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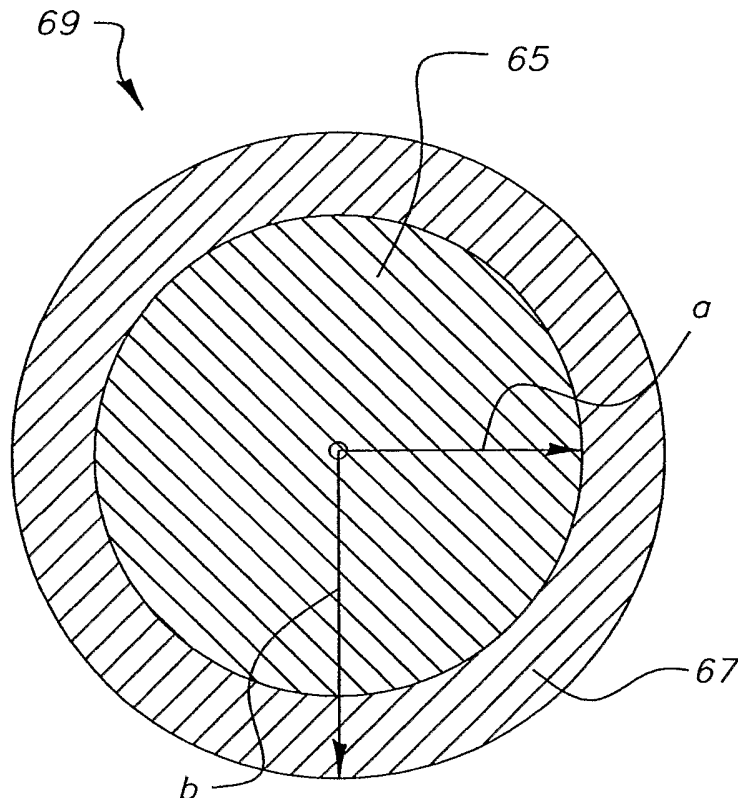
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[Continued on next page]

(54) Title: PLASTIC OPTICAL ARTICLES CONTAINING CORE SHELL COMPOSITE NANOPARTICLES



(57) Abstract: A nanocomposite optical plastic article has a plastic host material with a temperature sensitive optical vector (x) and a core shell nanoparticulate material dispersed into the plastic host material. The core shell nanoparticulate material is characterized by a core defined by a nanoparticulate material which has a temperature sensitive optical vector (x_1) and a shell defined by a coating material layer coated onto the core. It is important to the invention that temperature sensitive vector (x_1) is directionally opposed to the temperature sensitive optical vector (x) and $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$.



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PLASTIC OPTICAL ARTICLES CONTAINING CORE SHELL COMPOSITE NANOPARTICLES

FIELD OF THE INVENTION

The invention relates generally to the field of plastic optical articles
5 and films. More particularly, the invention concerns plastic optical materials and
articles containing core shell composite nanoparticles, such as plastic lenses,
optical films, and optical couplers, that must maintain stable performance
characteristics over a broad temperature range.

BACKGROUND OF THE INVENTION

10 Plastic lenses and glass lenses often perform the same function in
optical systems, such as in cameras, microscopes, telescopes, and ophthalmic wear.
The two main attributes that separate plastic lenses from glass lenses are cost and
optical stability. Plastic lenses typically cost 1/100th the price of a similar glass
lens while the stability of the refractive index of a glass lens with respect to
15 temperature and humidity is typically 100 times better than that of a plastic lens.

The difference in cost is due largely to the difference in
manufacturing processes that are required for the two materials and the relative
temperatures that the materials must be formed at. Plastic lenses are typically
produced at 230° C. using injection molding at cycle times that are 10 times faster
20 than glass lenses that are largely produced by grinding and polishing or
compression molding at 625° C. Grinding and polishing are labor intensive while
the high temperatures that glass must be formed at requires expensive mold
materials and extensive maintenance costs.

In contrast, the difference in optical stability between plastic and
25 glass is due to differences in their basic material properties. This difference in
optical stability results in substantially more variation in focus and image quality
in articles such as cameras when plastic lenses are used in place of glass. What is
desired, and a remaining challenge in the art, is a material with the optical stability
of glass that processes like a plastic.

30 While optical plastic materials such as cyclic olefins greatly
improve the refractive index stability with respect to humidity, improving the
refractive index stability with respect to temperature has remained an opportunity.

A study on the competing fundamental material characteristics that determine the sign and the magnitude of the dn/dT of glasses is available, for instance, by Lucien Prod'homme, "A New Approach To The Thermal Change In The Refractive Index Of Glasses," Physics and Chemistry of Glasses, Vol. 1, No. 4, 5 August 1960. There are two competing effects that determine the dn/dT in glasses. These are the density change, which produces a negative dn/dT and the electronic polarizability, which produces a positive dn/dT . The net dn/dT in a glass material depends on which effect dominates. In optical plastics however, the electronic polarizability is very small compared to the density change so that all 10 unfilled plastic materials have negative dn/dT values. Nonetheless, the article by Prod'homme does identify the possibility of using glass-like fillers with positive dn/dT values to substantially alter the dn/dT of a glass-plastic composite material.

Nanoparticulate fillers have been used to modify the index of refraction of optical plastics. By using a nanoparticulate filler small enough that it 15 is well below the wavelength of visible light (400-700 nm), light scattering from the nanoparticles is reduced and the filled plastic can remain transparent. WIPO Patent No. WO97/10527 by John S. Toeppen, published March 20, 1997, titled "Structured Index Optics And Ophthalmic Lenses For Vision Correction" describes the use of nanoparticles to increase the refractive index of plastics for 20 ophthalmic applications. In addition, technical references that describe the addition of nanoparticles to increase the refractive index of plastics include: "Optical And Thermomechanical Investigations On Thermoplastic Nanocomposites With Surface Modified Silica Nanoparticles" by C. Becker et al., SPIE Conference, Vol. 3469, pp. 88-98, July 1998; and "Tantalum Oxide Nanomers For Optical 25 Applications" by B. Braune et al., SPIE Conference, Vol. 3469, pp. 124-132, July 1998. While these references disclose the use of nanoparticles to modify refractive index of optical plastics they do not discuss the issue of refractive index stability with respect to temperature which requires a different set of characteristics in the nanoparticle.

30 U.S. Patent No. 6,441,077 titled "Polysulfone Nanocomposite Optical Plastic Article And Method Of Making Same" issued August 27, 2002 to Border et al., discloses the use of a nanoparticulate filler in an optical plastic,

where the nanoparticulate filler has been chosen with a positive dn/dT to counteract the negative dn/dT of the plastic host material such that the overall dn/dT of the nanocomposite has a substantially reduced magnitude of dn/dT , where dn is the change in refractive index of the nanocomposite material produced
5 by a change in temperature dT .

