

622682

COMMONWEALTH of AUSTRALIA
Patents Act 1952

APPLICATION FOR A STANDARD PATENT

I/We

Courtaulds Films & Packaging (Holdings) Ltd

of

Bath Road, Bridgwater, Somerset, TA6 4PA, United Kingdom

hereby apply for the grant of a Standard Patent for an invention entitled:

Polymeric films

which is described in the accompanying complete specification.

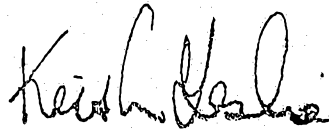
Details of basic application(s):-

<u>Number</u>	<u>Convention Country</u>	<u>Date</u>
8815983.5	United Kingdom	5 July 1988

The address for service is care of DAVIES & COLLISON, Patent Attorneys, of 1 Little Collins Street, Melbourne, in the State of Victoria, Commonwealth of Australia.

DATED this FOURTH day of JULY 1989

To: THE COMMISSIONER OF PATENTS



.....
a member of the firm of
DAVIES & COLLISON for
and on behalf of the
applicant(s)

Davies & Collison, Melbourne

M 010457 040789

COMMONWEALTH OF AUSTRALIA
PATENTS ACT 1952

DECLARATION IN SUPPORT OF CONVENTION OR
NON-CONVENTION APPLICATION FOR A PATENT

Insert title of invention.

In support of the Application made for a patent for an invention
entitled: polymeric films

Insert full name(s) and address(es)
of declarant(s) being the appli-
cant(s) or person(s) authorized to
sign on behalf of an applicant
company.

I ~~We~~ John Anthony CLAISSE of
Courtaulds Films & Packaging (Holdings) Ltd
Bath Road
Bridgwater
Somerset TA6 4PA
England

Cross out whichever of paragraphs
1(a) or 1(b) does not apply

1(a) relates to application made
by individual(s)

1(b) relates to application made
by company; insert name of
applicant company.

do solemnly and sincerely declare as follows :-

~~1. (a) I am~~ the applicant..... for the patent
~~We are~~

or (b) I am authorized by Courtaulds Films & Packaging (Holdings) Ltd

Cross out whichever of paragraphs
2(a) or 2(b) does not apply

2(a) relates to application made
by inventor(s)

2(b) relates to application made
by company(s) or person(s) who
are not inventor(s); insert full
name(s) and address(es) of inven-
tors.

the applicant..... for the patent to make this declaration on ^{its} ~~their~~ behalf.

~~2. (a) I am~~ the actual inventor..... of the invention
~~We are~~

or (b) Janette Mary SULLIVAN of 13 Greenhill Close,
Narborough, Leicester LE9 5WD, England

~~is~~ the actual inventor..... of the invention and the facts upon which the applicant.....
~~are~~
~~is~~ entitled to make the application are as follows :-

State manner in which applicant(s)
derive title from inventor(s)

By virtue of a contract of employment.
The nominated person is entitled to claim priority
from the basic application listed because the
nominated person made the basic application.

Cross out paragraphs 3 and 4
for non-convention applications.
For convention applications,
insert basic country(s) followed
by date(s) and basic applicant(s).

3. The basic application..... as defined by Section 141 of the Act ^{was} ~~were~~ made
in United Kingdom on the 5th July 1988
by Courtaulds Films & Packaging (Holdings) Ltd
in on the
by
in on the
by

4. The basic application..... referred to in paragraph 3 of this Declaration ^{was} ~~were~~
the first application..... made in a Convention country in respect of the invention the subject
of the application.

Insert place and date of signature.

Declared at Bridgwater this
Somerset, England

10th day of, December 1991

Signature of declarant(s) (no
attestation required)

J. A. Claisse
Dr J A Claisse
Authorised Signatory

Note: Initial all alterations.

DAVIES & COLLISON, MELBOURNE and CANBERRA.

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- (56) Prior Art Documents
AU 63102/86 B32B 27/08 B65D 65/40 30/08
AU 546747 80172/82 B29D 7/02 7/20 B29F 3/12
- (57) Claim

1. A polymeric film comprising a polymeric core having on each surface thereof a silicone coated heat-sealable layer containing an antiblock agent, the Davenport dynamic slip of the surface of one coated layer being different from that of the other.

