ester of a chain hydrocarbon diol and at least one fatty acid]

· UNISTAR H-208BRS, product of NOF Corp.

Neopentyl glycol di(2-ethylhexanoate), weight-average molecular weight: approximately 360

· COMPOL BL, product of NOF Corp.

5 Dodecanoic acid (C_{12}) monoester of butylene glycol, weight-average molecular weight: approximately 270

· COMPOL BS, product of NOF Corp.

Octadecanoic acid (C_{18}) monoester of butylene glycol, weight-average molecular weight: approximately 350

10 [0159]

[(C_2) Ester of a chain hydrocarbon tricarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 3 carboxyl groups, and at least one aliphatic monohydric alcohol]

15 · Tributyl O-acetylcitrate, product of Tokyo Kasei Kogyo Co., Ltd.

Weight-average molecular weight: approximately 400

· Tributyl citrate, product of Tokyo Kasei Kogyo Co., Ltd.

Weight-average molecular weight: approximately 360

20 [0160]

[(C_3) Ester of a chain hydrocarbon dicarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 2 carboxyl groups, and at least one aliphatic monohydric alcohol]

25 · Dioctyl adipate, product of Wako Pure Chemical Industries, Ltd.

Weight-average molecular weight: approximately 380

[0161]

[(d_3) Ester of a fatty acid and an aliphatic monohydric alcohol]

30 · ELECTOL WE20, product of NOF Corp.

Ester of dodecanoic acid (C_{12}) and dodecyl alcohol (C_{12}), weight-average molecular weight: approximately 360

· ELECTOL WE40, product of NOF Corp.

35 Ester of tetradecanoic acid (C_{14}) and dodecyl alcohol (C_{12}), weight-average molecular weight: approximately 390

[0162]

[(e_1) Polyoxy C_3 - C_6 alkylene glycol]

- UNIOL PB500, product of NOF Corp.

Polybutylene glycol, weight-average molecular weight: approximately 500

- UNIOL PB700, product of NOF Corp.

- 5 Polyoxybutylene polyoxypropylene glycol, weight-average molecular weight: approximately 700

[0163]

[(f₁) Chain alkane]

- PARLEAM 6, product of NOF Corp.

- 10 Branched chain hydrocarbon, produced by copolymerization of liquid isoparaffin, isobutene and n-butene followed by hydrogen addition, polymerization degree: approximately 5-10, weight-average molecular weight: approximately 330

15 [0164]

[Other materials]

- NA50, product of NOF Corp.

- 20 Glycerin and fatty acid triester obtained by addition of hydrogen to NA36 for reduced proportion of double bonds from unsaturated fatty acid starting material, weight-average molecular weight: approximately 880

- (Caprylic acid/capric acid) monoglyceride, product of NOF Corp.

- 25 Glycerin and fatty acid monoester, with octanoic acid (C₈) and decanoic acid (C₁₀) at a mass ratio of about 85:15, weight-average molecular weight: approximately 220

- Monomuls 90-L2 lauric acid monoglyceride, product of Cognis Japan

30 [0165]

- Isopropyl citrate, product of Tokyo Kasei Kogyo Co., Ltd.

Weight-average molecular weight: approximately 230

- Diisostearyl malate

- 35 Weight-average molecular weight: approximately 640

- UNIOL PB1000R, product of NOF Corp.

Polybutylene glycol, weight-average molecular

weight: approximately 1,000

· UNIOL D-250, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 250

5 [0166]

· UNIOL D-400, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 400

· UNIOL D-700, product of NOF Corp.

10 Polypropylene glycol, weight-average molecular
weight: approximately 700

· UNIOL D-1000, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 1,000

15 · UNIOL D-1200, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 1,160

[0167]

· UNIOL D-2000, product of NOF Corp.

20 Polypropylene glycol, weight-average molecular
weight: approximately 2,030

· UNIOL D-3000, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 3,000

25 · UNIOL D-4000, product of NOF Corp.

Polypropylene glycol, weight-average molecular
weight: approximately 4,000

[0168]

· PEG1500, product of NOF Corp.

30 Polyethylene glycol, weight-average molecular
weight: approximately 1,500-1,600

· WILBRITE cp9, product of NOF Corp.

Polybutylene glycol compound with OH groups at both
ends esterified by hexadecanoic acid (C₁₆), weight-average
molecular weight: approximately 1,150

35 · UNILUBE MS-70K, product of NOF Corp.

Stearyl ether of polypropylene glycol, approximately

15 repeating units, weight-average molecular weight:
approximately 1,140

[0169]

· NONION S-6, product of NOF Corp.

5 Polyoxyethylene monostearate, approximately 7
repeating units, weight-average molecular weight:
approximately 880

· UNILUBE 5TP-300KB

Polyoxyethylene polyoxypropylene pentaerythritol
10 ether, produced by addition of 5 mol of ethylene oxide
and 65 mol of propylene oxide to 1 mol of
pentaerythritol, weight-average molecular weight: 4,130

[0170]

· WILBRITE s753, product of NOF Corp.

15 Polyoxyethylene polyoxypropylene polyoxybutylene
glycerin, weight-average molecular weight: approximately
960

· UNIOL TG-330, product of NOF Corp.

Glyceryl ether of polypropylene glycol,
20 approximately 6 repeating units, weight-average molecular
weight: approximately 330

[0171]

· UNIOL TG-1000, product of NOF Corp.

Glyceryl ether of polypropylene glycol,
25 approximately 16 repeating units, weight-average
molecular weight: approximately 1,000

· UNIOL TG-3000, product of NOF Corp.

Glyceryl ether of polypropylene glycol,
approximately 16 repeating units, weight-average
30 molecular weight: approximately 3,000

· UNIOL TG-4000, product of NOF Corp.

Glyceryl ether of polypropylene glycol,
approximately 16 repeating units, weight-average
molecular weight: approximately 4,000

35 [0172]

· UNILUBE DGP-700, product of NOF Corp.

Diglyceryl ether of polypropylene glycol,

approximately 9 repeating units, weight-average molecular weight: approximately 700

· UNIOX HC60, product of NOF Corp.