In experiments and computer modeling done at Eastman Kodak, it has been noted that the addition of even very small nanoparticles (10-40nm) into plastic host materials can lead to low level light scattering or haze in the nanocomposite material which limits light transmission to less than 90%. The
10 haze is especially noticeable at loadings greater than 10% of the nanoparticles in the plastic host material, such as is required to substantially improve the thermal stability of the refractive index (reduced dn/dT) of the nanocomposite material. Haze is also particularly noticeable if the refractive index of the nanoparticles is substantially different from the refractive index of the plastic host material.

15 While there have been several attempts to modify properties of plastics using nanoparticles, none of these attempts have proven successful in producing optical plastic articles with improved temperature stable optical properties while retaining important processing characteristics and low levels of haze.

20 U.S. Patent No. 5,252,441 titled "Transparent Magnetic Recording Layers And Photographic Elements Containing The Same" issued October 12, 1993 to James et al. and U.S. Patent No. 5,217,804 titled "Magnetic Particles" issued June 8, 1993 to James et al. disclose a coating technique for magnetic particles that reduces the extinction coefficient by applying coatings to high
25 refractive index magnetic particles thereby reducing the haze of the material. Examples show how the coating of the particles with a low refractive index layer reduced the optical density (haze). However, James et al. do not contemplate the impact of the particles or the coating material on the overall thermal stability of the refractive index of the nanocomposite material.

30 U.S. Patent No. 5,985,173 titled "Phosphors Having A Semiconductor Host Surrounded By A Shell" issued November 16, 1999 to Gray et al. describes a technique for applying a shell coating to doped zinc sulfide

particles. However, the goal of this invention is related to establishing a bandgap to modify the surface electronic state of the doped host particle. This invention does not address the transparency of the nanocomposite material.

Likewise, in WIPO Publication Number WO99/21934 by
5 Mulvaney et al., published May 6, 1999 and titled "Stabilized Particles And Methods Of Preparation And Use Thereof," Mulvaney et al. disclose a method for stabilizing nanoparticles by coating them with an insulating, semiconducting and/or metallic coating. While Mulvaney et al., do disclose techniques for applying coatings to nanoparticles and some of the materials presented do have
10 positive dn/dT values such cadmium sulfide, zinc sulfide, zinc selenide, silica, and alumina, Mulvaney et al. only disclose the use of coated nanoparticles to improve fluorescence, electrofluorescence, and detection of an analyte. Mulvaney et al. do not anticipate the use of coated nanoparticles to modify the optical performance of optical articles such as lenses to improve thermal stability (dn/dT) of the refractive
15 index. In fact, in Example F on page 19, line 6, Mulvaney et al. make the statement that "the optical properties of the CdS particles are not significantly altered by silica deposition."

Therefore, a need persists in the art for optical plastic articles, such as lenses, and a method of making same that have improved temperature stable
20 optical properties with low levels of haze.

PROBLEM TO BE SOLVED BY THE INVENTION

The goal of the invention is to provide a nanocomposite material which processes like a plastic material but has a significantly reduced magnitude of dn/dT while also having low haze.

25 Border et al. (U.S. Patent No. 6,441,077) disclosed a technique in which plastic lens properties can be improved in respect to thermal stability by incorporating certain nanoparticles with a positive dn/dT into the plastic resins (which have a negative dn/dT) used for melt injection molding. As well as providing a positive contribution to dn/dT , the particles must be very small to
30 reduce light scatter and haze in the lens that would degrade image quality. However, if the particles are too big or the refractive index is too high relative to the plastic medium, the light scatter and haze becomes high. This is especially

true considering that the path length in lens applications is typically of the order of 1mm to 10 mm. Another factor is that the loading of nanoparticles required in the plastic host material is often greater than 20% to reduce the dn/dT by more than 20%.

5 Zinc sulfide (ZnS) is a particularly interesting material for improving thermal stability of optical properties because it has a rather high positive value of the parameter dn/dT (ca $+39 \sim +49 \text{E-}6/^{\circ}\text{C}$) so that less zinc sulfide nanoparticles have to be added to the plastic host material to achieve a level of dn/dT improvement of the nanocomposite. However, zinc sulfide also has a high
10 refractive index, e.g., $n \sim 2.39$ compared to the plastic host material which has a refractive index of approximately 1.5, which means that scattering and haze will be a serious problem. Scattering and haze are also a problem with other materials with a positive dn/dT , e.g., MgO, Al_2O_3 , AlOOH , SiO_2 , etc. as mentioned by Border et al., however, scattering and haze with these materials is not as
15 noticeable as with ZnS because the refractive indices of these materials are closer to the refractive index of the plastic host material. The greater the difference in refractive indices between the nanoparticle and the plastic host, the greater will be the haze produced in the nanocomposite material.

There is however a general, inventive trick in the optical science of
20 particles that can be applied to reduce scattering and essentially make nanoparticles invisible in virtually any plastic host material as long as the certain rules and material properties are obeyed.

For a composite or core shell particle as shown in Figure 6,
provided that:

25 $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$
where the core has real and imaginary optical constants: n_{core} , ($k_{\text{core}} \sim 0$)
the shell has real and imaginary optical constants: n_{shell} , ($k_{\text{shell}} \sim 0$)
and the plastic host has a refractive index: $n_{\text{plastic host}}$

30 Then there will be a particular value of the shell radius to core ratio or shell volume to core volume at which the scattering produced by the coated nanoparticle will tend to have very low values or become “essentially invisible” so

that the haze of the nanocomposite also tends to have very low levels and the transparency becomes very high. The optimum value for coating thickness can be calculated from an equation given by Milton Kerker in "The Scattering Of Light And Other Electromagnetic Radiation," Academic Press, New York, 1969 (page 192 Equation 5.1.25 and page 197 Equation 5.1.51) which when solved for the shell radius to core ratio becomes:

$$b/a = [(2m_2^2 + 1)(m_2^2 - m_1^2)/(m_2^2 - 1)(m_1^2 + 2m_2^2)]^{1/3}$$

Equation 1

where a is the radius of the core nanoparticle, b is the radius of the coated composite nanoparticle, m_1 is the ratio of the refractive indices of the core material to the plastic host material ($n_{\text{core}}/n_{\text{plastic host}}$) and m_2 is the ratio of the refractive indices of the shell material to the plastic host material ($n_{\text{shell}}/n_{\text{plastic host}}$).

This is a general rule that will hold true for all transparent materials that have a refractive index and do not have high values for the absorption coefficients or imaginary parts of the complex refractive index (k_{core} or k_{shell}).

This is the basis of the inventive idea for preparing highly filled plastic lenses with invisible nanoparticles.

The utility of this idea depends on the refractive index contrast between the core, shell, and plastic host. The lower the refractive index of the shell (n_{shell}), compared to the refractive index of the plastic host ($n_{\text{plastic host}}$), the thinner the shell can be and as a result, the less overall nanocomposite nanoparticles need to be added to the plastic host material to obtain the desired dn/dT .