2. A film according to claim 1, wherein the heat-sealable layers are of the same polymer, at least one surface of the film having been treated to increase its surface energy.

3. A film according to claim 1, wherein the heat-sealable layers are of different polymers having Vicat softening points differing by up to 10°C, at least one surface of the film having been treated to increase its surface energy.

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COMMONWEALTH OF AUSTRALIA
PATENTS ACT 1952
COMPLETE SPECIFICATION

**NAME & ADDRESS
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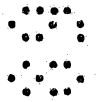
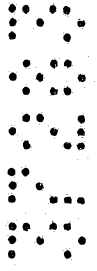
ADDRESS FOR SERVICE:

DAVIES & COLLISON
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1 Little Collins Street, Melbourne, 3000.

COMPLETE SPECIFICATION FOR THE INVENTION ENTITLED:

Polymeric films

The following statement is a full description of this invention, including the best method of performing it known to me/us:-



This invention concerns polymeric films, and in particular polymeric films having particular slip properties.

It is known to modify the slip of a polymeric film by the application of a coating of a silicone to one surface of the film. When the film is wound up, silicone is transferred from the coated surface to the uncoated surface of the film, and substantially identical slip properties are thereby

achieved for the two surfaces of the film. The increased degree of slip resulting from the use of increasing amounts of silicone can enable packaging machinery to operate at higher speeds. However, increased slip for the outer surface of the film is accompanied by increased slip for the inner surface of the film, and this is particularly disadvantageous for certain uses of such films since they slip to an unacceptable degree on the articles being packaged. The result is that the film is not sufficiently tight on the articles being packaged.

According to the present invention there is provided a polymeric film comprising a polymeric core having on each surface thereof a silicone coated heat-sealable layer containing an antiblock agent, the Davenport dynamic slip of the surface of one coated layer being different from that of the other.

Films of the present invention preferably have a difference in Davenport dynamic slip measured according to B.S. 2782 Method 824A of at least 0.6 units, more particularly at least 1.0 unit, and especially at least 2.0 units. Such films have been particularly advantageous in wrapping containers at high speed whilst combining low slip on the container with high slip between packaged containers.

The difference in Davenport dynamic slip can

be achieved by the use of the same polymer or different polymers for the heat-sealable layers and the inclusion of different amounts of antiblock agent in the two layers and/or by treating one surface of the film to increase its surface energy compared with that of the other film surface. Thus if the same polymer is used for the two heat-sealable layers, or if the difference in Vicat softening point between the two layers is not more than 10°C as measured by ASTM D1525, it is usually necessary to subject one surface of the film to a treatment to increase its surface energy in order to achieve the difference in Davenport dynamic slip. Treatments which can be used to increase the surface energy of the film include corona discharge treatment and flame treatment. This is usually the case whether the same or different amounts of antiblock agent are used in the two layers.

If the polymers used for the heat-sealable layers have Vicat softening points differing by more than 10°C as measured by ASTM D1525, a difference in Davenport dynamic slip can be achieved with the same or different amounts of antiblock agent in the respective layers, with or without a treatment to increase the surface energy of a surface of the film.

The polymers used to form the heat-sealable layers can be selected from those known in the art.

One preferred class of polymers which can be used contains units derived from at least two of ethylene, propylene and but-1-ene, a major amount in each case being derived from propylene. Another class of polymers which can be used is low density polyethylenes. It is generally preferred that at least one of the heat-seal layers is a propylene/ethylene copolymer, for example containing from 3 to 5, and preferably about 4, percent by weight of units derived from ethylene, especially where the same polymer is used for each layer. Where different polymers are used for the two layers, it is preferred that one is a propylene/ethylene copolymer and the other is a propylene/but-1-ene copolymer or a low density polyethylene, for example a linear low density polyethylene.

The absolute value of the Vicat softening points of the polymer or polymers should be such as not adversely to affect the properties of the resultant films. For example, satisfactory results have been achieved with polymers having Vicat softening points in the range of from 90 to 125°C. In all cases, the Vicat softening points of the polymers are as measured by ASTM D1525.