5 Polyoxyethylene hydrogenated castor oil, weight-average molecular weight: approximately 3,570

· Vaseline, product of Cognis Japan

Petroleum-derived hydrocarbon, semi-solid

[0173]

[Instance 1]

10 [Menstrual blood surface residue rate A, with absorption of large amount of blood]

A test was conducted to evaluate the absorption property of a sanitary napkin after one-time absorption of a large amount of blood.

15 There were prepared a top sheet, formed of a hydrophilic agent-treated air-through nonwoven fabric (composite fiber composed of polyester and polyethylene terephthalate, basis weight: 35 g/m²), a second sheet, formed of an air-through nonwoven fabric (composite fiber
20 composed of polyester and polyethylene terephthalate, basis weight: 30 g/m²), an absorbent body comprising pulp (basis weight: 150 to 450 g/m², increased at the center section), an acrylic super-absorbent polymer (basis weight: 15 g/m²) and tissue as a core wrap, a water-repellent agent-treated side sheet, and a back sheet
25 composed of a polyethylene film.

[0174]

The top sheet was a top sheet produced by the method described in Japanese Unexamined Patent Publication No.
30 2008-2034, having a ridge-furrow structure, with a ridge thickness of approximately 1.5 mm and a furrow thickness of approximately 0.4 mm, the pitch of the ridge-furrow structure (ridge width + furrow width) was approximately 4 mm, and open holes were formed in the furrows at an
35 open area of approximately 15%.

[0175]

UNISTAR H-408BRS (product of NOF Corp., tetraester

of pentaerythritol and fatty acid) was selected as the blood slipping agent, and it was coated onto the skin contact surface (ridge-furrow side) of the top sheet from a control seam HMA gun at room temperature, to a basis weight of 5.0 g/m². With an electron microscope it was confirmed that the H-408BRS was adhering onto the fiber surfaces as fine particulates.

A back sheet, an absorbent body, a second sheet, and a top sheet with the ridge-furrow side facing upward, were stacked in that order to form sanitary napkin No.1-1.

[0176]

Sanitary napkins No.1-2 to No.1-49 were produced, changing the blood slipping agent from UNISTAR H-408BRS to the ones listed in Table 2. Each blood slipping agent was used directly, when it was liquid at room temperature, or when the blood slipping agent was solid at room temperature it was heated to its melting point of +20°C, and then a control seam HMA gun was used for atomization of the blood slipping agent and coating onto the skin contact surface of the top sheet to a basis weight of about 5 g/m².

The blood slipping agent was coated onto essentially the entire skin contact surface of the top sheet, and on both the ridges and furrows.

[0177]

[Test methods]

After measuring the weight W_2 of the top sheet (the weight of the top sheet before the test), an acrylic board with an opened hole (200 mm × 100 mm, 125 g, with a 40 mm × 10 mm hole opened at the center) was placed on the top sheet, at the center section in the lengthwise direction and widthwise direction of the absorbent article, and 4.0 g of horse EDTA blood at 37±1°C (obtained by adding ethylenediaminetetraacetic acid (hereunder, "EDTA") to horse blood to prevent coagulation) was

dropped through the hole using a pipette.

[0178]

5 After dropping the horse EDTA blood, the acrylic board was immediately removed, the top sheet was taken off, the weight W_3 (weight of top sheet after the test) was measured and the "surface residue rate A (mass%)" was calculated by the following formula.

$$\text{Surface residue rate A (mass\%)} = 100 \times (W_3 - W_2) / 4.0$$

[0179]

10 The tack on the skin contact surface of the top sheet was measured at 35°C, and evaluated on the following scale.

G: No tack

F: Slight tack

15 P: Tack

[0180]

20 The surface residue rate A and tack of each absorbent article, and the properties of each blood slipping agent, are shown below in Table 2. Fig. 3 is an electron micrograph of the skin contact surface of a top sheet in a sanitary napkin wherein the top sheet comprises tri-C2L oil fatty acid glycerides.

[0181]

Table 2

No.	Blood slipping agent	Kinematic viscosity (mm ² /s, 40°C)	Water holding percentage (mass%)	Weight-average mol. wt.	IOB	Melting point (°C)	Surface residue rate A (mass%)	Tack
1-1	H-408ERS	45	0.7	640	0.13	<-5	0.8	G
1-2	H-2408ERS-22	22	0.8	520	0.18	<-5	0.8	G
1-3	Tri-C2L oil fatty acid glyceride	20	<1.0	570	0.27	37		G
1-4	Tri-CL oil fatty acid glyceride	15	<1.0	570	0.28	38		G
1-5	PANACET 810s	9	0.3	480	0.32	-5	0.8	G
1-6	PANACET 800	15	0.5	470	0.33	-5	1.8	G
1-7	PANACET 800B	20	<1.0	470	0.33	-5		G
1-8	NA36	40	<1.0	880	0.16	37		G
1-9	Tri-coconut oil fatty acid glyceride	25	<1.0	670	0.28	30		G
1-10	Capric acid diglyceride	25	2.7	340	0.58	<45	1.0	G
1-11	UNISTAR H-208ERS	8	0.7	360	0.24	<-5	0.5	G
1-12	COMPOL BL	10	1.6	270	0.50	2	1.3	G
1-13	COMPOL BS	35	0.3	350	0.36	37	2.5	G
1-14	Tributyl O-acetylcitrate	15	0.9	400	0.60	<45	0.5	G
1-15	Tributyl citrate	12	0.6	360	0.78	<45	1.8	G
1-16	Dioctyl adipate	7	0.4	380	0.27	<45	1.5	G
1-17	ELECTOL WE20	10	0.3	360	0.13	29	0.5	G
1-18	ELECTOL WE40	15	0.5	390	0.12	37	2.3	G
1-19	UNIOL PB500	40	3.6	500	0.44	<45	2.5	G
1-20	UNIOL PB700	50	2.3	700	0.49	-5	1.3	G
1-21	PARLEAM 6	5	0.06	330	0.00	-5	2.0	G

[0182]

Table 2 (cont.)

