



- (51) International Patent Classification:
A61F 13/15 (2006.01) *C08L 23/08* (2006.01)
- (21) International Application Number:
PCT/US2015/013851
- (22) International Filing Date:
30 January 2015 (30.01.2015)
- (25) Filing Language: English
- (26) Publication Language: English
- (71) Applicant: **KIMBERLY-CLARK WORLDWIDE, INC.**
[US/US]; 2300 Winchester Road, Neenah, WI 54956 (US).
- (72) Inventors: **CHEN, Feng**; 2300 Winchester Road, Neenah, WI 54956 (US). **FAULKES, Michael, J.**; 2300 Winchester Road, Neenah, WI 54956 (US). **MLEZIVA, Mark, M.**; 2300 Winchester Road, Neenah, WI 54956 (US). **WIDEMAN, Gregory, J.**; 2300 Winchester Road, Neenah, WI 54956 (US).
- (74) Agents: **JOHNSTON, Jason, W.** et al.; Dority & Manning, P.A., P.O. Box 1449, Greenville, SC 29602-1449 (US).
- (81) Designated States (*unless otherwise indicated, for every kind of national protection available*): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.
- (84) Designated States (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK,

[Continued on next page]

(54) Title: ABSORBENT ARTICLE PACKAGE WITH REDUCED NOISE

(57) Abstract: A package that contains a wrapper material that defines an interior cavity within which an absorbent article is removably positioned is provided. The wrapper material is formed from a film that includes a polymer composition containing at least one ethylene polymer and at least one nano-filler. The present inventors have discovered that through selective control over the particular type and concentration of these components, as well as the manner in which it is formed, the resulting package may generate a relatively low degree of noise when physically deformed.

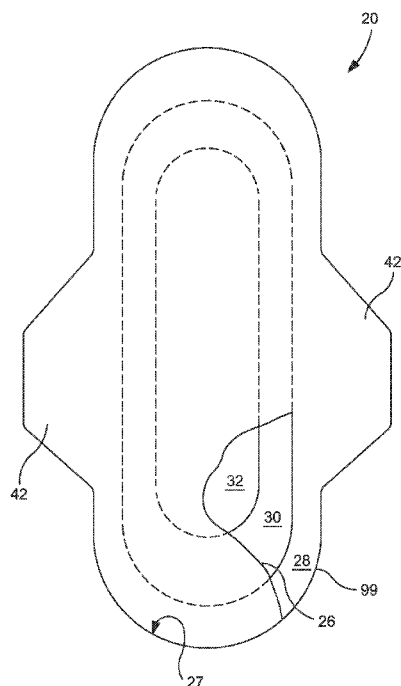


FIG. 2

WO 2016/122619 A1



SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, **Published:**
GW, KM, ML, MR, NE, SN, TD, TG).

— *with international search report (Art. 21(3))*

ABSORBENT ARTICLE PACKAGE WITH REDUCED NOISE**Background of the Invention**

Absorbent articles are generally constructed from an absorbent core that is positioned between a liquid-permeable topsheet, which defines a "body-facing" surface disposed toward the body, and a liquid-impermeable backsheet, which defines a "garment-facing" surface disposed away from the body. Feminine care absorbent articles, such as sanitary napkins, are often used during a woman's menstrual cycle or between menstrual cycles for light incontinence purposes. Because many of these articles are carried in a woman's purse or pocket prior to use, it is advantageous to wrap the article within a package to keep it clean and sanitary. Conventionally, the wrapping material includes a polyethylene film, which is folded around the absorbent article and then sealed. Many women value their personal privacy and prefer not to advertise to others that they are carrying or using feminine care products. Unfortunately, conventional wrapping materials and packages tend to exacerbate this problem as they are inherently "loud" when manipulated. In this regard, various technologies have been proposed to reduce the level of noise attributable to the film used in absorbent article. For example, attempts have been made to apply fibrous layers (e.g., nonwoven webs) to the film to help suppress the degree of noise it generates during use. In addition to being costly and overly complex, such materials can significantly increase the thickness of the wrapper material and also adversely impact certain of its mechanical properties, such as ductility.

As such, a need currently exists for improved absorbent article packages and wrapper materials that can generate a reduced amount of noise.

Summary of the Invention

In accordance with one embodiment of the present invention, a package is disclosed that comprises a wrapper material that defines an interior cavity and an absorbent article removably positioned within the cavity. The wrapper material is formed from a film having a thickness of from about 0.5 to about 80 micrometers. The film contains a polymer composition that includes one or more ethylene polymers in an amount of from about 75 wt.% to about 99 wt.% of the polymer composition and one or more nanofillers in an amount of from about 0.5 wt.% to about 20 wt.% of the polymer composition. The package exhibits a normalized noise level of about 3.0 or less as determined at a frequency of 4,000 Hz, the normalized noise level being determined by dividing the noise level of the package by the noise level of an ambient environment.

Other features and aspects of the present invention are described in more detail below.

Brief Description of the Drawings

A full and enabling disclosure of the present invention, including the best mode thereof, directed to one of ordinary skill in the art, is set forth more particularly in the remainder of the specification, which makes reference to the appended figures in which:

5 Fig. 1 is a schematic illustration of one embodiment of a process that can be used to form a film for use in the package of the present invention;

Fig. 2 is a perspective view of one embodiment of the absorbent article that can be employed in the package of the present invention;

10 Fig. 3 is a perspective view of one embodiment of an absorbent article package that may be formed in accordance with the present invention;

Fig. 4 is a perspective view of the package of Fig. 3 shown in its opened state;

Fig. 5 is a cross-sectional view of the package taken along the lines indicated in Fig. 3;

Fig. 6 is an X-ray diffraction graph for Example 8;

15 Fig. 7 are transmission electron microphotographs of Example 8 in which Fig. 7(a) is at a lower magnification (depicted scale of 2 μm) and Fig. 7(b) is at a higher magnification (depicted scale of 200 nm);

Fig. 8 is a graph that depicts logarithmic storage modulus (G') vs. logarithmic frequency (γ) for the samples of Example 8;

20 Fig. 9 is a graph that depicts complex viscosity η^* vs. logarithmic frequency (γ) for the samples of Example 8;

Fig. 10 is a graph that depicts $\log G'$ (storage modulus) and $\log G''$ (loss modulus) vs. logarithmic frequency (γ) for the samples of Example 8;

Fig. 11 is a perspective view of a testing apparatus that may be used to evaluate noise levels, with the apparatus door open;

25 Fig. 12 is a perspective view of the testing apparatus of Fig. 11, with the apparatus door closed; and

Fig. 13 is a plan view of the apparatus of Fig. 11 taken along arrow 190.

Repeat use of reference characters in the present specification and drawings is intended to represent same or analogous features or elements of the invention.

Detailed Description of Representative Embodiments

30 Reference now will be made in detail to various embodiments of the invention, one or more examples of which are set forth below. Each example is provided by way of explanation, not limitation of the invention. In fact, it will be apparent to those skilled in the art that various modifications and variations may be made in the present invention without departing from the scope or spirit of the

invention. For instance, features illustrated or described as part of one embodiment, may be used on another embodiment to yield a still further embodiment. Thus, it is intended that the present invention cover such modifications and variations.

Generally speaking, the present invention is directed to a package that contains a wrapper material that defines an interior cavity within which an absorbent article is removably positioned. The wrapper material is formed from a film that includes a polymer composition containing at least one ethylene polymer and at least one nanofiller. The present inventors have discovered that through selective control over the particular type and concentration of these components, as well as the manner in which it is formed, the resulting package may generate a relatively low degree of noise when physically deformed. When subjected to physical deformation for two (2) minutes, for instance, the noise level of the package may be about 50 decibels (dB) or less, in some embodiments about 48 dB or less, and in some embodiments, from about 20 dB to about 47 dB, such as determined at a frequency of 2,000 Hz or 4,000 Hz. The "normalized noise level" of the package, which is determined by dividing the noise level of the film that is generated while the package is subjected to physical deformation for two (2) minutes by the noise level generated by an ambient environment, may likewise be about 3.0 or less, in some embodiments about 2.9 or less, and in some embodiments, from about 1.5 to about 2.8 such as determined at a frequency of 2,000 Hz or 4,000 Hz. Further, the noise level of the wrapper material may also be about 45 decibels (dB) or less, in some embodiments about 42 dB or less, and in some embodiments, from about 20 dB to about 40 dB, such as determined at a frequency of 2,000 Hz or 4,000 Hz. The "normalized noise level" of the film, which is determined by dividing the noise level of the film that is generated while the film is subjected to physical deformation for two (2) minutes by the noise level generated by an ambient environment, may likewise be about 2.5 or less, in some embodiments about 2.4 or less, and in some embodiments, from about 1.5 to about 2.3, such as determined at a frequency of 2,000 Hz or 4,000 Hz.

One beneficial aspect of the present invention is that the low noise production may be provided without the need for a noise-reducing fibrous material (e.g., nonwoven web) on the film. In fact, the film may be entirely free of such a material. Among other things, this allows the film and resulting wrapper material to have a small thickness, such as from about 0.5 to about 80 micrometers, in some embodiments from about 1 to about 50 micrometers, in some embodiments from about 5 to about 35 micrometers, and in some embodiments, from about 10 to about 30 micrometers. Nevertheless, due to its unique structure and the manner in which it is formed, the film of the present invention can still maintain good mechanical properties, such as a high ductility. One parameter that is indicative of good ductility is the peak elongation of the film in the machine direction ("MD") and/or cross-machine direction ("CD"). For example, the film typically exhibits a peak elongation in the

machine direction of about 400% or more, in some embodiments about 500% or more, in some embodiments about 550% or more, and in some embodiments, from about 600% to about 2000%.

The film may likewise exhibit a peak elongation in the cross-machine direction of about 750% or more, in some embodiments about 800% or more, in some embodiments about 800% or more, and in some
5 embodiments, from about 850% to about 2500%. Despite having such good ductility, the film of the present invention is nevertheless able to retain good mechanical strength. For example, the film of the present invention may exhibit an ultimate tensile strength in the machine direction and/or cross-machine direction of from about 20 to about 150 Megapascals (MPa), in some embodiments from about 25 to about 100 MPa, and in some embodiments, from about 30 to about 80 MPa. The Young's
10 modulus of elasticity of the film, which is equal to the ratio of the tensile stress to the tensile strain and is determined from the slope of a stress-strain curve, may also be good. For example, the film typically exhibits a Young's modulus in the machine direction and/or cross-machine direction of from about 50 to about 500 MPa, in some embodiments from about 100 to about 400 MPa, and in some embodiments, from about 150 to about 350 MPa.

Surprisingly, the good ductility and other mechanical properties can be achieved even though the film has a very low thickness. In this regard, the normalized mechanical properties, which are determined by dividing a particular mechanical value (e.g., Young's modulus, tensile strength, or peak elongation) by the average film thickness (μm), may also be improved. For example, the film may exhibit a normalized peak elongation in the machine direction of about 30%/ μm or more, in some
20 embodiments about 40%/ μm or more, and in some embodiments, from about 50%/ μm to about 150%/ μm . The film may likewise exhibit a normalized peak elongation in the cross-machine direction of about 40%/ μm or more, in some embodiments about 50%/ μm or more, and in some embodiments, from about 60%/ μm to about 200%/ μm . The film may exhibit a normalized ultimate tensile strength in the machine direction and/or cross-machine direction of from about 0.5 to about 20 MPa/ μm , in some
25 embodiments from about 1 to about 12 MPa/ μm , and in some embodiments, from about 2 to about 8 MPa/ μm . The normalized Young's modulus in the machine direction and/or cross-machine direction may also be from about 5 to about 50 MPa/ μm , in some embodiments from about 10 to about 40 MPa/ μm , and in some embodiments, from about 15 to about 35 MPa/ μm .

Various embodiments of the present invention will now be described in more detail.

I. Polymer Composition

A. Ethylene Polymer

Ethylene polymers typically constitute from about 80 wt.% to about 99.9 wt.%, in some embodiments from about 90 wt.% to about 99.5 wt.%, and in some embodiments, from about 95 wt.% to 98 wt.% of the polymer content of the polymer composition. Likewise, ethylene polymers may

constitute from about 75 wt.% to about 99 wt.%, in some embodiments from about 80 wt.% to about 98 wt.%, and in some embodiments, from about 85 wt.% to 95 wt.% of the entire polymer composition.

Any of a variety of ethylene polymers may generally be employed in the present invention. In one embodiment, for instance, the ethylene polymer may be a copolymer of ethylene and an α -olefin, such as a C₃-C₂₀ α -olefin or C₃-C₁₂ α -olefin. Suitable α -olefins may be linear or branched (e.g., one or more C₁-C₃ alkyl branches, or an aryl group). Specific examples include 1-butene; 3-methyl-1-butene; 3,3-dimethyl-1-butene; 1-pentene; 1-pentene with one or more methyl, ethyl or propyl substituents; 1-hexene with one or more methyl, ethyl or propyl substituents; 1-heptene with one or more methyl, ethyl or propyl substituents; 1-octene with one or more methyl, ethyl or propyl substituents; 1-nonene with one or more methyl, ethyl or propyl substituents; ethyl, methyl or dimethyl-substituted 1-decene; 1-dodecene; and styrene. Particularly desired α -olefin comonomers are 1-butene, 1-hexene and 1-octene. The ethylene content of such copolymers may be from about 60 mole% to about 99 mole%, in some embodiments from about 80 mole% to about 98.5 mole%, and in some embodiments, from about 87 mole% to about 97.5 mole%. The α -olefin content may likewise range from about 1 mole% to about 40 mole%, in some embodiments from about 1.5 mole% to about 15 mole%, and in some embodiments, from about 2.5 mole% to about 13 mole%. The density of the polyethylene may vary depending on the type of polymer employed, but generally ranges from about 0.85 to about 0.96 grams per cubic centimeter (g/cm³). Polyethylene "plastomers", for instance, may have a density in the range of from about 0.85 to about 0.91 g/cm³. Likewise, "linear low density polyethylene" (LLDPE) may have a density in the range of from about 0.91 to about 0.940 g/cm³; "low density polyethylene" (LDPE) may have a density in the range of from about 0.910 to about 0.940 g/cm³; and "high density polyethylene" (HDPE) may have density in the range of from about 0.940 to about 0.960 g/cm³, such as determined in accordance with ASTM D792.

In certain embodiments, an ethylene polymer may be employed that has a relatively low density in the range of about 0.94 g/cm³ or less, in some embodiments from about 0.85 to about 0.94 g/cm³, and in some embodiments, from about 0.90 to about 0.935 g/cm³. One or more polymers may be employed in the composition that has these density characteristics. Linear low density polyethylene ("LLDPE") and/or low density polyethylene ("LDPE") are particularly suitable. The low density ethylene polymer may have a relatively low melting temperature and modulus of elasticity, which can provide the resulting film with a relatively soft and ductile feel. For example, the low density ethylene polymer may have a melting temperature of from about 50°C to about 145°C, in some embodiments from about 75°C to about 140°C, and in some embodiments, from about 100°C to about 135°C, and a modulus of elasticity of from about 50 to about 700 MPa, in some embodiments from about 75 to about 600 MPa, and in some embodiments, from about 100 to about 500 MPa, as determined in accordance

with ASTM D638-10. The low density ethylene polymer may also have a melt flow index of from about 0.1 to about 100 grams per 10 minutes, in some embodiments from about 0.5 to about 50 grams per 10 minutes, and in some embodiments, from about 1 to about 40 grams per 10 minutes, determined at a load of 2160 grams and at 190°C, as determined in accordance with ASTM D1238-13 (or ISO 1133).

5 If desired, low density ethylene polymers may constitute a substantial majority of the polymers employed in the composition. For example, low density ethylene polymers may constitute about 80 wt.% or more, in some embodiments about 85 wt.% or more, and in some embodiments, from about 90 wt.% to 100 wt.% of the polymers employed in the composition. Of course, in other embodiments, high density ethylene polymers may also be employed. For example, low density ethylene polymers
10 may constitute from about 5 wt.% to about 90 wt.%, in some embodiments from about 10 wt.% to about 80 wt.%, and in some embodiments, from about 20 wt.% to 70 wt.% of the polymer composition and high density ethylene polymers may constitute from about 5 wt.% to about 90 wt.%, in some embodiments from about 10 wt.% to about 80 wt.%, and in some embodiments, from about 20 wt.% to 70 wt.% of the polymer composition.