The amount of antiblock agent present in each layer will usually be from 0.01 to 0.30 percent by

weight of the heat-sealable layer. While the amount of antiblock agent present in each layer can be adjusted to provide differential slip between the two faces of the film, e.g. when the difference in Vicat softening point of the two layers is less than 10°C, one layer will usually contain at least 0.10 percent, preferably at least 0.15 percent and more particularly about 0.20 percent by weight of the antiblock agent. Where the two layers contain different amounts of antiblock agent, whether or not the difference in Vicat softening point between the two polymers is less than 10°C, the layer containing the lesser amount will usually contain from 0.01 to 0.15 percent by weight of the agent, while the layer containing the greater amount will usually contain from 0.15 to 0.30 percent by weight of the agent.

When different amounts of antiblock agent are present in the two heat-sealable layers, the difference is preferably at least 1000 ppm, and more preferably at least 1500 ppm.

The antiblock agent used in each layer can be selected from those known in the art, the preferred agent being silica. The particle size of the antiblock agent is preferably from 2 to 10 microns, more preferably from 4 to 6 microns, and especially about 5 microns.

The thickness of the polymeric core, which is preferably of a propylene homopolymer, can be, for example, from 15 to 50 microns. The thickness of the heat-sealable layers will usually be from 0.5 to 3.0 microns, and more particularly from 0.8 to 1.5 microns. Typically, a film of the present invention can be about 21 microns thick although thicker or thinner films can be produced according to the intended end use.

The polymeric core can contain materials used in the art to modify the physical properties of the film, for example the polymeric core can contain an anti-static agent, e.g. a bis-(2-hydroxyethyl)-amine.

The silicone coating on each surface of the films of the invention will usually have a viscosity of at least 10,000 centistokes (cs). preferably at least 30,000 cs and advantageously about 100,000 cs. In general, a coating of a silicone will be applied to one surface of the film, and this coating will then be transferred to the other surface of the film when the film is wound into a roll.

It is preferred to treat one surface of the films of the invention to increase the surface energy of the film and thereby provide and/or increase the difference in Davenport dynamic slip between the surfaces of the film, preferably using a known method

such as corona discharge treatment or flame treatment. Where the same polymer is used for each heat-sealable layer, and the same amount of antiblock agent is present in each layer, it is essential to treat one surface of the film to increase its surface energy and thereby provide a difference in Davenport dynamic slip between the surfaces of the film. It is also usually necessary to effect such a treatment with such polymers even if different amounts of antiblock agent are present in the two heat-sealable layers. Where the two heat-sealable layers contain different amounts of antiblock agent, it is generally preferred that the layer containing the lesser amount of antiblock agent be subjected to a treatment to increase its surface energy.

It is generally preferred that the surface of the film having a silicone coating applied thereto is not subjected to a treatment to increase its surface energy, but of course silicone will be transferred from the silicone coated surface to the treated surface when the film is wound up.

Films of the present invention can be manufactured in known manner and they will usually be biaxially oriented, for example sequentially or using a bubble technique.

The following Examples are given by way of

illustration only. All parts are by weight unless stated otherwise.

Example 1

A three-layer polymeric web was produced by coextruding a core layer of a propylene homopolymer with a layer of a propylene/ethylene copolymer (4% ethylene) containing 0.20 percent by weight of silica of 5 micron particle size on one side, and a layer of the same copolymer but containing 0.10 percent by weight of the silica on the other side.

The web was stretched 4.5 times in the direction of extrusion, and the layer containing 0.20 percent by weight of silica was coated with a layer of a silicone slip agent. The resulting coated film was then stretched 10 times in the transverse direction, and wound up. After approximately twenty one days, when silicone from the coating on one face of the film had transferred to the other face of the film, the Davenport dynamic slip of each face of the film to itself was then measured according to BS.2782 Method 824A. The thickness of the silicone coating on each surface of the film after this time was 11-12 Angstroms as measured by X-ray photo electron spectroscopy. The results are shown in the accompanying Table.

Example 2

The procedure of Example 1 was repeated, but with the surface not having the silicone coating applied thereto being corona discharge treated before wind-up of the film. The test results obtained with the film are shown in the accompanying Table.