No.	Blood slipping agent	Kinematic viscosity (mm ² /s, 40°C)	Water holding percentage (mass%)	Weight-average mol. wt.	IOB	Melting point (°C)	Surface residue rate A (mass%)	Tack
1-22	NA50	80<<	-	880	0.18	52	4.3	G
1-23	(Caprylic acid/Capric acid) monoglyceride	70	4.0<<	220	1.15	<45	5.0	G
1-24	90-12 Lauric acid monoglyceride	80<<	4.0<<	<1,000	0.87	58	5.0	G
1-25	Isopropyl citrate	120	4.0<<	230	1.56	<45	4.8	F
1-26	Diisostearyl malate	450	4.0<<	640	0.28	<45	3.3	F
1-27	UNIL PB1000R	70	5.5	1000	0.40	<45	2.5	F
1-28	UNIL D-250	20	4.0<<	250	-	<45	3.8	G
1-29	UNIL D-400	30	4.0<<	400	0.76	<45	4.8	G
1-30	UNIL D-700	50	34.6	700	0.58	<45	4.8	G
1-31	UNIL D-1000	70	26.7	1,000	0.51	<45	3.8	F
1-32	UNIL D-1200	90	16.2	1,160	0.48	<45	3.0	F
1-33	UNIL D-2000	160	-	2,030	-	<45	-	P
1-34	UNIL D-3000	-	0.6	3,000	0.39	<45	3.0	P
1-35	UNIL D-4000	450	0.5	4,000	0.38	<45	2.5	P
1-36	PEG1500	120	4.0<<	1,500-1,600	0.78	40	5.5	P
1-37	WILRITE CP9	120	0.6	1,150	0.21	35	6.8	P
1-38	UNILUBE MS-70K	50	2.8	1,140	0.30	<-10	1.5	F
1-39	NONION S-6	65	4.0<<	880	0.44	37	-	G
1-40	UNILUBE 5TP-300KB	310	3.9	4,130	0.39	<45	2.0	P
1-41	WILRITE s753	120	27.3	960	0.67	-5	3.5	F
1-42	UNIL TG-330	30	-	330	1.27	<45	-	G
1-43	UNIL TG-1000	100	21.2	1,000	0.61	<45	3.5	G
1-44	UNIL TG-3000	230	4.3	3,000	0.42	<45	1.0	P
1-45	UNIL TG-4000	300	2.4	4,000	0.40	<45	2.0	P
1-46	UNILUBE DGP-700	200	4.0<<	700	0.91	<0	3.5	F
1-47	UNIOX HC60	1150	-	3,570	0.46	33	-	P
1-48	Vaseline	80<<	0.0	<1,000	0.00	55	4.0	P
1-49	None	-	-	-	-	-	7.5	G

High viscosity, unmeasurable.

[0183]

With sanitary napkin No.1-49, which had no blood slipping agent, the surface residue rate A was 7.5 mass%, but with sanitary napkins No.1-1 to No.1-21 wherein the kinematic viscosity and water holding percentage were within the prescribed ranges, the surface residue rate A was 2.5 mass% or lower.

[0184]

With sanitary napkins No.1-1 to No.1-21, it was observed that the horse EDTA blood that was dropped onto the ridges of the top sheet slipped down from the ridges into the furrows, and was rapidly absorbed from the furrows into the absorbent body. However, with sanitary napkin No.1-49 which had no blood slipping agent, the dropped horse EDTA blood did not slip down into the furrows but slowly dripped down into the furrows, most of it remaining on the ridges of the top sheet. Also, with the absorbent articles with high water holding percentage, as with No.1-30, for example, the horse EDTA blood that was dropped onto the ridges of the top sheet did not slip down into the furrows but slowly dripped while partially remaining on the top sheet, and a portion thereof remained on the ridges.

[0185]

This suggests that sanitary napkins No.1-1 to No.1-21 allow rapid migration of menstrual blood from the top sheet into the absorbent body, when a large amount of menstrual blood has reached the top sheet at once.

[0186]

Next, several volunteer subjects were asked to wear sanitary napkins Nos. 1-1 to 1-49, and most of the obtained responses indicated that with the sanitary napkins comprising blood slipping agents Nos. 1-1 to 1-21, the top sheets had no sticky feel and the top sheets were smooth, even after absorption of menstrual blood.

[0187]

[Instance 2]

[Menstrual blood surface residue rate B, with absorption of small amount of blood]

A test was conducted to evaluate the absorption property of a sanitary napkin after absorption of a small amount of blood.

There were prepared a top sheet, formed of a hydrophilic agent-treated air-through nonwoven fabric (composite fiber composed of polyester and polyethylene terephthalate, basis weight: 35 g/m²) (hereunder also referred to as "top sheet with ridges-furrows"), a second sheet, formed of an air-through nonwoven fabric (composite fiber composed of polyester and polyethylene terephthalate, basis weight: 30 g/m²), an absorbent body comprising pulp (basis weight: 150 to 450 g/m², increased at the center section), an acrylic super-absorbent polymer (basis weight: 15 g/m²) and tissue as a core wrap, a water-repellent agent-treated side sheet, and a back sheet composed of a polyethylene film.

[0188]

The top sheet was a top sheet produced by the method described in Japanese Unexamined Patent Publication No. 2008-2034, having a ridge-furrow structure, with a ridge thickness of approximately 1.5 mm and a furrow thickness of approximately 0.4 mm, and the pitch of the ridge-furrow structure (ridge width + furrow width) was approximately 4 mm and open holes were formed in the furrows at an open area of approximately 15%.

[0189]

UNISTAR H-408BRS (product of NOF Corp., tetraester of pentaerythritol and fatty acid) was selected as the blood slipping agent, and it was coated onto the skin contact surface (ridge-furrow side) of the top sheet from a control seam HMA gun at room temperature, to a basis weight of 5.0 g/m². With an electron microscope it was confirmed that the H-408BRS was adhering onto the fiber surfaces as fine particulates.

A back sheet, an absorbent body, a second sheet, and

a top sheet with the ridge-furrow side facing upward, were stacked in that order to form sanitary napkin No.2-1(i).

[0190]

5 A sanitary napkin No.2-1(ii) was formed in the same manner as the sanitary napkin No.2-1(i), except that the top sheet was changed to a top sheet formed of a flat hydrophilic agent-treated air-through nonwoven fabric (composite fiber composed of polyester and polyethylene terephthalate, basis weight: 35 g/m²), without a ridge-furrow structure (hereunder also referred to as "flat top sheet").