15 The high density ethylene polymers typically have a density of greater than about 0.94 g/cm³, in some embodiments from about 0.945 to about 0.98 g/cm³, and in some embodiments, from about 0.95 to about 0.97 g/cm³. Once again, one or more polymers may be employed in the composition that has these characteristics. High density polyethylene ("HDPE") is particularly suitable. The high density ethylene polymers may have a relatively low melting temperature and high modulus of
20 elasticity. For example, the high density ethylene polymers may have a melting temperature of from about 70°C to about 160°C, in some embodiments from about 85°C to about 150°C, and in some embodiments, from about 110°C to about 145°C, and a modulus of elasticity of from about 700 to about 5,000 MPa, in some embodiments from about 750 to about 3,000 MPa, and in some embodiments, from about 1,000 to about 2,000 MPa, as determined in accordance with ASTM D638-
25 10. The high density ethylene polymers may also have a melt flow index of from about 0.1 to about 100 grams per 10 minutes, in some embodiments from about 0.5 to about 50 grams per 10 minutes, and in some embodiments, from about 1 to about 40 grams per 10 minutes, determined at a load of 2160 grams and at 190°C, as determined in accordance with ASTM D1238-13 (or ISO 1133).

Regardless of the particular type, various known techniques may generally be employed to
30 form the ethylene polymers. For instance, ethylene polymers may be formed using a free radical or a coordination catalyst (e.g., Ziegler-Natta). Typically, the ethylene polymer is formed from multi-site Ziegler-Natta catalysts, the resulting ethylene polymer has a broad molecular weight distribution with a polydispersity index (weight average molecular weight divided by number average molecular weight) of up to 20 or higher. The ethylene polymer made by a single-site coordination catalyst, such as a

metallocene catalyst, has a narrow molecular weight distribution. Such a catalyst system produces ethylene polymers in which a comonomer is randomly distributed within a molecular chain and uniformly distributed across the different molecular weight fractions. Metallocene-catalyzed polyolefins are described, for instance, in U.S. Patent Nos. 5,571,619 to McAlpin et al.; 5,322,728 to Davis et al.; 5,472,775 to Obijeski et al.; 5,272,236 to Lai et al.; and 6,090,325 to Wheat, et al. Examples of metallocene catalysts include bis(n-butylcyclopentadienyl)titanium dichloride, bis(n-butylcyclopentadienyl)zirconium dichloride, bis(cyclopentadienyl)scandium chloride, bis(indenyl)zirconium dichloride, bis(methylcyclopentadienyl)titanium dichloride, bis(methylcyclopentadienyl)zirconium dichloride, cobaltocene, cyclopentadienyltitanium trichloride, ferrocene, hafnocene dichloride, isopropyl(cyclopentadienyl,-1-flourenyl)zirconium dichloride, molybdocene dichloride, nickelocene, niobocene dichloride, ruthenocene, titanocene dichloride, zirconocene chloride hydride, zirconocene dichloride, and so forth. Polymers made using metallocene catalysts typically have a narrow molecular weight range. For instance, metallocene-catalyzed polymers may have polydispersity numbers (M_w/M_n) of below 4, controlled short chain branching distribution, and controlled isotacticity.

B. Nanofiller

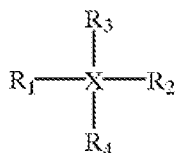
The nanofiller employed in the polymer composition typically has an average cross-sectional dimension (e.g., thickness or diameter) of from about 0.2 to about 100 nanometers, in some embodiments from about 0.5 to about 50 nanometers, and in some embodiments, from about 1 to about 20 nanometers. For example, the nanofiller may have a flake-like morphology in that it possesses a relatively flat or platelet shape. The platelets may, for example, have an average thickness within the ranges noted above. The "aspect ratio" of the platelets (i.e., the average length of the platelets divided by the average thickness) may also be relatively large, such as from about 20 to about 1000, in some embodiments from about 50 to about 80, in some embodiments, from about 100 to about 400. The average length (e.g., diameter) may, for instance, range from about 20 nanometers to about 10 micrometers, in some embodiments from about 100 nanometers to about 5 micrometers, and in some embodiments, from about 200 nanometers to about 4 micrometers.

The material used for the nanofiller is not particular critical. In certain embodiments, for example, the nanofiller may be carbon black, carbon nanotubes, carbon nanofibers, nanoclays, metal particles, silica, alumina, etc. Nanoclays are particularly suitable for use in the present invention. The term "nanoclay" generally refers to nanoparticles of a clay material (a naturally occurring mineral, an organically modified mineral, or a synthetic nanomaterial). The clay material may be formed from a phyllosilicate, such as a smectite clay mineral (e.g., bentonite, kaolinite, or montmorillonite, as well as salts thereof, such as sodium montmorillonite, magnesium montmorillonite, calcium montmorillonite,

etc.); nontronite; beidellite; volkonskoite; hectorite; saponite; sauconite; sobockite; stevensite; svinfordite; vermiculite; etc. Other useful nanoclays include micaceous minerals (e.g., illite) and mixed illite/smectite minerals, such as rectorite, tarosovite, ledikite and admixtures of illites with the clay minerals named above. Particularly suitable are montmorillonite (2:1 layered smectite clay structure), bentonite (aluminium phyllosilicate formed primarily of montmorillonite), kaolinite (1:1 aluminosilicate having a platy structure and empirical formula of $\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4$), halloysite (1:1 aluminosilicate having a tubular structure and empirical formula of $\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4$), etc.

If desired, the nanofiller may contain an organic surface treatment that enhances the hydrophobicity of the nanofiller. Without intending to be limited by theory, it is believed that the organic surface treatment can have a plastifying-like effect on the nanofiller, which can increase the compatibility between polyolefin and nanofiller and also reduce the degree of agglomeration of surface friction between the nanofiller phase and domains of the ethylene polymer when the composition is subjected to an elongational force. The surface treatment may also have a lubricating effect, which allows the macromolecular chains of the ethylene polymer to slip along the nanofiller surface without causing debonding. Each of these factors can help minimize the degree of noise that the film exhibits when subjected to physical deformation.

In one embodiment, the organic surface treatment may be formed from a quaternary onium (e.g., salt or ion), which may become intercalated via ion-exchange into the interlayer spaces between adjacent layered platelets. The quaternary onium ion may have the following structure:

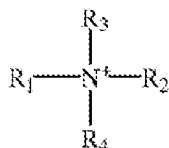


wherein

X is N, P, S, or O; and

R_1 , R_2 , R_3 and R_4 are independently hydrogen or organic moieties, such as linear or branched alkyl, aryl or aralkyl moieties having 1 to about 24 carbon atoms.

Particularly suitable quaternary ammonium ions are those having the structure below:



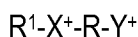
wherein

R_1 is a long chain alkyl moiety ranging from C_6 to C_{24} , straight or branched chain, including mixtures of long chain moieties, such as C_6 , C_8 , C_{10} , C_{12} , C_{14} , C_{16} , C_{18} , C_{20} , C_{22} and C_{24} , alone or in

any combination; and

R_2 , R_3 and R_4 are moieties, which may be the same or different, selected from the group consisting of H, alkyl, hydroxyalkyl, benzyl, substituted benzyl, e.g., straight or branched chain alkyl-substituted and halogen-substituted; ethoxylated or propoxylated alkyl; ethoxylated or propoxylated benzyl (e.g., 1-10 moles of ethoxylation or 1-10 moles of propoxylation).

Additional useful multi-charged spacing/coupling agents include for example, tetra-, tri-, and di-onium species such as tetra-ammonium, tri-ammonium, and di-ammonium (primary, secondary, tertiary, and quaternary), -phosphonium, -oxonium, or -sulfonium derivatives of aliphatic, aromatic or arylaliphatic amines, phosphines, esters, alcohols and sulfides. Illustrative of such materials are di-onium compounds of the formula:



where X^+ and Y^+ , are the same or different, and are ammonium, sulfonium, phosphonium, or oxonium radicals such as $-NH(CH_3)_2^+$, $-NH_2(CH_3)^+$, $-N(CH_3)_3^+$, $-N(CH_3)_2(CH_2CH_3)^+$, $-N(CH_3)(CH_2CH_3)_2^+$, $-S(CH_3)_2^+$, $-S(CH_3)_2^+$, $-P(CH_3)_3^+$, $-NH_3^+$, etc.;

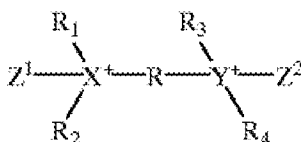
R is an organic spacing, backbone radical, straight or branched, such as those having from 2 to 24 carbon atoms, and in some embodiments from 3 to 10 carbon atoms, in a backbone organic spacing molecule covalently bonded at its ends to charged N^+ , P^+ , S^+ and/or O^+ cations;

R^1 can be hydrogen, or a linear or branched alkyl radical of 1 to 22 carbon atoms, linear or branched, and in some embodiments, 6 to 22 carbon atoms.

Illustrative of useful R groups are alkyls (e.g., methyl, ethyl, butyl, octyl, etc.); aryl (e.g., benzyl, phenylalkyl, etc.); alkenes (e.g., methylene, ethylene, octylene, nonylene, tert-butylene, neopentylene, isopropylene, sec-butylene, dodecylene, etc.); alkenylenes (e.g., 1-propenylenes, 1-butenylene, 1-pentylenes, 1-hexenylenes, 1-heptenylenes, 1-octenylenes, etc.); cycloalkenylenes (e.g., cyclohexenylenes, cyclopentylenes, etc.); hydroxyalkyl (e.g., hydroxymethyl, hydroxyethyl, hydroxyl-n-propyl, hydroxyisopropyl, hydroxyl-n-butyl, hydroxyl-iso-butyl, hydroxyl-tert-butyl, etc.), alkanoylalkylenes (e.g., butanoyl octadecylene, pentanoyl nonadecylene, octanoyl pentadecylene, ethanoyl undecylene, propanoyl hexadecylene, etc.); alkylaminoalkylenes (e.g., methylamino octadecylene, ethylamino pentadecylene, butylamino nonadecylene, etc.); dialkylaminoalkylene (e.g., dimethylamino octadecylene, methylethylamino nonadecylene, etc.); arylaminoalkylenes (e.g., phenylamino octadecylene, p-methylphenylamino nonadecylene, etc.); diarylaminoalkylenes (e.g., diphenylamino pentadecylene, p-nitrophenyl-p'-methylphenylamino octadecylene, etc.); alkylarylaminominoalkylenes (e.g., 2-phenyl-4-methylamino pentadecylene, etc.); alkylsulfinylenes, alkylsulfonylenes, alkylthio, arylthio, arylsulfinylenes, and arylsulfonylenes (e.g., butylthio octadecylene, neopentylthio pentadecylene, methylsulfinylnonadecylene, benzylsulfinyl

pentadecylene, phenylsulfinyl octadecylene, propylthiooctadecylene, octylthio pentadecylene, nonylsulfonyl nonadecylene, octylsulfonyl hexadecylene, methylthio nonadecylene, isopropylthio octadecylene, phenylsulfonyl pentadecylene, methylsulfonyl nonadecylene, nonylthio pentadecylene, phenylthio octadecylene, ethylthio nonadecylene, benzylthio undecylene, phenethylthio pentadecylene, sec-butylthio octadecylene, naphthylthio undecylene, etc.); alkoxyalkylalkylenes (e.g., methoxycarbonylene, ethoxycarbonylene, butoxycarbonylene, etc.); cycloalkylenes (e.g., cyclohexylene, cyclopentylene, cyclooctylene, cycloheptylene, etc.); alkoxyalkylenes (e.g., methoxymethylene, ethoxymethylene, butoxymethylene, propoxyethylene, pentoxybutylene, etc.); aryloxyalkylenes and aryloxyarylenes (e.g., phenoxyphenylene, phenoxyethylene, etc.); aryloryalkylenes (e.g., phenoxydecylene, phenoxyoctylene, etc.); arylalkylenes (e.g., benzylene, phenthylene, 8-phenyloctylene, 10-phenyldecylene, etc.); alkylarylenes (e.g., 3-decylphenylene, 4-octylphenylene, 4-nonylphenylene, etc.); and polypropylene glycol and polyethylene glycol substituents (e.g., ethylene, propylene, butylene, phenylene, benzylene, tolylene, p-styrylene, p-phenylmethylene, octylene, dodecylene, octadecylene, methoxyethylene, etc.), as well as combinations thereof. Such tetra-, tri-, and di-ammonium, -sulfonium, -phosphonium, -oxonium; ammonium/sulfonium; ammonium/phosphonium; ammonium/oxonium; phosphonium/oxonium; sulfonium/oxonium; and sulfonium/phosphonium radicals are well known in the art and can be derived from the corresponding amines, phosphines, alcohols or ethers, and sulfides.

Particularly suitable multi-charged spacing/coupling agent compounds are multi-onium ion compounds that include at least two primary, secondary, tertiary or quaternary ammonium, phosphonium, sulfonium, and/or oxonium ions having the following general formula:



wherein

R is an alkylene, aralkylene or substituted alkylene charged atom spacing moiety; and

Z₁, Z₂, R₁, R₂, R₃, and R₄ may be the same or different and selected from the group consisting of hydrogen, alkyl, aralkyl, benzyl, substituted benzyl (e.g., straight or branched chain alkyl-substituted and halogen-substituted); ethoxylated or propoxylated alkyl; ethoxylated or propoxylated benzyl (e.g., 1-10 moles of ethoxylation or 1-10 moles of propoxylation).

Particularly suitable organic cations may include, for instance, quaternary ammonium compounds, such as dimethyl bis[hydrogenated tallow] ammonium chloride (2M2HT), methyl benzyl bis[hydrogenated tallow] ammonium chloride (MB2HT), methyl tris[hydrogenated tallow alkyl] chloride (M3HT), etc. One example of a suitable nanofiller with such a surface treatment is Nanomer™ 1.44P,

which is a quaternary ammonium modified montmorillonite nanoclay and commercially available from Nanocor, Inc. Other suitable nanoclay additives include those available from Southern Clay Products, such as Cloisite™ 15A, Cloisite™ 30B, Cloisite™ 93A, and Cloisite™ Na⁺.

The onium ion may be introduced into (sorbed within) the interlayer spaces of the nanofiller in a number of ways. In one method, for example, the nanofiller is slurried in water, and the onium ion compound is dissolved therein. If necessary, the onium ion compound can be dissolved first in an organic solvent (e.g., propanol). If desired, the nanofiller may also be intercalated with an oligomer and/or polymer intercalant as is known in the art. For example, an olefin polymer or oligomer (e.g., ethylene polymer) intercalant may be employed. To intercalate an onium ion and an olefin intercalant between adjacent phyllosilicate platelets and optionally separate (exfoliate) the layered material into individual platelets, for example, the nanofiller may be first contacted with the onium ion and simultaneously or thereafter contacted with the melted oligomer/polymer intercalant to the onium ion-intercalated layered material. This may be accomplished, for instance, by directly compounding the materials in an extruder. Alternatively, the oligomer/polymer can be intercalated by an emulsion process by vigorously mixing with an emulsifier. If desired, a coupling agent (e.g., silane coupling agent) may also be employed to help bond the intercalant with the clay material. For example, the nanofiller may be initially treated with a coupling agent followed by ion-exchange of onium ions between the nanofiller, prior to or simultaneously with intercalation of the oligomer(s) or polymer(s). It should be understood that the oligomer or polymer intercalant(s) can also be intercalated between and complexed to the internal platelet faces by other well-known mechanisms, such as by dipole/dipole bonding (direct intercalation of the oligomer or polymer) as described in U.S. Patent Nos. 5,880,197 and 5,877,248, as well as by acidification with substitution with hydrogen (ion-exchanging the interlayer cations with hydrogen by use of an acid or ion-exchange resin) as described in U.S. Patent Nos. 5,102,948 and 5,853,886.

Nanofillers typically constitute from about 0.5 wt.% to about 20 wt.%, in some embodiments from about 1 wt.% to about 15 wt.%, and in some embodiments, from about 2 wt.% to about 10 wt.% of the polymer composition.