Examples 3-12

The procedure of Examples 1 and 2 were repeated using outer layers with different amounts of antiblock agents and/or different polymers, in some cases with and without corona discharge treatment. The results are shown in the accompanying Table.

A significant differential slip was noted where the difference in dynamic Davenport film-to-film slip values between outer to outer and inner to inner were greater than 0.6 slip units.

TABLE

Example	Outside Face	Inside Face	Davenport Slip Film-Film	
			Static O-O/I-I	Dynamic O-O/I-I
1	2000ppm, 5mym silica in PP-co-E	1000ppm, 5mym silica in PP-co-E	3.3/3.3	4.7/4.4
2	As Example 1	As Example 1 but corona treated to 42 dynes	3.1/3.0	4.4/3.6
3	As Example 1	500ppm, 5mym silica in PP-co-E	3.2/3.3	4.3/4.7
4	As Example 1	As Example 3 but corona treated to 42 dynes	3.0/2.2	5.2/2.9
5	As Example 1	500ppm, 2mym silica in PP-co-E	3.5/3.3	4.6/4.9
6	As Example 1	As Example 5 but corona treated to 42 dynes	3.2/2.2	4.8/3.4
7	As Example 1	100ppm, 5mym silica in PP-co-E	3.2/2.6	4.7/4.3
8	As Example 1	As Example 7 but corona treated to 42 dynes	3.5/2.3	4.7/3.8
9	As Example 1	100ppm, 2mym silica in PP-co-E	3.6/2.2	4.3/4.7
10	As Example 1	As Example 9 but corona treated to 42 dynes	3.3/2.1	5.0/3.7
11	As Example 1	500ppm, 5mym silica in PP-co-B (33% butene)	3.0/2.2	4.2/3.1
12	As Example 1	1000ppm, 5mym silica in LLDPE	3.6/2.6	5.6/4.3



THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS.

1. A polymeric film comprising a polymeric core having on each surface thereof a silicone coated heat-sealable layer containing an antiblock agent, the Davenport dynamic slip of the surface of one coated layer being different from that of the other.

2. A film according to claim 1, wherein the heat-sealable layers are of the same polymer, at least one surface of the film having been treated to increase its surface energy.

3. A film according to claim 1, wherein the heat-sealable layers are of different polymers having Vicat softening points differing by up to 10°C, at least one surface of the film having been treated to increase its surface energy.

4. A film according to claim 1, wherein the heat-sealable layers are of different polymers having Vicat softening points differing by at least 10°C.

5. A film according to any ^{one} of the preceding claims, wherein at least one of the heat-sealable layers is of a propylene/ethylene copolymer.

6. A film according to any ^{one} of the preceding claims, wherein at least one of the heat-sealable layers contains from 0.15 to 0.30 percent by weight of antiblock agent.



7. A film according to any^{one}₁ of the preceding claims, wherein one heat-sealable layer contains more antiblock agent than the other heat-sealable layer.

8. A film according to claim 7, wherein one of the heat-sealable layers contains from 0.15 to 0.30 percent by weight of antiblock agent and the other heat-sealable layer contains from 0.01 to 0.15 percent by weight of antiblock agent.

9. A film according to claim 8, wherein the polymer forming the heat-sealable layer containing from 0.15 to 0.30 percent by weight of antiblock agent has a Vicat softening point of at least 10°C higher than that of the other heat-sealable layer.

10. A film according to any^{one}₁ of the preceding claims, wherein the antiblock agent comprises silica.

11. A film according to any^{one}₁ of the preceding claims, wherein the antiblock agent has a mean particle size of from 2 to 10 microns.

12. A film according to any^{one}₁ of the preceding claims, wherein the polymeric core comprises polypropylene.



13. A film according to claim 1, substantially as hereinbefore described with reference to the examples.

~~14. The steps, features, compositions and compounds disclosed herein or referred to or indicated in the specification and/or claims of this application, individually or collectively, and any and all combinations of any two or more of said steps or features.~~

DATED this FOURTH day of JULY 1989

Courtaulds Films & Packaging (Holdings) Ltd

by DAVIES & COLLISON
Patent Attorneys for the applicant(s)