[0191]

15 Sanitary napkins No.2-2(i) to No.2-11(i) and No.2-2(ii) to No.2-11(ii) were produced, changing the blood slipping agent from UNISTAR H-408BRS to the ones listed in Table 3. Each blood slipping agent was used directly, when it was liquid at room temperature, or when the blood slipping agent was solid at room temperature it was

20 heated to its melting point of +20°C, and then a control seam HMA gun was used for atomization of the blood slipping agent and coating onto the skin contact surface of the top sheet to a basis weight of about 5 g/m².

25 The blood slipping agent was coated over essentially the entire skin contact surface of the top sheet, and on both the ridges and furrows of the top sheets with a ridge-furrow structure.

[0192]

[Test methods]

30 After measuring the weight W_1 of the top sheet (the weight of the top sheet before the test), approximately 0.25 g (2 drops) of horse EDTA blood at 37±1°C was added dropwise through a pipette, on the top sheet at the center in the lengthwise direction and widthwise

35 direction of the absorbent article. The horse EDTA blood was dropped onto the top parts of the ridges, in the top sheets with ridges-furrows.

[0193]

At 30 seconds after dropping, the top sheet was taken off, the weight W_5 (weight of top sheet after the test) was measured and the "surface residue rate B (mass%)" was calculated by the following formula.

$$\text{Surface residue rate B (mass\%)} = 100 \times (W_5 - W_4) / W_6$$

W_6 is the weight of the dropped horse EDTA blood, calculated from the weight of the pipette before and after dropping.

The results are shown in Table 3 below.

[0194]

Table 3

No.	Blood slipping agent	Surface residue rate B (mass%)	
		Top sheet with ridge-furrows	Flat top sheet
2-1	H-408BRS	4%	32%
2-2	PANACET 810S	8%	40%
2-3	Capric acid diglyceride	8%	24%
2-4	COMPOL BL	4%	32%
2-5	Tributyl O-acetylcitrate	8%	44%
2-6	Dioctyl adipate	8%	32%
2-7	ELECTOL WE40	8%	24%
2-8	UNIOL PB500	4%	68%
2-9	PARLEAM 6	4%	100%
2-10	UNIOL D-250	16%	48%
2-11	None	28%	28%

[0195]

Table 3 shows that when the blood slipping agent was H-408BRS, PANACET 810S, capric acid diglyceride, COMPOL BL, tributyl O-acetylcitrate, dioctyl adipate, ELECTOL WE40, UNIOL PB500 or PARLEAM 6, the surface residue rate B of the top sheet with ridges-furrows was low. This suggests that blood slipping agents having the prescribed properties cause rapid migration of small amounts of blood from the ridges to the furrows and into the absorbent body.

[0196]

[Example 3]

[Viscosity of blood containing blood slipping agent]

The viscosity of the blood slipping agent-containing blood was measured using a Rheometric Expansion System
5 ARES (Rheometric Scientific, Inc.). After adding 2 mass% of PANACET 810s to horse defibrinated blood, the mixture was gently agitated to form a sample, the sample was placed on a 50 mm-diameter parallel plate, with a gap of
10 100 μm , and the viscosity was measured at $37 \pm 0.5^\circ\text{C}$. The sample was not subjected to a uniform shear rate due to the parallel plate, but the average shear rate indicated by the device was 10 s^{-1} .

[0197]

The viscosity of the horse defibrinated blood
15 containing 2 mass% PANACET 810s was $5.9 \text{ mPa}\cdot\text{s}$, while the viscosity of the horse defibrinated blood containing no blood slipping agent was $50.4 \text{ mPa}\cdot\text{s}$. Thus, the horse defibrinated blood containing 2 mass% PANACET 810s clearly had an approximately 90% lower viscosity than the
20 blood containing no blood slipping agent.

[0198]

It is known that blood contains components, such as blood cells and has a thixotropic nature, and it is believed that the blood slipping agent of the present
25 disclosure has an effect of lowering the viscosity of blood, such as menstrual blood in the low viscosity range. Lowering the blood viscosity presumably allows absorbed menstrual blood to more easily migrate rapidly from the top sheet to the absorbent body.

30 [0199]

[Example 4]

[Photomicrograph of blood slipping agent-containing blood]

Menstrual blood was sampled from healthy volunteers
35 onto thin plastic wrap, and PANACET 810s dispersed in a 10-fold mass of phosphate-buffered saline was added to a portion thereof to a PANACET 810s concentration of 1

mass%. The menstrual blood was dropped onto a slide glass, a cover glass was placed thereover, and the state of the erythrocytes was observed with an optical microscope. A photomicrograph of menstrual blood containing no blood slipping agent is shown in Fig. 4(a), and a photomicrograph of menstrual blood containing PANACET 810s is shown in Fig. 4(b).

[0200]

From Fig. 7 it is seen that the erythrocytes formed aggregates, such as rouleaux in the menstrual blood containing no blood slipping agent, while the erythrocytes were stably dispersed in the menstrual blood containing PANACET 810s. This suggests that the blood slipping agent functions to stabilize erythrocytes in blood.

[0201]

[Example 5]

[Surface tension of blood containing blood slipping agent]

The surface tension of blood containing a blood slipping agent was measured by the pendant drop method, using a Drop Master500 contact angle meter by Kyowa Interface Science Co., Ltd. The surface tension was measured after adding a prescribed amount of blood slipping agent to sheep defibrinated blood, and thoroughly shaking.

The measurement was accomplished automatically with a device, and the surface tension γ was determined by the following formula (see Fig. 98).

[0202]

$$\gamma = g \times \rho \times (de)^2 \times 1/H$$

g : Gravitational constant

$1/H$: Correction factor determined from ds/de

ρ : Density

de : Maximum diameter

ds : Diameter at location of increase by de from

dropping edge

[0203]

The density ρ was measured at the temperatures listed in Table 4, according to JIS K 2249-1995, "Density test methods and density/mass/volume conversion tables", "5. Vibrating density test method".

The measurement was accomplished using a DA-505 by Kyoto Electronics Co., Ltd.

The results are shown in Table 4 below.

[0204]

Table 4

No.	Blood slipping agent		Measuring temperature (°C)	Surface tension (mN/m)
	Type	Amount (mass%)		
1	-	-	35	62.1
2	PANACET 810s	0.01	35	61.5
3		0.05	35	58.2
4		0.10	35	51.2
5	ELECTOL WE20	0.10	35	58.8
6	PARLEAM 6	0.10	35	57.5
7	-	-	50	56.3
8	WILBRITE cp9	0.10	50	49.1

[0205]

Based on Table 4 it is seen that the blood slipping agent has an effect of lowering the surface tension of blood.