C. Compatibilizer

Although not necessarily required, a compatibilizer may also be employed in the polymer composition, such as in an amount of from about 0.1 wt.% to about 10 wt.%, in some embodiments from about 0.2 wt.% to about 8 wt.%, and in some embodiments, from about 0.5 wt.% to about 5 wt.% of the polymer composition. The compatibilizer may be a polyolefin containing an olefin component and a polar component. The olefin component is non-polar and thus generally has an affinity for the ethylene polymer. The olefin component may generally be formed from any linear or branched α -olefin

monomer, oligomer, or polymer (including copolymers) derived from an α -olefin monomer. In one particular embodiment, for example, the compatibilizer includes at least one linear or branched α -olefin monomer, such as those having from 2 to 20 carbon atoms and preferably from 2 to 8 carbon atoms. Specific examples include ethylene, propylene, 1-butene; 3-methyl-1-butene; 3,3-dimethyl-1-butene; 1-pentene; 1-pentene with one or more methyl, ethyl or propyl substituents; 1-hexene with one or more methyl, ethyl or propyl substituents; 1-heptene with one or more methyl, ethyl or propyl substituents; 1-octene with one or more methyl, ethyl or propyl substituents; 1-nonene with one or more methyl, ethyl or propyl substituents; ethyl, methyl or dimethyl-substituted 1-decene; 1-dodecene; and styrene. Particularly desired α -olefin co-monomers are ethylene and propylene.

The polyolefin compatibilizer is also functionalized with a polar component, which can be grafted onto the polymer, incorporated as a monomeric constituent of the polymer (e.g., block or random copolymers), etc. When grafted onto a polymer backbone, particularly suitable polar groups are maleic anhydride, maleic acid, acrylic acid, methacrylic acid, fumaric acid, maleimide, maleic acid hydrazide, a reaction product of maleic anhydride and diamine, methyl nadic anhydride, dichloromaleic anhydride, maleic acid amide, etc. Maleic anhydride modified polyolefins are particularly suitable for use in the present invention. Such modified polyolefins are typically formed by grafting maleic anhydride onto a polymeric backbone material. Such maleated polyolefins are available from E. I. du Pont de Nemours and Company under the designation FUSABOND®, such as the P Series (chemically modified polypropylene), E Series (chemically modified polyethylene), C Series (chemically modified ethylene vinyl acetate), A Series (chemically modified ethylene acrylate copolymers or terpolymers), M Series (chemically modified polyethylene), or N Series (chemically modified ethylene-propylene, ethylene-propylene diene monomer ("EPDM") or ethylene-octene). Alternatively, modifier polyolefins are also available from Chemtura Corp. under the designation POLYBOND® (e.g., acrylic acid-modified polypropylene) and Eastman Chemical Company under the designation Eastman G series.

As noted above, the polar component may also be incorporated into the polyolefin compatibilizer as a monomer. For example, a (meth)acrylic monomeric component may be employed in certain embodiments. As used herein, the term "(meth)acrylic" includes acrylic and methacrylic monomers, as well as salts or esters thereof, such as acrylate and methacrylate monomers.

Examples of such (meth)acrylic monomers may include methyl acrylate, ethyl acrylate, n-propyl acrylate, i-propyl acrylate, n-butyl acrylate, s-butyl acrylate, i-butyl acrylate, t-butyl acrylate, n-amyl acrylate, i-amyl acrylate, isobornyl acrylate, n-hexyl acrylate, 2-ethylbutyl acrylate, 2-ethylhexyl acrylate, n-octyl acrylate, n-decyl acrylate, methylcyclohexyl acrylate, cyclopentyl acrylate, cyclohexyl acrylate, methyl methacrylate, ethyl methacrylate, 2-hydroxyethyl methacrylate, n-propyl methacrylate,

n-butyl methacrylate, i-propyl methacrylate, i-butyl methacrylate, n-amyl methacrylate, n-hexyl methacrylate, i-amyl methacrylate, s-butyl-methacrylate, t-butyl methacrylate, 2-ethylbutyl methacrylate, methylcyclohexyl methacrylate, cinnamyl methacrylate, crotyl methacrylate, cyclohexyl methacrylate, cyclopentyl methacrylate, 2-ethoxyethyl methacrylate, isobornyl methacrylate, etc., as well as combinations thereof. Other types of suitable polar monomers include ester monomers, amide monomers, etc.

D. Other Components

In addition to the components noted above, other additives may also be incorporated into the polymer composition, such as slip additives, melt stabilizers, processing stabilizers, heat stabilizers, light stabilizers, antioxidants, heat aging stabilizers, whitening agents, bonding agents, fillers, etc. Further, hindered phenols are commonly used as an antioxidant in the production of films. Some suitable hindered phenols include those available from Ciba Specialty Chemicals under the trade name "Irganox®", such as Irganox® 1076, 1010, or E 201. Moreover, bonding agents may also be added to the film to facilitate bonding of the film to additional materials (e.g., nonwoven webs). Examples of such bonding agents include hydrogenated hydrocarbon resins. Other suitable bonding agents are described in U.S. Patent Nos. 4,789,699 to Kieffer et al. and 5,695,868 to McCormack.

II. Film Construction

The film of the present invention may be mono- or multi-layered. Multilayer films may be prepared by co-extrusion of the layers, extrusion coating, or by any conventional layering process. Such multilayer films normally contain at least one base layer and at least one skin layer, but may contain any number of layers desired. For example, the multilayer film may be formed from a base layer and one or more skin layers. In one embodiment, for example, it may be desirable to employ two skin layers that sandwich a base layer. Regardless of the particular construction, the skin and/or base layers may be formed from the polymer composition of the present invention. In one embodiment, for example, the base layer is formed from the polymer composition of the present invention and the skin layer(s) are formed from the polymer composition or from an additional polymer material. Likewise, in other possible embodiments, one or more of the skin layers are formed from the polymer composition of the present invention and the base layer is formed from an additional polymer material. When employed, the additional material may include any type of polymer, such as polyolefins (e.g., polyethylene, polypropylene, etc.), polyesters, polyamides, styrenic copolymers, polyurethanes, polyvinyl acetate, polyvinyl alcohol, etc.

When multiple layers are employed, the base layer typically constitutes a substantial portion of the weight of the film, such as from about 50 wt.% to about 99 wt.%, in some embodiments from about 55 wt.% to about 90 wt.%, and in some embodiments, from about 60 wt.% to about 85 wt.% of the film.

The skin layer(s) may likewise constitute from about 1 wt.% to about 50 wt.%, in some embodiments from about 10 wt.% to about 45 wt.%, and in some embodiments, from about 15 wt.% to about 40 wt.% of the film. Each skin layer may also have a thickness of from about 0.1 to about 10 micrometers, in some embodiments from about 0.5 to about 5 micrometers, and in some
5 embodiments, from about 1 to about 2.5 micrometers. Likewise, the base layer may have a thickness of from about from about 1 to about 40 micrometers, in some embodiments from about 2 to about 25 micrometers, and in some embodiments, from about 5 to about 20 micrometers. As noted above, the total thickness of the film is typically about 50 micrometers or less, in some embodiments from about 1 to about 40 micrometers, in some embodiments from about 5 to about 35 micrometers, and in some
10 embodiments, from about 10 to about 30 micrometers.

Any of a variety of techniques may generally be employed to form the film of the present invention. In certain embodiments, for example, the components of the film (e.g., ethylene polymer, nanofiller, compatibilizer, etc.) may be individually supplied to a film forming system and blended together as the film is being formed. In such cases, the nanofiller may be in the form of a powder
15 containing particles, such as described above. Alternatively, however, it is may be desirable to pre-blend the ethylene polymer, nanofiller, and/or compatibilizer to form a masterbatch, which is then subsequently supplied to the film forming system. Without intending to be limited by theory, it is believed such a multi-step process can allow the nanofiller to be more uniformly oriented, thereby even further enhancing ductility. When supplied, the nanofiller may itself be in the form of a masterbatch,
20 which may contain nanofiller particles blended with a polymer (e.g., ethylene polymer), or in the form of a powder containing particles.

To form a masterbatch, for example, the components may initially be supplied to twin screw extruder that includes co-rotating screws rotatably mounted and received within a barrel (e.g., cylindrical barrel), which may be heated. The components are moved downstream from a feed end to
25 a discharge end by forces exerted by rotation of the screws. The ratio of the length to outer diameter ("L/D") of the screws may be selected to achieve an optimum balance between throughput and blend uniformity. For example, too large of an L/D value may increase the retention time to such an extent that the nanofiller degrades beyond the desired level or reduce the limit of torque along the twin screw, which may decrease the exfoliation and dispersion of nanofiller. On other hand, too low of an L/D
30 value may not result in the desired degree of blending due to the short residence time. Thus, the L/D value is typically from about 25 to about 60, in some embodiments from about 35 to about 55, and in some embodiments from about 40 to about 50. The speed of the screws may also be selected to achieve the desired residence time, shear rate, melt processing temperature, etc. Generally, an increase in product temperature is observed with increasing screw speed due to the additional

mechanical energy input into the system. The frictional energy results from the shear exerted by the turning screw on the materials within the extruder and results in the fracturing of large molecules. This results in lowering the apparent viscosity and increasing the melt flow rate of the finished material. For example, the screw speed may range from about 50 to about 400 revolutions per minute ("rpm"), in some embodiments from about 100 to about 300 rpm, and in some embodiments, from about 120 to about 280 rpm. As a result, melt processing may occur at a temperature of from about 100°C to about 500°C, in some embodiments, from about 150°C to about 350°C, and in some embodiments, from about 150°C to about 300°C. Typically, the apparent shear rate during melt processing may range from about 300 seconds⁻¹ to about 10,000 seconds⁻¹, in some embodiments from about 500 seconds⁻¹ to about 5000 seconds⁻¹, and in some embodiments, from about 800 seconds⁻¹ to about 1200 seconds⁻¹. The apparent shear rate is equal to $4Q/R^3$, where Q is the volumetric flow rate ("m³/s") of the polymer melt and R is the radius ("m") of the capillary (e.g., extruder die) through which the melted polymer flows. Of course, other variables, such as the residence time during melt processing, which is inversely proportional to throughput rate, may also be controlled to achieve the desired blending.

Once formed, the pre-blended masterbatch may be supplied to a film-forming system. Any known technique may be used to form a film from the compounded material, including blowing, casting, flat die extruding, etc. In one particular embodiment, the film may be formed by a blown process in which a gas (e.g., air) is used to expand a bubble of the extruded polymer blend through an annular die. The bubble is then collapsed and collected in flat film form. Processes for producing blown films are described, for instance, in U.S. Patent Nos. 3,354,506 to Raley; 3,650,649 to Schippers; and 3,801,429 to Schrenk et al., as well as U.S. Patent Application Publication Nos. 2005/0245162 to McCormack, et al. and 2003/0068951 to Boggs, et al. In yet another embodiment, however, the film is formed using a casting technique.

Referring to Fig. 1, for instance, one embodiment of a method for forming a cast film is shown. In this embodiment, the pre-blended masterbatch is supplied to an extruder 80 for melt processing. To help achieve good alignment and orientation of the nanofiller, it is typically desired to use a single screw extruder during film formation. Such single screw extruders are typically divided into three sections along the length of the screw. The first section is a feed section where the solid material is introduced to the screw. The second section is a melting section where a majority of the melting of the solid occurs. Within this section, the screw generally possesses a tapered diameter to enhance melting of the polymer. The third section is the mixing section, which delivers the molten material in a constant amount for extrusion. The L/D ratio for the screw is typically from about 5 to about 50, in some embodiments from about 10 to about 40, and in some embodiments from about 15 to about 35. Such L/D ratios may be readily achieved in a single screw extruder by using mixing section(s) that

constitute only a small portion of the length of the screw. The screw speed may likewise range from about 5 to about 150 rpm, in some embodiments from about 10 to about 100 rpm, and in some embodiments, from about 20 to about 80 rpm. As a result, melt processing may occur at a temperature of from about 100°C to about 500°C, in some embodiments, from about 150°C to about 350°C, and in some embodiments, from about 150°C to about 300°C.

Once formed, the extruded material may be immediately chilled and cut into pellet form. In the embodiment of Fig. 1, the extruded material is cast onto a casting roll 90 to form a single-layered precursor film 10a. If a multilayered film is to be produced, the multiple layers are co-extruded together onto the casting roll 90. The casting roll 90 may optionally be provided with embossing elements to impart a pattern to the film. Typically, the casting roll 90 is kept at temperature sufficient to solidify and quench the sheet 10a as it is formed, such as from about 20 to 60°C. If desired, a vacuum box may be positioned adjacent to the casting roll 90 to help keep the precursor film 10a close to the surface of the roll 90. Additionally, air knives or electrostatic pinners may help force the precursor film 10a against the surface of the casting roll 90 as it moves around a spinning roll. An air knife is a device known in the art that focuses a stream of air at a very high flow rate to pin the edges of the film.

Once cast, the film 10a may then be optionally oriented in one or more directions to further improve film uniformity and reduce thickness. In the case of sequential orientation, the “softened” film is drawn by rolls rotating at different speeds of rotation such that the sheet is stretched to the desired draw ratio in the longitudinal direction (machine direction). If desired, the uniaxially oriented film may also be oriented in the cross-machine direction to form a “biaxially oriented” film. For example, the film may be clamped at its lateral edges by chain clips and conveyed into a tenter oven. In the tenter oven, the film may be reheated and drawn in the cross-machine direction to the desired draw ratio by chain clips diverged in their forward travel.

Referring again to Fig. 1, for instance, one method for forming a uniaxially oriented film is shown. As illustrated, the precursor film 10a is directed to a film-orientation unit 100 or machine direction orienter (“MDO”), such as commercially available from Marshall and Willams, Co. of Providence, Rhode Island. The MDO has a plurality of stretching rolls (such as from 5 to 8) which progressively stretch and thin the film in the machine direction, which is the direction of travel of the film through the process as shown in Fig. 1. While the MDO 100 is illustrated with eight rolls, it should be understood that the number of rolls may be higher or lower, depending on the level of stretch that is desired and the degrees of stretching between each roll. The film may be stretched in either single or multiple discrete stretching operations. It should be noted that some of the rolls in an MDO apparatus may not be operating at progressively higher speeds. If desired, some of the rolls of the MDO 100 may act as preheat rolls. If present, these first few rolls heat the film 10a above room temperature.

The progressively faster speeds of adjacent rolls in the MDO act to stretch the film 10a. The rate at which the stretch rolls rotate determines the amount of stretch in the film and final film weight. The resulting film 10b may then be wound and stored on a take-up roll 60. While not shown here, various additional potential processing and/or finishing steps known in the art, such as embossing, slitting, treating, aperturing, printing graphics, or lamination of the film with other layers (e.g., nonwoven web materials), may be performed without departing from the spirit and scope of the invention.

In one embodiment, for example, the film may be embossed using any technique known in the art to form a pattern of embossed regions on one or more surfaces of the film. Suitable techniques include, for instance, the use of raised elements to impart the desired embossing pattern. Thermal and/or ultrasonic bonding techniques may be employed. For instance, a suitable process may involve thermal bonding wherein a layer is passed through two rolls (e.g., steel, rubber, etc.) in which one is engraved with an embossing pattern and the other is flat. One or both of the rolls may be heated. Regardless of the technique employed, the resulting embossed regions may have a relatively small depth such that the film is considered "micro-embossed." Micro-embossed films may, for instance, have embossed regions with a depth of from about 5 micrometers or less, and in some embodiments, from about 1 to about 4 micrometers. Of course, the embossed regions may also have a relatively large depth such that the film is considered "deep embossed." Such films may, for instance, have embossed regions with a depth of greater than about 5 micrometers, and in some embodiments, from about 5 to about 20 micrometers.

III. Package Configuration

The particular manner in which the wrapper material is configured to enclose an absorbent article and form a package is not particularly critical to the present invention. The absorbent article may be completely enclosed by the wrapper material, or only partially enclosed if so desired. Various suitable wrap configurations are disclosed, for instance, in U.S. Patent Nos. 6,716,203 to Sorebo, et al. and 6,380,445 to Moder, et al., as well as U.S. Patent Application Publication No. 2003/0116462 to Sorebo, et al.