SUMMARY OF THE INVENTION

It is, therefore, a first object of the invention to provide an optical nanocomposite material that can be formed into an optical article that has reduced temperature sensitivity.

Another object of the invention is to provide a method for reducing the haze in the nanocomposite material.

It is a feature of the optical article of the invention that a select nanoparticulate is dispersed into a plastic host material which has a temperature

sensitive optical vector that is directionally opposed to the temperature sensitive optical vector of the nanoparticulate filler.

Another feature of the optical article of the invention is that the nanoparticle dispersed in the plastic host material is a composite, core shell nanoparticle wherein the haze produced by the nanoparticle is reduced due the effects of the shell on the core when the materials have been selected such that their refractive indices follow the rule: $n_{\text{core}} > n_{\text{plastic host}} > n_{\text{shell}}$. Under particular combinations of shell and core geometry (e.g., radius ratio, or volume ratio or mass ratio), as determined by the refractive index contrasts of the core, shell, and plastic host, the scattering of light, or haze of the nanocomposite material in an optical article will be reduced to very low values.

To accomplish these and other objects, features and advantages of the invention, there is provided, in one aspect of the invention, a nanocomposite optical plastic article comprising a plastic host material having a temperature sensitive optical vector x ; a core shell nanoparticulate material dispersed into said plastic host material, said core shell nanoparticulate material having a core defined by a nanoparticulate material having a temperature sensitive optical vector x_1 , a shell defined by a coating material layer coated onto said core, said core shell having a temperature sensitive optical vector x_2 , and wherein said temperature sensitive optical vector x_1 is directionally opposed to said temperature sensitive optical vector x of said plastic host material.

The present invention has numerous advantageous effects over existing developments, including: the resulting nanocomposite has a significantly lower dn/dT (i.e., change in refractive index with temperature); lenses made with the nanocomposite material have more stable focal length over a given temperature range; low levels of dn/dT are achievable in the nanocomposite material with reduced loading of the nanoparticulate; the viscosity of the nanocomposite material is not significantly higher than the base plastic so that conventional plastic processing techniques can be used; light scattering and haze in the nanocomposite material and the optical article are reduced to the extent that transparency of the nanocomposite is similar to that of the

polymethylmethacrylate host material without the nanoparticles; and it has broad utility for making a variety of optical articles such as lenses, filters, film, or slabs.

BRIEF DESCRIPTION OF THE DRAWINGS

The above and other objects, features, and advantages of the present invention will become more apparent when taken in conjunction with the following description and drawings wherein identical reference numerals have been used, where possible, to designate identical features that are common to the figures, and wherein:

FIG. 1 is a prior art plastic lens showing a range of focal length variation produced by a change in temperature and the resulting change in refractive index;

FIG. 2a shows a lens made from a nanocomposite material that has improved stability of refractive index with respect to temperature and an associated reduced range of focal length variation produced by a change in temperature;

FIG. 2b shows a representative view of the nanocomposite material as a pellet before forming into an optical article;

FIG. 3 is a block diagram of the process for manufacturing a plastic optical article of the invention with improved refractive index stability;

FIG. 4 is a schematic diagram of a nanocomposite material making process of the invention based on compounding;

FIG. 5 is a schematic diagram of another nanocomposite material making process of the invention based on solvent dispersion; and

Fig. 6 is a diagram of a core shell composite nanoparticle used in the plastic article of the invention.

DETAILED DESCRIPTION OF THE INVENTION

Turning now to the drawings, and in particular to FIG. 1, prior art lens 1 is illustrated having focal length 5 of light rays 3 that varies significantly with changes in temperature (T) (shown schematically as a focal length range 7). The relationship between focal length 5 and refractive index (n) is given by the below equation:

$$f = R/(n-1)$$

Equation 2

wherein (f) is the focal length of the lens produced as incident light goes through the lens and is focused at a focal point, (R) is the radius of the lens surface, and (n) is the refractive index of the lens material.

In the case of a camera lens (not shown), the temperature range of operation can easily be 50° C. when used to photograph a tropical island and then later used to photograph a snowy mountain. As an example, a lens having a 10 mm radius and made, for instance, of polymethylmethacrylate, the index of refraction (n) at room temperature is 1.492 and the focal length (calculated from Equation 2 above) is 20.325 mm.

Referring again to FIG. 1, in a typical prior art camera lens, such as lens 1, comprising a plastic material selected from Table I, the change in refractive index (dn) over the temperature range of operation is 0.0055. Further, the change in focal point (shown as feature 7 in FIG. 1) of lens 1 from Equation 1 is 0.225 or 1%. Skilled artisans will appreciate that the image quality of images made with lens 1 will not be the same over the entire operating temperature range due to variations in focus quality.

Referring to FIGS. 2a and 2b, the reduced temperature sensitive, nanocomposite optical article, such as lens 10, of the invention is illustrated. According to FIG. 2a, the nanocomposite optical article or lens 10 is composed of a plastic host material 16 and a select nanoparticulate material 14 dispersed in the plastic host material 16 (shown clearly in FIG. 2b). The plastic host material 16 may be either a thermoplastic or thermoset material. It is important to the invention that the plastic host material 16 be selected based on a predetermined temperature sensitive optical vector x, for instance refractive index n. Similarly, the selection of the nanoparticulate material 14 dispersed in the plastic host material 16 must be based on a corresponding predetermined temperature sensitive optical vector x_1 , specifically refractive index. In this case, temperature sensitive optical vectors x, and x_1 are defined by a change in refractive index (dn) of the plastic host material 16 and the nanoparticulate material 14, respectively, with respect to a change in temperature (dT). It is also important to our invention

that x_1 is directionally opposed to x . By carefully selecting a nanoparticulate material 14 having a dn/dT , i.e., a rate of change of refractive index with respect to temperature, that has a sign that is directionally opposed to the dn/dT of the plastic host material 16, it is possible to significantly reduce the dn/dT of the resulting nanocomposite material 15 at relatively low loadings of the nanoparticulate material 14. As a result, the viscosity of the nanocomposite material 15 is not drastically increased and the processing characteristics will be similar to other optical plastics. Consequently, the resulting optical nanocomposite lens 10 has a focal length range 12 (FIG. 2a) over the operating temperature range that is much less than the focal length range 7 exhibited by the prior art lens 1 shown in FIG. 1.

According to Tables I and II, several select dn/dT values for plastic (polymeric) host materials and inorganic nanoparticulate fillers that comprise the nanocomposite material of the invention are illustrated.