Lowering the surface tension of blood presumably allows absorbed blood to rapidly migrate from the top sheet to the absorbent body, without being retained between the top sheet fibers.

[0206]

The present disclosure relates to the following J1 to J10.

[J1]

An absorbent article comprising a liquid-permeable top sheet, a liquid-impermeable back sheet and an absorbent body between the liquid-permeable top sheet and liquid-impermeable back sheet,

wherein the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on a skin contact surface thereof, and

5 the liquid-permeable top sheet comprises a blood slipping agent having a kinematic viscosity of 0.01 to 80 mm²/s at 40°C, a water holding percentage of 0.01 to 4.0 mass%, and a weight-average molecular weight of less than 1,000, on at least the projection in an excretory opening contact region.

10 [0207]

[J2]

The absorbent article according to J1, wherein the blood slipping agent further has an IOB of 0.00-0.60.

[0208]

15 [J3]

The absorbent article according to J1 or J2, wherein the blood slipping agent is selected from the group consisting of following items (i)-(iii), and any combination thereof:

20 (i) a hydrocarbon;

(ii) a compound having (ii-1) a hydrocarbon moiety, and (ii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

25 (iii) a compound having (iii-1) a hydrocarbon moiety, (iii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii-3) one or more, same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen of the hydrocarbon moiety;

35 with the proviso that when 2 or more oxy groups are inserted in the compound of (ii) or (iii), the oxy groups are not adjacent.

[0209]

[J4]

The absorbent article according to any one of J1 to J3, wherein the blood slipping agent is selected from the group consisting of following items (i')-(iii'), and any combination thereof:

(i') a hydrocarbon;

(ii') a compound having (ii'-1) a hydrocarbon moiety, and (ii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii') a compound having (iii'-1) a hydrocarbon moiety, (iii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii'-3) one or more, same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen on the hydrocarbon moiety;

with the proviso that when 2 or more same or different bonds are inserted in a compound of (ii') or (iii'), the bonds are not adjacent.

[0210]

[J5]

The absorbent article according to any one of J1 to J4, wherein the blood slipping agent is selected from the group consisting of following items (A)-(F), and any combination thereof:

(A) an ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(B) an ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(C) an ester of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (C2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(D) a compound having a chain hydrocarbon moiety and one bond selected from the group consisting of ether bonds (-O-), carbonyl bonds (-CO-), ester bonds (-COO-) and carbonate bonds (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety;

(E) a polyoxy C₃-C₆ alkylene glycol, or ester or ether thereof; and

(F) a chain hydrocarbon.

[0211]

[J6]

The absorbent article according to any one of J1 to J5, wherein the blood slipping agent is selected from the group consisting of (a₁) an ester of a chain hydrocarbon tetraol and at least one fatty acid, (a₂) an ester of a chain hydrocarbon triol and at least one fatty acid, (a₃) an ester of a chain hydrocarbon diol and at least one fatty acid, (b₁) an ether of a chain hydrocarbon tetraol and at least one aliphatic monohydric alcohol, (b₂) an ether of a chain hydrocarbon triol and at least one aliphatic monohydric alcohol, (b₃) an ether of a chain hydrocarbon diol and at least one aliphatic monohydric alcohol, (c₁) an ester of a chain hydrocarbon tetracarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 4 carboxyl groups, and at least one

aliphatic monohydric alcohol, (c₂) an ester of a chain hydrocarbon tricarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 3 carboxyl groups, and at least one aliphatic monohydric alcohol, (c₃) an ester of a chain hydrocarbon dicarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 2 carboxyl groups, and at least one aliphatic monohydric alcohol, (d₁) an ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol, (d₂) a dialkyl ketone, (d₃) an ester of a fatty acid and an aliphatic monohydric alcohol, (d₄) a dialkyl carbonate, (e₁) a polyoxy C₃-C₆ alkylene glycol, (e₂) an ester of a polyoxy C₃-C₆ alkylene glycols and at least one fatty acid, (e₃) an ether of a polyoxy C₃-C₆ alkylene glycol and at least one aliphatic monohydric alcohol, and (f₁) a chain alkane, and any combination thereof.

[0212]

[J7]

The absorbent article according to any one of J1 to J6, wherein the liquid-permeable top sheet has on the skin contact surface a ridge-furrow structure including a plurality of ridges and a plurality of furrows, with at least the ridges containing the blood slipping agent.

[0213]

[J8]

The absorbent article according to any one of J1 to J7, wherein the liquid-permeable top sheet has embossed sections formed by embossing at least the liquid-permeable top sheet.

[J9]

The absorbent article according to any one of J1 to J8, wherein the liquid-permeable top sheet is a nonwoven fabric or woven fabric, and the blood slipping agent is attached to the surfaces of the fibers of the nonwoven fabric or woven fabric.

[J10]

The absorbent article according to any one of J1 to J9, wherein the absorbent article is a sanitary napkin or

panty liner.

[References Signs List]

[0214]

- | | |
|----|-------------------------------|
| 5 | 1 Sanitary napkin |
| | 2 Top sheet |
| | 3 Absorbent body |
| | 4 Side sheet |
| | 5 Embossed section |
| 10 | 6 Back sheet |
| | 7 Projection |
| | 8 Recess |
| | 9 Skin contact surface |
| | 10 Blood slipping agent |
| 15 | 11, 11', 11'' Menstrual blood |

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

Claim 1

An absorbent article comprising a liquid-permeable top sheet, a liquid-impermeable back sheet and an absorbent body between the liquid-permeable top sheet and liquid-impermeable back sheet,

wherein the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on a skin contact surface thereof, and

the liquid-permeable top sheet comprises a blood slipping agent having a kinematic viscosity of 0.01 to 80 mm²/s at 40°C, a water holding percentage of 0.01 to 4.0 mass%, and a weight-average molecular weight of less than 1,000, on at least a projection in an excretory opening contact region, and a coating is configured to slip down from the liquid-permeable top sheet to the absorbent body along with menstrual blood.

Claim 2

The absorbent article according to claim 1, wherein the blood slipping agent further has an inorganic Organic Balance (IOB) of 0.00-0.60.