Referring to Figs. 3-5, for instance, one particular embodiment of a package 110 is shown that contains a wrapper material 114 that defines an interior cavity or pouch 115 within which an absorbent article 112 is removably positioned. The wrapper material 114 may be formed from the film of the present invention and define an exterior surface 138 of the package. In this embodiment, the wrapper material 114 may be folded and bonded into the pouch configuration. For example, the wrapper material 114 may be an elongated rectangular piece having a first end 126, an opposite second end 128, and generally parallel longitudinal sides 133 and 135 extending between the ends 126 and 128. As shown in Fig. 4, the wrapper material 114 may be folded around the absorbent article 112 (or

multiple absorbent articles) such that a pouch 115 is formed around the article. The wrapper material 114 is first folded at a first fold axis 130 such that the first end 126 is folded towards but spaced from the second end 128, as can be seen particularly in Figs 4-5. The distance between the first end 126 and second end 128 may vary depending on the desired length of a resulting flap 120, as described below. The aligned longitudinal sides of the wrapper material 114 define sides 134 and 136 of the pouch 115. The second end 128 of the wrapper material is then folded at a second fold axis 132 so as to extend back over the first end 126 and thus defines the flap 120 that closes off the pouch 115, as particularly seen in Fig. 3. The flap 120 has longitudinal sides 122 and 124 that align with the material sides 133 and 135 and pouch sides 134 and 136. The sides of the package 110 are then bonded in a conventional manner, for example with a heat/pressure embossing roll. The flap sides 122 and 124 are bonded to the material sides 133 and 135 and pouch sides 134 and 136 in a single pass operation. It may be the case that the first end 126 of the wrapper material 114 extends essentially to the second fold axis 132 and, thus, the flap sides 122 and 124 would be bonded along their entire length to pouch sides 134 and 136.

As shown in Fig. 4, the absorbent article 112 may be placed on an inner major surface 140 of the wrapper material 114, and then the material may be folded at the first axis 130 and then at the second axis 132 to define the pouch 115 and flap 120. The sides of the pouch and flap may then be bonded or sealed together. These seals or bonds can be formed by heat, heat and pressure, pressure, adhesive, ultrasonic bonding, or other types of bonding techniques known to those skilled in the art. The seals can be made to be a "permanent" seal, which means that the wrapper material adjacent to the seal will tear or break before the sealed layers separate. Alternately, the seals may be "frangible" seals, which means that the sealed layers will separate or pull apart. In a desirable embodiment of the package 110, the seal between the flap sides 122 and 124 and the pouch sides 134 and 136 is a frangible seal formed by simultaneously sealing the pouch sides and flap sides in a single sealing operation with a heated/pressure embossing roll. To open the pouch, a consumer may simply insert a finger between the flap 120 and front surface of the pouch 115 and then pull the flap away from the pouch to break the sealed sides of the flap. In some cases, the noise that is typically associated with the opening procedure can also be reduced in accordance with the present invention.

Regardless of the particular configuration, the film and wrapper material can enclose a single absorbent or even multiple absorbent articles if so desired. An "absorbent article" generally refers to any article capable of absorbing water or other fluids. Examples of some absorbent articles include, but are not limited to, personal care absorbent articles, such as diapers, training pants, absorbent underpants, incontinence articles, feminine care articles (e.g., sanitary napkins, pantliners, etc.), swim wear, baby wipes, and so forth; medical absorbent articles, such as garments, fenestration materials,

underpads, bedpads, bandages, absorbent drapes, and medical wipes; food service wipers; clothing articles; and so forth. Several examples of such absorbent articles are described in U.S. Patent Nos. 5,649,916 to DiPalma, et al.; 6,110,158 to Kielpikowski; 6,663,611 to Blaney, et al. Still other suitable articles are described in U.S. Patent Application Publication No. 2004/0060112 A1 to Fell et al., as well as U.S. Patent Nos. 4,886,512 to Damico et al.; 5,558,659 to Sherrod et al.; 6,888,044 to Fell et al.; and 6,511,465 to Freiburger et al. Materials and processes suitable for forming such absorbent articles are well known to those skilled in the art. Typically, absorbent articles include a generally liquid-impermeable layer (e.g., backsheet), a liquid-permeable layer (e.g., topsheet, surge layer, etc.), and an absorbent core.

Various embodiments of an absorbent article that may be formed according to the present invention will now be described in more detail. Referring to Fig. 2, for instance, one embodiment of a one particular embodiment of a feminine care absorbent article 20 that can be packaged in accordance with the present invention will now be described in more detail. As shown, the feminine care absorbent article 20 (e.g., feminine care pad or napkin) includes a topsheet 26, a backsheet 28, and an absorbent core 30 positioned between the topsheet 26 and the backsheet 28. In certain embodiments, the topsheet 26 and/or backsheet 28 may contain the film of the present invention. The topsheet 26 defines a body-facing surface 27 of the absorbent article 20. The absorbent core 30 is positioned inwardly from the outer periphery of the absorbent article 20 and includes a body-facing side positioned adjacent the topsheet 26 and a garment-facing surface positioned adjacent the backsheet 28. Typically, the topsheet 26 and the backsheet 28 are joined by adhesive bonding, ultrasonic bonding, or any other suitable joining method known in the art, the sealed edges defining an overall sealed peripheral edge 99 of the article 20. The article 20 may take on various geometries but will generally have opposite lateral sides and longitudinal ends.

The topsheet 26 is generally designed to contact the body of the user and is liquid-permeable. The liquid permeable topsheet 26 has an outwardly facing surface that may contact the body of the wearer and receive aqueous fluids from the body. The topsheet 26 is provided for comfort and conformability and functions to direct bodily exudates away from the body, through the topsheet 26 and toward the absorbent core 30. The topsheet 26 retains little or no liquid in its structure so that it provides a relatively comfortable and non-irritating surface next to the tissues within the vestibule of a female wearer. In certain embodiments, the topsheet 26 may contain a polyolefin (e.g., polyethylene) film. The topsheet 26 may also contain a woven or nonwoven material that is easily penetrated by bodily exudates contacting the surface of the backsheet. Examples of suitable materials include rayon, bonded carded webs of polyester, polypropylene, polyethylene, nylon, or other heat-bondable fibers, polyolefins, such as copolymers of polypropylene and polyethylene, linear low-density

polyethylene, and aliphatic esters such as polylactic acid. A specific example of a suitable topsheet material is a bonded carded web made of polypropylene and polyethylene such as that used as topsheet stock for KOTEX® pantliners. The topsheet 26 may also contain a plurality of apertures (not shown) formed therethrough to permit body fluid to pass more readily into the absorbent core 30. The apertures may be randomly or uniformly arranged throughout the topsheet 26, or they may be located only in the narrow longitudinal band or strip arranged along the longitudinal axis of the absorbent article 20. The apertures permit rapid penetration of body fluid down into the absorbent core 30. The size, shape, diameter and number of apertures may be varied to suit one's particular needs.

The backsheet 28 is generally liquid-impermeable and designed to face the inner surface, i.e., the crotch portion of an undergarment (not shown). The backsheet 28 may permit a passage of air or vapor out of the absorbent article 20, while still blocking the passage of liquids. Any liquid-impermeable material may generally be utilized to form the backsheet 28. For example, the backsheet 28 may contain a conventional polyolefin film, such as a polyethylene film such as that used in KOTEX® pantliners.

As indicated above, an absorbent core 30 is positioned between the topsheet 26 and the backsheet 28 that provides capacity to absorb and retain bodily exudates. The absorbent core 30 may be formed from a variety of different materials and contain any number of desired layers. For example, the core 30 typically includes one or more layers of an absorbent web material of cellulosic fibers (e.g., wood pulp fibers), other natural fibers, synthetic fibers, woven or nonwoven sheets, scrim netting or other stabilizing structures, superabsorbent material, binder materials, surfactants, selected hydrophobic and hydrophilic materials, pigments, lotions, odor control agents or the like, as well as combinations thereof. In a particular embodiment, the absorbent web material includes a matrix of cellulosic fluff, and may also include superabsorbent material. The cellulosic fluff may comprise a blend of wood pulp fluff. One preferred type of fluff is identified with the trade designation NB 416, available from Weyerhaeuser Corp., and is a bleached, highly absorbent wood pulp containing primarily soft wood fibers. The absorbent materials may be formed into a web structure by employing various conventional methods and techniques. For example, the absorbent web may be formed with a dry-forming technique, an air forming technique, a wet-forming technique, a foam-forming technique, or the like, as well as combinations thereof. A coform nonwoven material may also be employed. Methods and apparatus for carrying out such techniques are well known in the art.

The topsheet 26 may be maintained in secured relation with the absorbent core 30 by bonding all or a portion of the adjacent surfaces to one another. A variety of bonding mechanisms known to one of skill in the art may be utilized to achieve any such secured relation. Examples of such mechanisms include, but are not limited to, the application of adhesives in a variety of patterns

between the two adjoining surfaces, entangling at least portions of the adjacent surface of the absorbent with portions of the adjacent surface of the cover, or fusing at least portions of the adjacent surface of the cover to portions of the adjacent surface of the absorbent. The topsheet 26 typically extends over the upper, bodyside surface of the absorbent core 30, but can alternatively extend
5 around the article to partially or entirely, surround or enclose the absorbent core. Alternatively, the topsheet 26 and the backsheet 28 can have peripheral margins that extend outwardly beyond the terminal, peripheral edges of the absorbent core 30, and the extending margins can be joined together to partially or entirely, surround or enclose the absorbent core.

Although not required, the absorbent article 20 may also contain other additional layers as is
10 known in the art. In Fig. 2, for example, a liquid-permeable intake layer 32 is positioned vertically between the topsheet 26 and the absorbent core 30. The intake layer 32 may be made of a material that is capable of rapidly transferring, in the z-direction, body fluid that is delivered to the topsheet 26. The intake layer 32 may generally have any shape and/or size desired. In one embodiment, the intake layer 32 has a rectangular shape, with a length equal to or less than the overall length of the absorbent
15 article 20, and a width less than the width of the absorbent article 20. For example, a length of between about 150 mm to about 300 mm and a width of between about 10 mm to about 60 mm may be utilized. Any of a variety of different materials are capable of being used for the intake layer 32 to accomplish the above-mentioned functions. The material may be synthetic, cellulosic, or a combination of synthetic and cellulosic materials. For example, airlaid cellulosic tissues may be
20 suitable for use in the intake layer 32. The airlaid cellulosic tissue may have a basis weight ranging from about 10 grams per square meter (gsm) to about 300 gsm, and in some embodiments, between about 40 gsm to about 150 gsm. The airlaid tissue may be formed from hardwood and/or softwood fibers. The airlaid tissue has a fine pore structure and provides an excellent wicking capacity, especially for menses.

25 The absorbent article 20 may also contain a transfer delay layer (not shown) positioned between the intake layer 32 and the absorbent core 30. The transfer delay layer may contain a material that is substantially hydrophobic, such as a nonwoven web composed of polypropylene, polyethylene, polyester, etc. One example of a material suitable for the transfer delay layer is a spunbond web composed of polypropylene, multi-lobal fibers. Further examples of suitable transfer
30 delay layer materials include spunbond webs composed of polypropylene fibers, which may be round, tri-lobal or poly-lobal in cross-sectional shape and which may be hollow or solid in structure. Typically the webs are bonded, such as by thermal bonding, over about 3% to about 30% of the web area. Other examples of suitable materials that may be used for the transfer delay layer 36 are described in U.S. Patent Nos. 4,798,603 to Meyer, et al. and 5,248,309 to Serbiak, et al. To adjust performance,

the transfer delay layer may also be treated with a selected amount of surfactant to increase its initial wettability. The transfer delay layer typically has a basis weight less than that of the other absorbent members. For example, the basis weight of the transfer delay layer 36 is typically less than about 250 grams per square meter (gsm), and in some embodiments, between about 40 gsm to about 200 gsm.

5 The absorbent article 20 may also include laterally extending wing portions 42 that may be integrally connected to side regions along the intermediate portion of the article. The wing portions 42 may be a separately provided material that is subsequently attached or otherwise operatively joined to the intermediate portion of the article. In other configurations, the wing portions 42 may be unitarily formed with one or more components of the article. As representatively shown in Fig. 1, for example, 10 either or both wing portions 42 may be formed from a corresponding, operative extension of the material employed to form the topsheet 26. Alternatively, either or both wing portions 42 may be formed from a corresponding, operative extension of the material employed to form the backsheet 28.

Although various configurations of a feminine care article have been described above, it should be understood that other configurations are also included within the scope of the present 15 invention. In addition, the present invention is by no means limited to feminine care articles. In fact, any other absorbent article may be formed in accordance with the present invention, including, but not limited to, other personal care absorbent articles, such as training pants, absorbent underpants, adult incontinence products, diapers, swim wear, baby wipes, and so forth; medical absorbent articles, such as garments, fenestration materials, underpads, bandages, absorbent drapes, and medical wipes; food 20 service wipers; clothing articles; and so forth. The film may also be used in applications other than absorbent articles.

The present invention may be better understood with reference to the following examples.

Test Methods

Mechanical Properties:

25 The strip tensile strength values were determined in substantial accordance with ASTM Standard D638-99. A constant-rate-of-extension type of tensile tester was employed. The tensile testing system was an MTS Synergy 200 tensile frame. The tensile tester was equipped with TESTWORKS 4.08B software from MTS Corporation to support the testing. The load cell was 100 Newtons. The film samples were initially cut into dog-bone shapes with a center width of 3.0 mm 30 before testing. The samples were held between grips having a front and back face measuring 25.4 millimeters x 76 millimeters. The grip faces were rubberized, and the longer dimension of the grip was perpendicular to the direction of pull. The grip pressure was pneumatically maintained at a pressure of 40 pounds per square inch. The tensile test was run using a gauge length of 18.0 millimeters and a break sensitivity of 40%. Five samples were tested by applying the test load along the machine-

direction and five samples were tested by applying the test load along the cross direction. During the test, samples were stretched at a crosshead speed of about 127 millimeters per minute until breakage occurred. The modulus, peak stress, peak elongation (percent strain at break), and energy per volume at break were measured in the machine direction ("MD") and cross-machine directions ("CD").

5 X-Ray Diffraction:

Several film samples were analyzed using X-ray diffraction on a Rigaku D-max Rapid instrument, equipped with a 2-D position sensitive detector. The 2-D patterns were corrected for background, absorption and air scattering and converted to 1-D curves by using azimuthal integration. The WAXS detector covers interatomic distances in the range from 2.8 Å to 59 Å (0.28 – 5.9 nm; 2° is the diffraction angle).

10 Transmission Electron Microscopy:

Transmission electron microscopy ("TEM") was used to obtain images of film samples in accordance with PSI Method 902 Revision 2. Prior to testing, a portion of each sample was cut away with a razor blade. The samples were then sputter coated in accordance with PSI Method ID 14203 Revision 1 with a layer of gold to increase the chances of finding the sample while performing TEM. The samples were then molded in EpoFix slow cure epoxy. The samples were cross-sectionally cryomicrotomed in accordance with PSI Method ID 904 Revision 3 to produce sections of about 70 nm thickness. The samples were cut parallel to the extruded direction of each sample in order to provide images perpendicular to the extruded direction of the samples.

20 Noise Level:

Noise levels of film and package samples may be tested in an apparatus comprised of a test chamber, a control chamber, and a sound level meter. The purpose of the apparatus is to manipulate an article in a controlled noise environment, and to accurately quantify the noise produced by the movement of the sample. In general terms, a sample is physically deformed within the test apparatus to generate a noise level. As used herein, the "noise level" refers to the equivalent continuous noise level (referred to as "L_{EQ}" or "L_{AT}"), which is the time average sound level (expressed in units of dB) as determined according to the following equation:

$$L_{eq} = 10 \log \left[\frac{1}{t_2 - t_1} \int_{t_1}^{t_2} \frac{p_A^2}{p_0^2} dt \right]$$

p_0 is a reference pressure level (typically 20 μPa);

p_A is the acquired sound pressure;

t is time;

t_1 is the start time for measurement; and

t_2 is the end time for measurement.

This value is also described in IEC 61672-1 (2013).