15

Table I
Approximate dn/dT for Various Optical Plastics

Plastic	dn/dT ($10^{-6}/^{\circ}\text{C}$)
Polymethylmethacrylate	-105
Polystyrene	-127
Polycarbonate	-114
Cyclic olefin copolymer	-102
Polysulfone	-100
Kodak KTFR Photoresist (liquid)	-436
Olin HIPR 6505 Photoresist (liquid)	-230
Olin HIPR 6512 Photoresist (liquid)	-268

Table II
Approximate dn/dT for Various Inorganic Materials with Transmission
Bands in Visible Wavelengths

Material	dn/dT ($10^{-6}/^{\circ}\text{C}$)
Barium fluoride	-16
Aluminum oxide	14
ALON	12
Beryllium oxide	10
BBO	-16
Diamond	10
Calcium carbonate	7
Calcium fluoride	-10
Cesium bromide	-85
Cesium iodide	-99
Potassium bromide	-42
Potassium chloride	-36
Potassium fluoride	-23
Potassium iodide	-45
Potassium titano phosphate	12
Lithium borate	-7
Lithium fluoride	-17
Lithium iodate	-80
Magnesium aluminate	9
Magnesium oxide	19
Sodium bromide	-40
Sodium chloride	-35
Sodium fluoride	-13
Sodium iodide	-50
Silicon oxide	-5
Silica	12
Tellurium oxide	9
Titanium dioxide	-1
Yttrium oxide	8
Zinc Sulfide	49
Zinc Selenide	91
Cadmium Sulfide	80
Magnesium fluoride	1
Barium fluoride	-16

Referring again to FIGS. 2a and 2b, in addition to the plastic host material 16 and the nanoparticulate material 14 having directionally opposed dn/dT , the invention contemplates other qualifications for the nanoparticulate material 14 to make the useful, novel and unobvious optical nanocomposite material 15 of the invention.

5 For instance, the nanoparticulate material 14 must be transparent in the wavelength region of interest to maintain high optical transmission levels. Moreover, the nanoparticulate material or nanoparticles 14 must be available in a particle size range that is less than 40 nm to avoid scattering light. Most preferred is a particle size range below 20 nm. Further, it must be possible to disperse the
10 nanoparticles 14 into the base or host plastic such that no significant amounts of agglomerates and/or voids larger than 40 nm occur which would scatter light. FIG. 2b shows a representative view of the nanoparticles 14 dispersed into the plastic host material 16. The nanoparticles 14 are shown dispersed evenly throughout the plastic host material 16. The nanoparticles 14 as shown do not
15 have any larger agglomerates or voids associated with them. Furthermore, the cost of the nanoparticles 14 and any associated surface treatments of the nanoparticles 14 to improve dispersibility or reduce reactivity, etc. must be low enough that the total cost of the molded optical article is significantly less than a glass article.

20 As illustrated in Tables I and II, there exists a number of inorganic materials that have dn/dT values with an opposite sign compared to plastic host materials. Thus, a nanocomposite material with significantly improved refractive index stability with respect to temperature can be formulated by dispersing a select nanoparticulate material into a plastic host material that have directionally
25 opposed (or opposite signs) dn/dT .

According to another aspect of the invention, a method of manufacturing a reduced temperature sensitive optical article or lens 10 (as described above) includes the step of selecting a plastic host material 16, such as one described in Table I. According to the invention, the selected plastic host
30 material 16 has a temperature sensitive optical vector x or dn/dT , as described above. A nanoparticulate material 14 (Table II) is selected for dispersing in the plastic host material 16. The select nanoparticulate material 14, according to the

invention, is required to have a corresponding temperature sensitive optical vector x_1 . Moreover, it is further important to the invention that x_1 is directionally opposed to x , i.e., one of the two must be negative and the other positive. Once the nanoparticulate material 14 is selected, it is then dispersed in the plastic host material 16 using suitable dispersion techniques, such as compounding or solvent dispersion. Once the nanoparticulate material 14 is dispersed into the plastic host material 16, a nanocomposite material 15 is formed. The nanocomposite material 15 can then be used to form an array of optical articles such as the lens 10 of the invention having reduced temperature sensitivity.

Referring to FIGS. 2a, 2b and 6, in the event that the index of refraction n of the selected nanoparticulate material 14 differs from the index of refraction of the plastic host material 16 by more than approximately 0.035, and if the nanoparticles 14 are larger than approximately 3 nm in size, the haze of the resulting nanocomposite material 15 can be significantly reduced if a composite nanoparticulate material 69 (FIG. 6) is used. As illustrated in FIG. 6, composite nanoparticle 69 is composed of a core nanoparticle 65 that has been coated to form a shell 67 around the core nanoparticle 65. According to our invention, the indices of refraction of the materials should follow the rule that: $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$ for the coating to reduce the haze of nanocomposite material 15. Usually the refractive index of the core, e.g., ZnS (e.g. 2.39) will be greater than the plastic material (1.5 to 1.6) and the shell 67 must be less than the plastic material (this includes any non-absorbing, low refractive index material, e.g., amorphous silica, 1.45 to 1.47; fluoropolymer, ca 1.4; magnesium fluoride, 1.38; silsesquioxane materials, 1.47). Preferably, the shell material will also have a temperature sensitive vector x_2 that is directionally opposed to the temperature sensitive vector x of the plastic host material.

Referring to FIG. 3, a diagram 20 of the method for making the reduced dn/dT nanocomposite material 15 for optical articles, such as lens 10, is depicted. First the plastic host material 16 (FIGS. 2a, 2b) is selected (Step 22) based on optical, structural, and thermal design considerations such as % transmission, % haze, index of refraction, yield strength at a temperature, impact strength, scratch resistance, glass transition temperature, etc. Second, the

nanoparticulate material 14 (FIGS. 2a, 2b) is preferably selected (Step 24) based on dn/dT , transparency in the wavelength region of interest, index of refraction, particle size, cost, and availability. As disclosed in this invention, selecting suitable nanoparticulate materials 14 requires selecting a material that has a dn/dT that has a sign that is opposite to the plastic host material 16 being used and an average particle size less than about 40 nm. Third, the nanoparticles are coated (Step 25) to create a core shell composite nanoparticle. The coating should be selected such that: $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$ to reduce the haze in the final nanocomposite material. Fourth, the composite nanoparticles 14 are preferably dispersed (Step 26) into the plastic host material 16 although other mixing processes could be used, such as roll milling. Dispersion step 26 can be accomplished through preferably compounding (refer to FIG. 4) even though solvent dispersion (refer to FIG. 5) can be used with good results. Fifth, the optically modified material 15 is formed (Step 28) into an optical article or lens of the invention.

Referring to FIGS. 4 and 5, two methods of dispersing (Step 26) the nanoparticles 14 into the plastic host material 16 are schematically illustrated. According to FIG. 4, an outline of the process 32 for dispersion (Step 26) through compounding (Step 40) is depicted. As in Fig 3, the plastic host material 16 is selected (Step 34), the nanoparticulate is selected (Step 36) and the nanoparticles are coated (Step 37) to create composite nanoparticles. In a preferable step, the composite nanoparticles are then surface treated (Step 38) to improve the compatibility with the plastic host material 16 thereby making it easier to fully disperse the nanoparticles. Skilled artisans will appreciate that this treatment (Step 38 in FIG. 4 and Step 52 in FIG. 5) could be applied to the nanoparticles 14 directly or added as an additive to the compounder (Step 40) along with the nanoparticles 14 and the plastic host material 16. In compounding, the nanoparticles 14 are then fed into a compounder (Step 40), such as a twin screw extruder or a Farrell continuous mixer, along with pellets of the selected plastic host material 16. After compounding (Step 40), the optically modified material 15 (nanocomposite material) is pelletized (Step 42) for use in an injection molding machine (not shown).