Claim 3

The absorbent article according to claim 1 or 2, wherein the blood slipping agent is selected from the group consisting of following items (i)-(iii), and any combination thereof:

(i) a hydrocarbon;

(ii) a compound having (ii-1) a hydrocarbon moiety, and (ii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii) a compound having (iii-1) a hydrocarbon moiety, (iii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii-3) one or more,

same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen of the hydrocarbon moiety;

with the proviso that when 2 or more oxy groups are inserted in the compound of (ii) or (iii), the oxy groups are not adjacent.

Claim 4

The absorbent article according to any one of claims 1 to 3, wherein the blood slipping agent is selected from the group consisting of following items (i')-(iii'), and any combination thereof:

(i') a hydrocarbon;

(ii') a compound having (ii'-1) a hydrocarbon moiety, and (ii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii') a compound having (iii'-1) a hydrocarbon moiety, (iii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii'-3) one or more, same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen on the hydrocarbon moiety;

with the proviso that when 2 or more same or different bonds are inserted in a compound of (ii') or (iii'), the bonds are not adjacent.

Claim 5

The absorbent article according to any one of claims 1 to 4, wherein the blood slipping agent is selected from the group consisting of following items (A)-(F), and any combination thereof:

(A) an ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the

chain hydrocarbon moiety, and (A2) a compound having a chain hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(B) an ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(C) an ester of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (C2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(D) a compound having a chain hydrocarbon moiety and one bond selected from the group consisting of ether bonds (-O-) , carbonyl bonds (-CO-) , ester bonds (-COO-) and carbonate bonds (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety;

(E) a polyoxy C₃-C₆ alkylene glycol, or ester or ether thereof; and

(F) a chain hydrocarbon.

Claim 6

The absorbent article according to any one of claims 1 to 5, wherein the blood slipping agent is selected from the group consisting of (a₁) an ester of a chain hydrocarbon tetraol and at least one fatty acid, (a₂) an ester of a chain hydrocarbon triol and at least one fatty acid, (a₃) an ester of a chain hydrocarbon diol and at least one fatty acid, (b₁) an ether of a chain hydrocarbon tetraol and at least one aliphatic monohydric alcohol, (b₂) an ether of a chain hydrocarbon triol and at least one aliphatic monohydric alcohol, (b₃) an ether of a chain hydrocarbon diol and at least one aliphatic monohydric alcohol, (c₁) an ester of a chain hydrocarbon tetracarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 4 carboxyl groups, and at least one aliphatic monohydric alcohol, (c₂) an ester of a chain hydrocarbon tricarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 3 carboxyl groups, and at least one

aliphatic monohydric alcohol, (c₃) an ester of a chain hydrocarbon dicarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 2 carboxyl groups, and at least one aliphatic monohydric alcohol, (d₁) an ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol, (d₂) a dialkyl ketone, (d₃) an ester of a fatty acid and an aliphatic monohydric alcohol, (d₄) a dialkyl carbonate, (e₁) a polyoxy C₃-C₆ alkylene glycol, (e₂) an ester of a polyoxy C₃-C₆ alkylene glycols and at least one fatty acid, (e₃) an ether of a polyoxy C₃-C₆ alkylene glycol and at least one aliphatic monohydric alcohol, and (f₁) a chain alkane, and any combination thereof.

Claim 7

The absorbent article according to any one of claims 1 to 6, wherein the at least one protection is a plurality of ridges, and the at least one recess is a plurality of furrows.

Claim 8

The absorbent article according to any one of claims 1 to 7, wherein the liquid-permeable top sheet has embossed sections formed by embossing at least the liquid-permeable top sheet.

Claim 9

The absorbent article according to any one of claims 1 to 8, wherein the liquid-permeable top sheet is a nonwoven fabric or woven fabric, and the blood slipping agent is attached to the surfaces of the fibers of the nonwoven fabric or woven fabric.

Claim 10

The absorbent article according to any one of claims 1 to 9, wherein the absorbent article is a sanitary napkin or panty liner.

Claim 11

The use of an agent in a liquid permeable top sheet in an absorbent article for modifying blood on the liquid permeable top sheet, the absorbent article comprising a liquid-permeable top sheet, a liquid-impermeable back sheet and an absorbent body between the liquid-permeable top sheet and liquid-impermeable back sheet,

wherein the liquid-permeable top sheet has an uneven structure comprising at least one projection and at least one recess on a skin contact surface thereof, and

the liquid-permeable top sheet comprises a blood slipping agent having a kinematic viscosity of 0.01 to 80 mm²/s at 40°C, a water holding percentage of 0.01 to 4.0 mass%, and a weight-average molecular weight of less than 1,000, on at least a projection in an excretory opening contact region, and a coating is configured to slip down from the liquid-permeable top sheet to the absorbent body along with menstrual blood.

Claim 12

The use of an agent according to claim 11 wherein the blood slipping agent further has an inorganic Organic Balance (IOB) of 0.00-0.60.

Claim 13

The use of an agent according to claim 12 wherein the blood slipping agent is selected from the group consisting of following items (i)-(iii), and any combination thereof:

(i) a hydrocarbon;

(ii) a compound having (ii-1) a hydrocarbon moiety, and (ii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii) a compound having (iii-1) a hydrocarbon moiety, (iii-2) one or more, same or different groups selected from the group consisting of carbonyl group (-CO-) and oxy group (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii-3) one or more,

same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen of the hydrocarbon moiety;

with the proviso that when 2 or more oxy groups are inserted in the compound of (ii) or (iii), the oxy groups are not adjacent.

Claim 14

The use of an agent according to claim 13 wherein the blood slipping agent is selected from the group consisting of following items (i')-(iii'), and any combination thereof:

(i') a hydrocarbon;

(ii') a compound having (ii'-1) a hydrocarbon moiety, and (ii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety; and

(iii') a compound having (iii'-1) a hydrocarbon moiety, (iii'-2) one or more, same or different bonds selected from the group consisting of carbonyl bond (-CO-), ester bond (-COO-), carbonate bond (-OCOO-), and ether bond (-O-) inserted between a C-C single bond of the hydrocarbon moiety, and (iii'-3) one or more, same or different groups selected from the group consisting of carboxyl group (-COOH) and hydroxyl group (-OH) substituting for a hydrogen on the hydrocarbon moiety;

with the proviso that when 2 or more same or different bonds are inserted in a compound of (ii') or (iii'), the bonds are not adjacent.