The testing apparatus is illustrated in Figs. 11-13. The testing apparatus 200 includes a test chamber 201 and a control chamber 202. The test chamber 201 includes a door 203, a top wall 204, a bottom wall 205, two side walls 206 and 207, and a rear wall 208. The door and each wall are constructed of 0.25-inch (0.635 cm) thick 6061 grade anodized aluminum. The door 203 and rear wall 208 are each 36 inches (91.4 cm) in height and 24 inches (61.0 cm) in width. The test chamber side walls 206 and 207 are each 36 inches (91.4 cm) high and 18 inches (45.7 cm) wide. The test chamber top and bottom panels are each 24 inches wide (61.0 cm) and 18 inches (45.7 cm) long. The interior surface of the door 203 and each wall 204-208 has applied thereto two-inch thick polyurethane sound-dampening foam 209, available from Illbruck Inc. under the brand name SONEX and stock number SOC-2. As shown, a sound level meter support 216 extends perpendicularly outward from side wall 206 just below a microphone orifice 217. The microphone orifice 217 is positioned 14.5 centimeters above the floor of the bottom wall 205, and is further centered between the door 203 and the rear wall 208. The sound level meter support 216 is constructed of aluminum and is bolted (not shown) to side wall 206. The control chamber 202 includes a front wall 230, two side walls 231 and 232, a top wall 233, and a bottom wall 234. Each wall is constructed of 0.125-inch (0.3175 cm) thick 6061 grade anodized aluminum. The front wall 230 is 36 inches (91.4 cm) high and 24 inches (61.0 cm) wide. The control chamber side walls 231 and 232 are each 36 inches high (91.4 cm) and 12 inches (30.5 cm) wide. The control chamber top and bottom walls 233 and 234 are each 24 inches (61.0 cm) wide and 12 inches (30.5 cm) long. The control chamber 202 is bolted (not shown) to the outer surface of rear wall 208 along seam 270 (Fig. 12). The outer surface of the rear wall 208, and the front wall 230, two side walls 231 and 232, top wall 233, and bottom wall 234 of the control chamber 202 are each coated with 0.600-inch (1.524 cm) thick sound insulating material, part number NYC-600BE, available from Small Parts, Inc. The testing apparatus 200 further includes a sound level meter 220 (Fig. 13), such as a model 1900, equipped with a model OB-100 octave filter set, both available from Quest Technologies, a company having offices in Oconomowoc, Wisconsin. The sound level meter is supported by a model QC-20 calibrator and QuestSuite master module software, each also available from Quest Technologies. The software is installed on a personal computer (not shown). During operation of the testing apparatus, the sound level meter 220 rests in the sound level meter support 216. The sound level meter includes a microphone 221 extending 4.75 inches (12 centimeters) therefrom.

Although by no means required, the apparatus may also contain features for automatically deforming a sample during a test. For example, the apparatus may contain a lower slide bracket 210,

a six-inch (15.24 cm) high Series A1500 Model available from Velmex, Inc., which extends from the bottom wall 205 into the test chamber 201, and a lower clamp 211 that is affixed to the lower slide bracket 210. An eyelet 212 (Fig. 13) may optionally extend from the top wall 204 into the test chamber 201, and an optional lanyard 213 extends through the eyelet 212. One end of the lanyard 213 extends
5 into the test chamber 201, and has an upper clamp 214 affixed thereto. The other end of the lanyard 213 extends into the control chamber 202 through a lanyard orifice 215, which is 5/8 inch (16 mm) in diameter. The lanyard may be a premium-braid, 80-lb test Spiderwire®, part number SB80G-300, manufactured by Johnson Worldwide Associates (JWA), Inc.

Prior to testing a specimen using the testing apparatus 200, the following steps are followed:

- 10 1. Calibrate the sound level meter 220 following the instructions in the manufacturer's manual.
2. Insert the full length of the microphone 221 into the testing chamber 201 (it should extend past the wall and sound dampening material approximately 2.5 inches (6.35 cm)), positioned at a 90-degree angle to side wall 206. Allow the sound level meter 220 to rest in the sound level meter support 216.
- 15 3. Activate the sound level meter per the manufacturer's instruction manual. This will collect the ambient noise inside the cavity of the test chamber 200.
4. Set the octave filter to 2,000 or 4,000 Hz and take a reading for each test conducted by activating the sound level meter until the testing has been completed.

Having calibrated the testing apparatus 200 and having identified the ambient noise, five (5)
20 specimens of a sample (film or package) may then be physically deformed approximately 15 to 20 centimeters from the microphone within the test apparatus. The film and package samples in the Examples below were manually deformed as follows:

1. Open the dominant hand with the palm facing upward;
2. Place the sample in the palm of the dominant hand;
- 25 3. Compress the test specimen by making a gentle fist;
4. Quickly open the hand and release the test specimen; and
5. Repeat this four (4) more times to equate to five (5) sample "crumples."

The tests ranged between 1 to 2 seconds in duration. All tests involved starting the sound level meter, completing the respective product manipulation, and then stopping the sound level meter.
30 The chamber door was allowed to remain open during testing to accommodate the arms and hands of the tester, as well as the motions of the product manipulation protocols. The ambient noise outside of the apparatus was quiet and was the same during calibration (including measuring the ambient noise level) and testing of the sample.

CONTROL EXAMPLE 1

A commercial film was tested that is produced by Quanxing Plastics, Inc. The film had a basis weight of 18 grams per square meters and was cast extruded from a blend containing 35 to 45 wt.% high density polyethylene (HDPE 5070), 20 to 30 wt.% low density polyethylene (LDPE LD100AC), 25 to 35 wt.% linear low density polyethylene (LLDPE 7050), and 5 to 10 wt.% of a titanium dioxide masterbatch (1605H). A micro-embossed pattern having a depth of 1 to 5 micrometers was formed on a surface of the film.

CONTROL EXAMPLE 2

A package was formed that contained a wrapper material formed from the film of Control Example 1 and a feminine pad product. The feminine care product was formed that contained a backsheet having a basis weight of 22 grams per square meter and cast extruded from a blend containing 25 to 35 wt.% high density polyethylene (HDPE 5070), 20 to 30wt.% low density polyethylene (LDPE LD100AC), 15 to 25 wt.% linear low density polyethylene (LLDPE 7050), and 5 to 10 wt.% of a titanium dioxide masterbatch (1605H). A deep embossed pattern having a depth of 5 to 15 micrometers was formed on a surface of the film.

EXAMPLE 1

A film was formed from a blend containing 93.5 wt.% of a polyethylene masterbatch, 4.5 wt.% of Clayton™ HY, and 2 wt.% of Fusabond® E-528, which is a graft copolymer of polyethylene and maleic anhydride available from DuPont. The polyethylene masterbatch contained 35 to 45 wt.% high density polyethylene (HDPE 5070), 20 to 30 wt.% low density polyethylene (LDPE LD100AC), 25 to 35 wt.% linear low density polyethylene (LLDPE 7050), and 5 to 10 wt.% of a titanium dioxide masterbatch (1605H). Clayton™ HY is an organoclay powder treated with quaternary ammonium solution and is available from BYK Additives, Inc. (Gonzales TX). The blend was formed using a co-rotating, twin screw extruder available from Entek®. The extruder had 14 processing barrels, with 13 heated barrels. Three barrels are open barrels. The outer diameter of the screws was 53 mm. The processing temperatures along the extruder were set as 175°C and the compounding speed in the twin extruder was set as 700 rpm. The resultant strands were cooled in a water bath with 15 feet length in total. The cooled strand was then pelletized and collected for the following film processing.

A film was thereafter produced from the resulting blend using a single-screw extruder. The extruder has 8 processing barrels, with 7 heated barrels with temperatures ranging from 175-210°C. The screw speed was adjusted between 66.5 and 68.6 rpm depending on the required film thickness. The corresponding line speed was adjusted between 85 and 90 meters per minute. The resulting film had a basis weight of 13.5 grams per square meter. A micro-embossed pattern having a depth of 1 to 5 micrometers was also formed on the surface of the film.

EXAMPLE 2

A film was formed as described in Example 1, except that a deep embossing pattern was formed on the surface.

The films of Control Example 1, Example 1, and Example 2 were then tested for noise level as described herein. The tested films were rectangular and had a size of 13.5 centimeters by 28.0 centimeters. The results are shown in Table 1 below.

Table 1: Noise Level of Films at 4,000 Hz

	Noise Level (dB)	Normalized Noise Level
Control Example 1	46.0	2.6
Example 1	39.6	2.2
Example 2	38.0	2.1
Ambient	17.7	1.0

As indicated above, the film of Examples 1 and 2 showed considerable reduction in noise level compared with the control sample.

EXAMPLE 3

A film was formed from a blend containing 93.5 wt.% of a polyethylene masterbatch, 4.5 wt.% of Clayton™ HY, and 2 wt.% of Fusabond® E-528, which is a graft copolymer of polyethylene and maleic anhydride available from DuPont. The polyethylene masterbatch contained 25 to 35 wt.% high density polyethylene (HDPE 5070), 20 to 30 wt.% low density polyethylene (LDPE LD100AC), 15 to 25 wt.% linear low density polyethylene (LLDPE 7042), 15 to 25 wt.% linear low density polyethylene (LLDPE 7050), and 5 to 10 wt.% of a titanium dioxide masterbatch (1605H). The blend was formed using a co-rotating, twin screw extruder available from Entek. The extruder had 14 processing barrels, with 13 heated barrels. Three barrels are open barrels. The outer diameter of the screws was 53 mm. The processing temperatures along the extruder were set as 175°C and the compounding speed in the twin extruder was set as 700 rpm. The resultant strands were cooled in a water bath with 15 feet length in total. The cooled strand was then pelletized and collected for the following film processing.

A film was thereafter produced from the resulting blend using a single-screw extruder. The extruder has 8 processing barrels, with 7 heated barrels with temperatures ranging from 175-210°C. The screw speed was adjusted between 66.5 and 68.6 rpm depending on the required film thickness. The corresponding line speed was adjusted between 85 and 90 meters per minute. The resulting film had a basis weight of 17.0 grams per square meter. A micro-embossed pattern having a depth of 1 to 5 micrometers was also formed on the surface of the film.

EXAMPLE 4

A film was formed as described in Example 1, except that a deep embossing pattern was formed on the surface. The resulting film had a basis weight of 12.0 grams per square meter.

EXAMPLE 5

A feminine care absorbent article was produced on the commercial-scale equipment used to form KOTEX® pads (Kimberly-Clark). The absorbent article was placed inside a pouch and sealed to form a package. The film of Example 1 was used to form the pouch and the film of Example 3 was used to form the backsheet of the absorbent article.

EXAMPLE 6

A feminine care absorbent article was produced on the commercial-scale equipment used to form KOTEX® pads (Kimberly-Clark). The absorbent article was placed inside a pouch and sealed to form a package. The film of Example 1 was used to form the pouch and the film of Example 4 was used to form the backsheet of the absorbent article.

EXAMPLE 7

A feminine care absorbent article was produced on the commercial-scale equipment used to form KOTEX® pads (Kimberly-Clark). The absorbent article was placed inside a pouch and sealed to form a package. The film of Example 2 was used to form the pouch and the film of Example 4 was used to form the backsheet of the absorbent article.

The packages of Example 5-7 were then manually tested for noise level as set forth above. The results are set forth in Table 2 below.

Table 2: Noise Level of Packages at 4,000 Hz

	Noise Level (dB)	Normalized Noise Level
Control Example 2	54.4	3.1
Example 5	47.5	2.7
Example 6	45.7	2.6
Example 7	46.8	2.6
Ambient	17.7	1.0

CONTROL EXAMPLE 3

A film was formed from Dowlex™ EG 2244G using a HAAKE Rheomex® 252p single screw extruder. Dowlex™ EG2244G is a linear low density polyethylene having a melt flow index of 1.0 g/10 minutes at 190°C (Dow Chemical). The screw had a diameter of 19.05 mm and an L/D of 25 (L is the screw length). A chill roll was used to cool and flatten the polymer as it exited the cast film die. The screw speed was maintained at 60 rpm for a target thickness of 27.94 micrometers. The four controlled temperature zones from the first heating zone to the die adaptor were set at 190°C, 190°C, 190°C, and 190°C, respectively. The residence time was about 1 minute during extrusion.

CONTROL EXAMPLE 4

A film was formed as described in Control Example 3, except that the target thickness was 12.7 micrometers and the screw speed was about 30 rpm.

EXAMPLE 8

Film layers were formed from blends containing various percentages of LLDPE (Dowlex™ EG 2244G) and a nanoclay masterbatch (Nanocor™ available from Nanocor, Inc.), as reflected below in Table 3. The nanoclay masterbatch contained 50 wt.% Nanomer™ nanoclay (quaternary ammonium surface-modified montmorillonite) and 30 wt.% low density polyethylene and 20% maleic anhydride grafted polyethylene. The blends were formed using a Werner & Pfleiderer (W&P) ZSK-30 co-rotating, twin screw extruder. The extruder had 14 processing barrels, with 13 heated barrels. Three barrels are open barrels. The outer diameters of the screws were 30 mm and the inner screw diameters were 21.3 mm. The lengths of the screws were 1328 mm and the total processing section length was 1338 mm. The zones had a processing temperature of 170°C, 180°C, 190°C, 190°C, 190°C, 190°C, and 180°C, respectively. The melt temperature was about 202°C and the pressure was about 60-80 psi. The compounding speed in the twin screw extruder was set as 250 rpm.

Once formed, the blends were formed into a film layer having a target thickness of 28 micrometers using a HAAKE single screw extruder as described in Control Example 3. The resulting samples were then conditioned overnight at 23 ± 2°C and 50 ± 5% RH and subjected to mechanical testing as described above. The results are set forth below in Table 3.

Table 3: Mechanical Properties of the Films of Example 8

	LLDPE (wt.%)	Nanoclay Masterbatch (wt.%)	Avg. Peak Stress (MPa)		Avg. Peak Elongation (%)		Avg. Modulus (MPa)		Avg. Energy Per Volume at Break (J/cm ³)	
			MD	CD	MD	CD	MD	CD	MD	CD
Control Example 3	100	-	36	26	544	759	61	55	76	87
Example 8	96	4	41	31	625	859	116	115	99	111
	90	10	40	34	591	903	128	143	105	140
	84	16	40	33	581	873	165	154	112	138

As indicated, the mechanical properties (e.g., peak elongation) generally improved with the incorporation of nanoclay. In the machine direction (MD), a higher amount of nanoclay led to a slightly lower strain-at-break and higher elastic modulus due to the rigid nature of nanoclay, but the elongation in MD and CD are still higher than the control film without nanoclay, although the peak stress was approximately the same.

The film of Example 8 (10 wt.% nanoclay masterbatch) was also analyzed using X-ray diffraction and transmission electron microscopy. The results are shown in Figs. 6-7. As depicted in Fig. 6, only a small peak was observed at an angle of about 7°, which indicates that only a small portion of the nanoclay remained unexfoliated or undispersed. This is further evidenced by the

transmission electron microphotographs shown in Fig. 7. Namely, Fig. 7(a) shows that the nanoclay was well dispersed in the film and Fig. 7(b) shows that the nanoclay was exfoliated into single platelets (Fig. 7(b)) and dispersed in orientation. Minor nanoclay clusters were also seen in the film as represented by the circles.

Dynamic rheology testing was also performed on Control Example 5 and the samples containing 4 wt.%, 10 wt.%, and 16 wt.% of the nanoclay masterbatch. The results are shown in Figs. 8-10. In the terminal (low frequency) zone, the neat LLDPE melt demonstrated a typical liquid like behavior – logarithmic storage modulus (G') vs. logarithmic frequency (γ) showed a smooth linear relationship (lower curve in Fig. 8). The remaining films containing the nanoclay masterbatch not only exhibited much higher G' and complex viscosity (η^*) values than neat LLDPE, but also demonstrated drastically different terminal behaviors. Irrespective of frequency, G' increased monotonically with increasing nanoclay content in the blends (Fig. 8). The viscoelastic response of the blends was mainly altered at low frequencies, where G' exhibited weak frequency dependences (Fig. 8). The small slope of $\log(G')$ vs. $\log(\gamma)$ in the films containing 10 wt.% nanoclay masterbatch (labeled as “90/10”) and 16 wt.% masterbatch (labeled as “84/16”) indicates a significant pseudo-solid-like behavior of the melt. This result suggests that an interconnected network structure of anisometric filler – a characteristic solid or gel-like structure had formed in the system. From Fig. 8, it can be seen that the reduction of the slope of $\log(G')$ vs. $\log(\gamma)$ with respect to the neat polymer became phenomenal at 10% nanoclay masterbatch (“90/10”), suggesting the critical content of nanoclay for the percolation threshold.