According to FIG. 5, in the solvent-based dispersion process (Step 44), the plastic host material 16 is selected 46 and then dissolved in a solvent (Step 50). The nanoparticles are selected (Step 48), coated to create core shell composite nanoparticles (Step 49) and then preferably surface treated (Step 52) to
5 improve compatibility with the plastic host material 16. The treated composite nanoparticles are then dispersed in a solvent (Step 54) which is compatible with the solvent that the plastic host material 16 has been dissolved in (Step 50). The two solvent solutions are then mixed together (Step 56). After mixing the two solvent solutions together in Step 56, the solvents are removed in Step 58 and the
10 optically modified material 15 (nanocomposite material) is pelletized (Step 60) for use in an injection molding machine (not shown).

Referring to FIGS. 4 and 5, by following either method depicted in process diagrams 32 and 44 for making the optically modified material 15, the end result is typically plastic pellets which contain fully dispersed nanoparticles 14,
15 such as that shown in FIG. 2b. The resultant plastic pellets contain the composite nanoparticles present in sufficient quantity to deliver the reduced dn/dT desired while preserving a low level of haze.

Injection molding, compression molding, and casting are the three preferred techniques for forming the optical article 10 (refer to FIG. 3 step 28) of
20 the invention. For the case of compression molding, it may be more advantageous to use optically modified material in the form of sheet or film. For the case of casting, the optically modified material may be in the form of a liquid.

In a preferred embodiment, the nanocomposite optical article of manufacture 10 is comprised of a plastic host material 16 selected from the group
25 consisting of thermoplastic materials and thermoset materials. Thermoplastic materials used in optical articles include the following materials: polymethylmethacrylate, polycarbonate, polystyrene, polysulfone, polyether sulfone, cyclic olefins, and blends and copolymers of those listed. Thermoset materials used in optical articles include the following materials: diallyl
30 glycolcarbonate, epoxides, and thermoset polyesters.

Typically the reduced dn/dT articles of manufacture produced within the contemplation of the invention are simple lenses, an array of lenses,

ophthalmic lenses, window glazing, optical fibers, optical couplers, cover glasses for digital imagers, microlenses on digital imagers, and other optical devices of the like. Typically, all of these articles of manufacture require a low level of haze.

5 Skilled artisans will appreciate that modification of the optical properties of the host material is achieved, in accordance with the method of the invention, by reducing the dn/dT of the nanocomposite material and reducing haze. In our preferred embodiment, this is achieved by dispersing a nanoparticulate material filler having a dn/dT with a sign that is opposite that of the base plastic in which the nanoparticulate is a core shell composite nanoparticle
10 with indices of refraction $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$.

EXEMPLARY EXAMPLES

EXAMPLE 1

An example of the aforementioned method for reducing dn/dT of an optical plastic while retaining excellent transparency follows:

15 Three (3) samples were prepared using a polymethylmethacrylate plastic base material ($n = 1.49$) and nanoparticle fillers of approximately 15 nm size. The nanoparticles were all solvent dispersed at approximately 4% loading. The first sample was made with silica nanoparticles ($n = 1.45$) and it was very clear. This result is due to the fact that the silica nanoparticles have nearly the
20 same index of refraction as the polymethylmethacrylate plastic host material (1.49 vs 1.45) and light scattering does not occur to an appreciable extent. The second sample was made with magnesium oxide nanoparticles ($n = 1.72$) and it was quite hazy. The third sample was made with zinc sulfide nanoparticles ($n = 2.35$) and it was nearly opaque.

25 Observations of relative levels of haze produced in the 3 samples were supported by calculated values from a light scattering model based on the Debye form of Rayleigh-Mie Theory (Debye, P, Ann. d. Phys. 30, 57, 1909) in which the haze ($\text{haze} = [100 - \text{transmitted light \%} + \text{reflected light \%}]$) was calculated to be 1%, 16% and 53%, respectively.

30 A second set of calculations done by solving the light scattering model for each interface on core shell composite nanoparticles shows that by applying coatings of silica ($n = 1.45$) to the magnesium oxide and zinc sulfide

nanoparticles of 7.5 nm and 14.8 nm thicknesses respectively, the haze was reduced to 1.5 % for both cases.

EXAMPLE 2

A further example shows the importance of the coating thickness on the level of haze produced by composite nanoparticles. Again, the light scattering model was used to evaluate the haze levels produced by composite nanoparticles with various shell thicknesses. Three (3) coating thicknesses were evaluated with 4% zinc sulfide ($n = 2.35$) nanoparticles in polymethylmethacrylate ($n = 1.49$). The zinc sulfide nanoparticles were 20 nm and coated with silica ($n = 1.45$) with the coating thicknesses being 5 nm, 8 nm and 17 nm where the 17 nm thick coating is the optimum thickness as calculated by Equation 1. The results showed haze levels of 44%, 29%, and 2% respectively which shows that coating thickness is important to produce the lowest haze level.

EXAMPLE 3

Another example shows the impact of different types of coatings on optimum coating thickness. The light scattering model was again used to evaluate material combinations and the effect on haze. In this case, 10 nm zinc sulfide ($n = 2.35$) nanoparticles coated with 3 nm of magnesium fluoride ($n = 1.38$) and loaded to 4% in a polycarbonate ($n = 1.59$) plastic host. The result was a haze level of only 1%. In this case, the refractive index of the coating is quite different from the plastic host refractive index, 1.38 vs 1.59. Consequently the coating thickness does not need to be as thick as the case examined in Example 2 where the coating had to be 5 times as thick (17 nm) because the refractive index of the coating was close to that of the plastic host, 1.45 vs 1.49.

The invention has been described with reference to a preferred embodiment. However, it will be appreciated that variations and modifications can be effected by a person of ordinary skill in the art without departing from the scope of the invention.