Claim 15

The use of an agent according to claim 14 wherein the blood slipping agent is selected from the group consisting of following items (A)-(F), and any combination thereof:

(A) an ester of (A1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (A2) a compound having a chain

hydrocarbon moiety and 1 carboxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(B) an ether of (B1) a compound having a chain hydrocarbon moiety and 2-4 hydroxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (B2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(C) an ester of (C1) a carboxylic acid, hydroxy acid, alkoxy acid or oxoacid comprising a chain hydrocarbon moiety and 2-4 carboxyl groups substituting for hydrogens on the chain hydrocarbon moiety, and (C2) a compound having a chain hydrocarbon moiety and 1 hydroxyl group substituting for a hydrogen on the chain hydrocarbon moiety;

(D) a compound having a chain hydrocarbon moiety and one bond selected from the group consisting of ether bonds (-O-), carbonyl bonds (-CO-), ester bonds (-COO-) and carbonate bonds (-OCOO-) inserted between a C-C single bond of the chain hydrocarbon moiety;

(E) a polyoxy C₃-C₆ alkylene glycol, or ester or ether thereof; and

(F) a chain hydrocarbon.

Claim 16

The use of an agent according to claim 15, wherein the blood slipping agent is selected from the group consisting of (a₁) an ester of a chain hydrocarbon tetraol and at least one fatty acid, (a₂) an ester of a chain hydrocarbon triol and at least one fatty acid, (a₃) an ester of a chain hydrocarbon diol and at least one fatty acid, (b₁) an ether of a chain hydrocarbon tetraol and at least one aliphatic monohydric alcohol, (b₂) an ether of a chain hydrocarbon triol and at least one aliphatic monohydric alcohol, (b₃) an ether of a chain hydrocarbon diol and at least one aliphatic monohydric alcohol, (c₁) an ester of a chain hydrocarbon tetracarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 4 carboxyl groups, and at least one aliphatic monohydric alcohol, (c₂) an ester of a chain hydrocarbon tricarboxylic acid, hydroxy acid, alkoxy acid or oxoacid with 3 carboxyl groups, and at least one aliphatic monohydric alcohol, (c₃) an ester of a chain hydrocarbon dicarboxylic acid,

hydroxy acid, alkoxy acid or oxoacid with 2 carboxyl groups, and at least one aliphatic monohydric alcohol, (d₁) an ether of an aliphatic monohydric alcohol and an aliphatic monohydric alcohol, (d₂) a dialkyl ketone, (d₃) an ester of a fatty acid and an aliphatic monohydric alcohol, (d₄) a dialkyl carbonate, (e₁) a polyoxy C₃-C₆ alkylene glycol, (e₂) an ester of a polyoxy C₃-C₆ alkylene glycols and at least one fatty acid, (e₃) an ether of a polyoxy C₃-C₆ alkylene glycol and at least one aliphatic monohydric alcohol, and (f₁) a chain alkane, and any combination thereof.

Claim 17

The use of an agent according to claim 16 wherein the at least one protection is a plurality of ridges, and the at least one recess is a plurality of furrows.

Claim 18

The use of an agent according to claim 17 wherein the liquid-permeable top sheet has embossed sections formed by embossing at least the liquid-permeable top sheet.

Claim 19

The use of an agent according to claim 18 wherein the liquid-permeable top sheet is a nonwoven fabric or woven fabric, and the blood slipping agent is attached to the surfaces of the fibers of the nonwoven fabric or woven fabric.

Claim 20

The use of an agent according to claim 19 wherein the absorbent article is a sanitary napkin or panty liner.