On the other hand, in the terminal zone the plots of $\log(\eta^*)$ vs. $\log(\gamma)$ changed from a Newtonian (primary) plateau (lower curve in Fig. 9) for the neat LLDPE to a clear shear-thinning behavior for the blends, providing more evidence of elastic behavior due to the solid network structure of nanoclay. Further evidence of the formation of a pseudo-solid-like network of percolated threads is also noted in Fig. 10, where variations of storage modulus (G') and loss modulus (G'') of the blend films vs. γ are compared. The nanofilms with a low nanoclay content (e.g., 4 wt.% nanoclay, “96/4”) displayed a lower G' than G'' over the whole frequency range. However, with the buildup of network structure, in the terminal zone G' exceeded G'' due to the pseudo-solid like behavior. At higher frequencies, crossover of G' and G'' was noted, which was probably due to the destruction of the network structure at high shear rates. Expectedly, the crossover of G' and G'' was observed for all those nanofilms with nanoclay contents equal to or higher than 10%. It is worth mentioning that the crossover point shifted to higher frequency with more nanoclay in the blends, as can be seen by the blue arrows in Fig. 10.

EXAMPLE 9

Film layers were formed as described in Example 8, except that the target thickness was 12.7

micrometers. The results are set forth below in Table 4.

Table 4: Mechanical Properties of the Films of Example 9

	LLDPE (wt.%)	Nanoclay Masterbatch (wt.%)	Avg. Peak Stress (MPa)		Avg. Peak Elongation (%)		Avg. Modulus (MPa)		Avg. Energy Per Volume at Break (J/cm ³)	
			MD	CD	MD	CD	MD	CD	MD	CD
Control Example 4	100	-	35	26	391	818	103	39	53	75
Example 9	96	4	37	29	452	809	137	129	68	91
	90	10	44	28	404	852	185	133	80	97
	84	16	41	24	435	810	146	144	82	84
	78	22	61	25	351	765	245	200	119	106
	72	28	51	22	294	631	291	233	88	80
	66	34	52	19	254	540	396	250	88	67

As indicated, the mechanical properties (e.g., peak elongation) generally improved with the incorporation of nanoclay. In the machine direction (MD), a higher amount of nanoclay led to a slightly lower strain-at-break and higher elastic modulus due to the rigid nature of nanoclay, although the peak stress was approximately the same.

EXAMPLE 10

Film layers were formed from a blend containing 96 wt.% Dowlex™ EG 2244G and 4 wt.% of a Nanocor™ masterbatch. The blend was formed using a Werner & Pfleiderer (W&P) ZSK-30 co-rotating, twin screw extruder as described in Example 1, except that the screw speed was 150 rpm. The blend was formed into film having a target thickness of 27.94 micrometers and 12.7 micrometers using a HAAKE single screw extruder as described in Control Examples 5 and 6. The resulting samples were then conditioned overnight at 23 ± 2°C and 50 ± 5% RH and subjected to mechanical testing as described above. The results are set forth below in Table 5.

Table 5: Mechanical Properties of the Films of Example 10

	Thickness (μm)	LLDPE (wt.%)	Nanoclay Masterbatch (wt.%)	Avg. Peak Stress (MPa)		Avg. Peak Elongation (%)		Avg. Modulus (MPa)		Avg. Energy Per Volume at Break (J/cm ³)	
				MD	CD	MD	CD	MD	CD	MD	CD
Control Example 3	27.94	100	-	36	26	544	759	61	55	76	87
Control Example 4	12.70	100	-	35	26	391	818	103	39	53	75

Example 10	27.94	96	4	41	33	641	912	121	120	98	127
	12.70	96	4	41	29	473	815	154	122	72	99

While the invention has been described in detail with respect to the specific embodiments thereof, it will be appreciated that those skilled in the art, upon attaining an understanding of the foregoing, may readily conceive of alterations to, variations of, and equivalents to these embodiments.

5 Accordingly, the scope of the present invention should be assessed as that of the appended claims and any equivalents thereto.

WHAT IS CLAIMED IS:

1. A package comprising a wrapper material that defines an interior cavity and an absorbent article removably positioned within the cavity, wherein the wrapper material is formed from a film having a thickness of from about 0.5 to about 80 micrometers, wherein the film contains a polymer composition that includes one or more ethylene polymers in an amount of from about 75 wt.% to about 99 wt.% of the polymer composition and one or more nanofillers in an amount of from about 0.5 wt.% to about 20 wt.% of the polymer composition, and wherein the package exhibits a normalized noise level of about 3.0 or less as determined at a frequency of 4,000 Hz, the normalized noise level being determined by dividing the noise level of the package by the noise level of an ambient environment.
2. The package of claim 1, wherein the package exhibits a normalized noise level of about 3.0 or less as determined at a frequency of 2,000 Hz.
3. The package of claim 1, wherein the package exhibits a noise level of about 50 dB or less as determined at a frequency of 4,000 Hz.
4. The package of claim 1, wherein the ethylene polymer is a copolymer of ethylene and an α -olefin.
5. The package of claim 4, wherein the ethylene polymer is linear low density polyethylene.
6. The package of claim 1, wherein the polymer composition includes linear low density polyethylene, low density polyethylene, high density polyethylene, or a combination thereof.
7. The package of claim 1, wherein the nanofiller has an average cross-sectional dimension of from about 0.2 to about 100 nanometers.
8. The package of claim 7, wherein the nanofiller is in the form of platelets having an aspect ratio of from about 20 to about 1,000.
9. The package of claim 1, wherein the nanofiller includes carbon black, carbon nanotubes, carbon nanofibers, nanoclays, metal particles, silica, alumina, or a combination thereof.
10. The package of claim 1, wherein the nanofiller includes a nanoclay, wherein the nanoclay includes a phyllosilicate.
11. The package of claim 10, wherein the phyllosilicate includes a montmorillonite or a salt thereof.
12. The package of claim 1, wherein the nanofiller includes an organic surface treatment.
13. The package of claim 12, wherein the organic surface treatment includes a quaternary onium.
14. The package of claim 1, wherein the polymer composition further comprises a polyolefin compatibilizer that contains an olefin component and a polar component.
15. The package of claim 14, wherein the polar component of the compatibilizer includes

maleic anhydride.

16. The package of claim 1, wherein the film is multi-layered and contains a base layer and/or skin layer formed from the polymer composition.

17. The package of claim 1, wherein a surface of the film is embossed.

18. The package of claim 1, wherein the wrapper material consists of the film.

19. The package of claim 1, wherein the absorbent article comprises a liquid permeable layer, a liquid-impermeable layer, and an absorbent core positioned between the liquid permeable layer and the liquid-impermeable layer