PARTS LIST:

- | | | |
|----|----|---|
| | 1 | prior art lens |
| | 3 | incident light rays |
| | 5 | focal length of lens 1 |
| 5 | 7 | range of change of the focal length produced by a change in temperature of the lens |
| | 10 | nanocomposite lens |
| | 12 | reduced range of change of the focal length produced by a change in temperature of the nanocomposite lens |
| 10 | 14 | nanoparticulate material |
| | 15 | nanocomposite material (representation of nanoparticles dispersed into a plastic host material) |
| | 16 | plastic host material |
| | 20 | schematic of method for making reduced dn/dT article |
| 15 | 22 | step of selecting plastic host material |
| | 24 | step of selecting the nanoparticulate material |
| | 25 | step of coating the nanoparticles to create core shell composite nanoparticles |
| | 26 | step of dispersion |
| 20 | 28 | step of forming an optical article |
| | 32 | dispersion through compounding process |
| | 34 | step of selecting host material |
| | 36 | step of selecting nanoparticulate material |
| | 37 | step of coating the nanoparticles to create core shell composite nanoparticles |
| 25 | | nanoparticles |
| | 38 | step of surface treating nanoparticles |
| | 40 | step of compounding nanoparticles |
| | 42 | step of forming the nanocomposite into a useable form such as pelletizing |
| | 44 | solvent based dispersion process |
| 30 | 46 | step of selecting host material |
| | 48 | step of selecting nanoparticles |

Parts List - - continued

- 49 step of coating the nanoparticles to create core shell composite nanoparticles
- 5 50 step of dissolving host material in solvent
- 52 step of surface treating nanoparticles
- 54 step of dispersing nanoparticles in solvent
- 56 step of mixing together products of steps 50 and 54
- 58 step of removing the solvent
- 10 60 step of forming a useable material form
- 65 core material of composite nanoparticle
- 67 shell material of composite nanoparticle
- 69 composite nanoparticle

CLAIMS:

1. A nanocomposite optical plastic article, comprising:
a plastic host material having a temperature sensitive optical vector
x;
5 a core shell nanoparticulate material dispersed into said plastic host material, said
core shell nanoparticulate material having a core defined by a nanoparticulate
material having a temperature sensitive optical vector x_1 , a shell defined by a
coating material layer coated onto said core, said shell having a temperature
sensitive optical vector x_2 and wherein said temperature sensitive optical vector x_1
10 is directionally opposed to said temperature sensitive optical vector x of said
plastic host material.
2. The nanocomposite optical plastic article recited in claim 1
wherein said temperature sensitive optical vector x is defined as an index of
15 refraction $n_{\text{plastic host}}$, said temperature sensitive optical vector x_1 is defined as an
index of refraction n_{core} , and wherein said temperature sensitive optical vector x_2 is
defined as an index of refraction of the shell n_{shell} , wherein $n_{\text{shell}} < n_{\text{plastic host}} < n_{\text{core}}$.
3. The nanocomposite optical plastic article recited in claim 1
20 wherein said plastic host material comprises a material selected from the group
consisting of: polymethylmethacrylate, polystyrene, polycarbonate, cyclic olefin
polymer, polysulfone, polyethersulfone, diallyl glycolcarbonate, epoxides,
thermoset polyesters, and blends and copolymers of those listed.
- 25 4. The nanocomposite optical plastic article recited in claim 1
wherein said nanoparticulate material has a particle size of about 15nm.
5. The nanocomposite optical plastic article recited in claim 1
wherein said plastic host material is polymethylmethacrylate.
30
6. The nanocomposite optical plastic article recited in claim 4
wherein said nanoparticulate material comprises silica nanoparticles.

7. The nanocomposite optical plastic article recited in claim 4 wherein said nanoparticulate material comprises magnesium oxide nanoparticles.

8. The nanocomposite optical plastic article recited in claim 4
5 wherein said nanoparticulate material comprises zinc sulfide nanoparticles.

9. The nanocomposite optical plastic article recited in claim 8 wherein said zinc sulfide nanoparticles have a particle size of about 20nm, said zinc sulfide nanoparticles being provided with a silica coating layer, said silica
10 coating layer having a thickness in the range from about 5nm to about 17nm.

10. The nanocomposite optical plastic article recited in claim 8 wherein said zinc sulfide nanoparticles have a particle size of about 10nm, said zinc sulfide nanoparticles being provided with a 3nm thick coating layer of
15 magnesium fluoride forming a core shell nanoparticulate material, said core shell nanoparticulate material being dispersed at 5 to 50 wt-% in a polycarbonate plastic host material.

11. The nanocomposite optical plastic article recited in claim 1
20 wherein said nanoparticulate material comprises a material selected from the group consisting of: ALON, aluminum oxide, calcium carbonate, magnesium aluminate, yttrium oxide, zinc sulfide, cadmium sulfide, and potassium titano phosphate.

12. The nanocomposite optical plastic article recited in claim 1
25 wherein said nanoparticulate material has a particle size less than about 40 nm.

13. The nanocomposite optical plastic article recited in claim 1 wherein said nanoparticulate material has a particle size less than about 20 nm.

14. The nanocomposite optical plastic article recited in claim 1
30 wherein said coating material layer has a temperature sensitive optical vector x_2 ,

wherein x_2 is directionally opposed to said temperature sensitive optical vector x of said plastic host material.

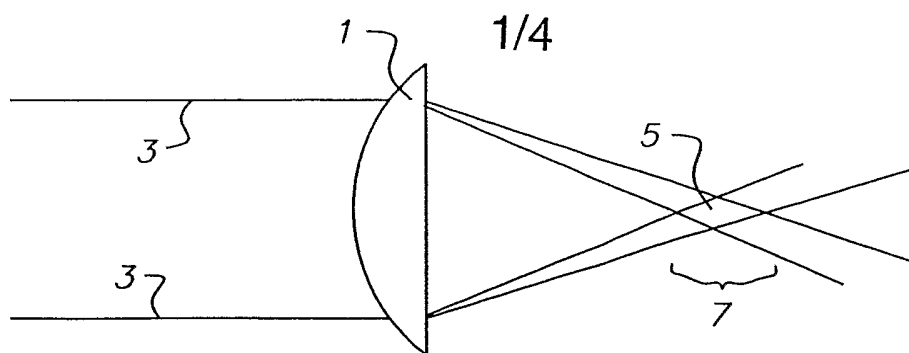


FIG. 1
(PRIOR ART)