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Fig.1

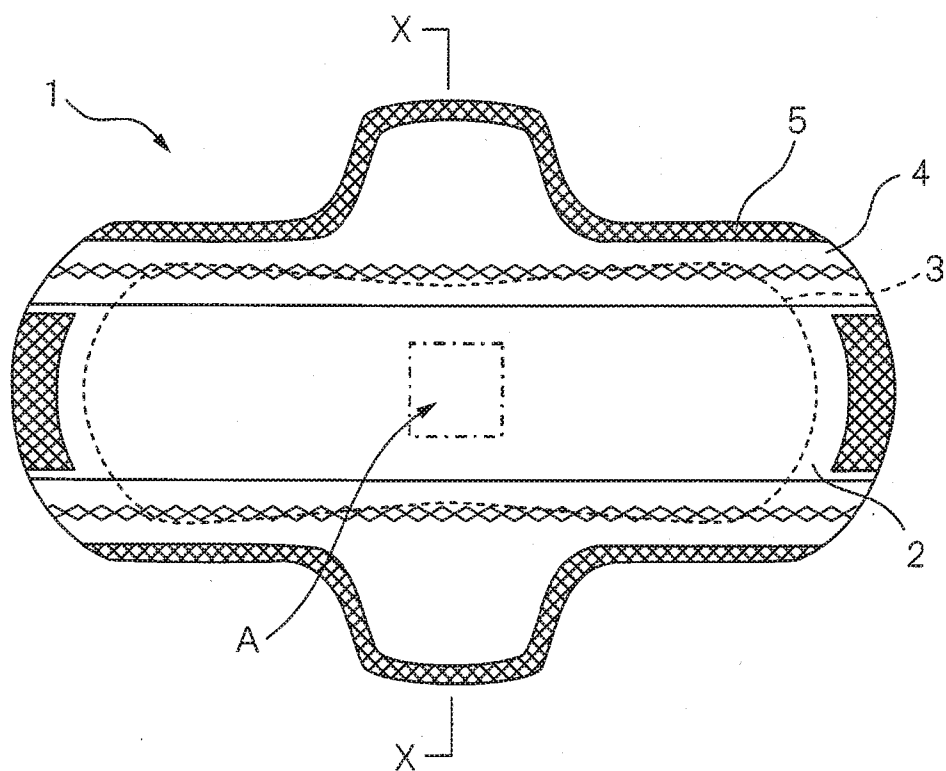
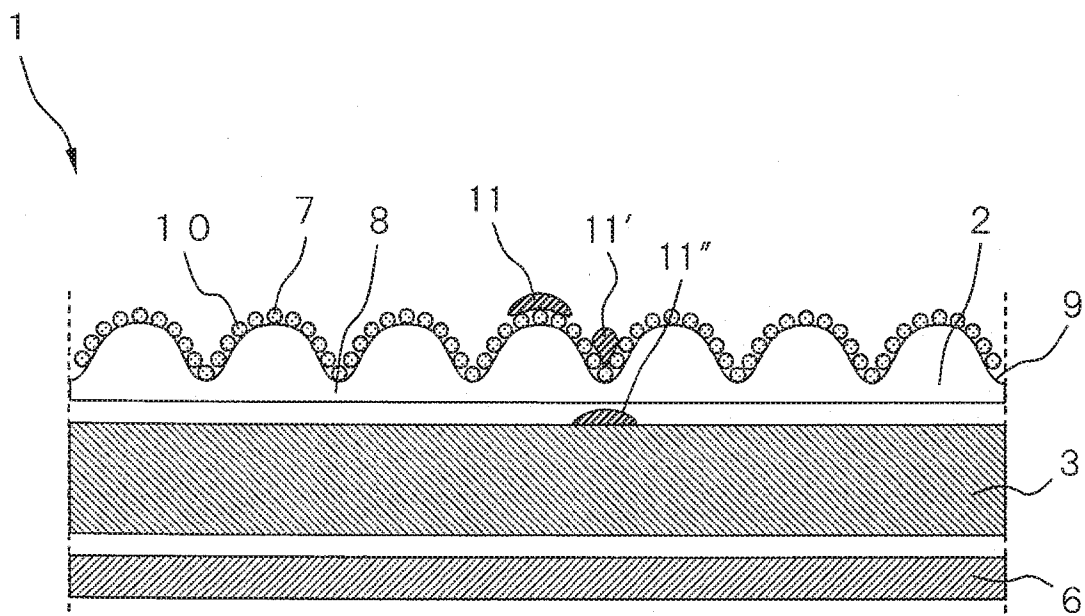
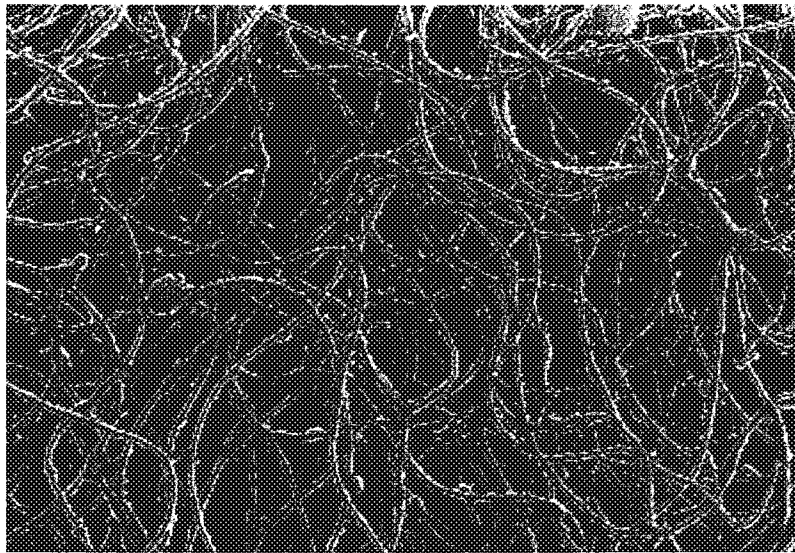


Fig.2



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Fig.3

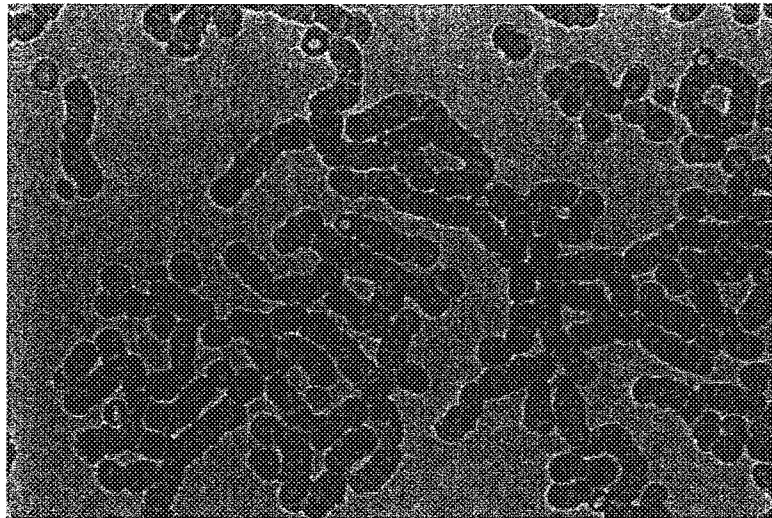


200 μ m

3/4

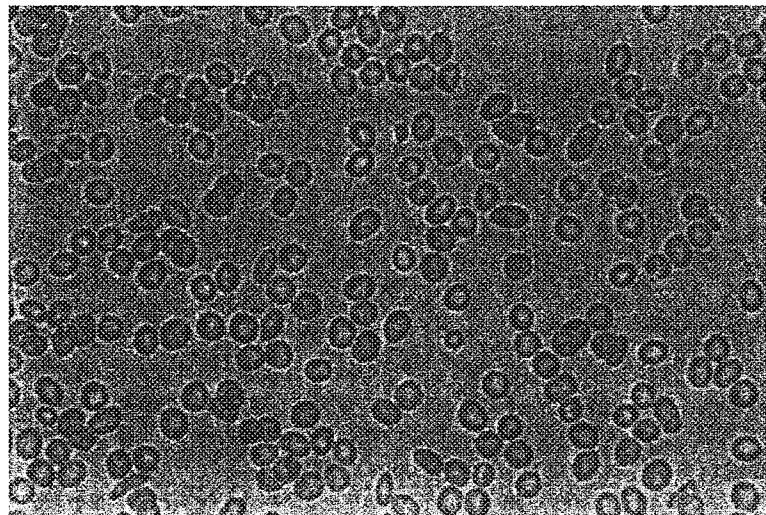
Fig.4

(a)



50 μ m

(b)



50 μ m

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Fig.5