20. The package of claim 1, wherein the absorbent article is a feminine care product.

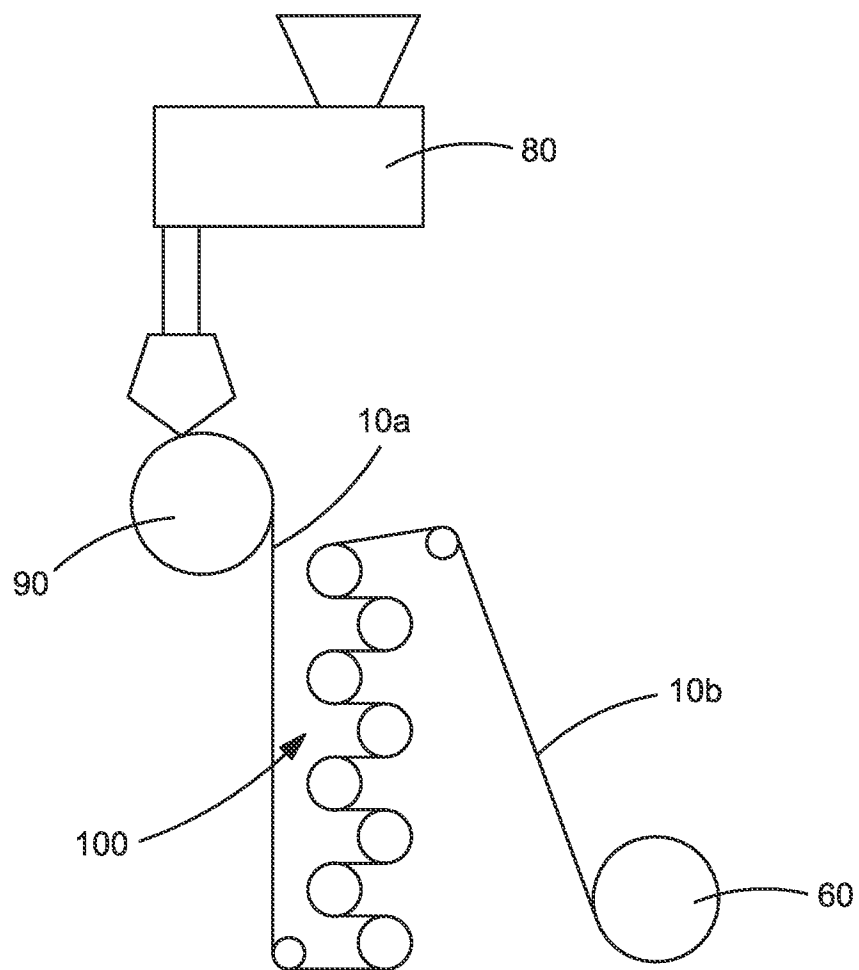


FIG. 1

2/11

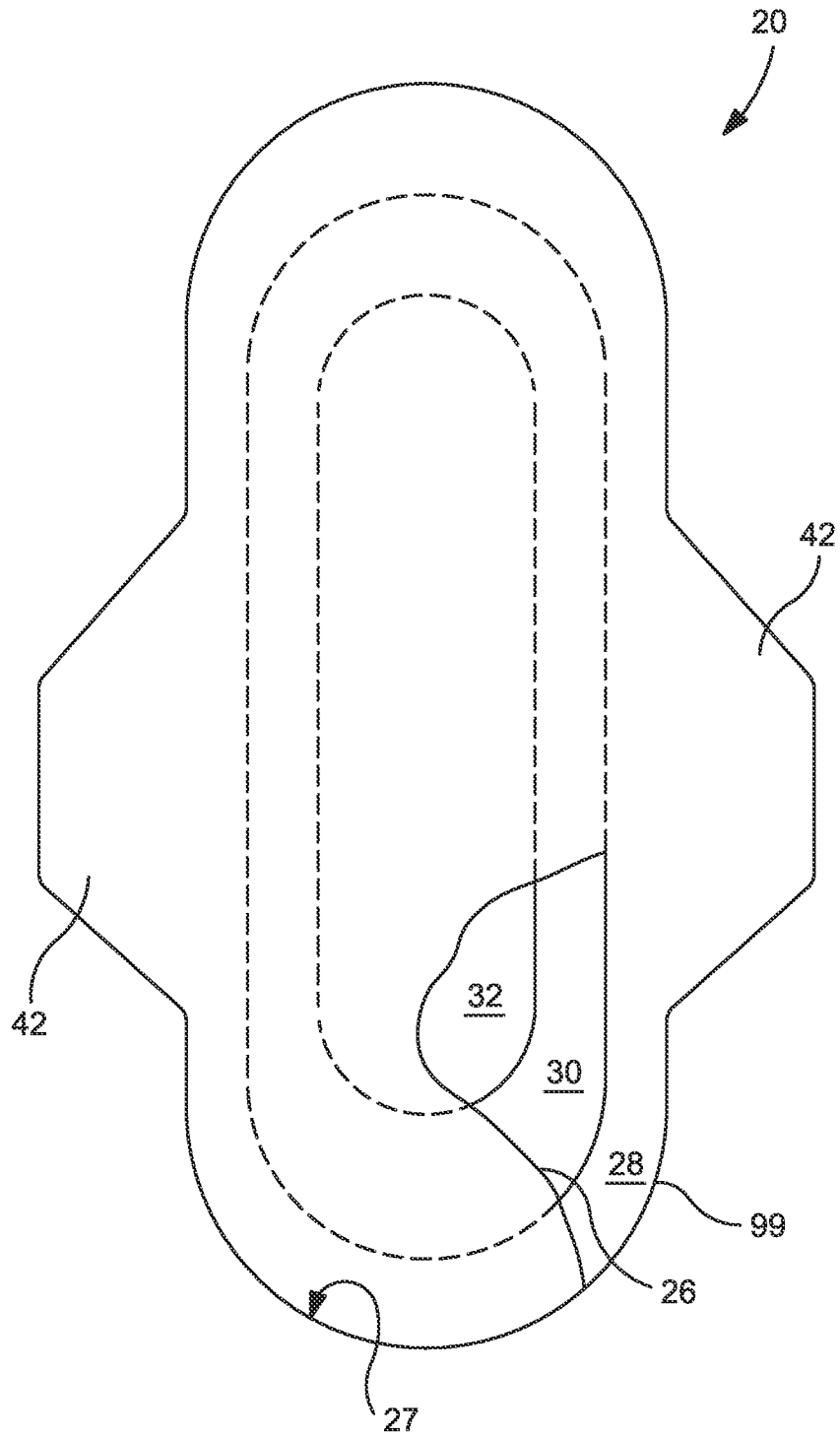


FIG. 2

3/11

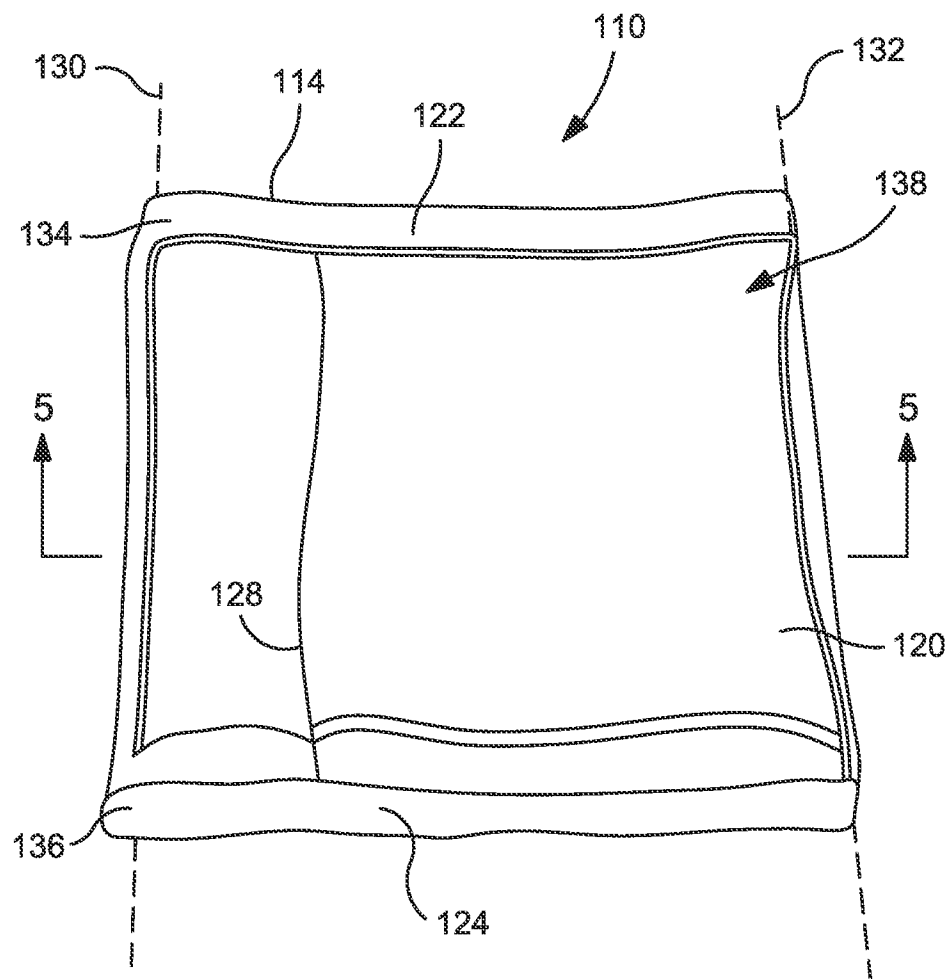
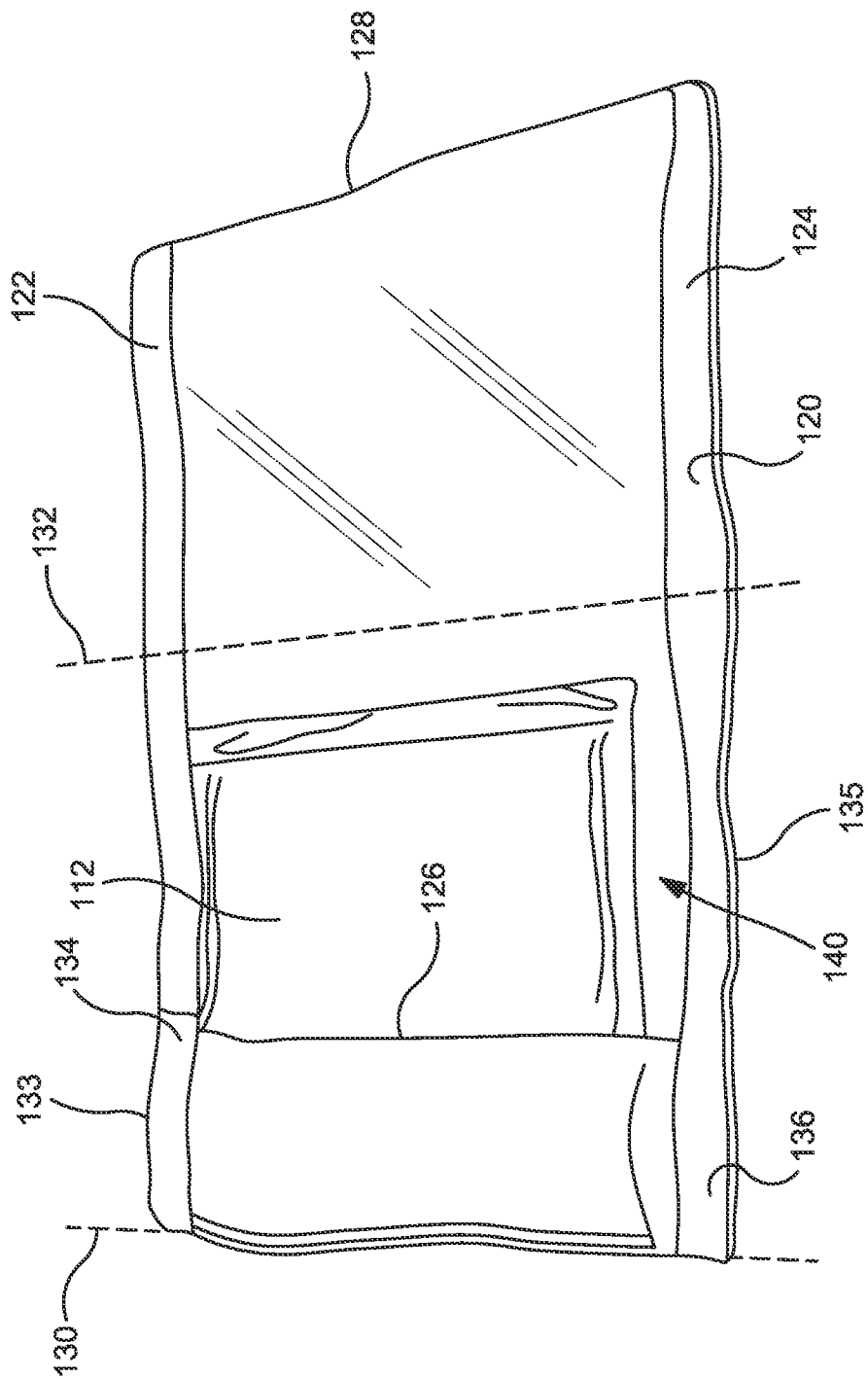


FIG. 3



ଶ୍ରୀ

5/11

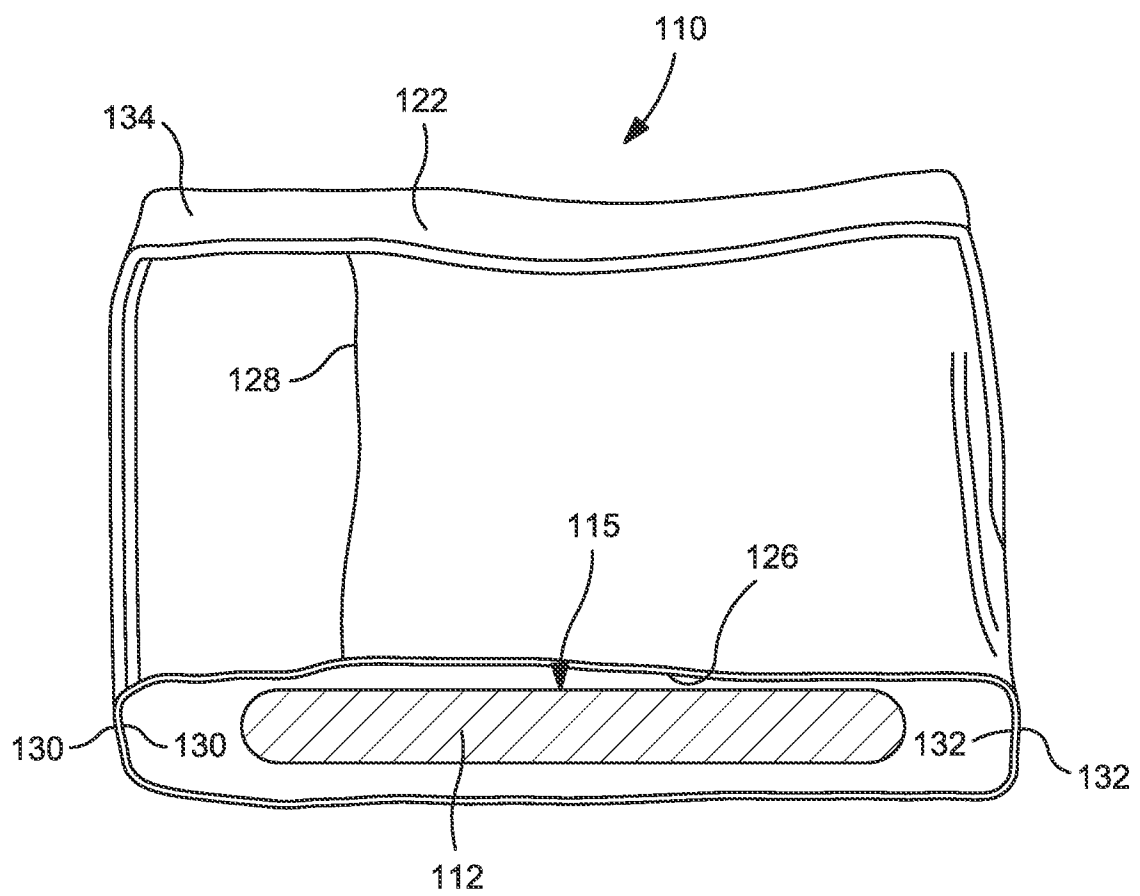
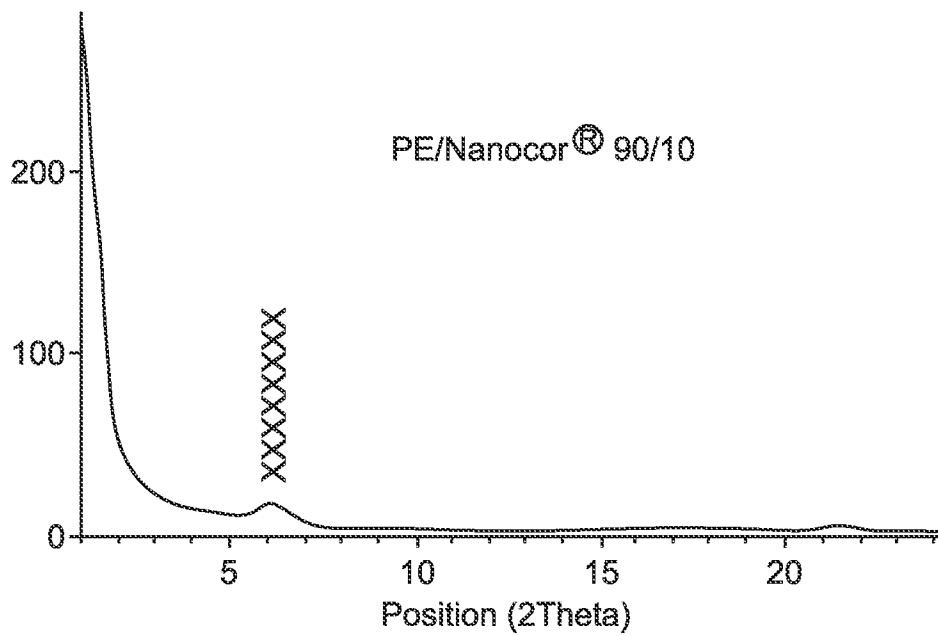
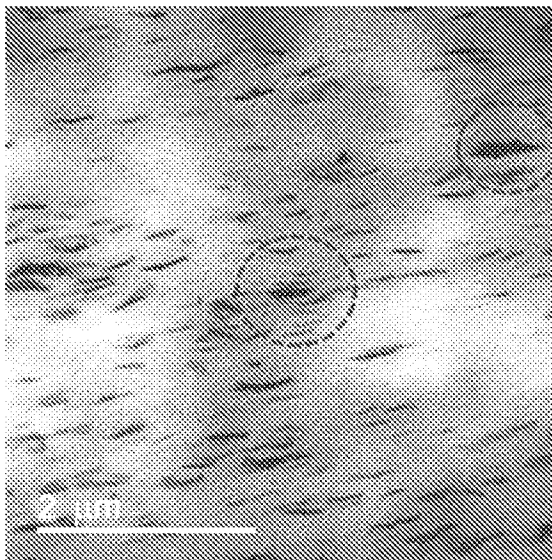
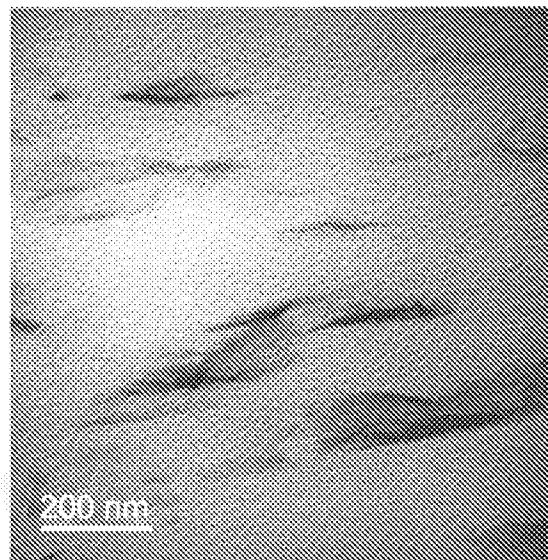
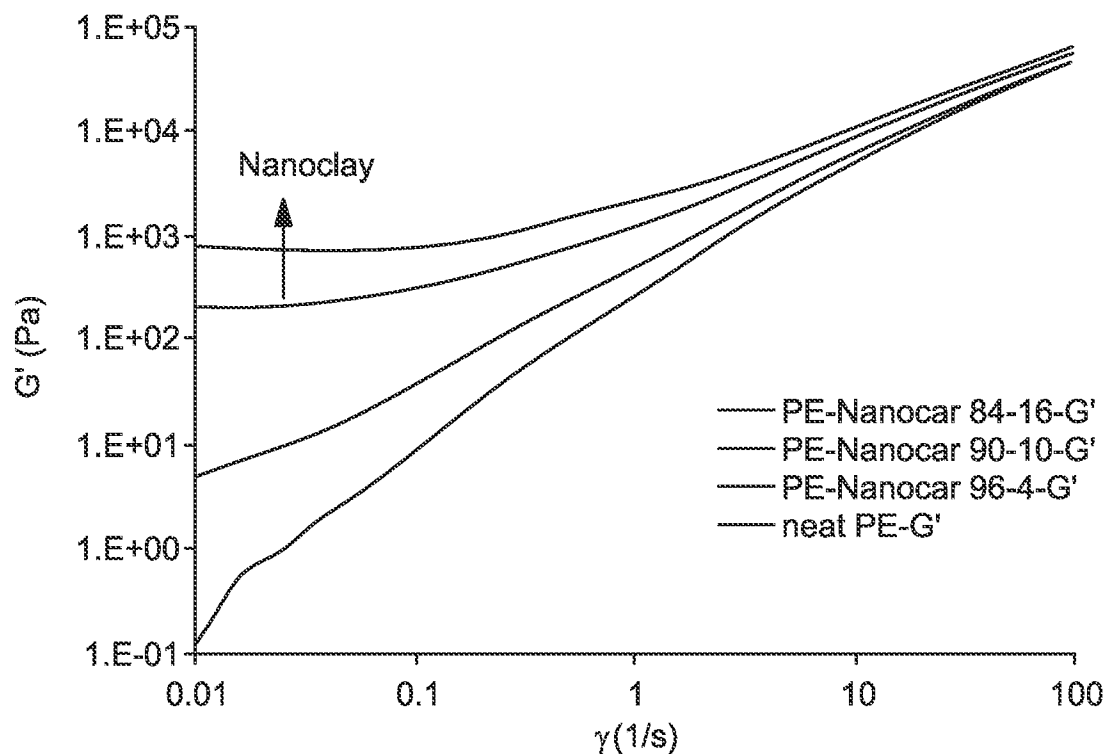
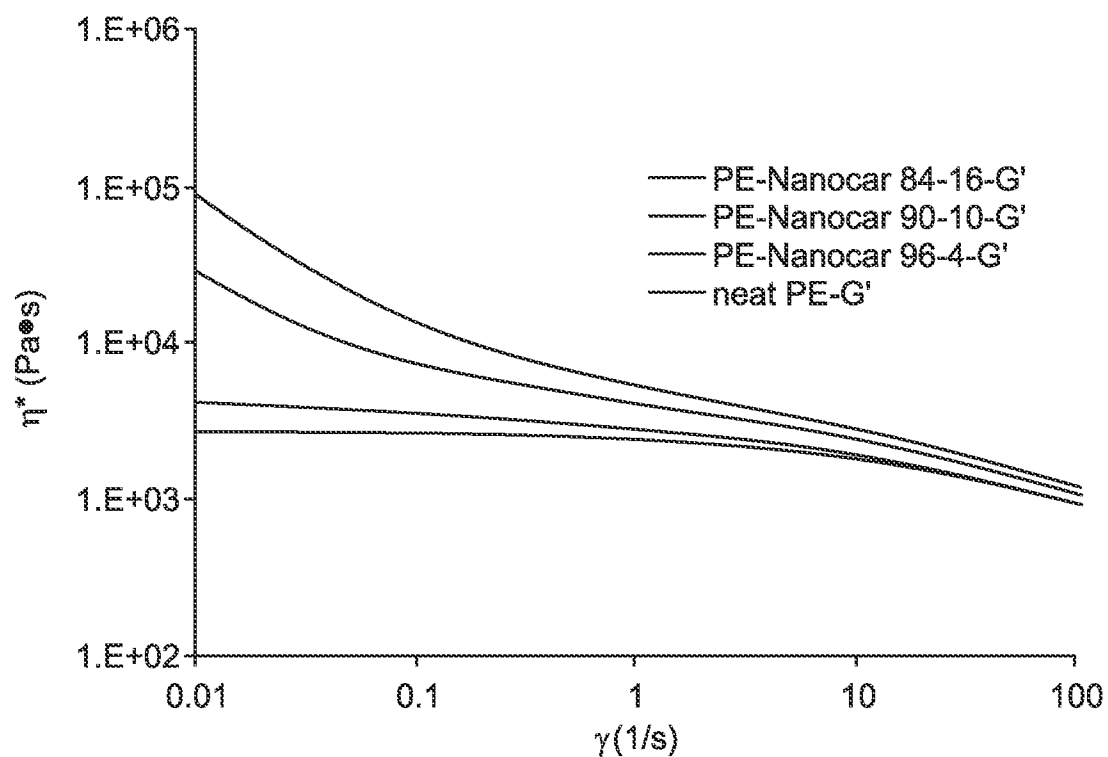