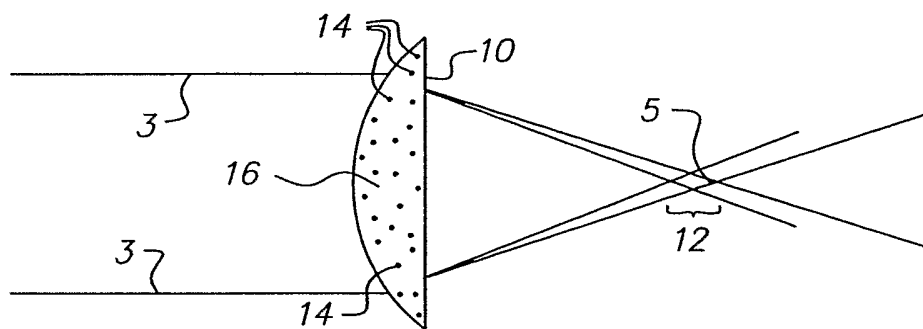


FIG. 2a

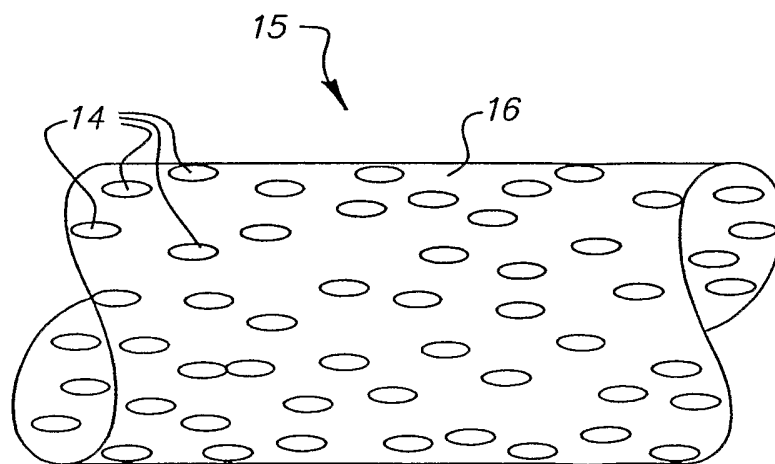
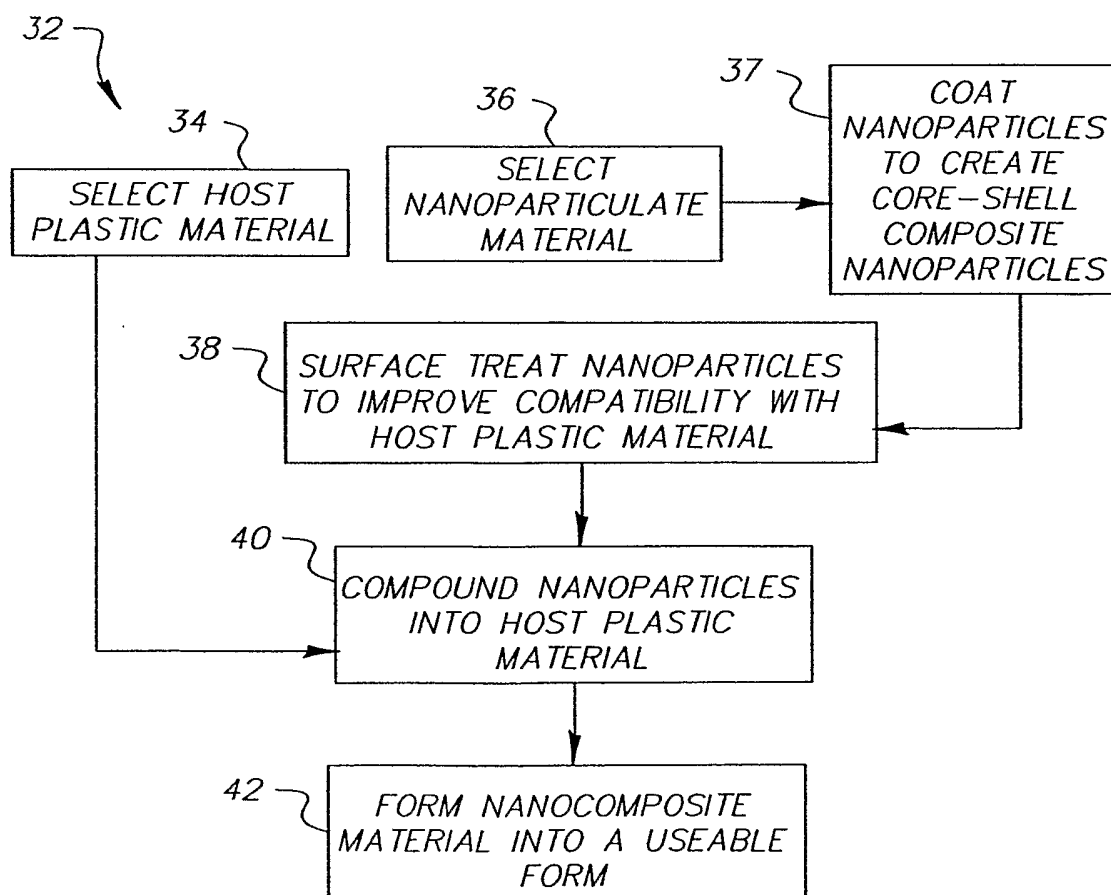
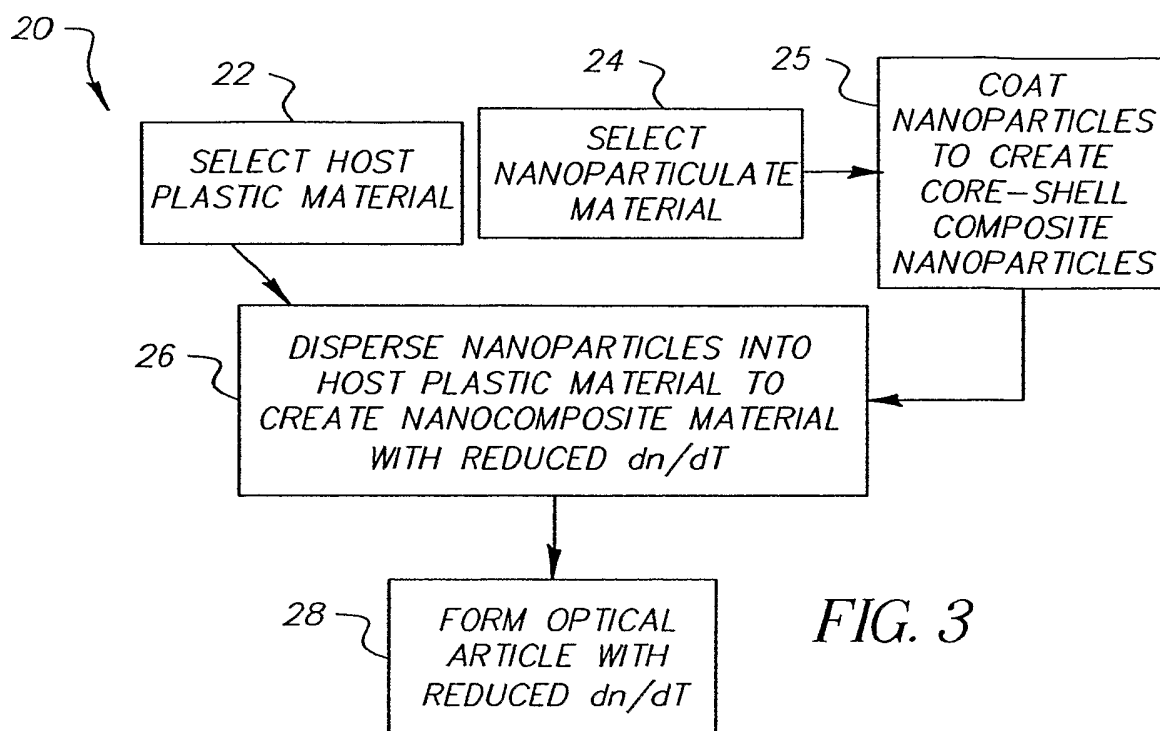