FIG. 5

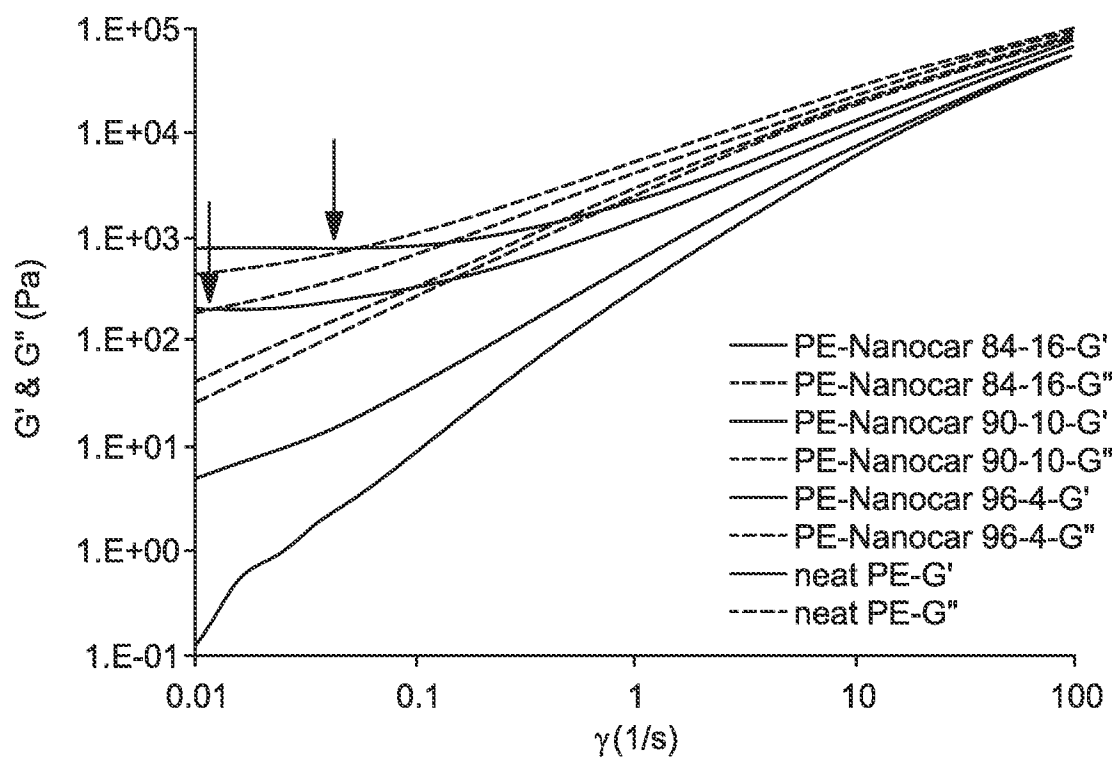
6/11

**FIG. 6****FIG. 7a****FIG. 7b**

7/11

**FIG. 8****FIG. 9**

8/11

**FIG. 10**

9/11

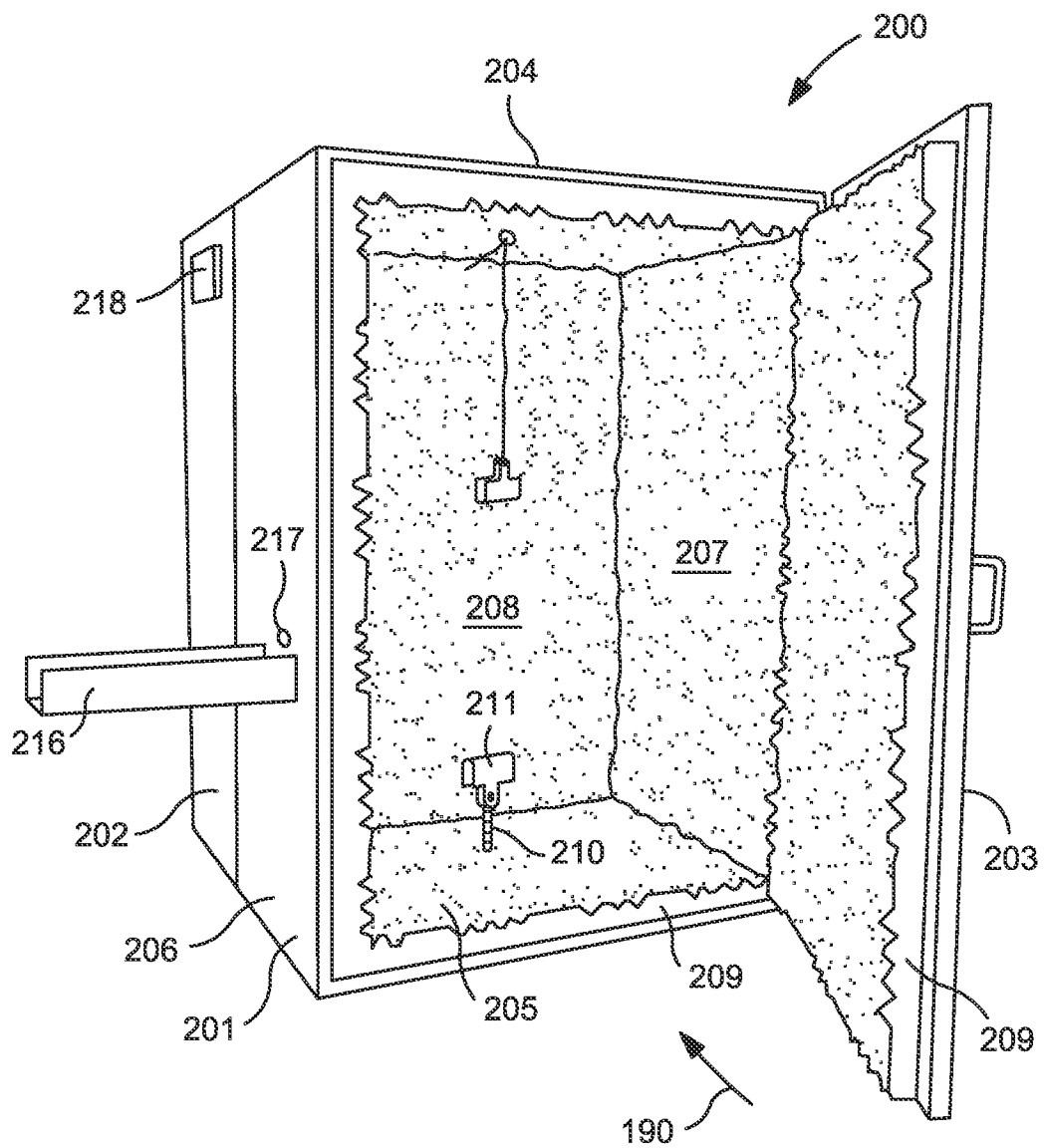
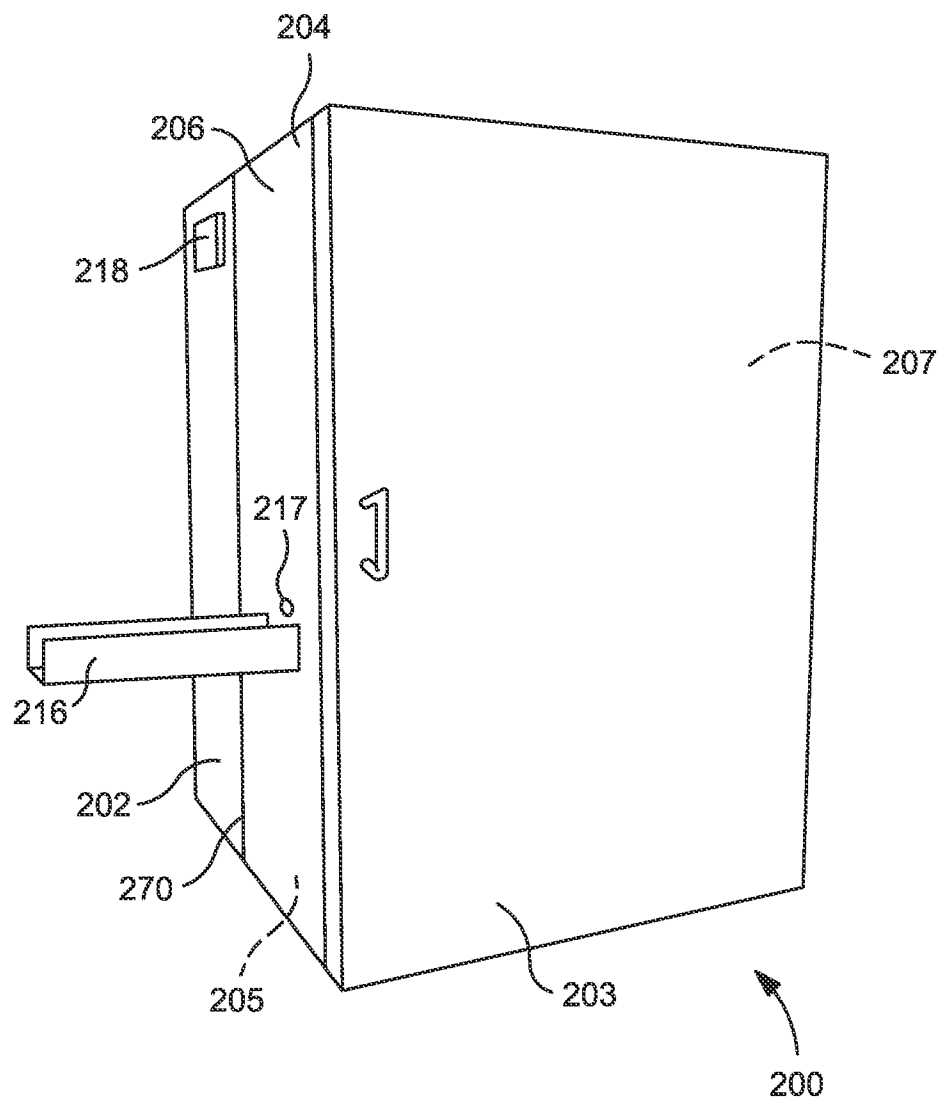


FIG. 11

10/11

**FIG. 12**

11/11

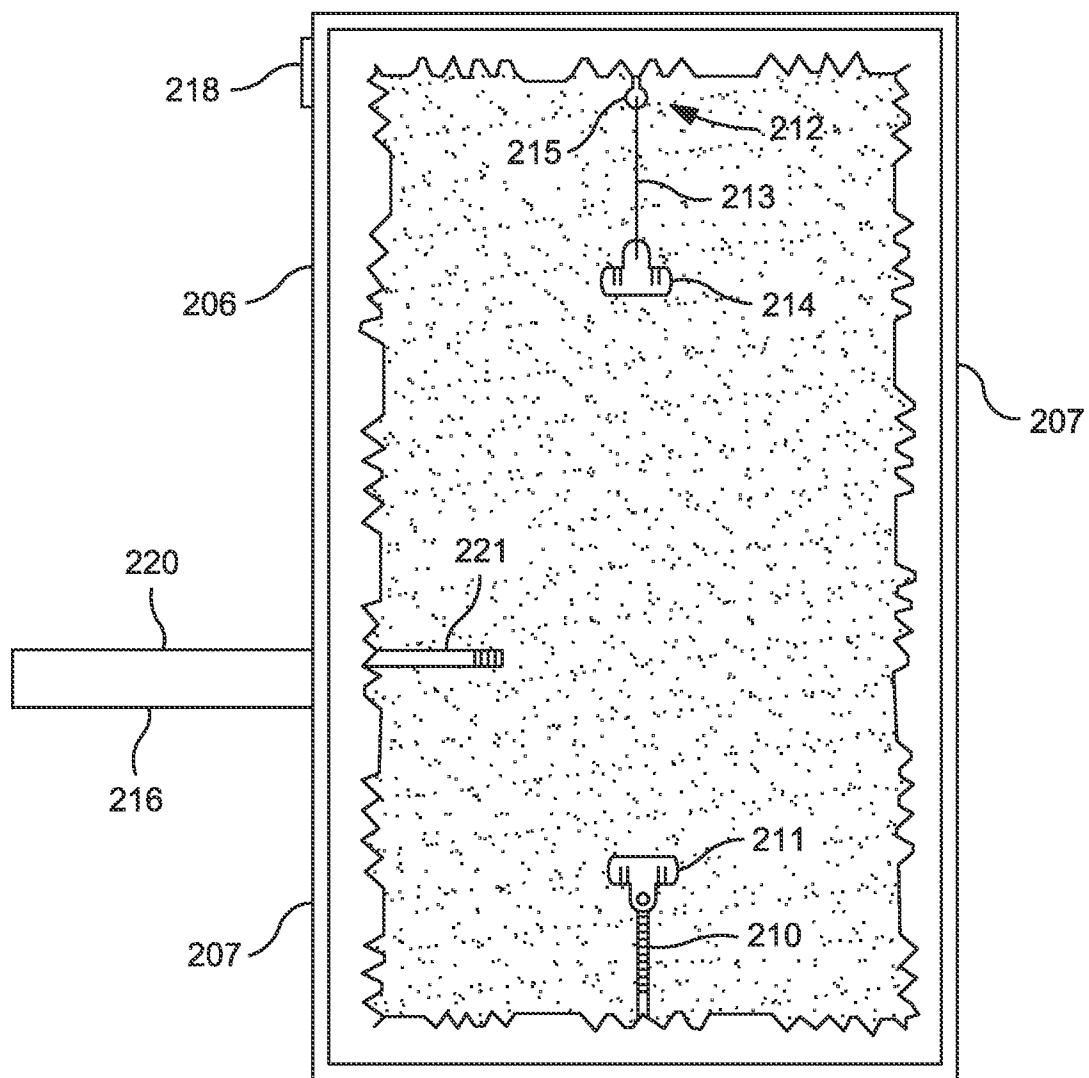


FIG. 13

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2015/013851**A. CLASSIFICATION OF SUBJECT MATTER****A61F 13/15(2006.01)i, C08L 23/08(2006.01)i**

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61F 13/15; B32B 27/30; B32B 27/08; A61F 13/20; C08L 23/08

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

eKOMPASS(KIPO internal) & keywords: package, film, absorbent article, noise level, ethylene, nanofiller

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 6455161 B1 (REGNIER, F. J. F. et al.) 24 September 2002 See abstract; claims 1-3, 11; column 7, lines 28-32.	1-20
A	US 2009-0030395 A1 (DALAL, U. P. et al.) 29 January 2009 See entire document.	1-20
A	US 7872169 B2 (RUIZ, O. A. et al.) 18 January 2011 See entire document.	1-20
A	US 5254111 A (CANCIO, L. V. et al.) 19 October 1993 See entire document.	1-20
A	US 2004-0078015 A1 (COPAT, M. S. et al.) 22 April 2004 See entire document.	1-20



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

26 August 2015 (26.08.2015)

Date of mailing of the international search report

05 October 2015 (05.10.2015)

Name and mailing address of the ISA/KR

International Application Division
Korean Intellectual Property Office
189 Cheongsu-ro, Seo-gu, Daejeon Metropolitan City, 302-701,
Republic of Korea

Facsimile No. +82-42-472-7140

Authorized officer

Han, Inho

Telephone No. +82-42-481-3362



INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2015/013851

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 6455161 B1	24/09/2002	EP 1196482 A2 EP 1196482 B1 JP 2003-503572 A JP 2011-174072 A US 6620474 B1 WO 01-00716 A2	17/04/2002 16/01/2008 28/01/2003 08/09/2011 16/09/2003 04/01/2001
US 2009-0030395 A1	29/01/2009	EP 2173304 A2 US 8323258 B2 WO 2009-018133 A2	14/04/2010 04/12/2012 05/02/2009
US 7872169 B2	18/01/2011	EP 1962761 A2 JP 2009-521347 A US 2007-0149938 A1 WO 2007-092104 A2	03/09/2008 04/06/2009 28/06/2007 16/08/2007
US 5254111 A	19/10/1993	EP 0515501 A1 EP 0515501 B1 US 5202173 A US 5296184 A WO 91-12125 A1	02/12/1992 01/06/1994 13/04/1993 22/03/1994 22/08/1991
US 2004-0078015 A1	22/04/2004	EP 1517783 A1 JP 2005-529768 A	30/03/2005 06/10/2005