FIG. 2b

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*FIG. 4*

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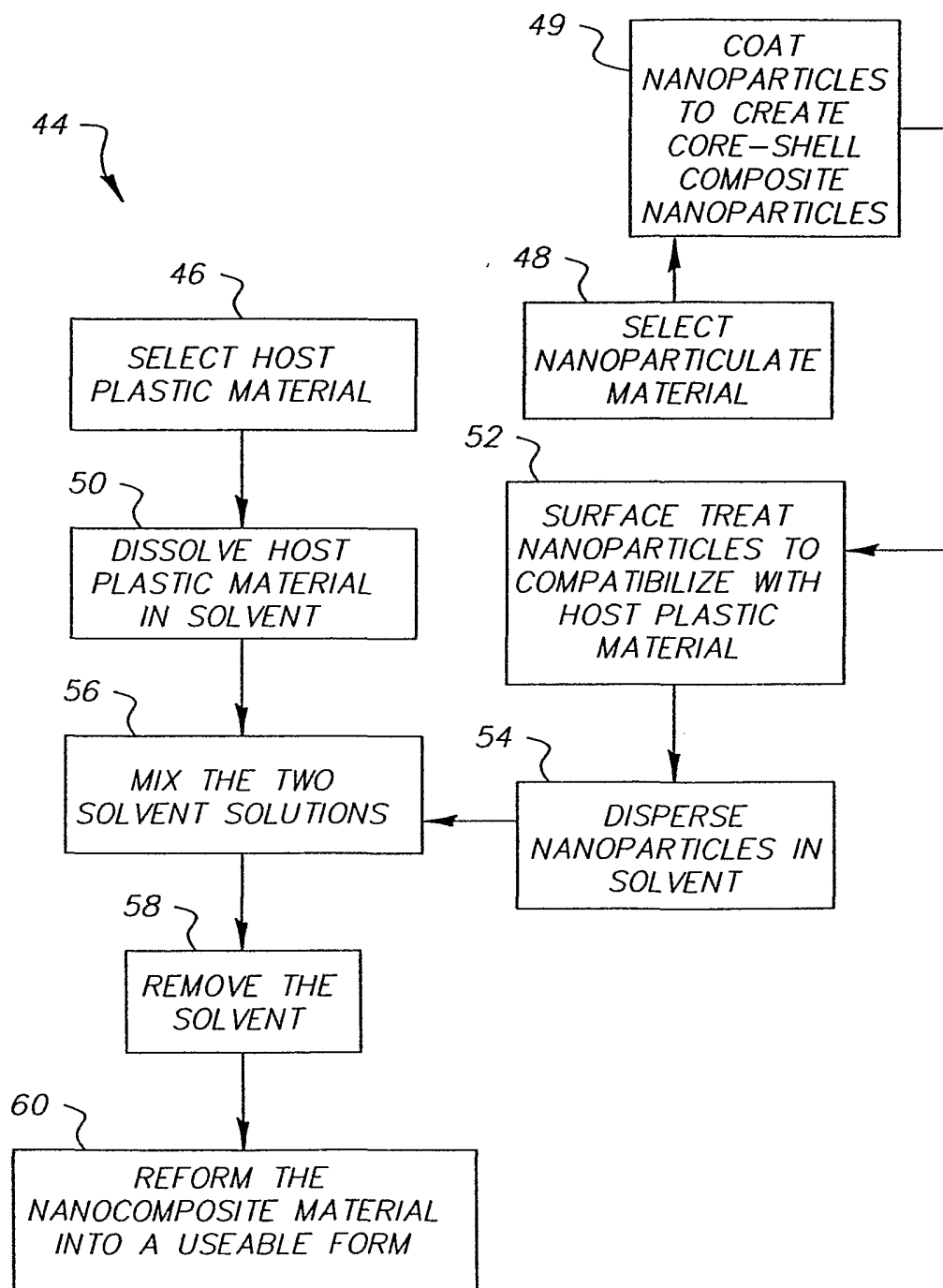


FIG. 5

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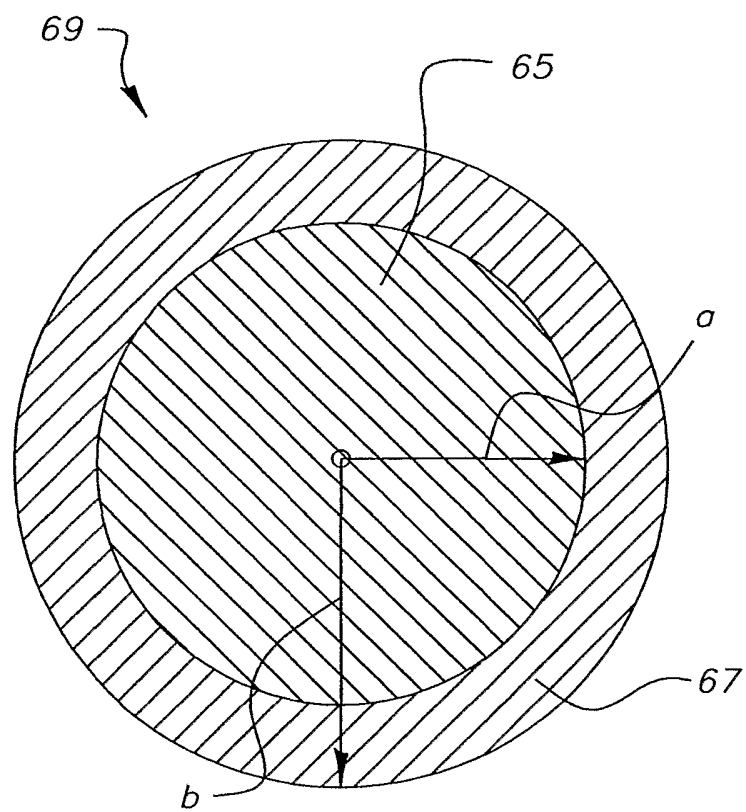


FIG. 6

INTERNATIONAL SEARCH REPORT

 International Application No
 PCT/US2004/025886

 A. CLASSIFICATION OF SUBJECT MATTER
 IPC 7 G02B1/04 C08K9/02

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)
 IPC 7 G02B C08K

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

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

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2003/119962 A1 (MCLAUGHLIN PAUL O ET AL) 26 June 2003 (2003-06-26) figure 4; examples 1,3,4 claims	1-14
A	US 5 910 522 A (JONSCHKER GERHARD ET AL) 8 June 1999 (1999-06-08) column 3, line 39 - column 4, line 67 claims 1,7	1-14
X	US 2003/136946 A1 (KUMACHEVA EUGENIA) 24 July 2003 (2003-07-24) paragraph '0005! figure 9 claims 1,10-14	1-14
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

° Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
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Date of the actual completion of the international search

5 November 2004

Date of mailing of the international search report

12/11/2004

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INTERNATIONAL SEARCH REPORT

International Application No
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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International Application No

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